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PAR
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**MÉCANISMES DE SOLIDIFICATION DES MAGMAS BASALTIQUES : ÉTUDE
QUANTITATIVE TEXTURALE ET GEOCHIMIQUE DES LAVES DU VOLCAN
KILAUEA, HAWAII**

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RÉSUMÉ

Le volcan Kilauea, Hawaï, est probablement le système magmatique basaltique actif le plus étudié sur Terre, et représente donc un site privilégié pour l'étude des processus de solidification basaltique en milieu naturel. Une meilleure compréhension de la solidification magmatique est d'importance majeure dans le raffinement de modèles expliquant le dynamisme des chambres magmatiques, et son étude détaillée est susceptible de grandement améliorer notre connaissance de l'évolution globale des systèmes magmatiques. Dans ce contexte volcanique, les lacs de lave offrent une rare opportunité d'étudier directement la solidification magmatique et peuvent être considérés, en première approximation, comme des analogues superficiels de petites chambres magmatiques.

Le but premier de ce doctorat est de déterminer et quantifier les principaux processus de solidification magmatiques à l'œuvre dans la genèse des basaltes tholéitiques. Ce travail s'articule autour du minéral olivine comme composant central. Dans les deux premiers chapitres, l'approche est double, texture et géochimie, mais l'emphase porte sur l'aspect textural dont l'analyse de la distribution de la taille des cristaux (CSD) est la composante phare. Ce travail a été réalisé sur les laves produites par les éruptions de 1969-1974 (Mauna Ulu) et 1959 (Kilauea Iki) du volcan Kilauea. L'étude des coulées de lave produites par l'éruption du Mauna Ulu permet de mieux comprendre les processus actifs de solidification dans tout le système magmatique superficiel (la "tuyauterie") de l'édifice. L'étude du lac de lave Kilauea Iki renseigne quant à elle sur la solidification en système semi-fermé en sub-surface. Dans un dernier temps, il est question d'évaluer plus en détail l'influence de la déformation magmatique sur la structure interne des olivines, et de la quantifier, en utilisant une technique *in situ* récente de micro-diffraction des rayons X. Chacun des trois chapitres de cette thèse est un article publié ou destiné à la publication dans une revue scientifique internationale.

L'article 1 présente les résultats en éléments majeurs et traces (roche totale), les compositions de l'olivine, et les CSDs de 11 échantillons de laves du Mauna Ulu. Les variations chimiques en roche totale sont interprétées comme étant partiellement produites par addition d'olivine dans le système magmatique. Les profils CSD suggèrent qu'au moins deux populations d'olivines interviennent : (1) une population d'âges 3-40 ans, caractérisée par une faible densité de "gros" cristaux et des pentes CSD relativement faibles ; et (2) une population d'âges 1,5-15 ans, marquée par une forte densité de petits cristaux et des pentes CSD plus fortes. La gamme de compositions de l'olivine suggère que ces cristaux se sont formés à partir de magmas différents, probablement reliés par cristallisation fractionnée. La présence d'olivines déformées de toutes tailles couvrant la totalité de la gamme de compositions, montre que la population 1 provient principalement de la désintégration et assimilation d'un cumulat déformé. Cette population d'olivines représente un composant magmatique cumulatif précoce qui a subi du mûrissement textural. A l'inverse, la population 2 représente un composant magmatique tardif formé dans la région sommitale de stockage de magma. Nos résultats sont en accord avec l'hypothèse que ces deux composants magmatiques ont suivi deux trajets différents avant d'alimenter l'éruption du Mauna Ulu. Le magma contenant les olivines déformées aurait transité le long du décollement basal sous le Kilauea, puis remonté verticalement par des conduits de type "*pipe*" sous le rift du Mauna Ulu. Le magma contenant la plupart des olivines non déformées aurait quant à lui transité

vers le réservoir sommital à travers le conduit magmatique principal, puis le long de la rift zone où les magmas se seraient finalement mélangés dans de petites chambres magmatiques satellites. La présence de fines zonalités inverses à la bordure de certains cristaux suggère que le mélange s'est fait juste avant l'éruption.

L'article 2 présente les compositions et CSDs d'olivine provenant de scories et d'échantillons de forage (0-90 m de profondeur) du lac de lave Kilauea Iki. Trois populations d'olivines sont distinguées sur la base de leur composition en forstérite (Fo) : (1) une population riche en Fo (Fo₈₅₋₈₈) ; (2) une population intermédiaire (Fo₇₇₋₈₁) ; et (3) une population mineure appauvrie en Fo (Fo₇₂₋₇₆). Les populations 1 et 2 sont composées à la fois de cristaux déformés et non déformés. La troisième population pourrait résulter d'une phase de recroissance tardive. Dans les 60 derniers mètres du lac, l'olivine est moins riche en Fo et la proportion de cristaux déformés augmente. Ces observations laissent penser à l'existence d'une stratification minéralogique et chimique verticale dans le lac de lave. L'analyse CSD a permis d'estimer les temps de résidence des olivines dans le magma, 1-60 ans, valeurs qui sont en accord avec les estimations préexistantes. Les CSDs sont globalement uniformes eu égard à la profondeur. Cependant, certaines caractéristiques spécifiques ressortent. Ainsi, les CSDs courbées sont considérées comme évidence de mélange de magmas ou de cristaux. L'inversion de pente aux petites tailles de la plupart des CSDs du lac de lave est interprétée comme résultant du mûrissement. Les résultats de la modélisation CSD suggèrent que la décantation / sédimentation des olivines et la convection à grande échelle ne sont pas significatives dans l'évolution du lac de lave. Enfin, la stratification verticale du lac peut être expliquée de différentes façons. Il peut s'agir d'une caractéristique originelle, résultat de la stratification de la chambre magmatique source. Cependant, plusieurs évidences montrent que le magma du lac a été fortement brassé pendant toute la durée de l'éruption ; cette première hypothèse n'est donc pas crédible. Le remplissage par la base du lac durant l'éruption serait une autre hypothèse à même d'expliquer cette stratification. Cependant, il nous manque encore de quoi définitivement valider cette théorie.

L'article 3 présente l'analyse microstructurale *in situ* par micro-diffraction des rayons X (μ XRD) d'olivines déformées et non déformées provenant d'une sélection d'échantillons préalablement étudiés dans les articles 1 et 2. Cette étude utilise une technique innovante, non destructive, peu coûteuse et rapide à mettre en œuvre permettant de recueillir des informations sur la structure interne des cristaux, ainsi que le mode et l'intensité de déformation. Les résultats ont permis de valider les observations pétrographiques de déformation faites à l'aide du microscope. Cette analyse μ XRD a aussi permis de confirmer la présence de déformation pour toutes les tailles de grains d'olivine, sans corrélation simple avec leur chimie, et de quantifier cette déformation. Cette technique ne permet cependant pas une estimation simple des conditions pression-température de déformation ou de formation des cristaux, ni d'apporter d'informations sur l'histoire magmatique. Il a cependant été possible de fixer un seuil quantitatif au-delà duquel toute olivine est déformée de façon significative : *full width at half maximum* (FWHM) > 1°.

Mots-clés : Kilauea ; Hawaii ; solidification magmatique ; basalte tholéiitique ; olivine ; distribution de la taille des cristaux (CSD) ; géochimie ; déformation cristalline ; Mauna Ulu ; Kilauea Iki ; lac de lave.

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« N'allez pas là où le chemin peut mener.
Allez là où il n'y a pas de chemin et laissez une trace ».

[Ralph Waldo Emerson]

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INTRODUCTION

La solidification du magma est un des problèmes fondamentaux en pétrologie ignée et volcanologie ; la surface de la Terre, qui est en perpétuelle évolution, est composée en partie de roches solidifiées dérivées d'un magma. La solidification comprend principalement la cristallisation, mais peut aussi inclure la dissolution des cristaux. Bien comprendre la solidification du magma est importante pour le raffinement de modèles concernant le dynamisme des chambres magmatiques, par exemple. Son étude détaillée peut améliorer notre connaissance de l'évolution des systèmes magmatiques, depuis la génération de magma au niveau de la frontière croûte-manteau ou dans le manteau plus profond, jusqu'à l'éruption volcanique en surface suite à la remontée de magma. Il y a solidification du magma en profondeur, dans le système magmatique de l'édifice volcanique, ou bien en surface, lors de l'éruption volcanique ainsi que la mise en place et le refroidissement des laves. Les roches volcaniques et leurs cristaux apportent ainsi de l'information sur les conditions de solidification du magma. L'information peut être recueillie à partir de la texture d'une roche (par exemple, la taille, type et distribution de ses cristaux), de sa composition chimique et minéralogique, mais aussi à partir de la composition chimique des cristaux. L'étude de la solidification du magma peut aussi aider à reconstruire la dynamique de transport et les trajets empruntés par le magma à travers l'ensemble du système magmatique sous-édifice.

L'information recueillie sur la solidification magmatique permet une meilleure détermination des principaux mécanismes physiques influençant le refroidissement, la solidification et la différenciation chimique d'un corps magmatique ; il s'agit d'un grand défi en

pétrologie ignée (e.g. Jellinek and Kerr, 2001). Les processus de refroidissement et de cristallisation sont par ailleurs les clés de la compréhension de l'évolution crustale continentale et océanique (e.g. Rupke and Hort, 2004). La solidification magmatique est cependant un processus complexe qui demeure mal compris et peu quantifié aujourd'hui, malgré les nombreuses études entreprises depuis quelques décennies. Son étude en milieu naturel, bien que généralement problématique, mérite une attention toute particulière.

La solidification magmatique comporte de nombreux processus intrinsèques, variables en nombre comme en importance suivant les conditions et l'environnement de refroidissement (e.g. Cashman, 1993). Le développement des textures ignées est grandement influencé par ces processus qui peuvent être de nature cinétique, mécanique ou de rééquilibrage. Des processus cinétiques produisent des populations cristallines qui peuvent ensuite être modifiées par des processus mécaniques, puis se rééquilibrer partiellement lorsque les forces cinétiques sont réduites (Fig. 1). La texture finale d'une roche ignée est donc le résultat de la combinaison entre effets cinétiques et effets liés au rééquilibrage (Higgins, 2006), s'il n'y a aucune réaction minéralogique subséquente. Les mécanismes spécifiques impliqués dans ces trois grands groupes se résument comme suit : (1) pour les processus cinétiques, il s'agit de la nucléation et croissance cristallines (e.g. Bonin, 1998; Dowty, 1980a; Lofgren, 1980) ; (2) pour les processus de rééquilibrage, le mûrissement textural (e.g. Baronnet, 1984; Cabane *et al.*, 2005; Higgins, 1998; Higgins and Roberge, 2003; Holness *et al.*, 2007; Hunter, 1996; Park and Hanson, 1999; Voorhees, 1992) ; et (3) pour les processus mécaniques, correspondant pour la plupart à la séparation entre liquide silicaté et cristaux, il s'agit de la décantation gravitaire des cristaux (Helz, 1980; e.g. Helz, 1987a; Higgins, 2002; Hort *et al.*, 1993; Hoshida *et al.*, 2006; Jellinek and Kerr, 2001; Marsh, 1988, 1996; Martin and Nokes, 1988, 1989; Rudman, 1992), l'agrégation (e.g. Schwindinger, 1999;

Schwindinger and Anderson, 1989), et la compaction et formation de chaînes de cristaux (e.g. Gray *et al.*, 2003; Jerram *et al.*, 2003; Philpotts *et al.*, 1999; Philpotts and Philpotts, 2005). La convection (ou advection) dans le corps magmatique (e.g. Bartlett, 1969; Brandeis and Jaupart, 1986; Campbell, 1996; Couch *et al.*, 2001; Davaille and Jaupart, 1993; Hort *et al.*, 1999; Huppert and Sparks, 1984; Jaupart and Tait, 1995; Martin *et al.*, 1987; Neri, 1998; Turner and Campbell, 1986; Worster *et al.*, 1993) et le mélange de magmas (e.g. Clague *et al.*, 1995; Helz, 1987b; Jellinek and Kerr, 2001; Martin *et al.*, 2006; Rhodes *et al.*, 1979; Shamberger and Garcia, 2007; Thornber, 2003; Worster *et al.*, 1993; Wright, 1973; Wright *et al.*, 1975), peuvent également jouer un rôle important en tant que processus mécaniques.

Ces processus peuvent marquer de leur empreinte la texture et/ou la chimie des roches volcaniques qui en résultent. L'approche est donc double dans ce travail de doctorat, texture et géochimie, mais l'emphase porte sur l'aspect textural.

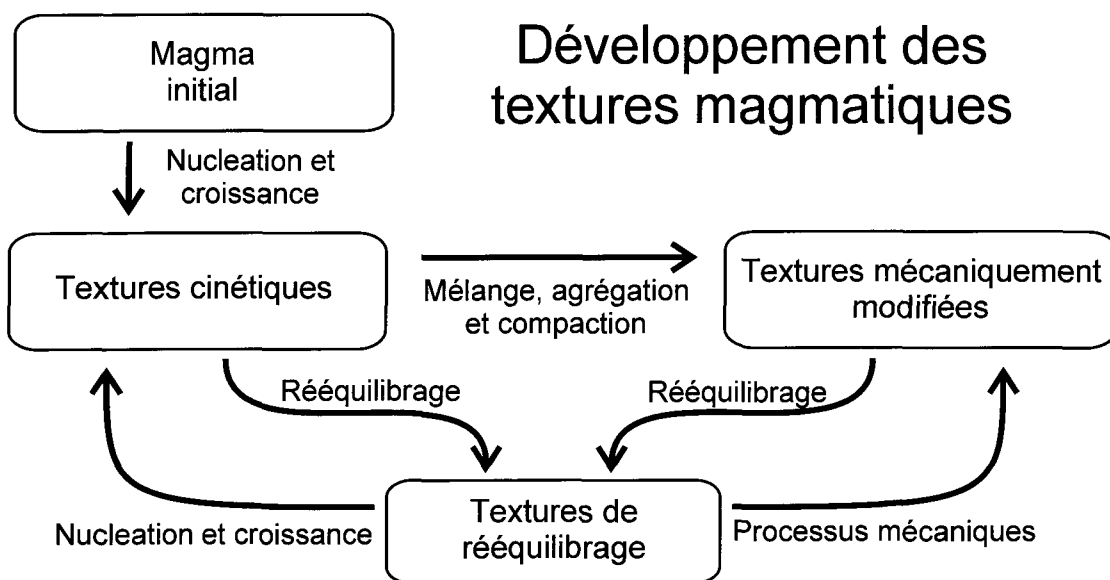


Figure 1 : Développement des textures magmatiques.

Rôle des processus cinétiques, mécaniques et de rééquilibrage. Modifiée d'après Higgins (2006)

1 Problématique

Cette thèse s'intéresse spécifiquement à la solidification des magmas basaltiques (*sensu lato*), et s'articule autour d'un composant central : l'olivine. Il s'agit de mieux comprendre (et quantifier) les processus de solidification des magmas basaltiques par le biais d'une étude texturale et géochimique quantitative appliquée en milieu naturel, les laves du volcan Kilauea, Hawaïi. Dans un premier temps, je traiterai des différents mécanismes de solidification magmatique, puis de quelques modèles de solidification des magmas basaltiques, et enfin de la déformation de l'olivine.

1.1 Mécanismes physiques de solidification magmatique

La solidification magmatique se caractérise, comme pour tous les processus de pétrologie ignée, par un transfert d'énergie et éventuellement de masse, de façon à atteindre un état d'équilibre à un niveau d'énergie plus faible (Bonin, 1998). La solidification d'un magma est en général une conséquence directe de son refroidissement. Les mécanismes physiques intervenant lors de la solidification des magmas basaltiques peuvent se décomposer en trois grandes catégories (fig. 1) : (1) les processus cinétiques ; (2) les processus de rééquilibrage ; et (3) les processus de séparation mécanique entre liquide silicaté et cristaux. A l'intérieur du corps magmatique basaltique en cours de refroidissement, deux contextes physiques de solidification peuvent être différenciés : (i) les mécanismes au niveau des fronts de solidification (Marsh, 1996), couches limites partiellement cristallisées ; et (ii) les mécanismes ayant lieu à l'intérieur même du corps magmatique, dans la partie dominée par le liquide silicaté. Suivant l'environnement

magmatique considéré – chambre magmatique, lac de lave, ou coulée de lave, par exemple – tous ces processus peuvent varier en nature comme en importance.

1.1.1 Processus cinétiques : nucléation et croissance cristallines

Les processus de solidification majoritairement contrôlés par des effets cinétiques, dépendent directement du degré de surfusion ("*undercooling*") du système (e.g. Dowty, 1980a; Lofgren, 1980). Le paramètre de surfusion correspond à un état de déséquilibre du système qui est généralement retranscrit en terme de différence ΔT entre température du liquidus et température réelle du magma (Baronnet, 1984; Bonin, 1998; Brandeis and Jaupart, 1986; Brandeis *et al.*, 1984; Dowty, 1980a; Lofgren, 1980). Le passage de l'état de liquide silicaté à celui de solide cristallisé nécessite deux étapes (Bonin, 1998) : la nucléation puis la croissance cristallines.

Nucléation des cristaux

Deux types de nucléation sont communément distingués (Dowty, 1980a; Lofgren, 1980) : (i) la nucléation homogène, un phénomène spontané induit par des fluctuations thermiques aléatoires dans le liquide silicaté ; et (ii) la nucléation hétérogène, qui prend naissance à la surface de particules "étrangères" préexistantes en contact avec le liquide silicaté. Ces particules étrangères peuvent prendre la forme de grains hôtes, de bulles, de défauts cristallins, d'impuretés, de parois, etc.

La formation d'un nucléus est directement fonction de l'état énergétique du système magmatique ; l'état énergétique du système est exprimé par la variation d'énergie libre totale ΔG_0 (Fig. 2a). Considérant une particule de rayon r , le maximum est atteint pour $\Delta G_0 = \Delta G^*$ et $r = r^*$

(Bonin, 1998); ΔG^* est appelé l'énergie libre critique (ou énergie d'activation), et r^* , le rayon critique. L'énergie libre totale d'une particule est la somme des énergies de surface et de volume (elle-même reliée à l'énergie de liaisons), toutes deux ayant un effet énergétique opposé (Tiller, 1991). Dans le cas de la *nucléation homogène*, il y a formation des nucléi (et début de croissance) pour $\Delta G_0 \geq \Delta G^*$, soit pour $r_{\text{nucléi}} \geq r^*$ (e.g. Dowty, 1980a). Pour un degré de surfusion trop faible, la barrière énergétique ΔG^* ne peut être franchie. Le rayon critique r^* n'est pas atteint. Le taux de nucléation est alors nul, et aucune nucléation n'est possible. Lorsque le degré de surfusion augmente, le taux de nucléation augmente également (Fig. 2b).

La *nucléation hétérogène* est associée à une réduction de l'énergie de surface du nucléus. Ceci implique une réduction du degré de surfusion nécessaire pour le début de cristallisation, et donc une diminution du rayon critique et de la barrière énergétique (Dowty, 1980a). Le degré de surfusion est alors relativement faible, généralement de l'ordre de quelques degrés Celsius (Bonin, 1998). Il est fonction du degré de similarités cristallographiques entre la phase hôte et la nouvelle phase. Ce degré de similarité est matérialisé par les angles diédraux θ (Tiller, 1991). Plus les angles diédraux sont faibles ($\theta < 90^\circ$), plus le degré de surfusion nécessaire à la nucléation sera faible (Fig. 2b). La nucléation hétérogène est donc possible à une température inférieure à la température de nucléation homogène.

En résumé, la nucléation dépend principalement du taux de refroidissement, de la surfusion et du temps d'incubation – temps requis pour que la nucléation ait lieu une fois que $\Delta T > 0$ (Lofgren, 1980). Le temps d'incubation influence ainsi le nombre de sites de nucléation produits.

Croissance cristalline et dissolution

Il y a croissance des cristaux s'il y a réduction de l'énergie libre totale du système en cours de cristallisation. De façon similaire, les cristaux peuvent également se dissoudre partiellement ou totalement si cela induit une réduction de l'énergie totale. Le mécanisme de croissance cristalline se fait par addition d'atomes ou molécules à la surface du cristal. Le taux d'addition atomique, et donc le taux de croissance, peut être contrôlé par deux processus de croissance principaux (e.g. Baronnet, 1984), conduisant à des morphologies différentes (Bonin, 1998) : (i) La croissance contrôlée par la cinétique d'attachement à l'interface (Dowty, 1980a; Tiller, 1991), ce qui produit des cristaux idiomorphes ; et (ii) la croissance contrôlée par le transfert de matière et de chaleur (diffusion et advection), ce qui produit des cristaux aux habitus variés (Lofgren, 1980) et peut par exemple générer la formation de zonalités.

Ces différents types de croissance s'observent pour des surfusions différentes selon la nature du minéral et sa place dans la séquence de cristallisation (Bonin, 1998). Pour un faible taux de surfusion, le taux de croissance sera contrôlé par les processus à l'interface. A l'inverse, pour un degré de surfusion élevé, les processus de transfert de matière et de chaleur deviennent prépondérants. Le taux de cristallisation, directement relié aux taux de nucléation et de croissance, est également contrôlé par ce degré de surfusion (Fig. 3 ; zone grisée). Ce taux de cristallisation est aussi fonction, entre autres, de la composition chimique du magma, donc du système considéré.

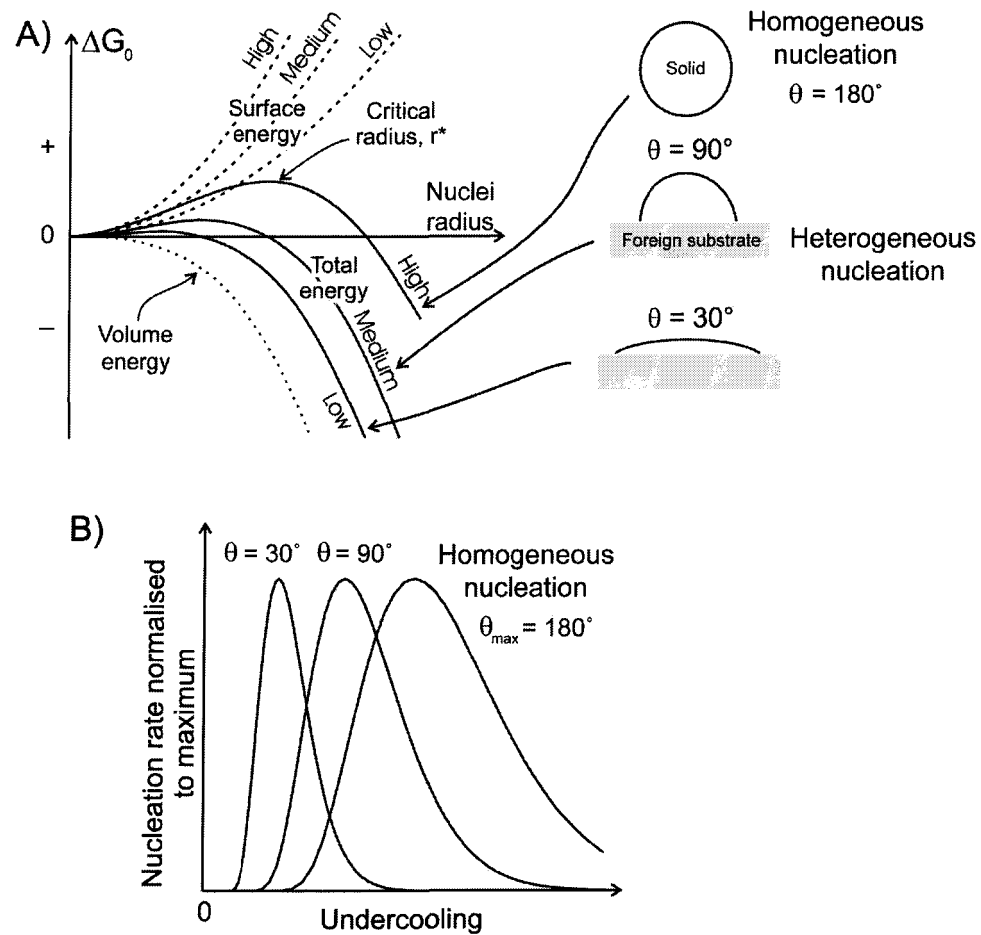


Figure 2 : Énergie libre totale et degré de surfusion vs. Nucléation.

(A) Énergie libre totale ΔG_0 de formation d'un nucléus en fonction de son rayon. Cette énergie totale est la somme de l'énergie de surface et de l'énergie de volume (liée à l'énergie de liaisons). (B) Taux de nucléation en fonction du degré de surfusion du système. La nucléation hétérogène (angles diédraux $\theta \neq 90^\circ$) réduit le degré de surfusion nécessaire pour la nucléation. Une barrière énergétique ΔG^* (non montrée ici) plus basse implique un θ et donc un degré de surfusion plus faible. Modifiée d'après Higgins (2006)

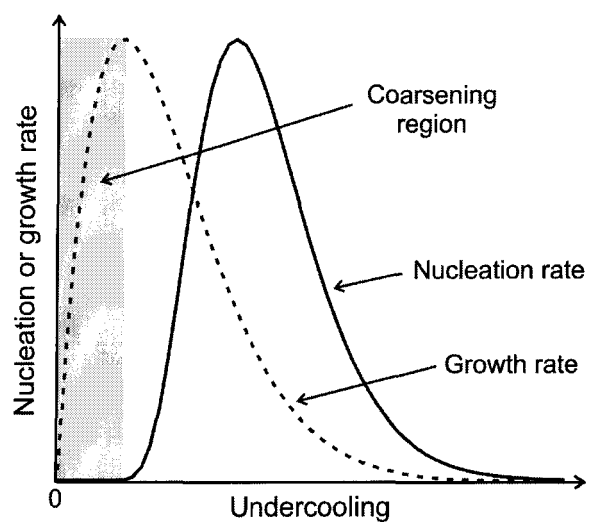


Figure 3 : Taux de nucléation ou de croissance en fonction du degré de surfusion.
 Dans les roches ignées, le mûrissement textural a lieu pour des taux de surfusion faibles (zone grisée), proches du liquidus, où le taux de nucléation est quasi nul mais le taux de croissance significatif. Modifiée d'après Higgins (2006)

1.1.2 Processus de rééquilibrage : mûrissement textural

Une fois les forces motrices cinétiques réduites, les populations de cristaux entrent en phase de rééquilibrage (Higgins, 2006). La texture cristalline répond à l'effet du rééquilibrage qui participe à la minimisation de l'énergie totale du système. Bien que l'équilibre complet ne soit jamais atteint, il y a homogénéisation de la distribution de la taille des cristaux, phénomène inexistant lors de processus purement cinétiques.

Dans le cas idéal d'un système magmatique fermé (solide + liquide silicaté), celui-ci atteindra son équilibre chimique et textural seulement si la contribution de surface à l'énergie libre totale est minimisée (Baronnet, 1984). Cette étape est satisfaite par diminution du rapport interface/volume des phases déjà cristallisées. Ceci peut avoir lieu plus ou moins simultanément par le processus de mûrissement textural ("*textural coarsening*"), via le changement d'habitus de croissance vers une forme d'équilibre. Ce mûrissement textural agit en plusieurs étapes quasi-simultanées (Baronnet, 1984) (Fig. 4) : résorption des particules les plus petites, transfert de masse par diffusion à travers le milieu environnant ou l'interface cristal-cristal, et croissance des particules les plus grosses. Ce mécanisme a principalement été mis en évidence en contexte plutonique (e.g. Higgins, 1998, 2002; Hunter, 1996; McBirney and Nicolas, 1997), mais semble également jouer un rôle important en domaine volcanique (Higgins and Roberge, 2003; Park and Hanson, 1999; Pupier *et al.*, 2008).

Comme discuté précédemment, pour des systèmes ignés la variation des taux de nucléation et de croissance fluctue avec le degré de surfusion (Fig. 3). Le mûrissement textural ne peut ainsi avoir lieu que pour un taux de nucléation quasi nul et un taux de croissance significatif, à savoir proche du liquidus du minéral considéré.

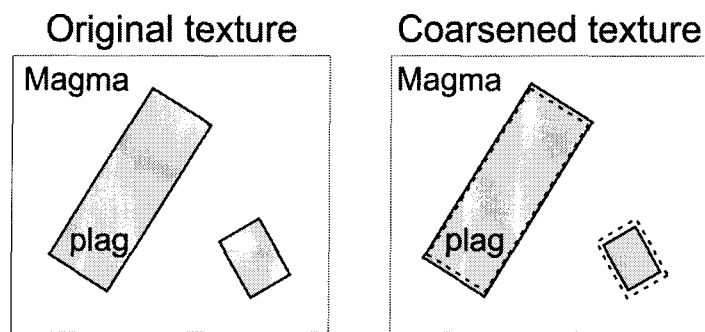


Figure 4 : Schéma de l'effet du mûrissement textural sur des cristaux de plagioclase immergés dans un magma basaltique. Modifiée d'après Higgins (2006)

1.1.3 Processus de séparation mécanique entre liquide silicaté et cristaux

Les caractéristiques cristallines comme la taille, la composition, l'orientation, et le nombre de cristaux, peuvent être modifiées par des processus mécaniques tels que le fractionnement et l'accumulation de cristaux, la formation de chaînes de cristaux, la compaction, et l'agrégation (Higgins, 2006). Certains de ces processus interviennent en systèmes fermés dans le sens où la masse totale du système reste inchangée. D'autres ont lieu en systèmes ouverts lorsqu'il y a addition/soustraction de cristaux ou de magma.

1.1.3.1 Fractionnement et accumulation de cristaux

Les processus de fractionnement et accumulation de cristaux (Fig. 5) correspondent à la séparation des cristaux du liquide silicaté (Bowen, 1928). Plusieurs mécanismes peuvent participer à cette séparation cristaux/liquide (Bonin, 1998) :

- l'exsolution des gaz ("*filter pressing*"), contrôlée par la formation de réseaux de cristaux, la viscosité du liquide, et le facteur temps ;
- la différenciation par fluage (ségrégation de flux ou effet *Bagnold*), contrôlée entre autres par la vitesse d'écoulement, la viscosité du liquide, la cristallinité, et le gradient de vitesse ;
- la décantation gravitaire, principalement contrôlée par le contraste de densité entre liquide et cristaux, la viscosité du liquide, et la taille et forme des cristaux ;
- la cristallisation aux parois.

Ces mécanismes sont associés à la distribution des phases solides dans le liquide, au niveau des fronts de solidification supérieur et inférieur du corps magmatique (Marsh, 1996) (Fig. 5). Ils dépendent aussi de la forme des grains et du comportement rhéologique du liquide silicaté, qui varie en fonction du degré de solidification (Bonin, 1998).

Les fronts de solidification constituent la masse de magma spatialement contenue entre les isothermes du *solidus* et du *liquidus*, et s'épaississent vers l'intérieur du corps magmatique au cours du refroidissement (Marsh, 1996). Ils se divisent en plusieurs zones rhéologiques distinctes suivant la cristallinité du magma (Fig. 6). (i) Pour une faible cristallinité, les cristaux sont en suspension dans le magma ; celui-ci se comporte alors comme un fluide visqueux Newtonien. Les phénomènes gravitaires comme la décantation des cristaux peuvent se produire, puisque les cristaux sont libres de bouger les uns par rapport aux autres. (ii) Pour une forte cristallinité, le magma se comporte comme un solide dont le réseau cristallin est plastique. Le liquide résiduel interstitiel ne peut s'extraire facilement, excepté s'il y a un gradient de pression. (iii) La zone intermédiaire se présente sous la forme d'un mélange cristaux-magma, une bouillie cristalline ("*mush*") ou suspension dense. Les cristaux restent en suspension dans le magma, mais leurs mouvements sont réduits puisqu'ils se touchent. En domaine basaltique, dans le cas (i) le niveau de cristallinité limite est d'environ 25% de cristaux, > 55% pour (ii), et 25-55% pour (iii).

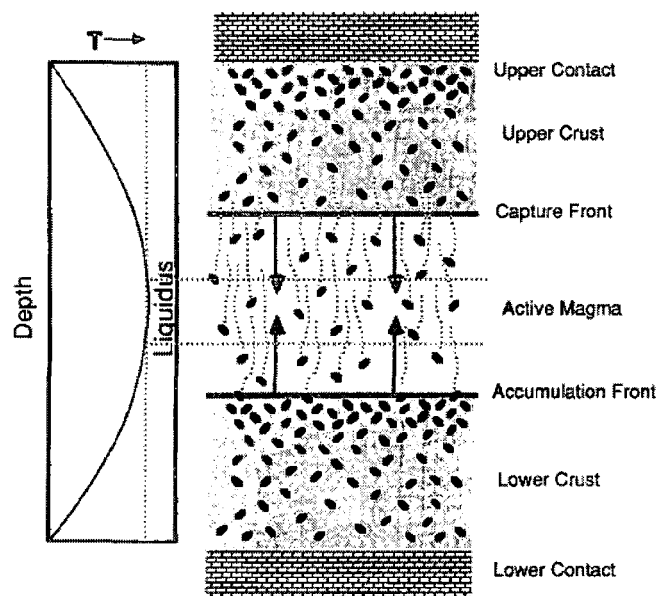


Figure 5 : Colonne de magma schématique.

Ce schéma illustre le fractionnement et l'accumulation de cristaux, ainsi que les fronts de solidification supérieur (front de capture) et inférieur (front d'accumulation). Reproduit de Marsh (1988)

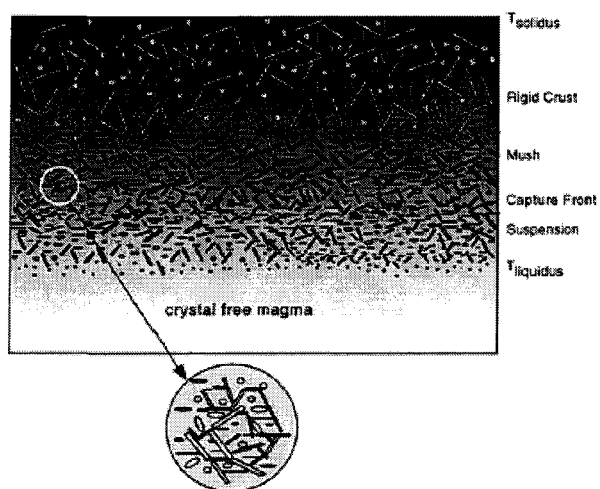


Figure 6 : Accumulation et capture de cristaux au niveau du front de solidification supérieur (front de capture). La limite supérieure du front est définie par le solidus, et la limite inférieure par le liquidus. Reproduit de Marsh (2002)

1.1.3.2 Formation de chaînes de cristaux, compaction et agrégation

Compaction et chaînes de cristaux

Le processus de compaction débute par une phase mécanique lors de laquelle un changement de l'orientation des grains permet un agencement plus dense des grains, par tassement avec expulsion du fluide interstitiel (Bowen, 1928). Cette compaction mécanique sans déformation interne des grains est limitée à un maximum (i.e. environ 50%, mais estimation complexe, fonction de plusieurs paramètres intrinsèques et de l'environnement magmatique considéré) du fait de la cohésion, ou inertie au changement, du magma (Higgins, 1991; Hunter, 1996; Marsh, 1989). Une compaction plus forte entraînera un changement de forme des grains par fracturation ou déformation plastique, permettant un tassement plus dense. On parle alors de compaction par pression-dissolution (Meurer and Boudreau, 1998; Nicolas and Ildefonse, 1996), un mécanisme de transfert de masse induit par la contrainte. Ce mécanisme est très commun dans les roches plutoniques et métamorphiques, mais assez marginal en domaine volcanique.

Le passage de l'état de suspension à l'état de bouillie cristalline dans un magma basaltique a généralement lieu à partir de 25% de cristallisation (Marsh, 1988; Philpotts *et al.*, 1998). La compaction est alors possible seulement après l'interconnexion des cristaux pour former un réseau 3-D continu (Philpotts *et al.*, 1998). Un cas typique est celui du plagioclase, initialement de flottabilité neutre (Scoates, 2000). Cependant, seule l'olivine est communément présente sous forme de phénocristaux dans les laves du Kilauea, parfois associée à un faible pourcentage de plagioclase. Par conséquent, une telle compaction par formation de chaînes de cristaux apparaît comme très peu probable ici, et ne sera pas traitée dans la suite de ce travail.

Agrégation et synneusis

Dans la plupart des contextes géologiques, la concentration des grains est suffisamment forte pour qu'ils interagissent entre eux et s'agglomèrent en amas, particulièrement lors de leur décantation gravitaire (Schwindinger, 1999). La force contrôlant ce processus est la réduction de l'énergie totale interfaciale des grains (e.g. Baronnet, 1984; Herring, 1951; Schwindinger and Anderson, 1989). Ces amas se comportent comme un grain unique plus dense, et sédimentent donc plus vite que chacun des grains originaux. La vitesse de refroidissement élevée dans les roches volcaniques permet de préserver les agrégats, ce qui paraît moins commun dans le cas de roches cumulatives plutoniques (Higgins, 2006). Dans certains agrégats, les cristaux idiomorphes peuvent s'assembler le long de leurs faces cristallographiques. Ce phénomène, communément appelé *synneusis* (Vance, 1969; Vance and Gilreath, 1967), a été en particulier mis en évidence pour les cristaux d'olivine de l'éruption de 1959 du Kilauea Iki (Schwindinger, 1999; Schwindinger and Anderson, 1989). L'ensemble de la structure ressemble à un cristal maclé, mais sans en avoir les propriétés cristallines (Dowty, 1980b).

1.1.4 Autres processus

1.1.4.1 Convection/advection

Il y a convection dans un corps magmatique lorsque la force d'entraînement est suffisamment élevée pour dépasser la résistance liée à la viscosité et la diffusivité thermique (e.g. Bartlett, 1969; Jaupart and Parsons, 1985; Shaw, 1965). Le critère de convection est alors donné par le nombre sans dimension de *Rayleigh* qui dépend principalement de la viscosité, diffusivité

thermique, température du liquide, et de l'épaisseur du corps magmatique (e.g. Brandeis and Jaupart, 1986; Turner and Campbell, 1986). Il y a convection pour un nombre de *Rayleigh* supérieur à la valeur critique (variable suivant les conditions), et simple conduction pour un nombre de *Rayleigh* inférieur à cette valeur.

Deux types de convection peuvent intervenir lors de la solidification d'une chambre magmatique ou d'un lac de lave basaltiques (Brandeis and Jaupart, 1986; Martin *et al.*, 1987) : (1) la convection compositionnelle, contrôlée par les contrastes de densité dus à la cristallisation fractionnée et/ou à l'exsolution des gaz (Chen and Turner, 1980; Huppert and Sparks, 1984; Turner and Campbell, 1986; Wright and Okamura, 1977) ; et (2) la convection thermique, contrôlée par les contrastes de température. Les mouvements convectifs peuvent se matérialiser sous la forme de panaches verticaux partiellement cristallisés, constitués de la bouillie cristalline formant les fronts de solidification (e.g. Bartlett, 1969; Marsh, 1988, 1996; Philpotts and Dickson, 2000). Ces panaches verticaux sont susceptibles d'entraîner la compaction, ainsi que la migration de liquides résiduels interstitiels sous la forme de veines de ségrégation par exemple.

1.1.4.2 Mélange de magmas

Le mélange de magmas est un processus très commun ; de nombreuses évidences existent autant en domaine plutonique que volcanique. Deux types de mélange peuvent être distingués : (1) le mélange complet ("*mixing*"), correspondant à l'extrême limite du processus dont le résultat est un produit chimiquement homogène ; et (2) le mélange "mécanique" ("*mingling*"), correspondant à un processus moins extrême où les deux composants peuvent être reconnus par

la pétrographie et/ou la chimie. Il s'agit en fait d'un problème d'échelle de mélange, en considérant tous les intermédiaires possibles entre l'absence de mélange et le mélange parfait.

Les différents processus décrits dans cette section 1.1. sont généralement pris en considération pour l'élaboration de modèles de solidification des magmas basaltiques que je traite maintenant.

1.2 Modèles de solidification des magmas basaltiques

De nombreux modèles de solidification magmatique existent, mais la majorité est basée sur l'expérimentation en laboratoire (e.g. Brandeis and Jaupart, 1986, 1987; Brandeis and Marsh, 1989; Huppert and Worster, 1985; Jaupart and Tait, 1995; Kerr et al., 1989; Worster et al., 1993), à partir de matériaux de synthèse (systèmes binaires) ou naturels (e.g. basaltes tholéitiques), et sur la modélisation numérique (e.g. Hort, 1997; Jaupart and Tait, 1995; Jellinek and Kerr, 2001; Mangan and Marsh, 1992; Marsh, 1988, 1989, 1996; Rupke and Hort, 2004). Dans ce travail de doctorat, nous avons distingué : (1) la solidification en système ouvert à relativement faible profondeur dans une chambre magmatique sommitale ; et (2) la solidification en système fermé en sub-surface dans un lac de lave.

1.2.1 Solidification en profondeur dans une chambre magmatique

Il est communément admis que le refroidissement d'une chambre magmatique implique les effets combinés de la convection et de la cristallisation fractionnée (Brandeis and Jaupart,

1986). Les travaux expérimentaux de Brandeis and Jaupart (1986) sur les chambres magmatiques de rapports de forme élevés (effets de bordure négligeables) montrent que, pour une faible viscosité typique de magmas basaltiques très fluides, la convection intervient avant le début de la cristallisation. Le temps de croissance des cristaux est trop court pour permettre une décantation (ou sédimentation) gravitaire significative. Aucun contraste de composition ne peut se développer. Pour des viscosités plus fortes typiques de magmas moins mafiques, des instabilités thermiques se développent dans un magma partiellement cristallisé, expliquant la génération de panaches verticaux chargés en cristaux. Cependant, ce dernier cas ne sera pas traité ici, puisqu'il concerne des magmas plus évolués.

La plupart des études sur les lacs de lave naturels supposent qu'ils se comportent comme de petites chambres magmatiques, à partir du moment où une croûte superficielle s'est formée. Par conséquent, il est possible d'appliquer, dans une certaine limite, les conclusions acquises sur l'étude des lacs de lave aux chambres magmatiques, pour mieux comprendre leur mode de solidification. La présente section n'est donc pas développée davantage, au profit de la suivante sur la solidification dans les lacs de lave.

1.2.2 Solidification en sub-surface dans un lac de lave

L'étude de la solidification des lacs de lave a été l'objet de nombreux travaux, autant basés sur l'expérimentation que sur des échantillons naturels ; ces deux approches sont souvent combinées à la modélisation numérique.

Worster *et al.* (1993) ont montré expérimentalement que la convection thermique, uniquement contrôlée par la surfusion de solidification, peut produire des variations

compositionnelles et texturales dans un corps lavique initialement uniforme (Fig. 7). Ceci est principalement valable dans le cas d'un refroidissement par le toit du corps lavique. Les mouvements convectifs réduisent le temps de solidification complète, et entraînent la cristallisation secondaire du liquide silicaté dans des régions éloignées des fronts de solidification. Ces travaux mettent en évidence l'effet de la viscosité et du dégazage (exsolution des gaz) sur l'histoire de la solidification.

Jellinek and Kerr (2001) suggèrent, via une modélisation numérique basée sur des échantillons naturels, que les forts taux de refroidissement au toit et à la base du lac de lave Kilauea Iki aboutissent à des contrastes thermiques et compositionnels, ainsi qu'à des variations dans la vésicularité et la cristallinité (variation de la quantité d'olivine par exemple). Les auteurs proposent que tout ceci engendre des mouvements convectifs chaotiques et vigoureux qui agitent le magma (Fig. 8), produisant un mélange thermique et chimique imparfait mais efficace sur presque toute l'histoire de solidification du lac de lave. Cet aspect dynamique de la solidification du Kilauea Iki est en désaccord avec les conclusions de Helz (1989) qui considère que les transferts diapiriques de matériel du toit vers la base du lac ont lieu sans mélange. À l'inverse, la vision de Jellinek and Kerr (2001) du mélange induit par la convection est en accord avec le modèle dynamique de Worster *et al.* (1993).

De plus, la comparaison entre la modélisation numérique et les données acquises sur les lacs de lave hawaïens, montre que le refroidissement par les parois latérales a un impact significatif sur la solidification des lacs de lave (Rupke and Hort, 2004).

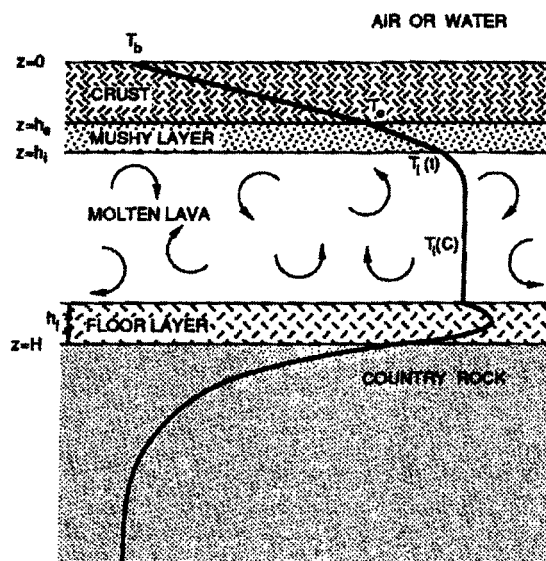


Figure 7 : Modèle simple d'un lac de lave.

Le refroidissement principal se fait par transfert convectif à travers le toit, et secondairement par transfert de chaleur par conduction vers les roches encaissantes basales. Une croûte solide composite croît depuis le toit vers la base du lac de lave. Une couche constituée d'une bouillie cristalline sépare la croûte de solidification et le liquide convectif.

T_b , température du toit du lac, supposée constante ; T_e , température de l'eutectique ; T_i , température à l'interface entre la bouillie cristalline et le liquide convectif ; $T_l(C)$, température d'équilibre du liquide convectif ; H , profondeur totale du lac.

Reproduit de Worster *et al.* (1993)

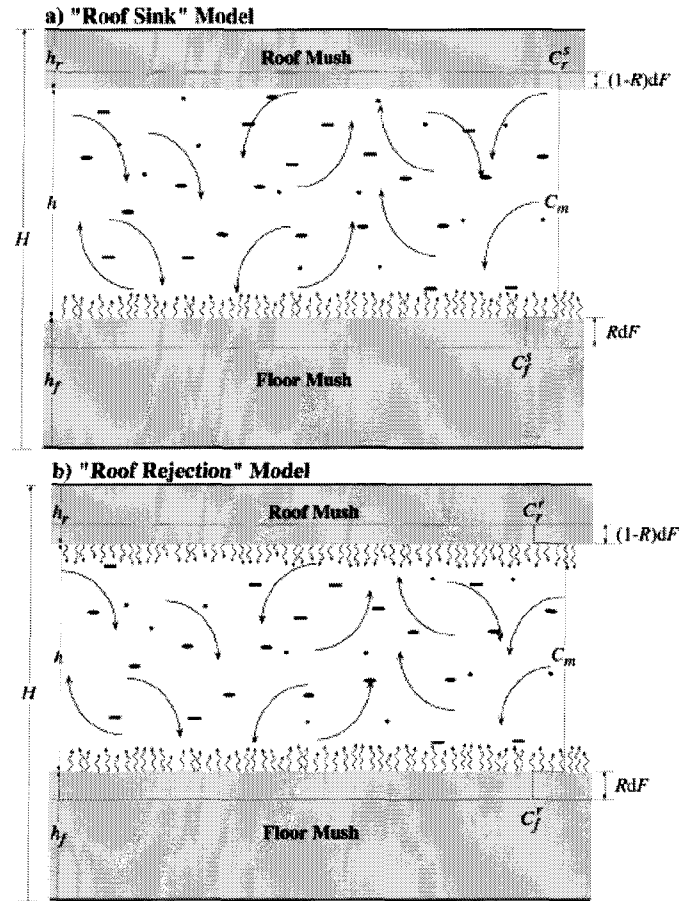


Figure 8 : Schémas descriptifs des modèles de solidification du lac de lave du Kilauea Iki.

Ces modèles sont basés sur le processus de fractionnement de Rayleigh, tout en supposant un mélange complet. La solidification progresse par cristallisation *in situ* d'olivine au niveau du toit et de la base du lac de lave, puis sédimentation des cristaux, ce qui provoque la diminution de la hauteur de magma, h . Dans le modèle (a) ("roof sink model"), il y a différenciation par (1) mélange convectif du liquide résiduel qui remonte depuis les cristaux de base de lac, et (2) par mélange du liquide résiduel restant issu de la décantation gravitaire des cristaux depuis l'intérieur convectif du lac. Le liquide résiduel produit au niveau du front de solidification supérieur reste à ce niveau et n'influence pas l'évolution chimique du magma sous-jacent. Dans le modèle (b) ("roof rejection model"), le liquide résiduel produit au niveau du front de solidification supérieur est incorporé et brassé dans le magma sous-jacent, puisque entraîné par des panaches thermiques descendants.

$F = h/H$; $R = h_f/(h_f + h_r)$; $(1-R)dF$: incrément de fractionnement au sommet du lac ; RdF : incrément de fractionnement en base de lac ; C_r : concentration globale du front de solidification supérieur ("r" pour "roof") ; C_f : concentration globale du front de solidification inférieur ("f" pour "floor") ; les exposants "s" et "r" sur C_r et C_f font référence aux modèles "Roof Sink" et "Roof Rejection", respectivement. Reproduit de Jellinek and Kerr (2001)

À partir de la modélisation numérique, certains auteurs ont mis en évidence plusieurs processus de solidification des lacs de lave hawaïiens (Jellinek and Kerr, 2001; Marsh, 1988; Moore and Evans, 1967). Par souci de simplification et de clarté, ces processus ne sont pas décrits en détail mais simplement listés ci-dessous :

- Mouvements de convection par entraînement vertical d'un panache, à la fois vers la base et le toit du lac de lave ;
- Stratification compositionnelle mineure au toit, et stratification thermique à la base du lac de lave, toutes deux causées par le mélange incomplet dû à la convection ;
- Nucléation, croissance et sédimentation gravitaire des cristaux d'olivine ;
- Flux ascendant de magma évolué (veines de ségrégation) dû à la convection de liquides interstitiels dans la bouillie cristalline ;
- Perte de chaleur accentuée par l'infiltration hydrothermale au niveau du toit et de la base du lac.

A défaut d'être exhaustive, cette liste synthétique apporte quelques informations utiles, parfois quantitatives, sur les processus de solidification qu'il est bon de prendre en considération en tant que base de travail dans cette thèse. Cette liste n'est cependant pas directement applicable en totalité dans le cadre d'une telle étude en milieu naturel que représente cette thèse ; certains de ces processus ne sont pas quantifiables ni même détectables par le biais de l'analyse texturale ou géochimique.

D'un intérêt certains pour cette thèse, les études précédentes en milieu naturel apportent une information complémentaire très utile comme base à ce travail. Ainsi, les études en systèmes naturels de certains lacs de lave du Kilauea (Kilauea Iki, picritique ; Makaopuhi, basaltique tholéiitique) ont mis en évidence les processus de solidification suivants :

- Redistribution des phénocristaux (d'olivine principalement) par décantation/sédimentation gravitaire (Cashman and Marsh, 1988; Helz, 1987a; Wright and Okamura, 1977) ;
- Migration et perte des bulles de gaz (vésicules) par dégazage du magma (Helz, 1987a) ;
- Différenciation du magma par redistribution des olivines, augites et plagioclases lors de la convection (Cashman and Marsh, 1988; Helz, 1987a) ;
- Formation de veines pauvres en olivine par ségrégation de liquides interstitiels résiduels (Cashman and Marsh, 1988; Helz, 1987a; Jellinek and Kerr, 2001; Richter and Moore, 1966; Wright and Okamura, 1977) ;
- Mélange de magmas (Helz, 1987b).

Cette liste n'est pas exhaustive non plus, mais elle recense les principaux processus de solidification observables et parfois quantifiables par l'étude en milieu naturel des textures et des compositions chimiques.

L'ensemble des modèles proposés dans la littérature, et synthétisés ici, ne tient pas compte de la présence de cristaux déformés et leur influence sur le mode de solidification. Dans cette thèse, l'emphasis est portée sur l'olivine puisqu'il s'agit du minéral le plus commun dans les magmas basaltiques, et le plus précoce dans la séquence de cristallisation.

1.3 Déformation des olivines

La présence d'olivines déformées est commune dans les roches mantelliques et plutoniques, mais rare en contexte volcanique. Les cas connus d'olivine volcanique déformée se

trouvent principalement dans les laves d'Hawaii, pour les volcans Mauna Kea et Kilauea (e.g. Clague and Denlinger, 1994; Clague *et al.*, 1995; Yang *et al.*, 1994), et dans les laves du Piton de la Fournaise, à La Réunion (Albarede *et al.*, 1997; Albarede and Tamagnan, 1988; Boivin and Bachelery, 2009; Famin *et al.*, 2009; Welsch *et al.*, 2009). Ces cristaux déformés proviennent en très grande majorité (voire en totalité) de cumulats magmatiques formés au sein même de l'édifice volcanique. Il s'agit donc ici de déformation magmatique.

La déformation magmatique des cristaux d'olivine peut être considérée comme un processus spécifique de la solidification des magmas basaltiques. De nombreuses études expérimentales ont été menées dans le but de démontrer l'influence de la déformation plastique des olivines sur la rhéologie du manteau (e.g. Langdon, 1985; Li *et al.*, 2004; Li *et al.*, 2006; Mercier, 1985; Raleigh, 1968; Raterron *et al.*, 2004). La raison principale en est qu'une meilleure compréhension du fluage plastique dans le manteau est essentielle dans la formation et le raffinement des modèles expliquant les processus géodynamiques et les propriétés géophysiques ou rhéologiques du manteau terrestre (le manteau supérieur en particulier) (e.g. Mercier, 1985).

La déformation des roches est influencée par de nombreux facteurs intrinsèques et extrinsèques (Hobbs, 1985); certains contrôlent la forme, l'orientation et la distribution des grains, et donc laissent une empreinte sur les microfabriques. La température est un paramètre important influençant la déformation. Plusieurs études basées sur l'expérimentation à des pressions et températures variées, ont proposé des équations constitutives pour décrire les mécanismes de dislocations intervenant dans les cristaux d'olivine, en tenant compte de toute une gamme de conditions thermodynamiques, thermochimiques et microstructurales (Bai and Kohlstedt, 1992; Bai *et al.*, 1991; Demouchy *et al.*, 2009; Durham and Goetze, 1977; Evans and Goetze, 1979; Kohlstedt and Goetze, 1974; Li *et al.*, 2004; Raterron *et al.*, 2004).

La déformation des olivines est maintenant connue en contexte volcanique. Cependant, ce processus demeure mal contraint, bien que la présence d'olivines déformées dans certaines roches volcaniques soit fréquente et que leur importance dans l'histoire magmatique d'édifices volcaniques tel le Kilauea soit parfaitement reconnue. Certes encore considérée comme une curiosité par certains, la déformation des olivines dans un tel contexte volcanique apparaît comme un point à ne pas négliger pour une bonne compréhension du système magmatique dans son ensemble, et de la solidification des magmas basaltiques en particulier.

Dans un tel contexte volcanique, la présence, la formation et l'histoire de solidification des cristaux déformés d'olivine sont des thématiques clés. L'étude de ces cristaux déformés est d'un grand intérêt pour l'évaluation de l'origine des différentes populations de cristaux présentes dans le système magmatique. Les cristaux déformés peuvent être de composition chimique similaire à celle des cristaux non déformés, mais traduisent une histoire de solidification plus complexe (Albarede et al., 1997; Bureau et al., 1998; Garcia, 1996). Dans la plupart des études antérieures, l'observation au microscope optique était le seul moyen de distinguer les cristaux déformés des cristaux non déformés.

En contexte volcanique, il n'y a pas de lien direct, voire aucun lien du tout, avec les thématiques de fluage du manteau. Les approches, techniques et interprétations classiques appliquées aux problèmes mantelliques ne sont pas envisageables ici. De plus, très peu de données existent concernant la déformation magmatique de l'olivine ; ce manque doit être comblé. La quantification de la déformation de l'olivine par micro-diffraction des rayons X peut aider à la reconnaissance de l'origine des olivines déformées en contexte volcanique. Idéalement, cette technique pourrait permettre de contraindre la pression (donc la profondeur), la température et le type de déformation (i.e. type de mécanisme de dislocation par exemple) de

l'olivine. L'étude de la déformation de l'olivine apporte également un moyen supplémentaire de discriminer les différentes populations d'olivine, tout comme la taille et la forme (i.e. idiomorphe vs. xénomorphe), et se mesure sur toutes tailles de grains, y compris les petits grains pour lesquels aucune évidence de déformation ne peut être reconnue optiquement. L'analyse *in situ* par micro-diffraction des rayons X est une technique récente, et peu de données quantitatives existent, limitant la mise en comparaison de nos données. Les études précédentes utilisant cette même technique ont examiné le quartz (Flemming, 2007), le zircon (Moser et al., 2009), l'orthopyroxène (Izawa et al., 2009, *in review*), le diamant (Smith et al., 2010), et l'olivine de météorites (McCausland et al., 2010); aucune étude n'a examiné l'olivine terrestre. Par ailleurs, la caractérisation *in situ* basée sur la diffraction a été utilisée pour l'étude d'autres types de roches ou minéraux : des roches métamorphiques de ultra haute pression et les minéraux de coésite et quartz associés, par exemple (Ikuta et al., 2007). D'autres techniques existent également pour l'étude de la microstructure cristalline et caractérisation de la déformation, telle que l'approche par diffraction Laue qui utilise le synchrotron comme source de rayons X (Ice et al., 2006; Ice et al., 2005).

Tout ceci a directement motivé l'intégration de la déformation des olivines dans les laves du Kilauea en tant que Chapitre 3 de ce travail de doctorat.

1.4 Synthèse

L'étude et la quantification de la solidification des magmas basaltiques sont cruciales pour une meilleure compréhension de notre planète en général, et du fonctionnement des systèmes

magmatiques basaltiques en particulier. La multitude et la diversité des travaux ne permettent pas vraiment de trouver un consensus qui rendrait compte à la fois des données expérimentales et de modélisation numérique, mais surtout des données correspondant aux systèmes basaltiques naturels. Le problème majeur de ce type d'études sur la solidification magmatique réside bien dans la représentativité et la cohérence des résultats appliqués aux milieux naturels.

Peu d'études proposent une réelle quantification, en fonction du type d'environnement magmatique, de la plupart des processus physiques mis en jeu. On peut donc se demander si les modèles proposés jusqu'à présent s'appliquent tous en domaine volcanique, et s'ils sont suffisamment généraux pour être étendus aux différents environnements de solidification des magmas basaltiques.

L'olivine est certes un minéral très commun et généralement abondant dans les basaltes, mais il se retrouve ici aussi sous la forme de cristaux déformés, d'origine magmatique. Ce point constitue clairement un autre aspect crucial de l'étude, la présence d'olivines déformées étant généralement une particularité des roches mantelliques et plutoniques, mais beaucoup plus rare en contexte volcanique.

2 Objectifs

L'objectif principal de ce travail de doctorat est de déterminer et quantifier par une approche double alliant texture et géochimie, les principaux processus de solidification à l'oeuvre en contexte volcanique, i.e. le volcan Kilauea, Hawaïi. L'analyse structurale des cristaux d'olivine déformés s'insère dans l'optique générale d'une meilleure compréhension de la déformation et du système magmatique hôte. Cette approche double est appliquée ici à la fois en système ouvert et

en système fermé, dans un but comparatif. Différents environnements de solidification sont représentés par les échantillons suivants : (i) en système ouvert, des échantillons de lave provenant de différentes coulées issues de l'éruption de 1969-1974 du Mauna Ulu ; et (ii) en système fermé, des échantillons de lave provenant du lac de lave du Kilauea Iki, né de l'éruption sommitale de 1959.

Le corollaire de cet objectif est d'analyser en milieu naturel les différents modèles de solidification magmatique présentés dans la problématique. Il s'agira ensuite d'élaborer de nouveaux modèles pour les magmas basaltiques, directement appliqués au milieu naturel : un premier modèle appliqué à des processus crustaux (0-12 km) qui ont lieu en système ouvert dans tout le système magmatique sous-édifice ; et un second modèle appliqué à des processus de surface qui ont lieu en système fermé à l'intérieur du lac de lave.

Plus spécifiquement, il s'agit de reconnaître l'influence des processus cités ci-dessous lors de la solidification magmatique, en analysant leur(s) effet(s) sur la texture et la chimie de l'olivine, et éventuellement la chimie en roche totale. L'analyse proposée prend la forme d'une comparaison entre modèles théoriques et réponses réelles en milieux naturels. Ce mode comparatif est particulièrement utile pour évaluer l'influence des différents processus sur la texture. Les principaux processus de solidification étudiés ici sont :

- La nucléation et croissance cristallines ;
- L'agrégation de cristaux ;
- Le mûrissement textural ;
- La séparation mécanique entre liquide silicaté et cristaux (e.g. la sédimentation gravitaire des cristaux) ;
- Le mélange de cristaux ou magmas ;

- La convection/advection ;
- La déformation des cristaux.

Un autre objectif est de proposer un modèle de transfert des magmas à travers la tuyauterie du Kilauea et à l'origine de l'éruption de 1969-1974 du Mauna Ulu. Ce modèle est censé s'appliquer sur toute l'épaisseur de l'édifice (i.e. 0-12 km de profondeur), et concerne uniquement le centre et toute la partie Est du Kilauea (*East Rift Zone*). Il permettra de mieux comprendre comment fonctionne le système magmatique du Kilauea lors d'éruptions latérales volumineuses.

Finalement, un dernier objectif de ce travail de doctorat est d'évaluer plus en détail le processus de déformation magmatique des cristaux d'olivine et son influence sur la structure interne du cristal, par une analyse structurale *in situ* via la micro-diffraction des rayons X. Il s'agit de :

- (i) Quantifier la déformation de l'olivine en contexte volcanique par l'utilisation du paramètre FWHM (*Full Width at Half Maximum*), ce qui revient à mesurer la déformation dans des cristaux d'olivine orientés aléatoirement en tant qu'indicateur de la contrainte moyenne (ou maximale) ;
- (ii) Valider l'approche analytique par micro-diffraction des rayons X, et donc ;
- (iii) Reconnaître clairement la présence de déformation dans les cristaux d'olivine, même petits ;
- (iv) Quantifier la réponse cristalline à la déformation pour différents plans cristallographiques ;
- (v) Reconstruire les conditions de déformation initialement présente avant la formation de sous-grains.

Cette approche est supposée être facile à mettre à œuvre sur une large gamme de roches comportant de l'olivine. A termes, la communauté scientifique pourra disposer de valeurs références comme les données FWHM recueillies sur des olivines dont l'origine est bien connue.

Une méthodologie complète est proposée dans la section suivante afin de répondre à ces différents objectifs.

3 Méthodologie

Ce travail de doctorat est basé sur l'étude de systèmes naturels basaltiques. Des carottes de forage du lac de lave du Kilauea Iki, Hawaii, et des échantillons de laves collectés sur le terrain, constituent les matériaux de base.

La variété et la grande quantité des laves émises par le Kilauea font de ce volcan actif un site d'étude privilégié. De plus, son activité historique, puisque très bien documentée, fournit une excellente opportunité de contraindre les processus de solidification mis en jeu au niveau du réservoir magmatique sommital, des lacs de lave, et de certaines coulées de lave. Le Kilauea bénéficie également d'un archivage important et facile d'accès de carottes de lave provenant des différents forages de certains lacs de lave en cours de refroidissement, tel que le Kilauea Iki.

L'étude des coulées de lave comme celles produites par l'éruption de 1969-1974 du Mauna Ulu (Fig. 9a-b), permet de mieux comprendre les processus actifs de solidification dans tout le système magmatique superficiel (la "tuyauterie") de l'édifice. Pour simplifier, nous pouvons affirmer que la plupart des processus de solidification pré-éruption ont lieu au niveau du réservoir magmatique sommital. Le lien entre échantillons de lave et solidification dans cette tuyauterie

s'applique principalement dans le cas d'échantillons à matrice vitreuse, témoignant d'un refroidissement très rapide (trempe). L'information texturale acquise avant l'éruption est donc préservée. Les échantillons de lave à matrice microlitique, témoignant d'un refroidissement un peu plus lent, sont aussi considérés en première approximation comme représentatifs des processus pré-éruption.

Il est aussi question ici de mieux comprendre la solidification en système fermé en subsurface à travers l'étude des lacs de lave du Kilauea, tel que le Kilauea Iki (Fig. 9c). Ces lacs de lave sont formés par le remplissage de *pit craters* préexistants lors d'éruptions sommitales ou latérales (Wright and Okamura, 1977). Ils peuvent constituer de bons analogues des chambres magmatiques basaltiques quant à leur mode de solidification, particulièrement pour les premiers stades du processus. Le Kilauea Iki est privilégié dans ce travail puisqu'il est très étudié depuis plusieurs décennies, et qu'il est facilement accessible aujourd'hui. Une brève étude de la distribution de la taille des cristaux (CSD) sur les téphras de cette éruption de 1959 a été publiée (Mangan, 1990), mais le lac de lave lui-même n'a pas été étudié en utilisant cette approche texturale quantitative.

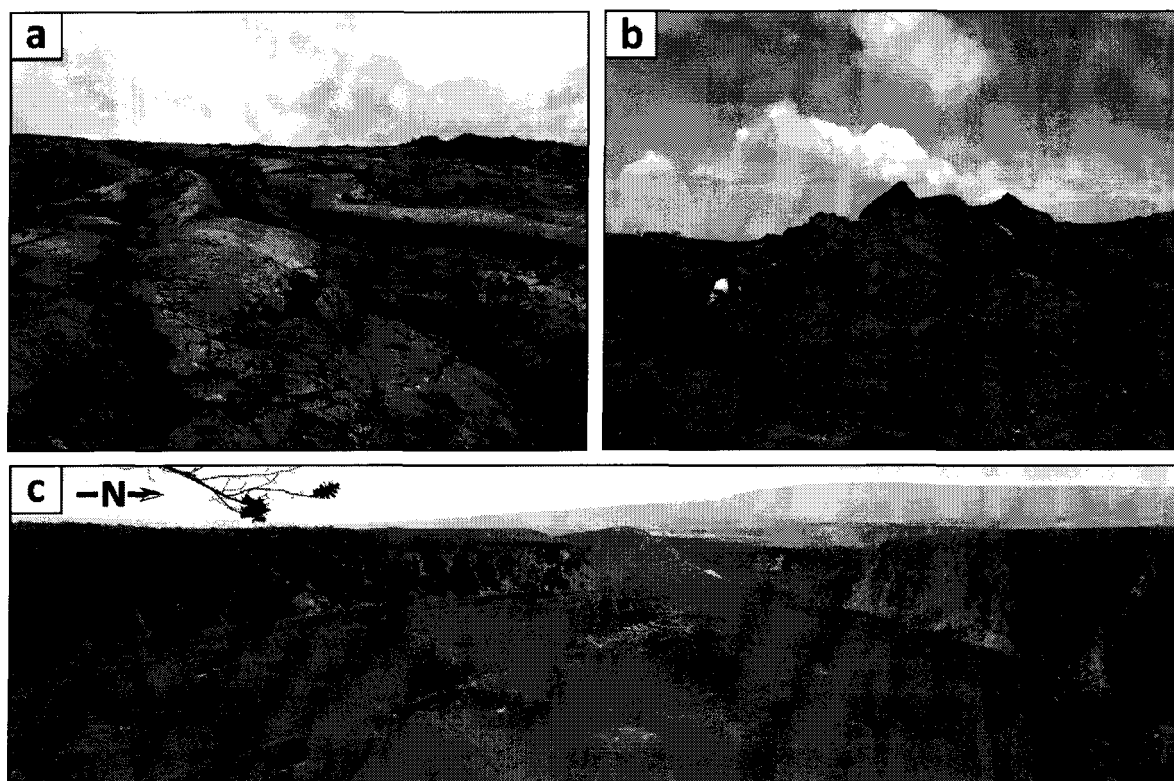


Figure 9 : Photographies des zones étudiées : Mauna Ulu et Kilauea Iki, Hawaii.

(a) Champ de lave sur les pentes du Mauna Ulu, mettant en évidence un chenal de lave solidifié ; (b) Échantillonnage d'un bloc de lave au niveau d'un hornito situé dans le champ de lave du Mauna Ulu, proche du cratère Makaopuhi ; (c) Vue d'ensemble du cratère du Kilauea Iki et son lac de lave solidifié, avec en arrière-plan la silhouette du Mauna Loa. Photos prises en Juin 2006.

L'aspect innovateur de ce travail réside dans l'étude quantitative combinée des textures de laves naturelles et des compositions chimiques des laves et cristaux d'olivine. L'analyse de la distribution de la taille des cristaux (*crystal size distribution* : CSD) est la technique principale de cette analyse texturale quantitative. Elle a été uniquement menée sur les cristaux d'olivine, puisque cette phase minérale est la plus abondante dans nos échantillons, voire la seule phase présente en phénocristaux.

Mesures texturales quantitatives

Les données texturales fournissent une information unique sur les conditions de cristallisation (Cashman et al., 1999), et par extension de solidification, d'un système magmatique. Cette approche est d'autant plus intéressante que les mesures texturales s'appliquent particulièrement bien aux systèmes naturels. L'approche texturale, complémentaire de l'approche géochimique classique, permet de mettre en évidence des processus magmatiques non observables ni quantifiables autrement.

Nombre de processus mis en jeu lors de la solidification des magmas basaltiques peuvent produire des CSDs différentes, ou modifier les CSDs préexistantes. Les CSDs théoriques attendues diffèrent selon le type de processus considéré. Une revue de ces différents profils CSD ainsi que le détail des méthodes analytiques sont présentés dans les Chapitres 1 et 2.

Imagerie électronique des olivines

L'imagerie électronique constitue une méthode complémentaire pour visualiser les textures, et donc d'en retirer une information plus complète. Cette approche qualitative permet

principalement d'identifier les cristaux d'olivine ayant subi un ou des épisodes de rééquilibrage (recristallisation) et/ou déstabilisation (résorption/dissolution) lors de la solidification magmatique, et de déterminer s'il existe une ou plusieurs populations de cristaux. Les éventuelles zonalités à l'intérieur des cristaux pourront ainsi être caractérisées en détail. En pratique, l'imagerie des olivines se fait par microsonde électronique en mode "électrons rétro-diffusés" (Fig. 10).

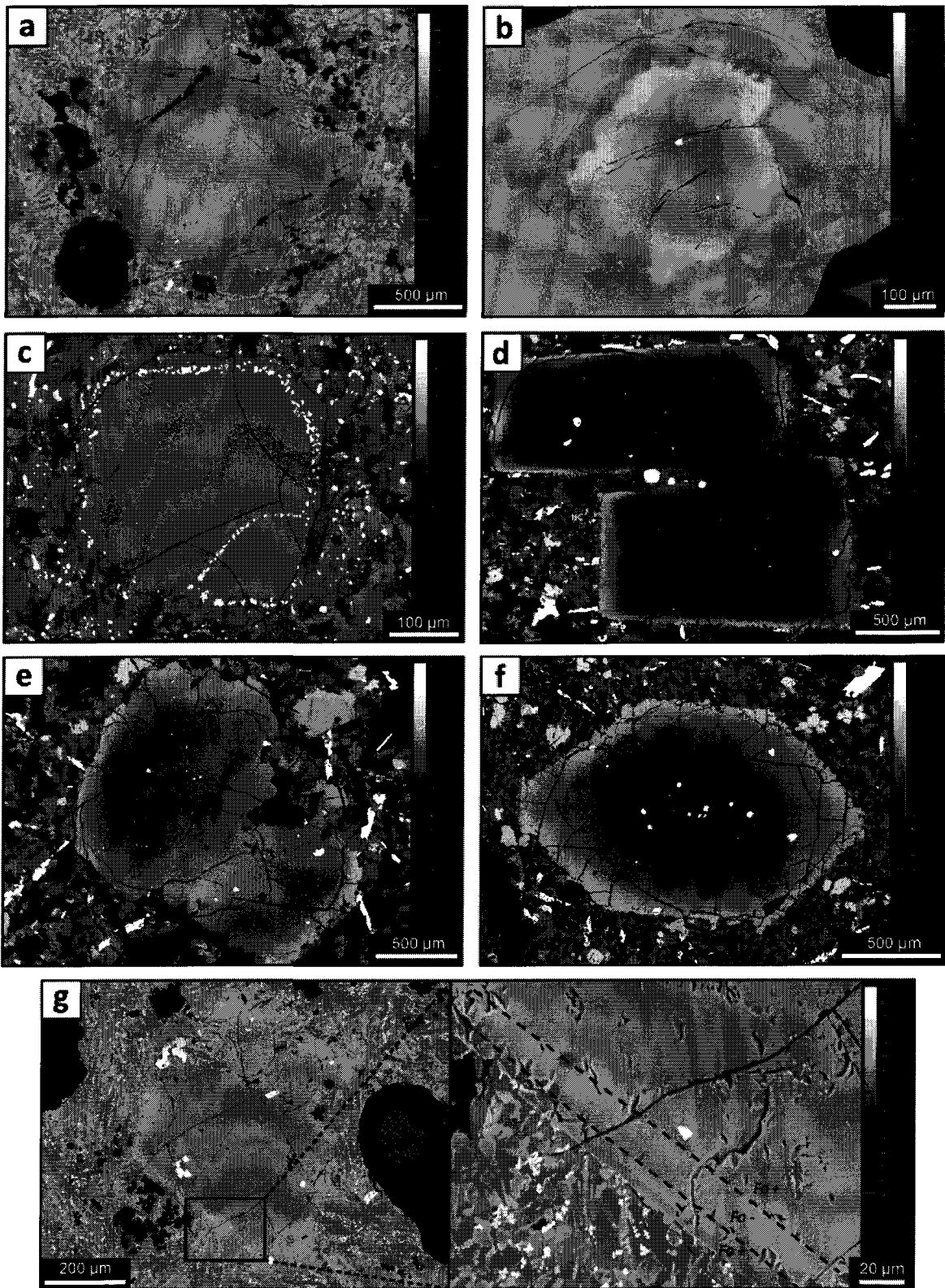


Figure 10 : Exemples généraux illustrant les différents types de zonalité chimique de l'olivine et autres observations rencontrés dans ce travail.

Photographies prises à la microsonde en mode électrons rétrodiffusés (15 kV ; 20 nA). (a) Zonalité inverse ; MU69-1a, grain 9. (b) Zonalité complexe ; MU74-2a, grain V. (c) Bordure de chromites autour de l'olivine et mise en évidence d'un sous-grain ; KI67-1-60, grain m. (d) Zonalité (normale) vs agrégation ; KI75-1-11, grain N. (e) Zonalité normale vs sous-grains ; KI75-1-27, grain F. (f) Zonalité normale ; KI75-3-35, grain D. (g) Zonalité complexe : à gauche, le grain entier ; à droite, une vue détaillée de la bordure (Fo+, riche en forstérite ; Fo-, pauvre en forstérite) ; MU74-5a, grain 11.

Chimie des olivines et chimie en roche totale

Tout comme l'imagerie électronique, les analyses chimiques *in situ* des olivines par microsonde électronique (éléments majeurs et mineurs ; Annexes 1 et 2), ainsi que les analyses en roche totale (éléments majeurs, traces et Terres Rares ; Annexe 5), permettent de faciliter l'interprétation des textures. La chimie des olivines permet également de confirmer l'existence d'une ou plusieurs populations cristallines et minérales, et de quantifier la proportion de chaque population par balance de masse. Les analyses chimiques permettent aussi de caractériser les cristaux qui présentent des évidences chimiques de dissolution ou de recristallisation. De plus, l'approche géochimique est le seul moyen, ou un moyen complémentaire, de distinguer et quantifier certains mécanismes de solidification, tel que le mélange de magmas, pour ne citer que lui. À nouveau ici, le lecteur est invité à se référer aux Chapitres 1 et 2 pour de plus amples détails.

Certaines chromites et la matrice vitreuse de quelques échantillons ont également été analysées *in situ* par microsonde électronique (Annexes 3 et 4), mais ces résultats n'ont pas été utilisés dans les articles qui constituent cette thèse.

Micro-diffraction des rayons X (μ XRD) sur olivines

L'analyse structurale de l'olivine par μ XRD est une technique récemment mise au point par Dr. Roberta Flemming, *The University of Western Ontario*, London, Ontario (Canada). Cette technique permet, entre autres, la caractérisation *in situ* et rapide de la déformation au sein du minéral. Il est ainsi possible de vérifier les observations pétrographiques de déformation faites au microscope, et de quantifier l'intensité de la déformation par intégration du signal μ XRD. La description de la méthode est fournie au Chapitre 3.

4 Format de la thèse

Cette thèse est de type "recueil d'articles scientifiques" qui sont au moment du dépôt final, soit publiés, soit acceptés, soit soumis aux fins d'évaluation par un comité de lecture dans un journal international.

Le premier article a été publié dans la revue *Journal of Petrology* en Juin 2010 sous le titre : "*Magma solidification processes beneath Kilauea Volcano, Hawaii: A quantitative textural and geochemical study of the 1969-1974 Mauna Ulu lavas*". Cet article constitue le Chapitre 1 de cette thèse. Dans ce manuscrit, les principaux processus de solidification actifs à des profondeurs de 0 à 12 km à l'intérieur du volcan Kilauea, sont examinés et quantifiés via une approche double, géochimique et texturale, appliquée aux laves de l'éruption de 1969-1974 du Mauna Ulu (*East Rift Zone*). L'article met en évidence l'existence de deux populations différentes d'olivines magmatiques, cogénétiques par ailleurs. Aucune olivine mantellique n'est reconnue ici, malgré la présence de nombreux cristaux déformés. Les résultats montrent que les magmas à l'origine de l'éruption du Mauna Ulu ont suivi deux trajets différents : (1) transit du magma transportant la plupart des olivines déformées le long du décollement majeur sous l'édifice, puis remontée à travers des conduits verticaux jusqu'à la région sous Mauna Ulu ; et (2) remontée du magma transportant les olivines non déformées par le conduit magmatique principal jusqu'au réservoir sommital, puis transfert sous la *East Rift Zone* et mélange des magmas dans des réservoirs secondaires.

Le second article a été accepté et est en cours de révision pour la revue *Journal of Volcanology and Geothermal Research*, sous le titre : "*What can crystal size distributions and*

olivine compositions tell us about magma solidification processes inside Kilauea Iki lava lake, Hawaii?". Cet article constitue le Chapitre 2 de cette thèse. Dans ce manuscrit, l'approche combinée de la chimie *in situ* et des CSDs de l'olivine est appliquée à la fois à des échantillons provenant de la surface du lac de lave du Kilauea Iki ainsi qu'à une large gamme d'échantillons provenant de forages dans le lac, à différentes profondeurs (0-90 m). Trois populations d'olivines sont mises en évidence à partir de leur teneur en forstérite. Au moins deux populations sont retrouvées par l'analyse texturale quantitative. Nos résultats suggèrent l'influence majeure du mélange de magmas ou cristaux et du mûrissement textural lors du refroidissement du lac. La recroissance a aussi joué un rôle important. Au contraire, la modélisation CSD n'a fourni aucune évidence de sédimentation gravitaire des cristaux ni de convection. Ce travail met aussi l'accent sur l'existence dans le lac de lave d'une stratification verticale minéralogique et chimique, dont l'origine pourrait être attribuée au remplissage basal du lac en cours d'éruption.

Le troisième article a été accepté et re-soumis après révisions en Octobre 2010 dans la revue *American Mineralogist*, sous le titre initial : "*Crystal structure, mosaicity and strain analysis of Hawaiian olivines using in situ micro X-ray diffraction (μ XRD)*". Cet article constitue le Chapitre 3 de la thèse. Il présente l'analyse microstructurale *in situ* par μ XRD d'olivines déformées et non déformées provenant des laves du Mauna Ulu et Kilauea Iki. Cette courte étude utilise une technique innovante, non destructive, peu coûteuse et rapide à mettre en œuvre permettant de recueillir des informations sur la structure interne des cristaux, ainsi que le mode et l'intensité de déformation. Les résultats ont permis de valider les observations pétrographiques faites à l'aide du microscope, au sujet des évidences de déformation. Cette analyse μ XRD a aussi permis de confirmer la présence d'olivines déformées pour toutes les tailles de grains, sans corrélation simple avec leur chimie. Enfin, il a été possible de fixer un seuil quantitatif au-delà duquel toute

olivine est déformée de façon significative. J'ai mené ce travail au laboratoire de Microanalyse par Rayons X de *The University of Western Ontario*, London, Ontario (Canada), avec la collaboration de Dr. Roberta Flemming. J'ai recueilli tous les résultats alors que Dr. Flemming m'a aidé à correctement utiliser l'appareillage et les divers logiciels de traitement de données ; elle a par ailleurs rédigé la partie « *Methods* » de l'article.

Les Chapitres 1 et 2 de cette thèse ont également fait l'objet de conférences présentées lors des congrès internationaux et locaux suivants :

- Vitrine étudiante des Sciences de la Terre, UQAC, Chicoutimi, QC, Canada, Mars 2010 ;
- AGU Fall Meeting, San Francisco, CA, USA, Décembre 2009 ;
- IAVCEI General Assembly, Reykjavik, Islande, Août 2008 ;
- Frontiers in Mineral Sciences, Cambridge, UK, Juin 2007 ;
- Vitrine étudiante des Sciences de la Terre, UQAC, Avril 2007.

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CHAPITRE 1

MAGMA SOLIDIFICATION PROCESSES BENEATH KILAUEA VOLCANO, HAWAII: A QUANTITATIVE TEXTURAL AND GEOCHEMICAL STUDY OF THE 1969-1974 MAUNA ULU LAVAS

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1.1 ABSTRACT

Kilauea Volcano is a very intensively studied, active basaltic magmatic system and thus, represents an ideal location to study magma solidification processes in a natural environment. Understanding solidification is important in refining models of magma chamber dynamics, and its detailed study can improve our knowledge of magma system evolution. In this study, magma solidification processes are examined and quantified using samples from the 1969-1974 Mauna Ulu (MU) rift eruption. We have collected major and trace element whole-rock data plus *in situ* olivine compositions, along with crystal size distribution data on eleven lava samples. The observed whole-rock chemical variation was partly produced by olivine addition within the Kilauea edifice. At least two distinct olivine populations are inferred from quantitative textural analysis: (1) a 3-40 year old population characterized by a low crystal density, greater crystal length and flatter slopes of the crystal size distributions (CSDs); and (2) a 1.5-15 year old population marked by a high density of smaller crystals and steep CSD slopes. The range in olivine composition suggests that all these crystals grew from a range of different magmas, probably closely related by crystal fractionation. The ubiquitous presence of deformed olivine crystals shows that population 1 reflects a component that must have mostly originated by disruption of a deformed cumulate. This antecrystic olivine population represents an earlier-coarsened and aggregated, cumulate-forming magma component. In contrast, the phenocrystic population 2 represents a late magma component formed in the summit magma storage region. Our results are consistent with the hypothesis that the components of the MU magmas followed two different routes. The deformed-olivine bearing magma moved along the deep basal decollement then rose through vertical pipe-like conduits under MU rift. The undeformed-olivine-bearing magma rose via the main conduit to

the summit reservoir and then moved out along the rift zone, where the magmas mixed in small chambers. The presence of narrow, reversely-zoned rims suggests that the mixing occurred just prior to eruption.

KEY WORDS: *Kilauea (Hawaii); Mauna Ulu; crystal size distributions; olivine cumulates; solidification processes*

1.2 INTRODUCTION

A fundamental issue in igneous petrology is the understanding of magma solidification. Solidification is the result of a complex interplay between crystallisation and solution. Magma in a chamber can be reheated by the addition of a new hotter magma or by convective self-mixing. The partial reduction of pressure during magma ascent can result in an increase or a decrease in the rate of undercooling. Since direct examination of active basaltic magma chambers is impossible, we use samples from lava lakes or other volcanic products to study solidification processes and time scales. These volcanic products are commonly studied by quantitative chemical and isotopic analysis and by qualitative petrological observations of texture. However, it is also possible to approach the problem by quantitative analysis of rock textures. One of the most commonly employed techniques is the study of crystal size distribution (CSD).

CSD measurements can provide a complementary approach to chemical and experimental methods, of looking at solidification processes in magma chambers, lava flows and lava lakes (Cashman & Marsh, 1988; Marsh, 1988, 1998). Furthermore, the CSD approach offers the possibility of investigating solidification in natural dynamic systems and providing clues to the physical and chemical processes that have taken place. The CSD method yields valuable kinetic

information for examining the link between crystal nucleation and growth, and the overall cooling and magma dynamics (Marsh, 2007).

Kilauea Volcano is one of the most intensively studied and best understood magmatic systems on Earth (e.g. Tilling & Dvorak, 1993; Wright & Klein, 2008), due to the continuous monitoring of its activity by the Hawaiian Volcano Observatory since 1912. In addition, numerous petrographic studies of the lavas from Kilauea have documented the complexity of the magmatic processes (Garcia *et al.*, 2003; Garcia *et al.*, 1996; Helz, 1980, 1987a, b; Helz, 2009; Hofmann *et al.*, 1984; Richter & Murata, 1966; Scowen *et al.*, 1991; Thornber, 2001). Kilauea is composed of tholeiitic basalt with minor olivine-rich tholeiite. Olivine is the dominant – and generally the only – phenocrystic mineral, associated with accessory Cr-spinel microphenocrysts (Macdonald, 1949; Powers, 1955; Wright, 1971).

Kilauea's historical activity from ~1790 to present includes decades of quiescent lava effusion, interrupted by major collapses ($\geq 100\text{m}$) of the caldera floor, rare violent explosions, and brief to decade-long rift zone eruptions (Garcia *et al.*, 2008). From 1952 to 1982, there were mostly short-lived fissure eruptions in and near Halema'uma'u Crater and frequently in the East and Southwest Rift zones (Garcia *et al.*, 2008) (Fig. 1-1a). No summit eruptions have taken place since the start of the current Pu'u 'O'o eruption of 1983, East Rift Zone (e.g. Garcia *et al.*, 2000), until the opening of a new vent in Halema'uma'u Crater on March 19, 2008.

Mauna Ulu (*Growing Mountain* in Hawaiian) is located on the upper East Rift of Kilauea Volcano (Fig. 1-1) about 8-10 km from the rim of the summit caldera (Hofmann *et al.*, 1984). This spatter-and-lava basaltic shield grew during a nearly continuous eruption lasting from May 24, 1969 to July 22, 1974, with a pause between mid-October 1971 and early-February 1972 (Peterson *et al.*, 1976; Swanson, 1973; Swanson *et al.*, 1979; Swanson *et al.*, 1971; Tilling *et al.*, 1987; Wright

& Klein, 2008; Wright *et al.*, 1975). The eruptive activity is divided into two main parts: the 1969-1971 and the 1972-1974 periods. The Mauna Ulu eruption began with high fountains, like the ongoing Pu'u 'O'o eruption, and evolved into a passive, effusive eruption with tube-fed pahoehoe first entering the ocean in September 1970 (Peterson *et al.*, 1976; Swanson, 1973; Swanson *et al.*, 1979). The first half of the eruption produced an estimated 185 million cubic meters of lava, and the second half an additional 160 million cubic meters. The lavas erupted between 1969 and 1974 from Mauna Ulu and closely associated vents covered 45.6 km² and added almost about 1 km² of new land to Hawaii. Before the still on-going 1983 Pu'u 'O'o eruption, the 1969-1974 MU eruption was the longest, the most voluminous and most varied flank eruption in the recorded history of Kilauea. This was also the first eruption in the last 150 years that lasted longer than a few weeks (Swanson *et al.*, 1979).

The magmatic plumbing system of Kilauea Volcano has been the subject of vigorous debate (see summaries by Pietruszka & Garcia, 1999; Wright & Klein, 2008). Eaton & Murata (1960) proposed that mantle-generated melt migrates through discontinuous vertical conduits from a depth of 60-170 km to a shallow, 2-4 km deep, magma reservoir beneath the summit caldera prior to eruption. Eruptions are either at the volcano's summit by vertical propagation of dikes, or at the rift zones by lateral intrusion. Magma may be further stored in the lateral conduits prior to eruption in the rift zones. This model has been refined over the last four decades by geophysical studies, but is still subject to controversy (e.g. Decker, 1987; Delaney *et al.*, 1990; Tilling & Dvorak, 1993; Wright & Klein, 2006, 2008). Combined geophysical and petrologic data for the 1994-1999 eruptive interval support the model of storage and mixture of discrete magma bodies along the active east rift conduit (Thornber *et al.*, 2003). Mostly based on summit deformation

measurements for the Puu Oo-Kupaianaha eruption, several authors found evidence of magmatic continuity between the summit reservoir and rift vents (e.g. Cervelli *et al.*, 2002; Thornber, 2001).

The geometry of the shallow magma reservoir that lies under the summit region is of great importance. Two distinct models have been proposed based on geodetic data: a single, 'spherical' magma body (Yang *et al.*, 1992) versus a plexus of dikes and sills (Fiske & Kinoshita, 1969). Measurements of ground deformation associated with the volcano's frequent activity have been interpreted as resulting from the migration of the apparent locus of deformation over a large area up to 16 km² (see summary by Pietruszka & Garcia, 1999). The classic interpretation of Fiske & Kinoshita (1969) is related to a network of interconnected dikes and sills. However, some bias on the deformation center due to dike intrusion during periods of summit inflation, may lead to an apparent migration of the pressure center (Yang *et al.*, 1992). Removal of this effect results in the convergence of the inflation center, and thus seems consistent with the single, 'spherical' body model. Using lava chemistry, Pietruszka & Garcia (1999) also argued that the shape of the volcano's summit chamber is relatively simple, consistent with a single, 'spherical' body. Further, they refined the wide range of reported reservoir size estimates of 0.08 – 40 km³ to ~2-3 km³. In another geochemical study, Garcia *et al.* (2003) also concluded that the Kilauea summit magma reservoir is a well-mixed single body. Similarly, Thornber (2003) argued in favour of continuous input and turbulent mixing of primitive magma into the summit reservoir, based on chemical data. As a consequence of the continuous replenishment of the summit reservoir, the magma, mostly olivine-saturated, moves through the shallow rift zone, where it may interact prior to the eruption with cooler magma persisting in open- or closed-system rift reservoirs (Owen *et al.*, 2000; Thornber, 2001; Thornber *et al.*, 2003). The simple reservoir model was also favoured by Peltier *et*

al. (2009) in their combined geophysical and geochemical study of Piton de la Fournaise Volcano, La Réunion Island, which is very similar to Kilauea.

Although the lavas of Kilauea have been extensively studied using chemical and isotopic methods, few quantitative textural studies have been published. Cashman & Marsh (1988) worked mainly on plagioclase and oxides of tholeiitic basalt from the Makaopuhi lava lake, Kilauea. Mangan (1990) studied olivine in a few scoria samples of picritic basalt from the 1959 eruption of Kilauea Iki. These early papers were exploratory in nature and time has come for more detailed work with a greater number of samples, combining textural and geochemical approaches. Our aim is to characterize texturally the MU olivines in order to understand and quantify the solidification processes of tholeiitic magmas at depth via textures and chemical compositions.

This paper presents the results of textural and geochemical analyses from eleven lava samples and over 330 individual olivine crystals from the 1969-1974 MU eruption; all produced during a multiple, prolonged rift-zone eruption. These samples have significantly different chemical compositions and CSDs. Although olivine can re-equilibrate chemically rapidly at magmatic temperatures (e.g. Roeder & Emslie, 1970; Ulmer, 1989), analyses of olivine can provide evidence of magma mixing, crystal accumulation and other magmatic processes (e.g. Garcia, 1996). We consider that the following processes are likely to have participated in generating the petrologic diversities observed at Mauna Ulu: (1) simple crystal nucleation and growth; (2) crystal accumulation and fractionation; (3) textural coarsening; (4) crystal deformation; (5) aggregation and synneusis, and; (6) mixing of multiple parental magmas.

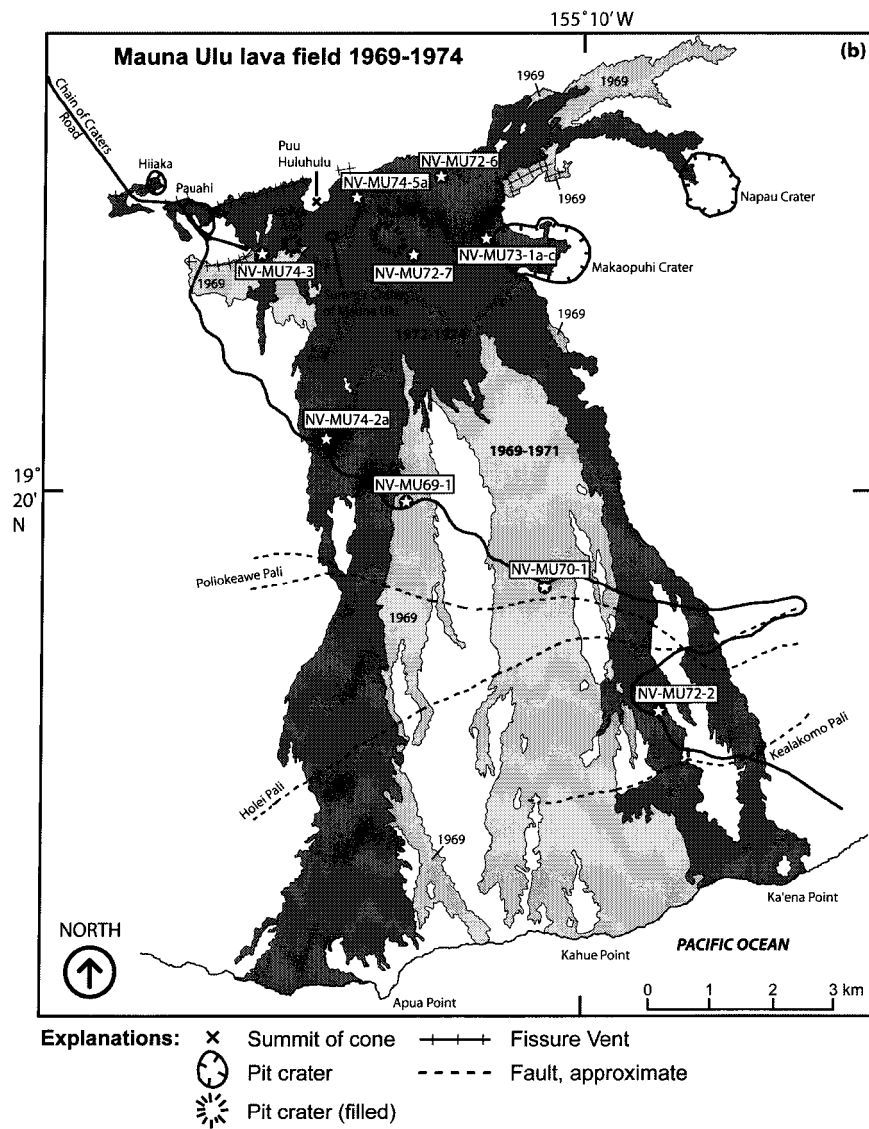
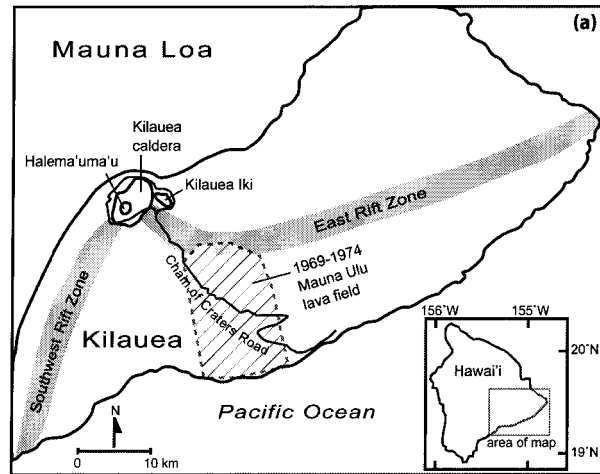


Figure 1-1. (a) Sketch map of Kilauea Volcano, showing the summit region, the two rift-zones and the Mauna Ulu lava field 1969-74. Location map of the island of Hawaii with the study area in inset; (b) Mauna Ulu lava sampling map, modified after the maps of Wolfe & Morris (1996) and Tilling *et al.* (1987). Pali means cliff in Hawaiian.

1.3 MINERALOGY AND PETROGRAPHY

1.3.1 Sampling

We sampled at least one lava flow per eruption year from the 1969-1974 MU eruption (Fig. 1-1; Table 1-1). Where possible, we collected a temporal suite from each lava flow, typically from the fissure to the end of the flow. We sampled either the flow terminations (mostly lava lobes) or the middle of the flow. This sampling method allowed us to study the chemical and textural evolution of individual eruptive episodes. Unfortunately only one sample in the middle of the flow was sampled for the 1969 and 1970 lava flows.

To minimize the effects of crystal growth during post-emplacement cooling, we collected scoria and/or spatter material from the fissure vent, lava blocks from the flow surface, or quenched blocks where possible from the flow tops (the 'glassy crust'). When neither quenched crust nor loose blocks were available, we collected lava samples from beneath the surface of the flow. In some cases, we sampled deeper 'slabs' seen in flow or tumulus cracks, or lava tree moulds that often constituted partly glassy, or finely crystallized matrix-bearing, material. The glassy material was also preferred as the subsequent creation of texture maps from the thin section images in the lab is more effective and contains less uncertainty (i.e. no microlites to increase the complexity of the image).

Table 1-1: Sample locations of Mauna Ulu lava flows

Sample	Flow type*	Eruption year	UTM zone 5Q (WGS 1984)		Latitude (N)°	Longitude (E)°	Altitude (m)
			mN	mE			
NV-MU69-1	aa	1969	2138519	0269830	19.327472°	-155.190793°	752
NV-MU70-1	aa	1970	2137341	0272323	19.317117°	-155.166936°	635
NV-MU72-2	phh tube	1972	2135063	0274074	19.296741°	-155.150011°	110
NV-MU72-6	aa	1972	2144031	0269960	19.327264°	-155.190221°	977
NV-MU72-7	phh	1972	2142573	0270118	19.364115°	-155.188541°	999
NV-MU73-1a-c	phh tube	1973	2142535	0271380	19.363916°	-155.176529°	918
NV-MU74-2a	phh tube	1974	2139594	0268533	19.337031°	-155.203261°	840
NV-MU74-3	aa	1974	2142602	0267451	19.364070°	-155.213921°	1000
NV-MU74-5a	phh (in channel)	1974	2143405	0269082	19.371510°	-155.198500°	1016

* 'a'a (aa) or pahoehoe (phh)

° Latitude and longitude are in decimal degrees

1.3.2 Macroscopic characteristics

The samples consist of moderately to strongly vesicular olivine tholeiitic lavas with no visible foliation. They contain 3-16 vol. % olivine in phenocrysts (>0.5 mm across) and microphenocrysts (>0.1 to <0.5 mm across) as the most common crystalline phase. In some samples the crystallinity can vary abruptly on the scale of centimetres, in terms of grain size, abundance of crystals or interstitial glassy matrix, or types of microlites. For example, a few thin sections have ~1 cm pockets or bands with a contrasting matrix. These also have fewer and smaller olivine phenocrysts and microphenocrysts. The olivine may also be dendritic.

The lavas can be classified into three categories based on the groundmass crystallinity. Samples NV-MU69-1, -MU72-7, -MU73-1c, and -MU74-3 have a microcrystalline groundmass of plagioclase, clinopyroxene and opaque minerals, mostly Ti-magnetite, with or without olivine. Samples NV-MU70-1, -MU72-2, -MU73-1a, -MU74-2a and -MU74-5a have a glassy groundmass. Samples NV-MU72-6 and -MU73-1b are quite difficult to classify in these two categories and were labelled as intermediate. MU73-1a-c are from the same sample site, although they show significant textural variations. No obvious control on the groundmass texture, such as top, bottom or middle of the flow, is noted.

Helz (1987b) observed and defined five classes of olivine crystals in the magmas of the 1959 Kilauea eruption, from both lava lake and eruption pumice samples:

- (1) Irregular blocky crystals with deformation features;
- (2) Equant, euhedral, skeletal or strongly elongate grains;
- (3) Round or strongly resorbed grains;
- (4) Fragmental grains ('angular or conchoidal fragments');

(5) Subhedral sulphide inclusion-bearing crystals.

To the list, two other types were added by Helz (1987b): the olivine aggregates and olivine megacrysts. She found in the lake samples only two examples of olivine megacrysts, of size larger than 20 mm by 10 mm. In our samples, no such megacrysts have been identified, but only rare megacrysts of 5-10 mm long. Those megacrysts were not sufficiently numerous to be included in the CSD analysis.

The five classes, along with the aggregate type, are also recognized in the MU lavas studied here (Fig. 1-2; class 5 not shown). Class 1 consists here of large grains, 1-10 mm long, that are present in all the samples in varying proportions ranging from ~1 to 35% of the olivine crystals present. It represents around 0.1 to 3 vol. % deformed olivine in total per sample. Deformation is observed optically as planar extinction discontinuities, undulose extinction, or subgrains in all the crystal sizes. Although deformed crystals less than 0.5 mm in size are more difficult to recognize, they were found in every sample. Edge rounding or resorption are common features of this class, as well. Class 2 olivine is the most abundant, and consists of smaller grains in average, present as both microphenocrysts and phenocrysts. Class 3 olivine is fairly rare, typically a few percent of the total amount of olivines, as all the resorbed crystals with deformation features have been placed in class 1. Classes 4 and 5 are rare and deformed grains are uncommon for both of those, even when the grains containing non-sulphide inclusion swarms are counted.

Most olivine crystals in every sample, and thus most of the Helz's classes appear to be part of larger olivine aggregates, as also observed by Helz (1987b), Schwindinger and Anderson (1989), and Schwindinger (1999), in the lavas from the 1959 Kilauea eruption. In our samples olivine aggregates are omnipresent in varying but high proportions. It is >20-50 % by volume of the total amount of olivine crystals in all our samples. The clusters typically contain 2-4 individual crystals. A

wide range of olivine aggregate styles are recognized in the MU lavas, especially in the way in which the grains are connected in each aggregate. They range from adjacent, separate grains that may be connected in the third dimension (Fig. 1-2a), to less common clumps of crystals that are completely attached. Finally some clusters are re-equilibrated with originally separate grains becoming subgrains (Fig. 1-2c). The olivines have either straight or curved edges, depending of the aggregate. No dihedral angles or triple junction angles (between only olivine grains) were measured, as only few aggregates contain three grains touching. Often multiple grains are attached in a single cluster, more or less as a small chain of crystals. When present, three touching olivine grains joined by a well-defined grain boundary have apparent triple junctions that are not 120° .

MU lavas also contain minor amounts of other minerals. Cr-spinel occurs within the olivine grains and as separate crystals, commonly adjacent to the olivine grains, as well as in the olivine aggregates. The Cr-spinel crystals are subhedral to euhedral and 10-100 μm in size. Very rare examples of clinopyroxene and plagioclase phenocrysts are present, but they are more common as microphenocrysts, 0.1 to 0.5 mm across. Some clinopyroxenes show deformation features such as undulose or sector extinction. Plagioclase, clinopyroxene and Cr-spinel were too small or not sufficiently abundant for their textures to be determined quantitatively.

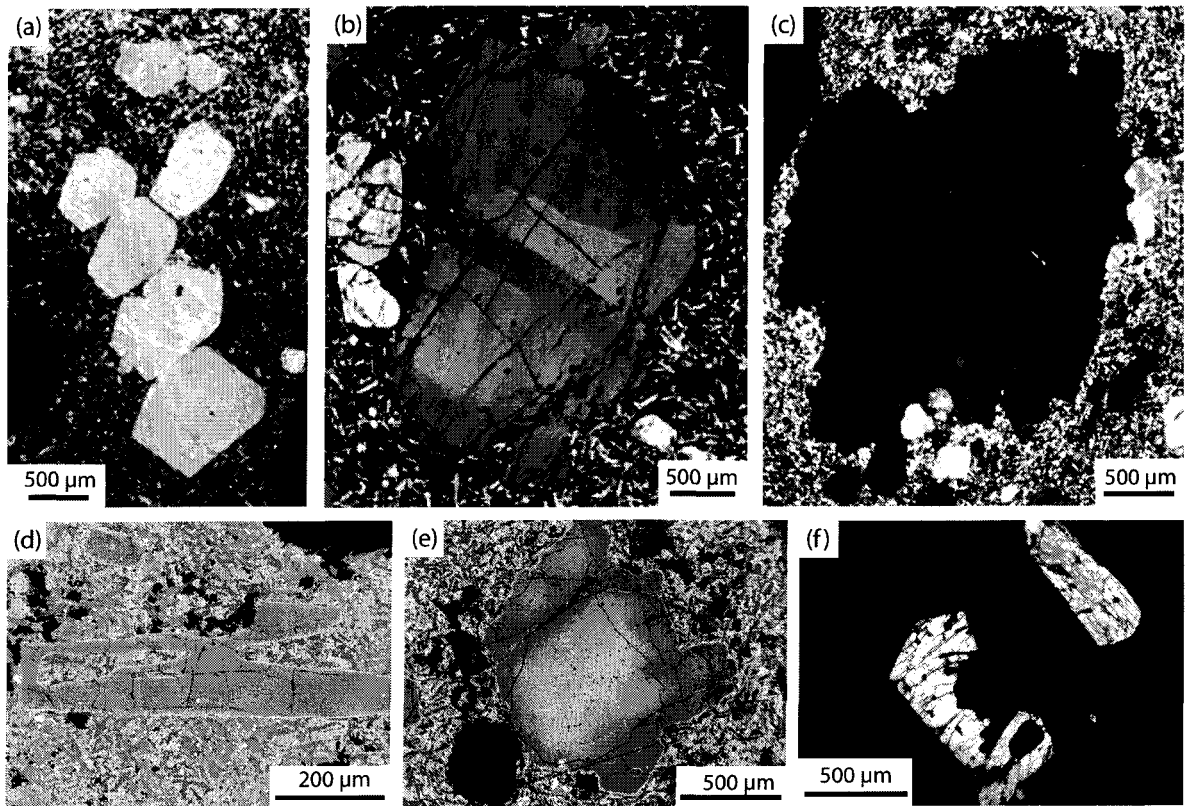


Figure 1-2. Representative photomicrographs of the different types of Mauna Ulu olivine crystals, with the class 5 olivine not shown here: (a) Aggregate of euhedral, class 2 olivine grains, NV-MU69-1; (b) Deformed, class 1 grain with kink-bands; (c) Deformed, unknown class grain with kink-bands, in aggregate of a different type than in (a), NV-MU69-1; (d) Skeletal, strongly elongate, class 2 grain, BSE image, NV-MU69-1; (e) Strongly resorbed and Fe-Mg chemically zoned, class 3 grain, BSE image, NV-MU69-1; (f) Fragmental, class 4 grains (lower left of the photo), NV-MU72-2.

1.4 CHEMICAL STUDIES

1.4.1 Analytical methods

Olivine and glass major element compositions were measured in thin sections using the five-spectrometer, Cameca SX-100 electron-microprobe at the Université Laval Microanalysis Laboratory, Québec city, Canada. Both olivine and glass analyses were done at 15 kV and 20 nA beam current, with a 5 μm spot size and using a wavelength dispersive spectra (WDS) mode. At least one spot both in the core and the rim of each olivine was analyzed to check for compositional zoning. Where the back-scattered electron (BSE) imagery showed the presence of complex zoning, more points on the rim of the selected grain were analyzed. Whole-rock major and minor element compositions were determined by X-ray fluorescence (XRF) spectrometry at the TEAL / Geochemical Laboratories of McGill University, Montréal, Canada. The precision for element analyses was better than 1% with the exception of SiO_2 , $\leq 0.5\%$.

1.4.2 Whole-rock geochemistry

1.4.2.1 Major elements

MgO contents of the MU samples range from 9.4 to 12.0% with a single sample at 16.2 wt % (Table 1-2). With the exception of Fe_2O_3 , all MU oxides display a negative linear trend when plotted against MgO, with little scatter for Na_2O , K_2O and P_2O_5 (Fig. 1-3). A positive trend with more scatter is observed for Fe_2O_3^* .

Our data is most easily understood when combined with that of earlier studies. Overall, the major element compositions and chemical behaviour of the MU lavas analysed in this study are similar to analyses of the 1969-1971 MU lavas (Wright *et al.*, 1975) and other historical Kilauea lavas (Garcia *et al.*, 2003; Garcia *et al.*, 2000). However, there is a slight offset of our data toward lower SiO₂ and higher Fe₂O₃* contents when compared with the 1969-1971 MU lavas (Wright *et al.*, 1975). This apparent discrepancy in the analysis is not an analytical artefact. Our analyses were calibrated using 39 geological Certified Reference Materials from a variety of sources (e.g., NIST, China, Japan, France). The whole-rock values presented here are thus believed reliable. Accordingly, the previous data on Mauna Ulu lavas might either reflect some uncertainty in Si and Fe, or be indicative of some unknown compositional discrepancy in the magma. The MgO composition of MU lavas is somewhat higher than other historical Kilauea lavas. MU samples are amongst the most Fe-rich of historical Kilauea lavas, for a given MgO content. In comparison with MU lavas, the Puu Oo field plots towards more evolved compositions (<10 wt% MgO) along with the related higher compositional variations in alkali and P, and in a minor extent, Ti and Ca, at a given MgO content.

In selected major-oxide variation diagrams such as MgO-FeO, CaO-Al₂O₃ and TiO₂-CaO (Fig. 1-4), an olivine control line was added, delineated by an arrow that points toward a mean composition. This olivine control line represents accumulation or subtraction of Fo₈₇ olivine. It appears that the chemical compositions of the MU and other historical Kilauea lavas fully, or at least partly, define a trend that matches this olivine control line. Hence, they are in most part olivine-controlled, especially the MU lavas. The MU lavas have constant CaO/Al₂O₃ values, around 0.83, along the range of MgO (Table 1-2), which further supports this idea of olivine control.

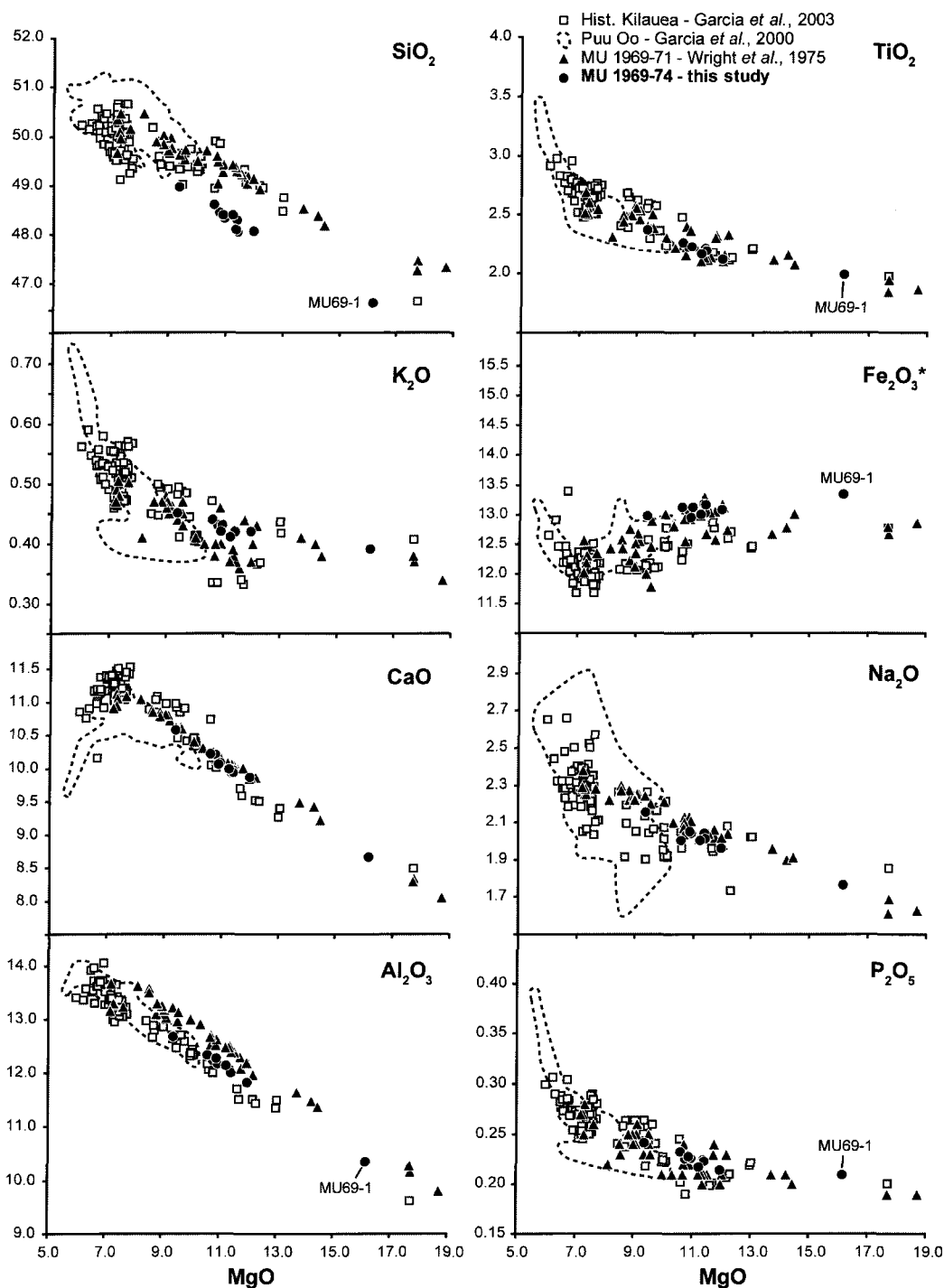


Figure 1-3. Whole-rock MgO variation diagrams for the Mauna Ulu (MU) lavas compared to other historical Kilauea lavas.

The slightly, but significant offset towards lower SiO₂ and higher Fe₂O₃* contents of the studied samples compared to other 1969-71 MU lavas (from Wright *et al.*, 1975) is not an analytical artefact of our data (see text for details). The oxide compositions are expressed in wt %. The 2σ errors are smaller than symbol size.

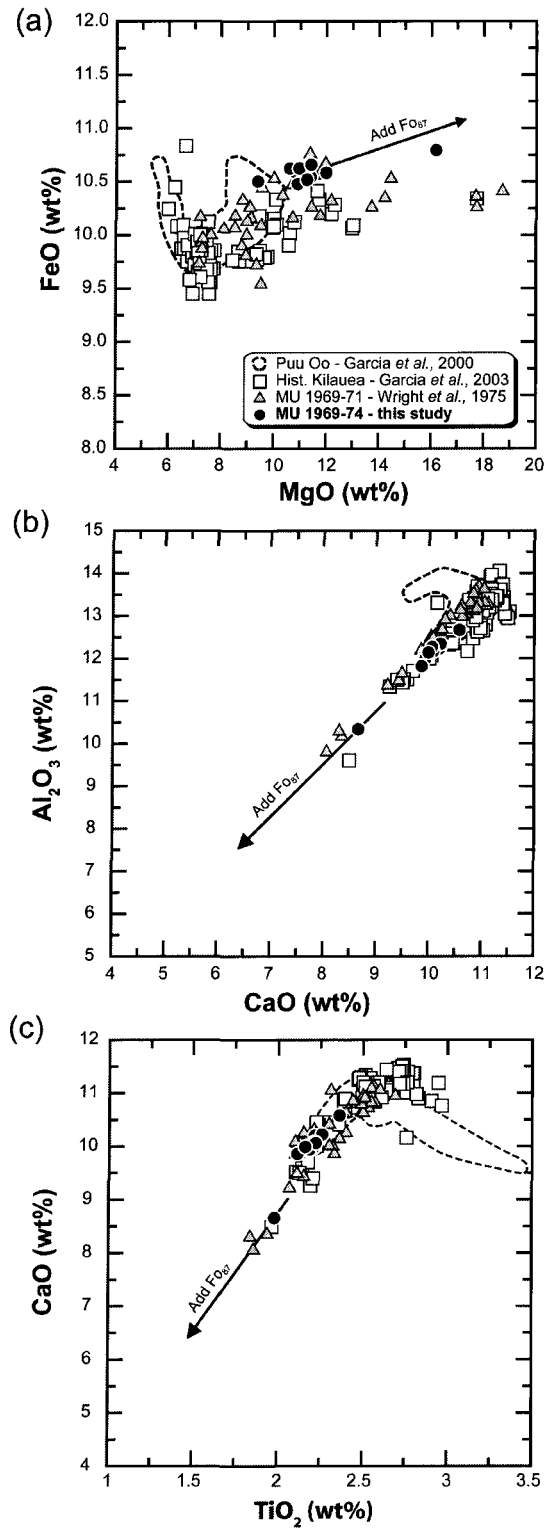


Figure 1-4. Selected major-oxide variation diagrams illustrating the compositional tendency that can result from addition of olivine Fo₈₇ to the whole-rock composition: (a) MgO vs FeO, (b) CaO vs Al₂O₃ and (c) TiO₂ vs CaO. Notice that the MU lavas, along with other Kilauea lavas, are clearly olivine-controlled.

Table 1-2: Whole-rock major-element, plus Ni and Cr compositions and olivine modal abundances of the Mauna Ulu samples

Sample	MU69-1	MU70-1	MU72-2	MU72-6	MU72-7	MU73-1a	MU73-1b	MU73-1c	MU74-2a	MU74-3	MU74-5a
<i>(XRF)</i>											
SiO ₂	46.61	48.11	48.46	48.31	48.62	48.33	48.06	48.97	48.08	48.40	48.40
TiO ₂	1.981	2.181	2.214	2.204	2.257	2.211	2.183	2.360	2.111	2.219	2.158
Al ₂ O ₃	10.34	12.06	12.23	12.05	12.34	12.17	12.01	12.67	11.82	12.27	12.14
Fe ₂ O ₃ *	13.33	13.17	13.08	13.02	13.12	13.12	13.16	12.97	13.07	12.94	12.99
MnO	0.177	0.177	0.179	0.177	0.177	0.177	0.176	0.180	0.177	0.176	0.176
MgO	16.17	11.35	10.79	11.39	10.60	10.96	11.41	9.39	11.98	10.90	11.25
CaO	8.66	9.92	10.20	9.97	10.22	10.08	9.94	10.58	9.86	10.06	9.99
Na ₂ O	1.76	2.02	2.03	2.04	2.00	2.04	2.01	2.15	1.96	2.05	2.00
K ₂ O	0.39	0.42	0.43	0.42	0.44	0.43	0.42	0.45	0.42	0.42	0.41
P ₂ O ₅	0.209	0.222	0.225	0.224	0.232	0.226	0.223	0.241	0.214	0.227	0.217
Cr	1190	740	717	744	681	706	751	571	825	714	751
Ni	695	339	297	365	289	308	331	227	388	336	342
<i>Total</i>	<i>99.63</i>	<i>99.63</i>	<i>99.84</i>	<i>99.81</i>	<i>100.01</i>	<i>99.74</i>	<i>99.59</i>	<i>99.96</i>	<i>99.69</i>	<i>99.66</i>	<i>99.73</i>
Mg#	0.728	0.655	0.645	0.658	0.640	0.648	0.656	0.614	0.669	0.650	0.656
CaO/Al ₂ O ₃	0.838	0.823	0.834	0.827	0.828	0.828	0.828	0.835	0.834	0.820	0.823
<i>CIPW Norms</i>											
Or	2.3	2.5	2.5	2.5	2.6	2.5	2.5	2.7	2.5	2.5	2.4
Ab	14.9	17.1	17.2	17.3	16.9	17.3	17.0	18.2	16.6	17.3	16.9
An	19.2	22.6	23.0	22.5	23.4	22.8	22.5	23.6	22.2	23.0	22.9
Di	18.1	20.5	21.3	20.8	21.0	21.0	20.6	22.4	20.6	20.7	20.5
Hy	11.5	16.2	17.1	16.4	18.5	16.7	16.0	18.1	16.1	17.3	17.8
Ol	27.5	14.2	12.1	13.8	10.8	12.9	14.3	8.1	15.3	12.2	12.7
Mt	1.9	1.9	1.9	1.9	1.9	1.9	1.9	1.9	1.9	1.9	1.9
Il	3.8	4.1	4.2	4.2	4.3	4.2	4.1	4.5	4.0	4.2	4.1
Ap	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.6	0.5	0.5	0.5
<i>Sum</i>	<i>99.6</i>	<i>99.6</i>	<i>99.9</i>	<i>99.8</i>	<i>100.0</i>	<i>99.8</i>	<i>99.6</i>	<i>100.0</i>	<i>99.7</i>	<i>99.7</i>	<i>99.7</i>
<i>Modal abundances of olivine (%)</i>											
Deformed	2.9	0.3	0.1	0.7	0.5	0.8	1.0	0.2	1.7	0.6	2.8
Undeformed	13.1	9.0	7.3	7.6	5.7	10.1	7.0	3.2	7.4	7.4	5.2

* Fe₂O₃ is total iron.

n.d. means 'Not Detected'.

Oxides are in wt %; Ni and Cr are in µg/g. Normative compositions are in mol. %; Mg# is atomic Mg/(Mg+Fe²⁺)

1.4.2.2 Ni and Cr content

Ni and Cr content is given in Table 1-2. The MU lavas of this study show great variation in Ni and Cr content for a small range in K like many historical Kilauea summit lavas (Fig. 1-5). However, the MU lavas have higher Ni and Cr contents, with MU69-1 displaying one of the highest Ni (695 ppm) and Cr (1190 ppm) contents published. Moreover, a direct correlation is observed between Ni, Cr and the amount of olivine in the rock (Fig. 1-6), with the exception of MU73-1a being a little off the trend. This may be related to the very compatible character of Ni in olivine and Cr in Cr-spinel. Again, it is likely that olivine and Cr-spinel are major factors of compositional control.

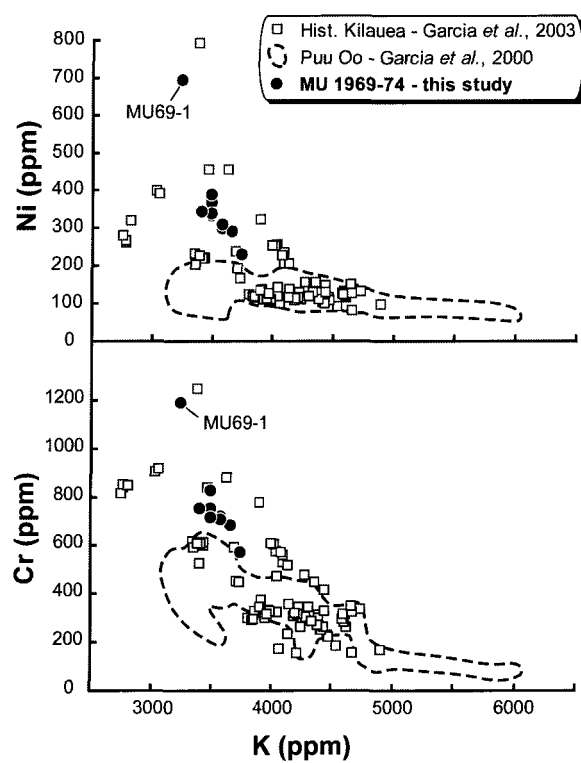


Figure 1-5. Variations of Ni and Cr compositions versus K for Mauna Ulu (MU) lavas compared to some other historical Kilauea lavas. The 2σ errors are smaller than symbol size.

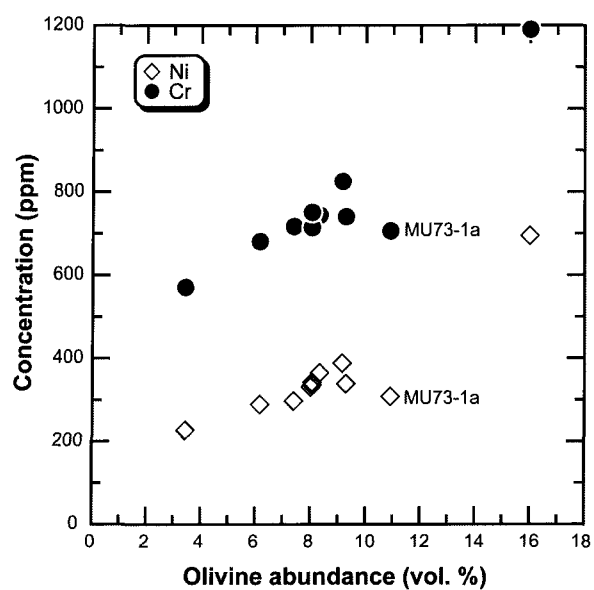


Figure 1-6. Ni and Cr whole-rock content versus olivine abundance for the Mauna Ulu lavas.

1.4.3 Olivine chemistry

Over 330 olivine phenocrysts and microphenocrysts were analysed from eleven MU samples. All analyses were spot analyses; no line traverses were done. Olivines were analysed to determine whether they were in equilibrium with their host magma and whether there was compositional zoning. Although olivine can re-equilibrate rapidly at magmatic temperatures (e.g. Roeder & Emslie, 1970; Ulmer, 1989), analyses of olivine can provide evidence of magma mixing, crystal accumulation and other magmatic processes (e.g. Garcia, 1996).

The forsterite (Fo) content of the olivine cores in MU lavas ranges from 78.2 to 90.2 mol % (Table 1-3). It seems that the highest Fo values generally increase with the whole-rock Mg-number (Fig. 1-7), whereas the lower Fo values show no systematic relationship to Mg-number. The CaO content of olivine is high to moderate and mildly increases from 0.20 to 0.35 wt % with a decrease in Fo content. These CaO contents are typical of magmatic values and consistent with low pressure crystallisation in the crust (Jurewicz & Watson, 1988; Kamenetsky *et al.*, 2006; Simkin & Smith, 1970; Stormer, 1973; Thompson & Gibson, 2000). Mantle values are significantly lower (Fig. 1-8). The NiO content ranges widely and shows a generally increasing trend from 0.16 to 0.51 wt % with increasing Fo content.

Most of the olivine crystals show moderate to strong normal zoning from core to rim (Table 1-3; Fig. 1-9), the range of Fo variation is fairly wide. The typical Fo variation of normal zoning in all the MU lavas is around 1-5% with a maximum of ~22% in MU69-1. Some crystals are unzoned, while others are reversely zoned with a typical Fo difference of 2%, with a maximum variation of up to ~9% in MU69-1. The reversed zone is only 10-50 μm wide. The core forsterite content of the reversely zoned olivines ranges from ~78 to 86%. Many of these reversely zoned olivines are

observed in MU72-6. It should be noted that the core compositions of the normally zoned or unzoned olivines in the early erupted MU69-1 lava display a narrow range of Fo values, 87-90% with an average of $Fo_{88.7}$. In the other MU lavas, these high-Fo olivines are not so abundant. Based on spot analyses (and not traverses through the entire grain), most, if not all, deformed olivine crystals appear to be normally zoned in Fo content.

Of great interest is the presence of rare, complexly zoned olivine crystals. They were recognised using BSE images. However, their number may be somewhat underestimated because the whole thin section was not imaged using BSE. Some of the complexly zoned grains have a high-Fo core (85-87%) and a bulk lower-Fo rim which can be divided as follows: a 10-50 μm wide inner rim with Fo 74-76%, and a 10-20 μm wide outer rim with Fo 79-81%. Very rarely, the complex zoning of olivine is expressed as a core forsterite composition around 81-82%, an inner rim (20-30 μm width) around 70-71% and an outer rim of 10-50 μm wide skeletal overgrowth around 82-83%.

All MU lavas have some olivines that appear in Fe^{II}/Mg equilibrium with their host rock which displays a wide range of Mg-numbers (61-73%; Fig. 1-7). With the exception of MU73-1c, all MU lavas also contain olivines with Fo contents too low to be in equilibrium with the bulk composition of their host rock. This is especially well marked for MU69-1 sample, which has the highest Mg-number. The lavas with Mg-numbers of 61-67% all contain euhedral or subhedral, apparently undeformed olivine crystals with Fo contents that are too high for equilibrium, up to Fo_{90} (Fig. 1-7). These high-Fo olivines have CaO contents similar to that of the olivines in equilibrium with the magma (Table 1-3).

There are no systematic compositional differences between undeformed and deformed olivines in Fo, CaO, NiO and even MnO content, in either microphenocrysts or phenocrysts (Fig. 1-8, CaO and Fo% only represented). In other words, only crystal plastic deformation features permit

us to distinguish the deformed crystals from the undeformed ones. This is in agreement with previous findings from lavas from the Hawaii Scientific Drilling Project (Garcia, 1996) and from other olivine-rich lavas (Norman & Garcia, 1999; Wilkinson & Hensel, 1988).

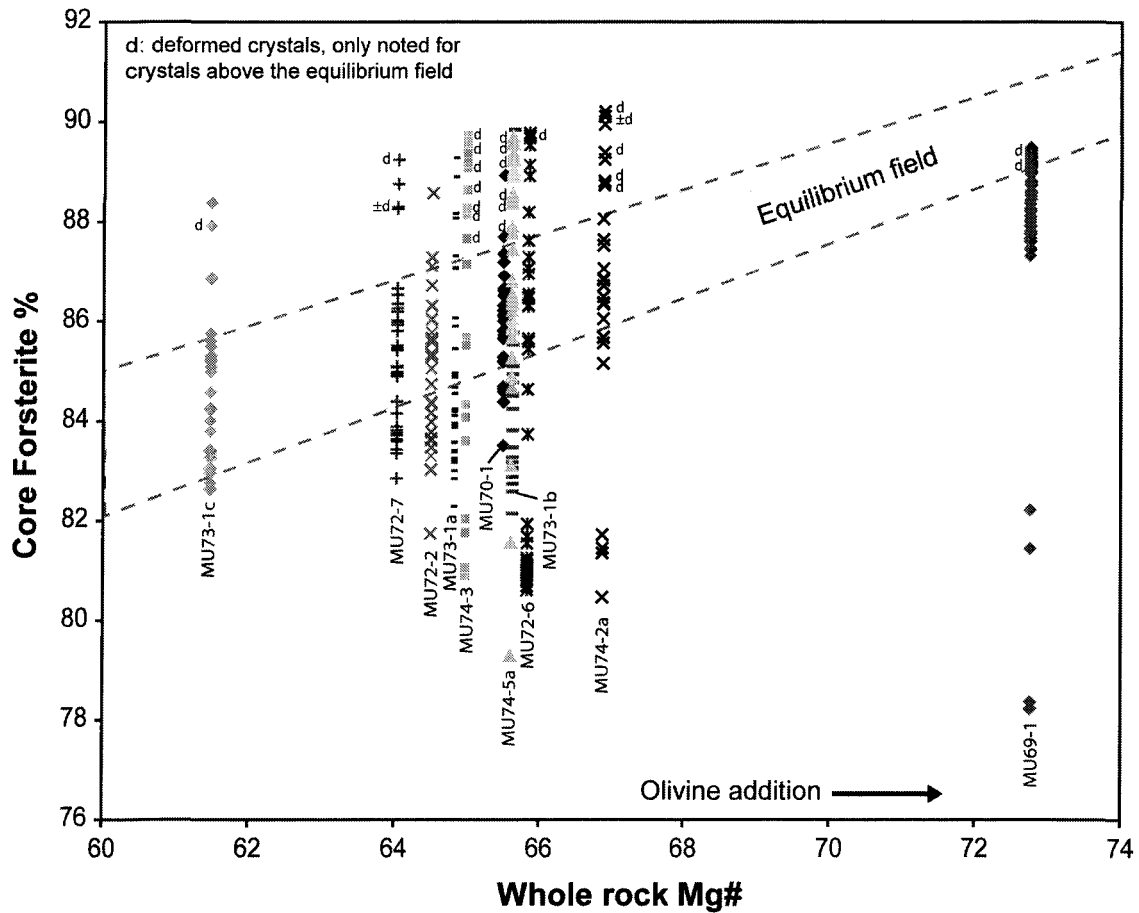


Figure 1-7. Whole-rock Mg-number (atomic $\text{Mg}/(\text{Mg} + \text{Fe}^{2+}) \times 100$) plotted against olivine core composition (forsterite %) for Mauna Ulu lavas. The Fe^{2+} is set to 90% of the total iron, as constrained by ferrous iron measurements (e.g., Herzberg *et al.*, 2007; Rhodes & Vollinger, 2004, 2005; Wright *et al.*, 1975). The diagonal field shows the range of equilibrium olivine compositions at 1 atm for a given Mg-number based on Roeder & Emslie (1970) and Ulmer (1989). Rocks with olivines that partly plot below the equilibrium field have probably experienced olivine accumulation. Representative analyses of olivine compositions given in Table 1-3.

d: deformed crystals (i.e., kink-bands, undulose extinction); only indicated in the high-Fo area of the plot.

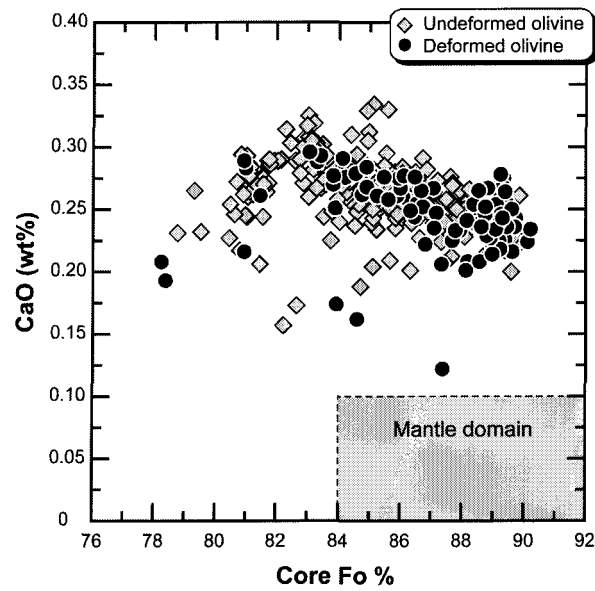


Figure 1-8. CaO content versus Fo content (%) of the core of both the undeformed and deformed olivine crystals analyzed in the Mauna Ulu lavas of this study. All olivines plot above the mantle domain indicated as the shaded area. Mantle domain after Kamenetsky *et al.* (2006), Simkin & Smith (1970) and Thompson & Gibson (2000).

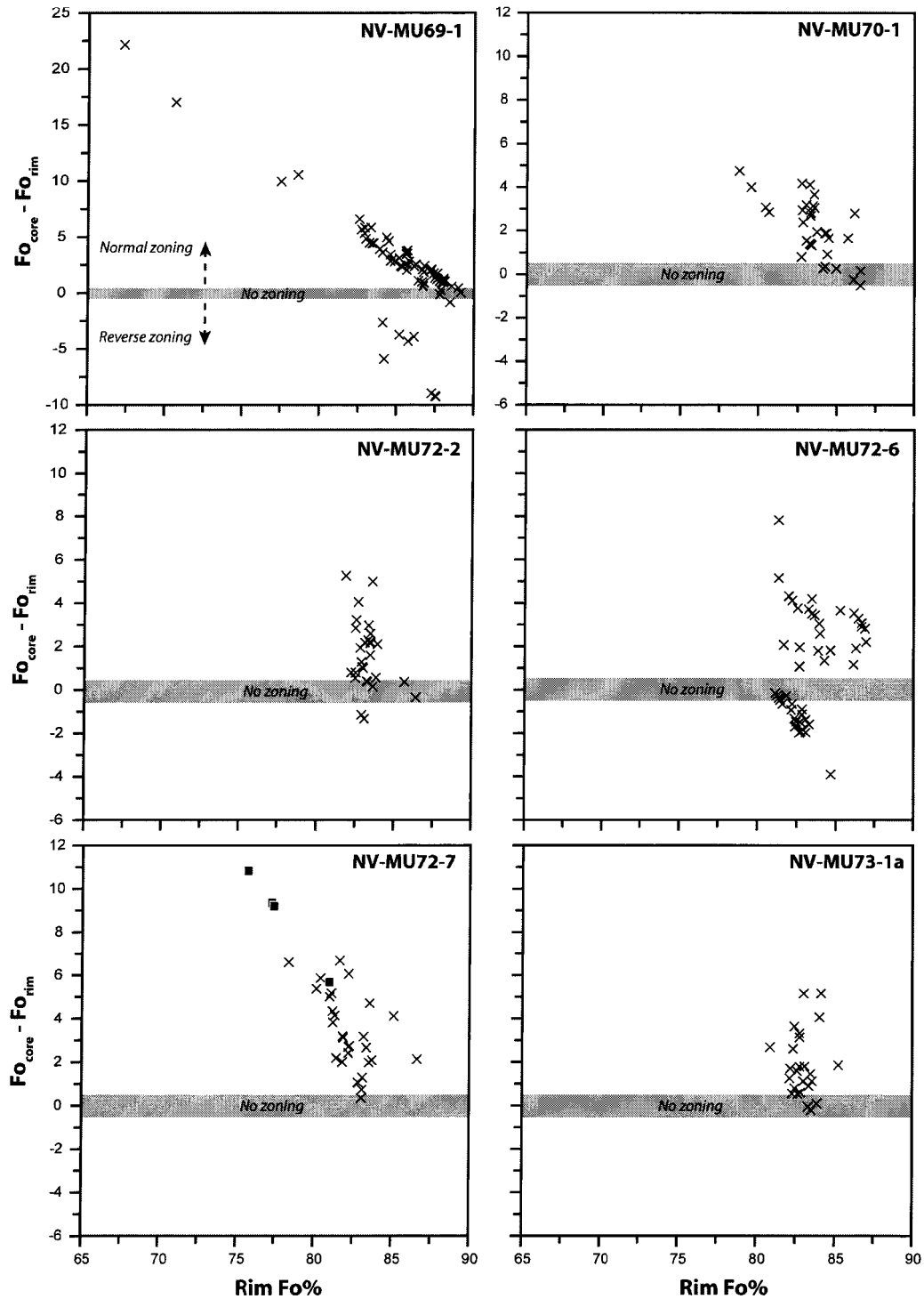


Figure 1-9. Variation of Fo content from core to rim versus Fo composition of the corresponding rim, for the olivine grains analyzed in different Mauna Ulu samples. The shaded area in each plot indicates the Fo composition range (around the variation value of 0) of no zoning. As represented in the first plot, grains normally zoned lie above this shaded area, and grains reversely zoned below it. Complexly zoned crystals are indicated by filled squares. Representative analyses given in Table 1-3.

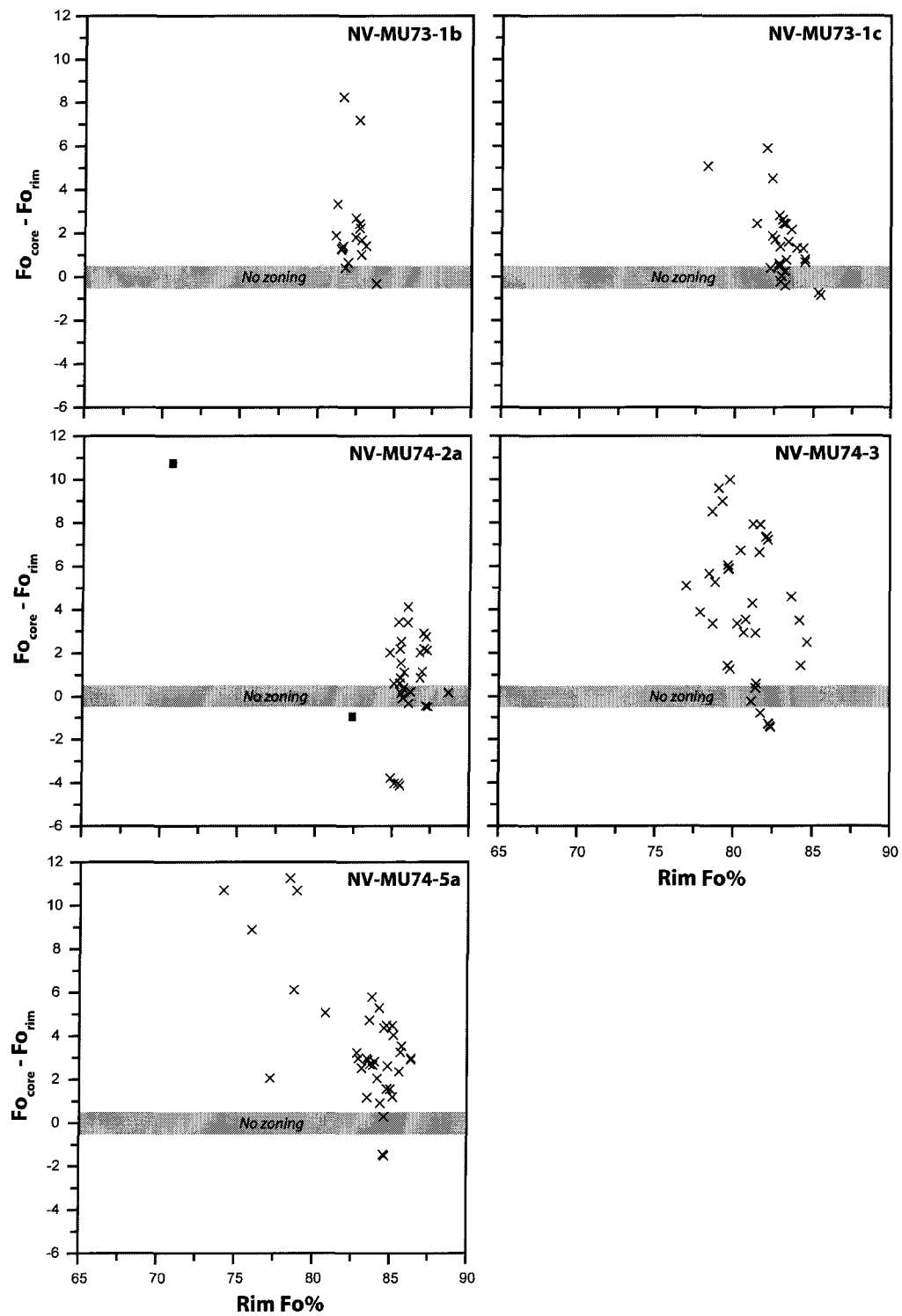


Figure 1-9 (continued)

Table 1-3: Representative microprobe analyses of olivine cores and rims from Mauna Ulu samples

Sample:	MU69-1												
Rock Mg-no.:	72.8												
Size:	ph-c	ph-r	ph-c	ph-r1	ph-r2	ph-c	ph-r1	ph-r2	ph-c*	ph-r*	ph-c	ph-r1	ph-r2
SiO ₂	40.08	36.12	37.98	38.84	39.48	39.87	40.20	39.26	39.50	39.29	39.45	38.94	36.78
MgO	47.64	31.73	39.42	43.73	46.16	47.50	47.61	45.07	46.06	45.63	46.38	44.56	34.36
CaO	0.24	0.29	0.21	0.22	0.26	0.26	0.24	0.29	0.21	0.28	0.27	0.26	0.29
MnO	0.08	0.22	0.30	0.24	0.22	0.16	0.11	0.14	0.15	0.29	0.19	0.23	0.31
FeO	11.10	30.53	21.71	16.25	13.32	11.41	11.47	14.92	13.24	13.85	12.90	15.25	28.24
NiO	0.44	0.15	0.24	0.38	0.33	0.34	0.45	0.31	0.35	0.28	0.30	0.32	0.17
Total	99.66	99.17	99.86	99.75	99.97	99.63	100.22	100.13	99.60	99.77	99.58	99.66	100.35
Fo%	89.5	67.3	78.2	84.2	87.3	89.2	89.2	85.7	87.3	86.7	87.7	85.3	70.7

Sample	MU70-1						MU72-2					
Rock Mg-no.	65.5						64.5					
Size:	mph-c*	mph-	ph-c	ph-r	ph-c	ph-r	ph-c	ph-r	ph-c	ph-r	ph-c	ph-r
SiO ₂	39.74	38.64	39.80	39.14	39.98	39.70	39.29	38.76	39.81	39.19	38.85	38.81
MgO	46.39	42.93	46.01	43.29	47.72	45.76	45.03	43.09	47.28	43.47	42.15	42.59
CaO	0.12	0.31	0.24	0.28	0.24	0.26	0.27	0.30	0.25	0.31	0.29	0.33
MnO	0.22	0.15	0.18	0.19	0.14	0.20	0.28	0.23	0.16	0.20	0.25	0.20
FeO	13.29	17.11	14.09	16.98	11.75	14.58	14.98	16.91	12.07	16.90	18.63	17.40
NiO	0.38	0.24	0.31	0.22	0.37	0.34	0.24	0.28	0.39	0.20	0.21	0.19
Total	100.26	99.46	100.80	100.21	100.35	101.09	100.22	99.68	100.14	100.40	100.45	99.63
Fo%	87.4	83.3	86.6	83.5	88.9	86.1	85.6	83.5	88.6	83.6	81.8	82.9

Sample	MU72-6						MU72-7					
Rock Mg-no.	65.8						64.0					
Size:	ph-c*	ph-r*	ph-c	ph-r	ph-c	ph-r	ph-c	ph-r	mph-c*	mph-	mph-	
SiO ₂	40.41	39.78	38.27	38.28	39.41	38.63	38.49	39.26	39.21	37.84	38.22	
MgO	48.52	45.51	41.42	42.51	45.77	42.29	41.56	44.51	44.28	38.37	42.85	
CaO	0.24	0.26	0.29	0.29	0.20	0.29	0.22	0.23	0.26	0.26	0.30	
MnO	0.15	0.21	0.27	0.26	0.22	0.27	0.23	0.24	0.24	0.31	0.30	
FeO	11.05	14.48	19.45	17.53	14.37	18.14	19.58	15.94	15.79	24.29	19.95	
NiO	0.51	0.31	0.16	0.29	0.43	0.21	0.25	0.30	0.25	0.17	0.22	
Total	101.10	100.65	99.97	99.40	100.52	99.95	100.50	100.58	100.16	101.35	101.92	
Fo%	89.7	86.2	80.8	82.8	86.3	82.2	80.8	84.7	84.7	75.8	81.0	

Table 1-3 (continued)

Sample	MU72-7				MU73-1a							
Rock Mg-no.	64.0				64.8							
Size	ph-c	ph-r	ph-c*	ph-r*	ph-c	ph-r	mph-c	mph-r	mph-c	mph-r	ph-c	ph-r
SiO ₂	39.33	38.70	39.56	39.19	40.16	39.09	39.18	38.95	39.07	38.87	39.31	38.75
MgO	44.48	42.08	47.55	44.53	47.58	43.91	43.32	42.94	42.83	42.32	44.68	42.43
CaO	0.28	0.34	0.25	0.28	0.25	0.26	0.27	0.31	0.27	0.31	0.31	0.33
MnO	0.24	0.21	0.14	0.17	0.09	0.24	0.27	0.33	0.21	0.28	0.17	0.26
FeO	15.42	18.51	11.35	15.40	11.31	16.41	17.05	17.64	17.56	18.02	15.66	18.00
NiO	0.26	0.21	0.38	0.31	0.43	0.29	0.25	0.21	0.23	0.20	0.27	0.22
Total	100.17	100.14	99.39	100.00	99.97	100.26	100.47	100.47	100.31	100.12	100.51	100.03
Fo%	85.1	81.8	89.2	85.1	89.3	84.1	83.4	82.8	82.9	82.3	85.0	82.4

Sample	MU73-1b				MU73-1c							
Rock Mg-no.	65.6				61.4							
Size:	ph-c	ph-r	ph-c	ph-r	mph-c	mph-r	ph-c*	ph-r*	ph-c	ph-r	mph-c	mph-r
SiO ₂	39.16	38.79	40.35	38.80	38.54	38.50	39.66	38.81	39.27	38.79	39.00	39.09
MgO	44.49	42.38	48.32	42.76	42.27	42.17	46.59	42.21	44.18	42.82	43.34	43.32
CaO	0.33	0.31	0.26	0.32	0.29	0.32	0.23	0.30	0.28	0.29	0.29	0.28
MnO	0.20	0.35	0.22	0.27	0.17	0.28	0.12	0.27	0.27	0.19	0.19	0.16
FeO	15.42	17.90	10.82	17.76	18.19	18.66	12.67	18.33	16.37	17.54	17.06	17.37
NiO	0.30	0.22	0.31	0.25	0.20	0.18	0.39	0.20	0.25	0.22	0.18	0.23
Total	99.98	100.02	100.31	100.28	99.78	100.27	99.84	100.25	100.68	100.12	100.24	100.55
Fo%	85.1	82.4	89.8	82.7	82.2	81.7	87.9	82.0	84.2	82.9	83.4	83.2

Sample	MU74-2a						
Rock Mg-no.	66.9						
Size:	ph-c*	ph-r1*	ph-r2*	ph-c*	ph-r*	mph-c	mph-r
SiO ₂	38.71	36.92	39.00	40.54	39.73	39.77	39.57
MgO	40.98	33.24	41.60	47.92	44.45	45.65	44.70
CaO	0.26	0.22	0.31	0.22	0.26	0.28	0.28
MnO	0.25	0.31	0.23	0.19	0.17	0.22	0.10
FeO	18.47	27.25	17.58	10.42	14.35	13.68	14.72
NiO	0.16	0.15	0.16	0.39	0.20	0.24	0.23
Total	98.94	98.11	99.10	99.86	99.43	100.04	99.78
Fo%	81.5	70.7	82.4	90.1	86.0	86.9	85.7

Table 1-3 (continued)

Sample	MU74-3						MU74-5a					
Rock Mg-no.	65.0						65.6					
Size:	ph-c	ph-r	ph-c	ph-r	ph-c*	ph-r*	ph-c*	ph-r*	ph-c	ph-r	mph-c	mph-r
SiO ₂	40.17	38.81	38.62	38.29	39.30	38.52	39.75	38.90	38.83	39.16	39.37	38.84
MgO	47.24	41.85	41.51	40.24	44.22	41.46	47.59	44.31	42.91	44.06	45.56	43.36
CaO	0.24	0.28	0.27	0.33	0.27	0.32	0.25	0.29	0.31	0.32	0.26	0.31
MnO	0.20	0.24	0.28	0.29	0.19	0.28	0.16	0.25	0.23	0.25	0.19	0.24
FeO	12.43	18.65	19.20	20.38	16.27	18.75	10.95	15.33	17.26	15.83	14.03	16.50
NiO	0.39	0.23	0.22	0.24	0.30	0.27	0.37	0.28	0.18	0.26	0.28	0.26
Total	100.85	100.20	100.23	100.00	100.66	99.71	99.24	99.46	99.77	100.12	99.76	99.60
Fo%	88.3	81.6	81.1	79.6	84.3	81.4	89.6	85.1	83.1	84.6	86.5	83.9

Rock Mg-no. is $[\text{Mg}/(\text{Fe}^{2+} + \text{Mg})] \times 100$ assuming 90% of the total iron is Fe^{2+} ; ph, phenocryst; mph, microphenocrysts; r, rim (r1, inner rim; r2, outer rim); c, core; oxides are in wt %; forsterite (Fo); * deformed (or \pm deformed) crystal.

1.5 QUANTITATIVE TEXTURAL MEASUREMENTS: CRYSTAL SIZE DISTRIBUTION (CSD)

1.5.1 CSD theory

The CSD analysis of a rock is a quantitative measure of the number of crystals of a mineral phase per unit volume in intervals of fixed size (Higgins, 2006a). CSD theory was initially developed in chemical engineering by Randolph & Larson (1971), among others, for understanding industrial crystallisation processes, and then applied to magmatic systems by Marsh (1988). Since then a number of studies have focused on the quantification of crystal populations in rocks using advanced textural analysis techniques (Jerram & Kent, 2006).

Marsh (1988, 1998) proposed two distinct CSD end-member models for igneous systems: (1) an open-system, steady-state model, and (2) a closed-system, batch model. The Marsh steady-state model describes an open system where there is balance between the number of crystals that nucleated and grew in the magma and the loss or gain of grains due to further mechanisms of transport (e.g., fractionation, mixing). Such an ideal system may be represented by a continuously tapped and refilled magma chamber, tapping and refilling occurring at the same rate. The continuous crystallisation typically results in a straight, or log-linear, CSD plot with a negative slope on a graph of $\ln(\text{population density})$ versus size, as described by the equation

$$n = n^{\circ} \cdot \exp\left(-\frac{L}{G\tau}\right)$$

Where n is the population density (mm^{-4}), n° is the nucleation density and represents the intercept on the y-axis, G is the mean crystal growth rate (mm/s), τ is the mean crystal growth time or

crystal residence time, and L is the crystal size (maximum length; mm). G and the nucleation rate are assumed to be constant. G is independent of L . One unique CSD exists in constant conditions. Log-linear or near log-linear CSDs are common in igneous rocks (Marsh, 1998).

In a closed-system batch model (Marsh, 1998), there is no input or output of crystals, and therefore the volume proportion of crystallizing magma decreases with time. With exponentially increasing nucleation rates and a constant growth rate, closed systems can also produce straight CSDs. The Makaopuhi lava lake, Kilauea, is an example of this in a volcanic system (Cashman & Marsh, 1988).

Marsh (1998) noted that CSDs from a wide range of igneous settings do not show typical steady-state or batch behaviour, but reflect hybrid aspects of the two end-member models. He also argued that CSDs in comagmatic sequences do not closely resemble CSDs expected in either batch or open systems, but seem to reflect the dynamic regime of the system. Hence, the steady-state model should be applied to near-surface crystallisation, and the batch model to crystallisation within a deep-seated cumulate body. The steady-state model is employed in this study as an overall proxy for quantifying most magma processes, since it yields effective values of crystal growth times and nucleation rates (J) from this commonly used CSD model (Marsh, 1988, 1998). These quantitative parameters may be extracted from the slope and intercept of a log-linear CSD plot:

$$\text{Slope} = -\frac{1}{G\tau} \text{ and } \text{Intercept} = \frac{J}{G}.$$

In an open-system end-member model, a straight or log-linear CSD might be the result of simple crystal nucleation and growth processes, and referred to as a kinetic texture (e.g. Higgins, 2006a). Nucleation and growth rates are controlled by the magnitude of the crystallisation driving

force. Variations in temperature and pressure will change the driving force of crystallisation. A decrease in temperature causes the appearance of a family of straight lines with the same slope but different intercepts (i.e. different nucleation densities) on a classic CSD diagram (Fig. 1-10a). An increase in residence time and a constant growth rate will favour an increase in larger crystals and hence a change in the CSD slope, but no change in the intercept (Fig. 1-10b).

Kinking or curvature in the CSD plot profile (Fig. 1-10c-e) has been attributed to processes such as crystal accumulation and fractionation (Marsh, 1998), compaction and compaction-driven recrystallisation (e.g. Boorman *et al.*, 2004), mixing of crystal populations or magmas (e.g. Higgins, 1996; Higgins & Roberge, 2007; Jerram *et al.*, 2003), textural coarsening (= Ostwald ripening) (e.g. Higgins, 2002b; Higgins & Roberge, 2003), and variation of cooling rate during ascent and emplacement of magma (e.g. Armienti *et al.*, 1994).

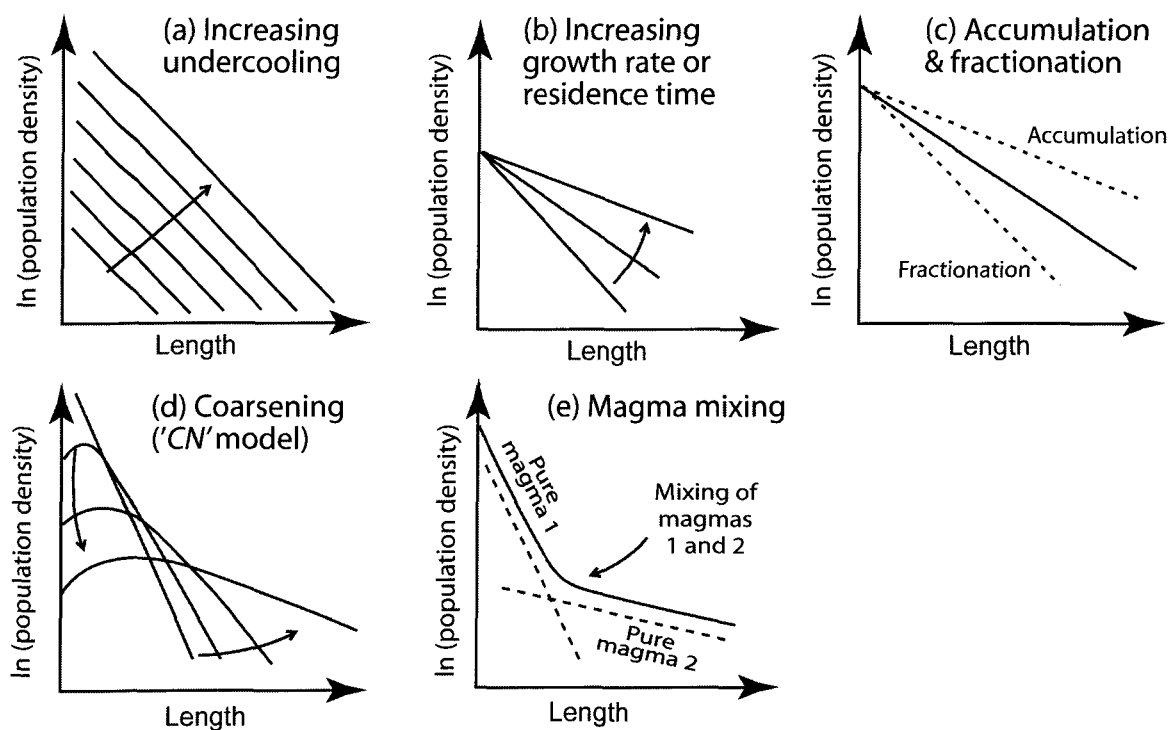


Figure 1-10. Schematic examples of processes that can influence the shape of the CSD (after Higgins, 2006a): (a) Increase in undercooling (or saturation); (b) Increase in residence time or growth rate; (c) Accumulation and fractionation of crystals; (d) Coarsening; (e) Magma mixing (or mixing of two crystal populations).

1.5.2 Analytical methods

To determine olivine CSDs, we stitched together high-resolution photomicrographs of each thin section to make a single image of each thin section (Fig. 1-11). Subsequently, individual crystals were identified and outlined by using a polarizing microscope and a vector graphics editor. Effort was made to outline and measure separately each individual crystal in a cluster. In addition, all optically discernible deformed grains were outlined in a separate layer in order to compare the CSDs of undeformed versus total olivine populations (Fig. 1-11b). The digitized olivine texture images were then edited into simple greyscale compressed tiff format before importing into the software *ImageJ* (*Image Processing and Analysis in Java*). Area, orientation and length of major and minor axes of a best fitted ellipse to crystal outlines were measured.

We measured 360 to >1000 olivine crystals in each thin section. Crystal size ranges are presented in Table 1-4. Intersection data based on the major axis of the fitted ellipse determined with *ImageJ* were converted to 3-D CSDs using the program *CSDCorrections 1.37* (Higgins, 2000, 2002a; Higgins, 2006a). Logarithmic length intervals were used, with each bin $10^{0.2}$ times the size of the previous bin. There were no gaps in the CSDs; all bins contained crystals. Bins with less than three crystals were removed from the CSD analysis as not significant. The lower limit of the CSD was 0.05 mm, the smallest crystal that could be measured consistently and not the smallest crystal in the rock.

To calculate CSDs from 2-D intersection measurements using *CSDCorrections 1.37*, we needed to specify the mean crystal shape as expressed by the crystal aspect ratio $S:I:L$ (short:intermediate:long dimensions). These aspect ratios were objectively estimated for each thin section, following Morgan & Jerram's (2006) method and their extensive database *CSDS/ice*. A bulk

fit value of the olivine shape, 1:1.2:1.7, was used for every CSD plot of all the samples from Mauna Ulu, in order to compare them adequately. This specific aspect ratio was chosen because, for every sample of this study, it appeared in every set of the five best fit values yielded as output results by *CSDSlice*. In contrast, all the individual CSD-derived textural parameters were obtained by using in the calculations the best fit individual aspect ratio (Table 1-5), among the set of the best five values estimated by *CSDSlice*. This gave greater accuracy in quantification. For the purposes of *CSDCorrections 1.37*, rock fabric was specified as massive; crystal roundness factor was estimated to be 0.5 in all cases. The complete list of all quantitative textural data from *CSDCorrections 1.37* is given in Table 1-5.

The CSD data were plotted on a $\ln(\text{population density})$ versus size diagram (Marsh, 1988) (Fig. 1-12). For CSDs only slightly curved on a standard CSD diagram (S-type distribution of Higgins, 2006b), meaningful values for the slope and intercept of the CSD were determined by linear regression following Higgins (2006b). The 'Goodness of fit', Q , is a parameter that describes how well the data fit to a straight line (Table 1-6), taking into account the error in each point. A value of Q greater than 0.1 indicates a very significant fit, but values greater than 0.01 are also acceptable (Higgins, 2006b). Strongly curved CSDs necessitated a different approach: the slope and intercept were calculated separately for the smallest and largest crystals using the mean of several CSD bins (Fig. 1-13).

The characteristic length (L_D) of the CSDs was then calculated as $-1 / \text{slope}$ for the entire pattern or a single segment. This parameter is equal to the mean crystal length for a straight CSD that extends to all crystal sizes (Marsh, 1988). The volumetric olivine proportion is obtained by dividing the total area of the olivine crystals by the total measured area. A plot of phase volumetric proportion versus L_D (Fig. 1-14) is more useful than the commonly used graph of slope

versus intercept, because of the correlation between these parameters produced by closure (pp71-74 in Higgins, 2006a).

The maximum 3-D crystal size is also an important parameter of the CSD, but no rigorous assessment is possible using two-dimensional data. An estimate of this characteristic is the maximum intersection size (L_{max} ; Table 1-5). This is not a very reliable measure because of the small number of large crystals present, which are not necessarily in the correct orientation. Here, we used the average of the intersection length of the three or four largest crystals for each sample as suggested by Marsh (1998) and O'Driscoll *et al.* (2007).

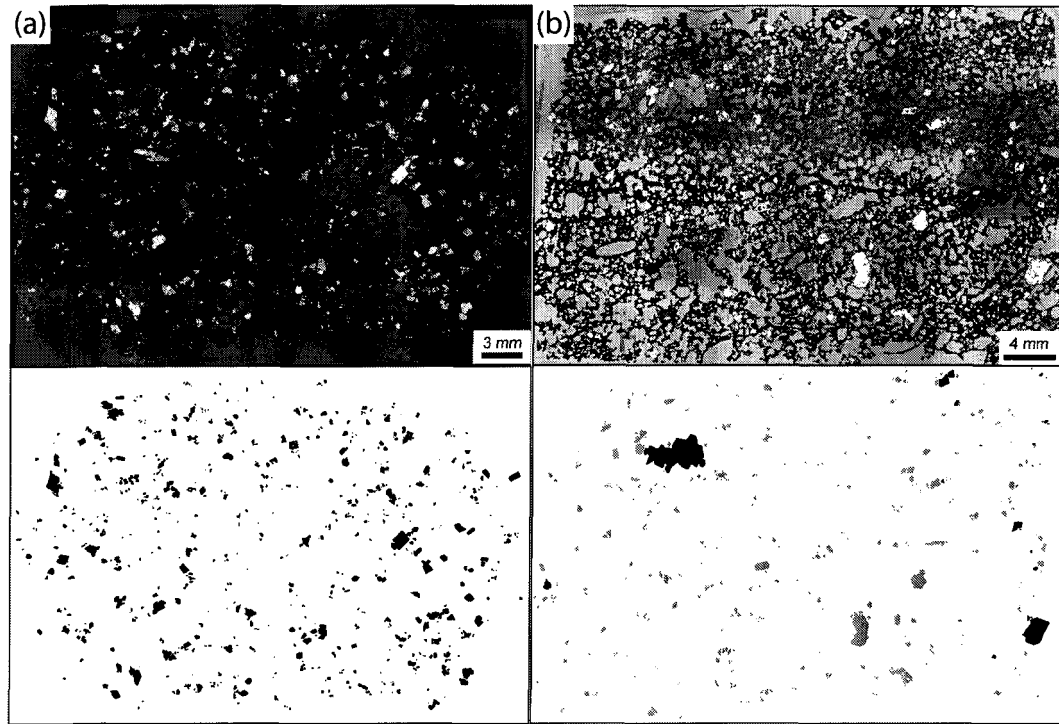


Figure 1-11. Stacked photomicrographs above and digitized olivine texture images below.
(a) Glassy lava NV-MU72-2, (b) Microcrystallized lava NV-MU74-5a (undeformed crystals are light grey, deformed crystals are black).

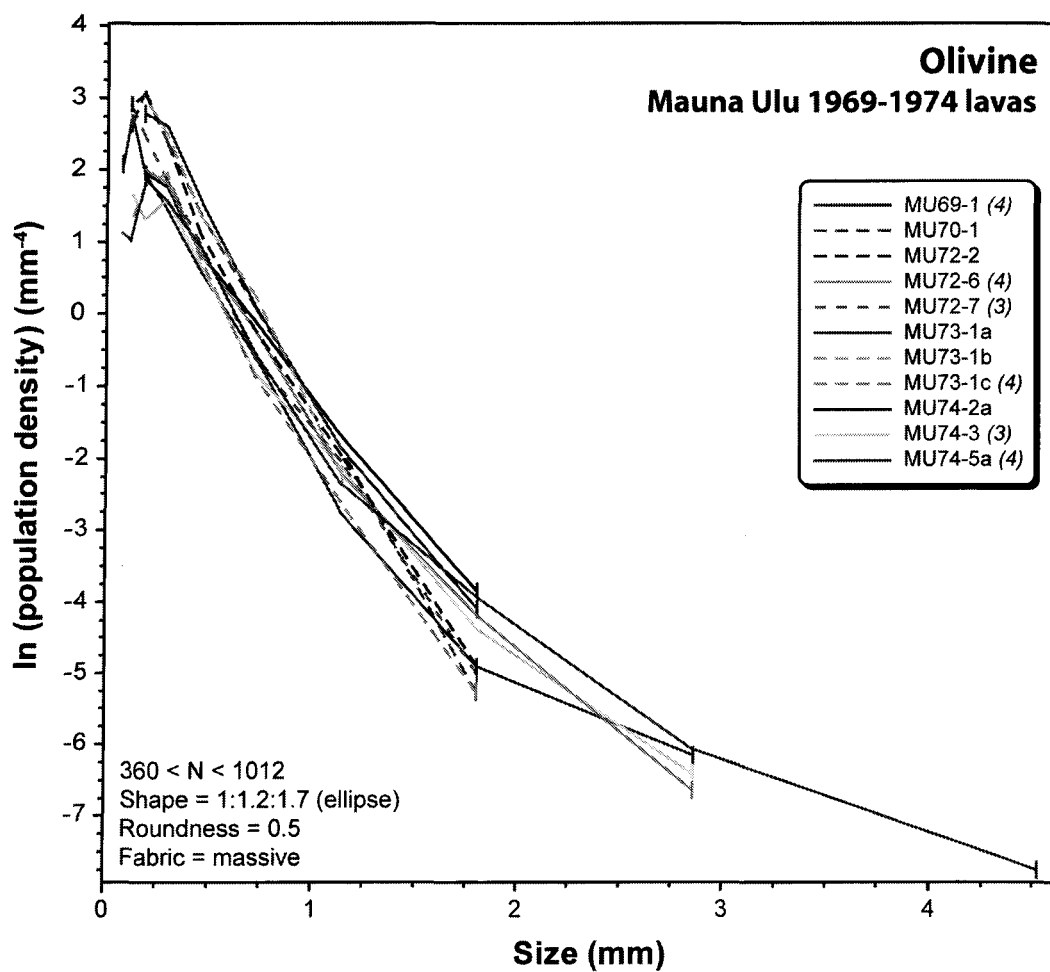


Figure 1-12. Olivine crystal size distributions of the Mauna Ulu lavas plotted following the conventions of Higgins (2006a). Numbers in parentheses next to some sample names are the number of crystals counted in the CSD analysis for the last bin, that is, for the largest grains.

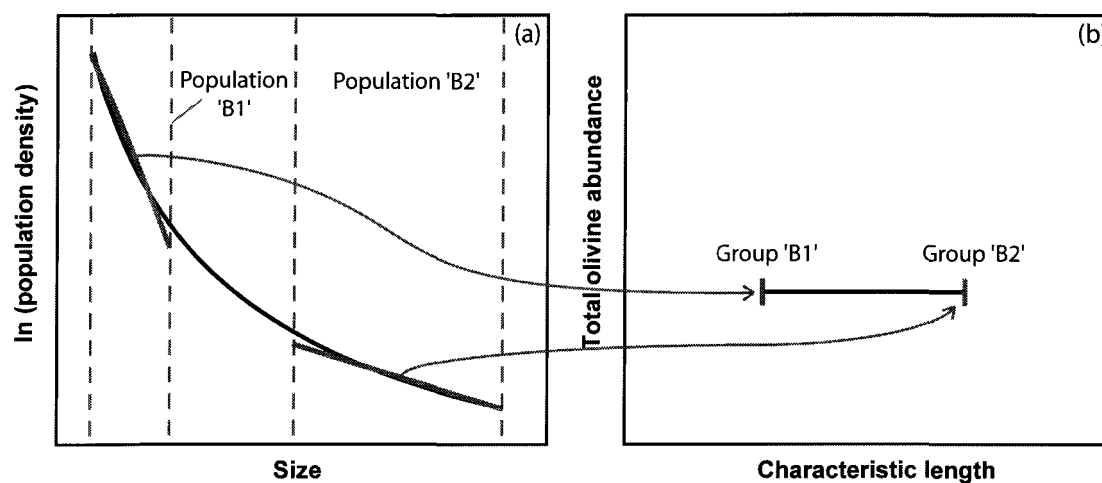


Figure 1-13. Schematic diagrams illustrating the segmentation of CSDs. (a) The original CSD curve is in black. The segments in grey both at the left and right sides of the CSD are for the olivine populations B1 and B2, respectively. Each segment (straight line) represents a mean of several CSD bins. (b) A plot of total sample olivine abundance vs. characteristic length for the two segments. The calculation inferred from the CSD segmentation permits placement of two points for the same sample of a given olivine abundance, representing here by the two small vertical bars in grey. The left bar corresponds to the population B1 segment and the right bar to the population B2 segment. The length of the horizontal line that links the vertical bars gives us information on the degree of curvature of the CSD, and thus on the level of mixing of at least two different crystal populations, for example.

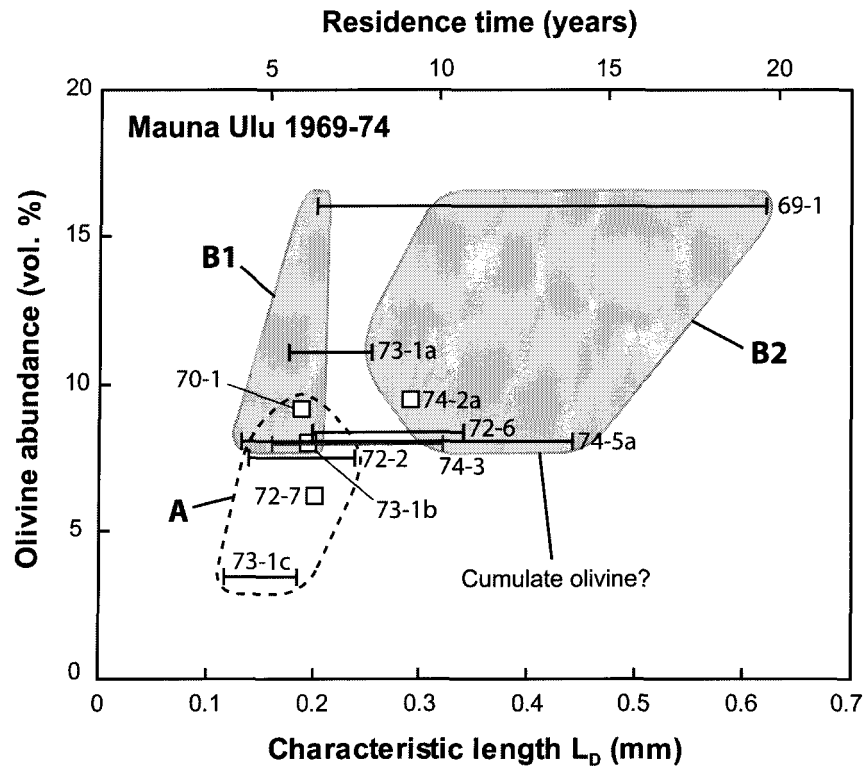


Figure 1-14. Characteristic length (L_D) versus olivine volumetric proportion in the Mauna Ulu lavas. The volume-percent olivine axis applies to the whole sample mentioned in the plot. Inferred from segmentation of the curved CSDs, the two extremities of the horizontal bar per sample correspond to the L_D of the extreme CSD segments. The dashed area is representative of group 'A', and the grey-filled areas of group 'B' (including populations B1 and B2) discussed in the text. The vertical ranges of the areas of populations B1 and B2 are the same. Also indicated in X-axis is the residence time of crystal, assuming a mean growth rate of 10^{-9} mm/s (estimate after Cashman, 1993; Mangan, 1990; Marsh, 1988, 1998; Wright & Fiske, 1971), instead of a range of values for the clarity of the diagram. This value of 10^{-9} mm/s is within the range defined in the text.

Table 1-4: Crystal size ranges expressed as the number of crystals per unit area

Samples		Number of crystals per crystal size interval used in the CSD analysis									
Sample	Area	6.31-	3.98-	2.51-	1.58-	1.00-	0.631-	0.398-	0.251-	0.158-	0.1000-
name	measured	3.98	2.51	1.58	1.00	0.631	0.398	0.251	0.158	0.100	0.0631
	(mm ²)	(mm)									
NV-MU69-1	1950	4	9	30	60	146	263	225	147	61	24
NV-MU70-1	965.9			4	46	133	209	240	208	90	
NV-MU72-2	853.6			5	38	89	142	204	177	89	
NV-MU72-6	1478			4	18	52	151	185	210	128	
NV-MU72-7	548.9			3	23	41	81	84	68	47	
NV-MU73-1a	1023			13	50	144	260	314	192		
NV-MU73-1b	880.1				29	141	187	242	185		
NV-MU73-1c	965.5				4	23	55	113	153	74	34
NV-MU74-2a	927.1			15	53	107	118	126	77		
NV-MU74-3	870.9		3	9	34	54	95	102	46	29	8
NV-MU74-5a	946.5		4	6	19	67	94	98	71	65	23

Table 1-5: Textural data (input and output) from CSDCorrections

Sample	Crystal morphology	R ² of shape best fit	Olivine vol. prop. (%)	No. of crystals	L _{max} (mm)
NV-MU69-1	rectangular prism (1:1.2:1.6)	0.930	16.0	974	5.68
NV-MU70-1	rectangular prism (1:1.15:1.6)	0.901	9.3	950	1.48
NV-MU72-2	rectangular prism (1:1.2:1.7)	0.928	7.4	754	1.39
NV-MU72-6	rectangular prism (1:1.15:1.5)	0.905	8.3	796	2.48
NV-MU72-7	rectangular prism (1:1.2:1.7)	0.916	6.2	360	1.17
NV-MU73-1a	rectangular prism (1:1.15:1.7)	0.919	10.9	1011	1.49
NV-MU73-1b	rectangular prism (1:1.2:1.8)	0.917	8.0	839	0.99
NV-MU73-1c	rectangular prism (1:1.4:1.5)	0.903	3.4	462	1.21
NV-MU74-2a	rectangular prism (1:1.15:1.8)	0.902	9.1	503	1.43
NV-MU74-3	rectangular prism (1:1.1:1.4)	0.872	8.0	381	2.30
NV-MU74-5a	rectangular prism (1:1.15:1.5)	0.924	8.0	452	3.23

Aspect ratios and R² calculated after Morgan & Jerram (2006).

L_{max} calculated from the three largest intersection lengths of each sample (after Marsh, 1998; O'Driscoll *et al.*, 2007).

Table 1-6: CSD results including slope and intercept data calculated by CSDCorrections for each sample

Sample	Matrix	Eruption year	Slope	error	Intercept $\ln(n_0)$	error	Crystal growth time (years)		Nucleation rate ($\times 10^{-9}$ mm ³ /s)		Characteristic length (mm)	Goodness of fit (Q)
				(1 σ)		(1 σ)	G1	G2	G1	G2		
NV-MU69-1	microlitic	1969	-4.95	0.17	2.88	0.09	2.6	12.8	7.2	1.44	0.20	0.05
			-1.61	0.22	-1.60	0.48	7.9	39.4	4.0	0.80	0.62	0.14
NV-MU70-1	glassy	1970	-5.30	0.15	3.82	0.08	2.4	12.0	9.6	1.91	0.19	0.14
NV-MU72-2	glassy	1972	-7.13	0.50	4.33	0.16	1.8	8.9	10.8	2.17	0.14	0.45
			-4.20	0.35	2.40	0.33	3.0	15.1	6.0	1.20	0.24	0.77
NV-MU72-6	intermediate	1972	-5.02	0.20	3.03	0.11	2.5	12.6	7.6	1.52	0.20	0.12
			-2.92	0.31	0.82	0.42	4.3	21.7	2.1	0.41	0.34	0.46
NV-MU72-7	microlitic	1972	-4.87	0.21	3.23	0.11	2.6	13.0	8.1	1.62	0.21	0.18
NV-MU73-1a	glassy	1973	-5.62	0.24	4.19	0.12	2.3	11.3	10.5	2.10	0.18	0.40
			-3.93	0.23	2.91	0.22	3.2	16.1	7.3	1.46	0.25	0.08
NV-MU73-1b	intermediate	1973	-5.08	0.16	3.89	0.09	2.5	12.5	9.7	1.95	0.20	0.06
NV-MU73-1c	microlitic	1973	-8.57	0.53	3.92	0.18	1.5	7.4	9.8	1.96	0.12	0.52
			-5.43	0.56	2.26	0.40	2.3	11.7	5.7	1.13	0.18	0.88
NV-MU74-2a	glassy	1974	-3.42	0.13	2.52	0.10	3.7	18.5	6.3	1.26	0.29	0.20
NV-MU74-3	microlitic	1974	-6.16	0.51	3.20	0.22	2.1	10.3	8.0	1.60	0.16	0.88
			-3.08	0.41	0.76	0.52	4.1	20.6	1.9	0.38	0.32	0.21
NV-MU74-5a	glassy	1974	-7.46	1.32	3.38	0.28	1.7	8.5	8.5	1.69	0.13	0.10
			-2.26	0.38	-0.52	0.55	5.6	28.1	1.3	0.26	0.44	0.14

Nucleation rates and crystal growth times (residence times) are calculated for olivine growth rates $G1 = 2.5 \times 10^{-9}$ mm/s and $G2 = 5 \times 10^{-10}$ mm/s (estimate after Cashman, 1993; Mangan, 1990; Marsh, 1988, 1998; Wright & Fiske, 1971). The "Goodness of fit", Q, is a parameter that describes the quality of fit of the linear regression (see text for details).

1.5.3 CSD results

Although all the MU CSD patterns are broadly similar, they show significant differences in size, nucleation density and curvature, depending on the approximate eruption date and the proportions of deformed crystals. The complete CSD results are presented in Table 1-6. The size range of each CSD is variable (Fig. 1-12). Most samples have CSDs that range from ~0.1 mm, the lower analytical limit, to 1.8 mm; MU72-6, MU74-3 and MU74-5a extend to ~2.8 mm; MU69-1 is over 2.8 mm, and MU73-1b only extends to ~1.2 mm in size. There is no simple link between CSDs and eruption date.

Samples collected from specific lava flows show significant inter-flow variability in their CSDs. However, CSDs show no significant in-flow variations, except for the 1973 flow. Both MU72-2 and MU72-7 sample the same flow and have similar CSDs and comparable olivine abundances. MU72-2 has a glassy matrix and was collected from the termination of the tube-fed flow. However, MU72-7 has a matrix with microlites and was taken from the top of the flow, much closer to the MU vent. Similarly, 1974 flow samples MU74-2a, MU74-3 and MU74-5a have relatively comparable CSDs and olivine proportions, but MU74-2a fits a short straight line not exceeding 1.8 mm in size. In contrast, samples from the 1972 lava flow exhibit significant variations compared to the 1974 flow samples, as illustrated in their CSDs (Figs. 1-12 and 1-15). The only example of in-flow variability is the 1973 flow. The three samples, MU73-1a, MU73-1b and MU73-1c, collected from the same tumulus, near the westernmost upper rim of Makaopuhi Crater, exhibit significant textural differences. The glassy MU73-1a and the intermediate MU73-1b are from a blue glassy pahoehoe surface and have similar CSDs. In contrast, the microlitic MU73-1c is from deeper in the tumulus and displays a parallel, but displaced downwards CSD.

The majority of the samples show a turndown of the CSD curves at smallest crystal sizes and it is important to determine if this is a measurement artefact. When the samples are plotted separately, all glassy samples show a turndown (Fig. 1-15a). Thus the turndown at the left side of the CSD is clearly not an artefact for all crystals are measured in glassy samples. The same holds true for some of the samples with a crystallized matrix that also have a turndown at smallest sizes (Fig. 1-15b) where only crystals >0.1 mm were measured. Therefore, the turndown is unlikely to be an artefact of measurement for the microlite MU lavas either and hence must reflect solidification processes.

Two groups of samples can be defined on the basis of their CSDs (Fig. 15). Group 'A' (Fig. 1-15c) consists mostly of samples with straight CSDs (MU70-1, MU72-7 and MU73-1b) and two with slightly curved CSDs (MU72-2, MU73-1c). All samples have steep negative slopes from -4.2 to -8.6 and corrected maximum crystal sizes no larger than 1.8 mm. CSD group 'A' has higher intercepts (2.3-4.3) and moderate characteristic lengths (0.12-0.24 mm). It should be noted that, although MU73-1c is thought to belong to group 'A' based on the textural parameters inferred from the CSD analysis, this sample has a significantly lower olivine volumetric proportion (~ 3.4 %) than all the samples studied here (Fig. 1-14). Thus, to a certain extent, MU73-1c is texturally unique.

Group 'B' (Fig. 1-15d) mostly consists of samples with highly curved CSDs (MU69-1, MU72-6, MU73-1a, MU74-3 and MU74-5a) except for MU74-2a which has a straight and short CSD. These curved CSDs were segmented so that they could be better understood: one segment for the smallest sizes, and the other for the largest sizes. The two olivine populations are referred to as 'B1' for the smallest crystals and 'B2' for the largest crystals. Two different grey-filled areas enclosing each of those two populations are represented in Figure 1-14. Population 'B1' shows higher intercepts of 2.5 to 4.2 and lower L_D (0.13-0.20 mm). This population has CSD similarities

with group 'A'. Population B2 has lower intercepts (<2.5 , except for MU73-1a with 2.9) and higher L_D values (0.25-0.62 mm).

Overall, there are significant differences in the maximum 3-D sizes (i.e., L_{max}) between the MU lavas. MU69-1 has the highest L_{max} (~5.7 mm), while most samples have L_{max} in the range of 1-1.5 mm, including all of group 'A' and some group 'B' samples (MU73-1a and 74-2a). Three group 'B' samples have intermediate values, ranging from 2.3 to 3.2 mm (MU74-3, MU72-6 and MU74-5a).

The CSDs of deformed crystals of olivine are very variable. It should be noted that the proportion of deformed crystals for each sample is generally underestimated since the petrologic evidence is not always clear, particularly when the grains are small. In many samples the number of deformed crystals was so low that separate CSDs could not be constructed. Hence, we have compared the CSD of the total number of crystals with that of the undeformed crystals only (Figs. 1-16 and 1-A1). First, no obvious relationship with glassy samples is noted. Second, the samples from group 'A' have, in general, CSDs of undeformed crystals identical to the total CSDs. MU72-7 is an exception, because it lacks large deformed crystals. Another exception is MU73-1b, which has an undeformed olivine CSD displaced downward relative to the total CSD. In the same way, undeformed crystal CSDs of several samples from group 'B' also show this small displacement downward, which is accentuated at larger sizes. Finally, the CSDs of the undeformed olivines of MU69-1 and MU74-3 from group 'B' are marked by the absence of the large deformed crystals compared to the total CSDs.

A plot of I/L versus S/I highlights the slight morphological differences in shapes of olivine crystals between each individual sample, with MU73-1c showing the highest I/L and lowest S/I

ratios (Fig. 1-17). This latter sample is thus characterized by a mean crystal shape clearly different compared to the other samples.

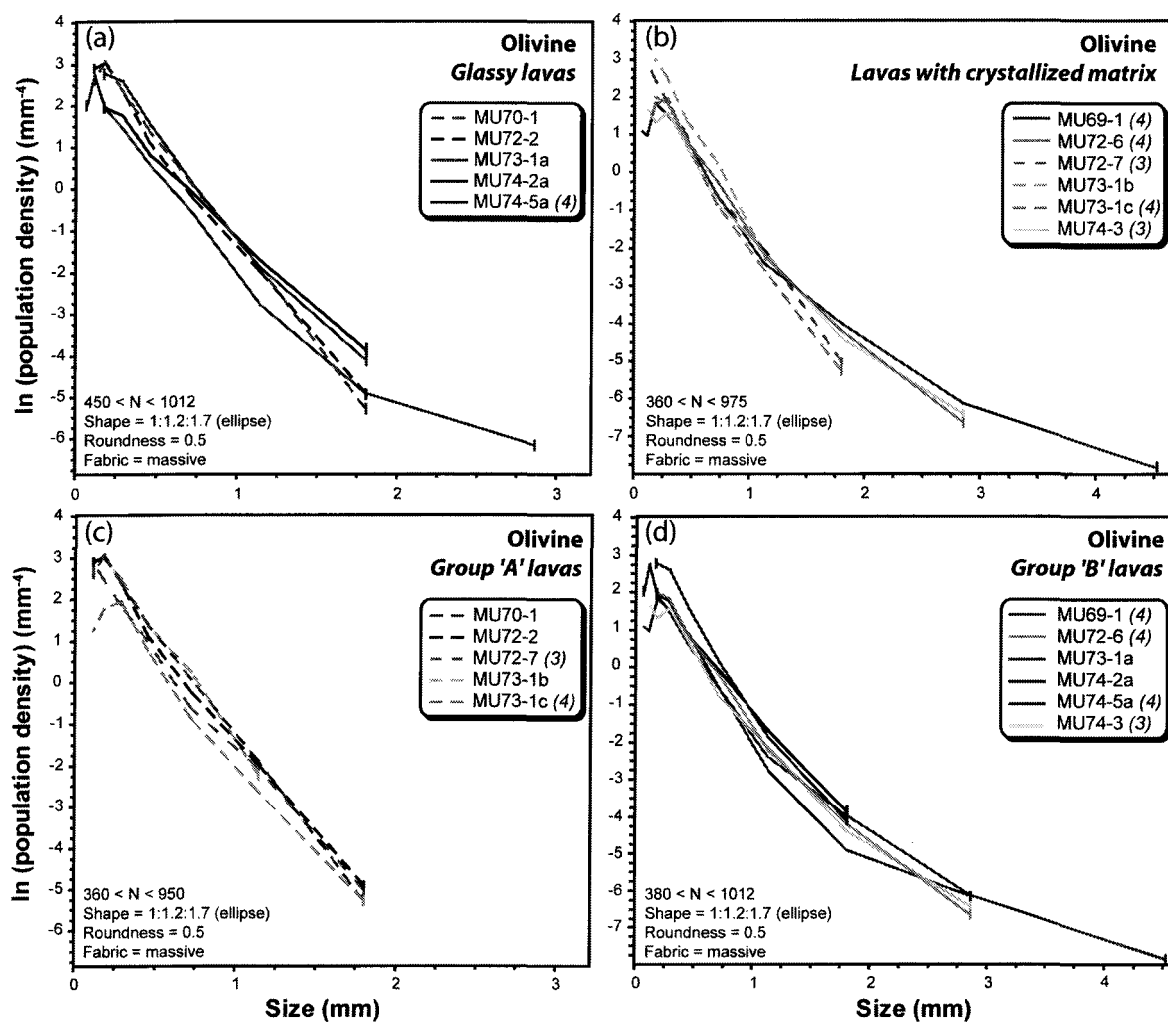


Figure 1-15. Olivine CSDs of Mauna Ulu lavas plotted separately as follows: (a) glassy samples, (b) samples with microlites, (c) group 'A' samples, (d) group 'B' samples. Numbers in parentheses next to some sample names are the number of crystals counted in the CSD analysis for the last bin.

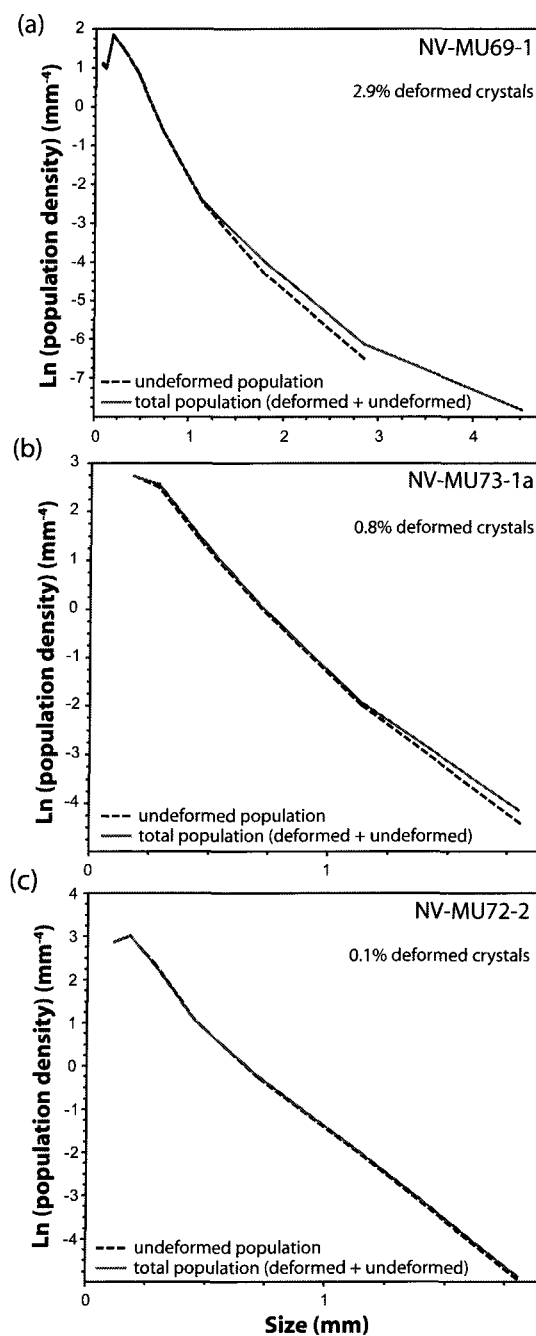


Figure 1-16. Effects of the presence of deformed crystals on the crystal size distribution for (a) NV-MU69-1 that displays significant differences in the CSD of undeformed versus total olivine populations, (b) NV-MU73-1a that shows very little CSD changes, and (c) NV-MU72-2 that has unchanged CSD because of the lack of deformed crystals. Quantitative estimate of the proportion of deformed crystals in each sample listed at the top right of the plot.

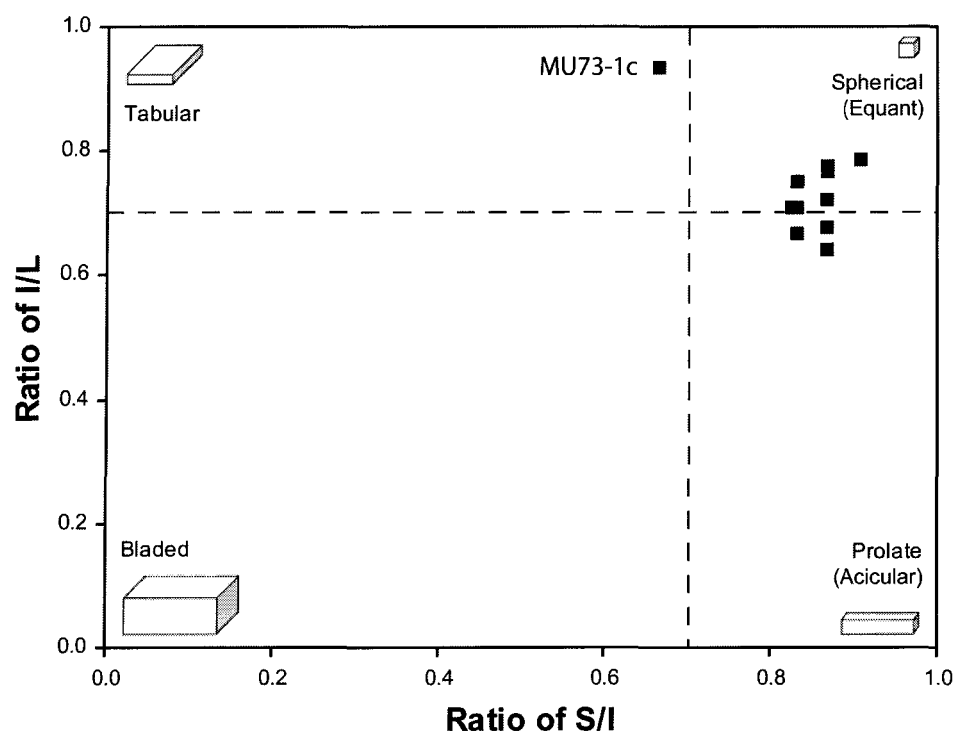


Figure 1-17. Plot of I/L versus S/I ratios for Mauna Ulu lavas, dimensions as calculated by the program *CSDSlice* (Morgan & Jerram, 2006). There is a slight morphological difference in olivine crystal shape between each individual sample. Notice the peculiarity of MU73-1c in exhibiting the highest I/L and lowest S/I ratios.

1.6 DISCUSSION

The MU samples produced during this multiple, prolonged rift-zone eruption, have significantly different chemical compositions and CSDs. These differences have been investigated in terms of the magmatic processes and plumbing, mostly from depth to the surface. The combined chemical and CSD characteristics are considered to document the origin, fractionation, accumulation and deformation of the different MU olivine populations. Our data also lead us to discuss textural coarsening, crystal-liquid and liquid-liquid mixing during magma storage or movement through the plumbing system.

1.6.1 Origin of Mauna Ulu olivine: mantle, crust or both?

To build a coherent story through the plumbing of Kilauea Volcano, the question of the mantle or crustal origin of the MU olivine populations must be addressed. The deformed olivine crystals observed in all MU lavas clearly have a different history from the undeformed crystals, even though they may have the same origin. It is commonly thought that the deformed crystals in Hawaii are unlikely to represent mantle xenocrysts or unrelated cumulates from different volcanoes (Baker *et al.*, 1996; Garcia, 1996; Norman & Garcia, 1999; Wilkinson & Hensel, 1988). This idea is consistent with:

- (1) the magmatic origin deciphered from the fairly high CaO values, mostly 0.20-0.35 wt. %, of all our olivines that may have formed in the crust (e.g., Norman & Garcia, 1999; Simkin & Smith, 1970; Thompson & Gibson, 2000);

- (2) the persistence of melt and Cr-spinel inclusions in both deformed and undeformed crystals;
- (3) the absence of compositional differences between deformed and undeformed olivines, even in minor components such as CaO, NiO and MnO, that suggests that the deformed olivines have a cognate origin with the undeformed ones, and are unrelated to xenocrysts;
- (4) the presence of numerous zoned grains, which is also indicative of a magmatic origin, since mantle xenocrysts would be expected to have homogeneous compositions, resulting from long equilibration times;
- (5) the large range of Fo, even in the lower-Ca olivines, which is uncommon for mantle xenocrysts and would require sampling from very different peridotite lithologies;
- (6) the occurrence of deformed olivines with Fo content less than 84%, thus lying outside the compositional range of mantle olivine (Kamenetsky *et al.*, 2001; Thompson & Gibson, 2000).

Sakyi *et al.* (2008) studied the dislocation microstructures in olivine grains from Kilauea. They found high densities of dislocations ($>10^6 \text{ cm}^{-2}$) and suggest that the olivines have experienced high-temperature plastic deformation. This interpretation is in agreement with the kink-banded olivines observed in our samples, given that kink-bands normally form upon migration of dislocations during thermal restoration at high temperature. The appearance of those kink-bands attests to a strong deformation regime (Nicolas & Poirier, 1976), in the solid state (e.g., Green II & Radcliffe, 1972).

The deformed olivine crystals from MU lavas can be considered as 'antecrysts', in the sense recently proposed by Jerram & Martin (2008): the term 'antecryst' is "becoming the accepted term used to define crystals that have been 'reincorporated' into the final magma", and are "directly

associated with the active magmatic system". These antecrysts are interpreted here to have started as cumulates from earlier Kilauea magmas, which were subsequently disrupted and incorporated into the later magmas. Thus, all the MU olivine crystals must have crystallized from magma within the Kilauea plumbing system. Undeformed and deformed olivines may have the same chemical provenance, coming from the same region of the plumbing system. However, our chemical and CSD data are not sufficient here to prove this shared origin.

Similarly, we propose that the total olivine population of the MU lavas is composed of an early cumulate component represented by the deformed grains, and a later phenocrystic component that crystallized directly from the latest magma(s) of the system. Both of those components are derived from the whole Kilauea magma system, and no mantle component is involved.

The composition of the undeformed olivines indicates that they have several different origins, based on major and minor element analyses. In terms of Fe-Mg systematics, many crystals appear to have been in equilibrium with their current host magma, whereas others are clearly out of equilibrium. We propose that the undeformed, high-Fo olivines (Fo_{88-90}) from MU lavas are relics from a less evolved, earlier magma. The high NiO content of the olivines (0.3-0.5 wt. %) at Fo_{88-90} is reliable evidence here, being strongly compatible, thus strongly resistant to later or even solid state changes, unlike Fe and Mg. The positive correlation between Ni and the amount of olivine in the rocks is in agreement with the compatible character of Ni in olivine. These olivine compositions suggest a parental magma with Mg-numbers $>69 \pm 1$ which is equivalent to >16 wt % MgO (Clague *et al.*, 1995; Garcia *et al.*, 2003; Garcia *et al.*, 2000; Maaloe *et al.*, 1988). This magma could have been similar in composition to MU69-1. Olivines that have compositions that fall below the equilibrium band must have crystallized from more evolved magmas. We will discuss later the

possibility that such magma was mixed with the main MU magma. It is now time to discuss the occurrence and influence of magma processes during solidification within the Kilauea plumbing system under Mauna Ulu.

1.6.2 Where and when did the crystallisation occur?

Understanding the location and the timing of crystallisation that produced the MU lavas is of great importance in the refining of a general plumbing model for Kilauea. Crystallisation of the MU lavas appears to occur at different times and locations or depths in Kilauea's magma system, as indicated by a significant proportion of deformed olivine crystals and their wide compositional range.

No accurate estimation of pressure, and hence depth of crystallisation, can be inferred from crystal and glass geochemistry. However, it is commonly accepted that most olivine fractionation and accumulation occurs within shallow magma reservoirs and conduits beneath the summit and upper parts of the East Rift Zone (e.g. Clague & Denlinger, 1994; Garcia *et al.*, 2003; Pietruszka & Garcia, 1999; Wright & Klein, 2006, 2008). Thus, as proposed in the literature and repeated here, the olivine crystallisation and growth are assumed to have mostly occurred within Kilauea's main conduit and summit reservoir at 3-12 km depth, and subsequently en-route to and within secondary shallow rift chambers as under MU region at a depth of about 3-4 km (see reviews by Wright & Klein, 2006, 2008).

The chemical and CSD characteristics of the MU lavas studied here suggest that the currently deformed olivine could have crystallized at different locations and then settled within the Kilauea edifice, forming a cumulate body deep in the edifice (depth of ca. 9-12 km), which was

subsequently tapped by ascending magma. Macroscopic (i.e. thin section) evidence of mingling within the matrix of some samples that have significantly curved CSDs (MU72-6, MU74-3 and MU74-5a), is consistent with the presence of several olivine populations of distinct origins or magma histories, and hence some sort of mixing. Mixing is also supported by the presence of resorbed and reversely zoned olivine crystals. In contrast, the presence of small euhedral, unzoned and undeformed crystals in equilibrium with their host melt (Table 1-3) is consistent with later crystallisation in the upper parts of the volcano, either in the summit magma chamber or in the conduits and/or adjacent magma pockets under the upper East Rift Zone, 3-4 km deep. Hence, crystallisation may have occurred at least at two different time periods and maybe at different levels too within the Kilauea edifice. The first took place early and participated in the formation of a cumulate body that was partially incorporated into later magma; the second was late and shallow in summit and/or rift storage regions. A mixing of the two distinct olivine populations took place before the eruption.

Magma residence times can be estimated from CSDs if a CSD model is chosen, such as that of Marsh (1988), and a growth rate defined. Here we used a growth rate range of 2.5×10^{-9} mm/s (G_1) to 5×10^{-10} mm/s (G_2) based on a survey of studies on similar rocks (Cashman, 1993, Mangan, 1990, Marsh, 1988, Marsh, 1998, Wright & Fiske, 1971). These two end-member values give an idea of uncertainty on the growth rate, and thus on timescale. The samples from group 'A' have mean crystal residence time ranging from 1.5 to 3.0 years when using G_1 , and from 7.4 to 15.1 years when using G_2 (Table 1-6). Calculated residence times for B1 samples show narrower variations and lie in the same range as the 'A' sample values ($\tau_1 = 1.7$ -2.6 years with G_1 ; $\tau_2 = 8.5$ -12.8 years with G_2). B2 samples have magma residence times that are significantly longer with values ranging from 3.2 to 7.9 years using G_1 in the calculations, and from 16.1 to 39.4 years using

G₂. Given that the population B2 is thought to have undergone postcumulus textural and chemical equilibration processes in the crystal mush, the calculated residence times for B2 are considered here as minimum residence times. In summary, the shorter times of 1.5-15 years estimated from 'A' and B1, are thought to represent the residence times of the populations of olivines crystallized within the shallow plumbing system of Kilauea, while B2 sample ages of 3-40 years reflect minimum residence times of olivine in the deep-seated cumulate body at the base of the edifice, 9-12 km deep. These residence times accord with that determined by Mangan (1990) from olivine CSDs. She found an average olivine residence time of $\sim 11 \pm 4$ years for the 1959 Kilauea Iki eruption, with a growth rate of 10^{-9} mm/s. However, we would like to compare our results with independent methods.

There are few published residence times for magmas erupted from Kilauea's rift zones. Pu'u 'O'o lavas were estimated to be 7-14 years old based on temporal changes in geochemistry (Garcia *et al.*, 1996). $^{226}\text{Ra}/^{230}\text{Th}$ disequilibria applied to the early stage of the 1955 Kilauea east rift eruption gave a mean crystal residence time of ~ 550 years (Cooper *et al.*, 2001). In contrast, Pietruszka & Garcia (1999) proposed an estimate of ~ 30 -40 years of storage in the summit reservoir for Kilauea's late 20th century summit lavas, based on the volcano's magma supply rates and the temporal geochemical fluctuations of the Nb/Y ratios. Furthermore, based on the coherence of the geochemical variation for Kilauea summit lavas, Garcia *et al.* (2003) proposed a relatively rapid, <10 years, period of mixing of new magma batches with the summit magma in the Kilauea reservoir. Based on geodetic data, Wright (1984) estimated ~ 100 -120 years for the same lavas. Average residence times of ~ 100 -3000 years have also been proposed for the entire magma reservoir system beneath Kilauea, based on geophysical estimates of reservoir size and magma

supply rate (Decker, 1987; Denlinger, 1997). The latter values may not be strictly relevant here, as they refer to magma transfer and not crystallisation.

In summary, the magma residence times of MU lavas determined here broadly accord with previous values estimated from Kilauea lavas, although they range widely between the authors and the approaches employed. This is especially true for the minimum residence times calculated for B2 samples. 'A' and B1 samples are significantly lower. This conclusion supports our results, even for the shortest residence time.

1.6.3 Addition and deformation of olivine

The occurrence of multiple populations of olivine is demonstrated by the common presence of deformed olivine crystals in our MU lavas. The whole-rock and olivine chemistry is indicative of olivine that was originally in equilibrium with different liquids, which also supports the idea of multiple populations. Here, the addition of olivine to host magma was treated as equivalent, as a first approximation, to accumulation.

Accumulation (and fractionation) influences overall CSD shapes. Both analytical and stochastic approaches have demonstrated that during crystal accumulation larger crystals sink faster than smaller crystals (Higgins, 2002b; Higgins, 2006a; Marsh, 1998). The resulting CSD slopes are gentler for the magma with accumulated crystals and steeper for the complementary fractionated magma, which retains proportionately more of the small crystals (Fig. 1-10c). At greater sizes, CSDs of population B2 has much gentler CSD slopes than either population B1 at smaller sizes or all lavas of group 'A'. This is interpreted as evidence of olivine accumulation, or more generally, addition, for population B2. The presence of steeper CSDs for MU73-1b and

MU73-1c compared to the flatter one of MU73-1a also suggests accumulation, or addition. But to discount other possible explanations like textural coarsening or mixing, we need to examine the chemical data for evidence of olivine addition.

We have observed chemical evidence for olivine addition in the MU samples: there is a slightly scattered, positive linear trend of whole-rock MgO content with respect to olivine abundance (Fig. 1-18) and the Fo content in a large number of olivines is low for the bulk MgO composition of the host (Fig. 1-7) (Garcia *et al.*, 2003; Marske *et al.*, 2008). Olivine addition is also characterized by a linear array of the whole-rock data on plot of Ni versus MgO (Fig. 1-18b) (Clague *et al.*, 1995; Hart & Davis, 1978). We estimated the composition of the added olivines to be Fo₈₇, the mean value from the MU deformed olivine data of this study. It also corresponds to the mean composition determined from the intersection of the regression lines through the whole-rock and deformed olivine compositions in a Ni-MgO plot (Fig. 1-19). This value is consistent with those presented in previous studies on Kilauea lavas (Clague *et al.*, 1995; 1959 Kilauea Iki olivines, Murata & Richter, 1966; summit lava olivines, Wright, 1971). Nevertheless, we cannot determine on chemical grounds the amount of olivine in disequilibrium with the host, and the amount of added olivine. To do so, we would have to have had prior knowledge of the exact parental magma composition and thus the amount of phenocrysts that fractionated from it. In the present case, we lack the necessary data.

The population of deformed olivine is undoubtedly derived from cumulates originated within Kilauea edifice. But, where and when did cumulate formation and subsequent deformation take place? Was it within the deep crustal cumulate region only, in the narrow conduits during transport to the surface, in the summit magma chamber, or in shallow reservoirs along the rift zone? One may also reasonably invoke a combination of all or part of these possibilities. Helz

(1987b) proposed that the deformed olivines of the 1959 Kilauea Iki eruption were derived originally from conduit walls, lined by deep crustal cumulates from earlier Kilauea tholeiites. Wilkinson & Hensel (1988) proposed that the deformed olivines may be cognate with their hosts and only reflect phenocryst deformation during magma flow through narrow feeders and conduits, at elevated (but unknown) pressures. Others argued in favour of a dunitic cumulate body formed at the base of, and partly under the shallow reservoirs, at a depth of 4-10 km within the edifice (Clague & Denlinger, 1994; Okubo *et al.*, 1997; Wright & Klein, 2006; Wyss *et al.*, 2001). The deformation of the olivine antecrysts may be a response to inelastic strain (Clague & Denlinger, 1994) caused by the seaward plastic flow of the hot and ductile dunite body beneath the south flank of Kilauea. This possibility is also mentioned by Peltier *et al.* (2009) to explain the formation of the dislocation lamellae in olivine crystals of Piton de la Fournaise volcano, La Réunion island. They proposed that the lamellae could be generated by “solid-state gravitational creep and slumping towards the unsupported eastern flank of the volcano edifice”. They argued in favour of dunitic complexes that may underlie the edifice, as well. In their model, deformed olivine antecrysts were derived from the walls of the summit magma chamber or from an olivine-enriched layer stored at the base of the reservoir. We propose that the deformed olivine found in the MU lavas formed in a similar way by volcano spreading.

Deformation affects all crystal sizes and more or less all final crystal shapes. In addition, crystals of clinopyroxene offer evidence of deformation. However, we lack the necessary data to constrain the strength and the P-T conditions of deformation. Consequently, deformation may have occurred at several places within the edifice, both in the various cumulates, shallow or deeper, and in conduits, with the cumulate-phase of deformation being more important. But what happened during and after the formation of cumulates, in terms of magma processes?

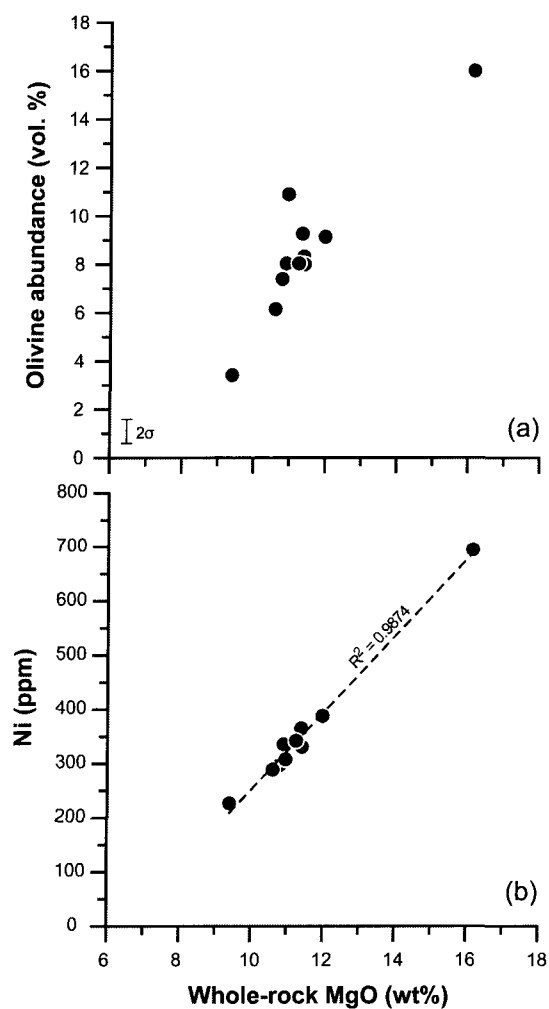


Figure 1-18. (a) Plot of olivine volumetric abundance versus whole-rock MgO content for the Mauna Ulu (MU) lavas. The overall positive trend of the bulk MU lavas likely indicates addition of olivine.

(b) Variations of Ni against MgO contents for Mauna Ulu lavas. The straight dashed line with the R-squared value is indicative of the positive linear correlation between Ni and MgO contents for the original MU lavas. This is consistent with addition of olivine. See details in the text.

The 2σ errors are smaller than symbol size, when not indicated.

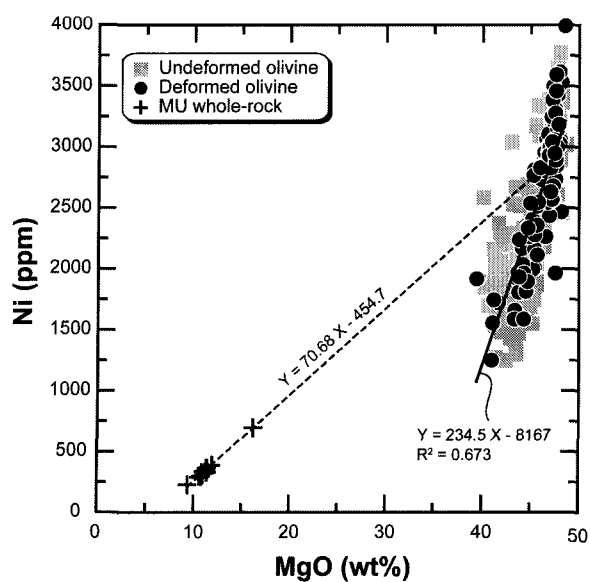


Figure 1-19. Plot of Ni versus MgO for Mauna Ulu (MU) lavas and both undeformed and deformed olivines. Linear regression lines along with their equation are shown for both whole-rock analyses (dashed line) and deformed olivine in the lavas (simple line). The intersection of those two regression lines yields the composition of the added olivine.

1.6.4 Aggregation and textural coarsening

Both aggregation and coarsening are closely related processes of importance in the formation of cumulates. Aggregation is a general term describing the clustering of grains in a melt. The concept of clusters of crystals (glomeroporphyritic texture) that originate in magmas by aggregation and attachment of individual grains has been introduced by many authors in different contexts (Vance, 1969). In some cases crystals appear to be attached along crystallographically similar faces, which assures that structural mismatch and interfacial energy are minimized. The term *synneusis* – the swimming together of crystals (Vogt, 1921) – is often used both for this texture and the associated mechanism. In contrast, it has also been proposed that growth is enhanced by annexation of small grains by larger crystals, followed by grain boundary migration (Marsh, 1998). This process can produce optically continuous minerals, and is considered by Marsh (1998) as a type of rapid coarsening process.

Textural coarsening (also known as Ostwald ripening, textural equilibration or maturation) is proposed as an important process in plutonic (e.g. Higgins, 1998, 2002b; Higgins, 2006a) and some volcanic rocks (e.g. Higgins & Roberge, 2003). In an igneous context, coarsening is the resorption of small crystals and the simultaneous growth of larger ones as a process of total surface (free) energy minimization (see review by Voorhees, 1992). This process is most important at temperature close to the mineral liquidus. Although no consensus exists about the exact process of textural coarsening, we used the Communicating Neighbours (CN) theory (Dehoff, 1991) instead of the more commonly used Lifshitz-Slyozov-Wagner (LSW) theory (Lifshitz & Slyozov, 1961; Voorhees, 1992), because it appears to be a better fit for igneous rocks (e.g. Higgins, 1998; Higgins

& Roberge, 2003, 2007). Recently, Simakin & Bindeman (2008) showed that coarsening can be driven more effectively by thermal oscillations in the magma.

Aggregation and textural coarsening are magma processes that can be recognized and quantified only by textural approaches, such as CSD analysis. Figure 1-10d shows the theoretical CSD evolution of magmas undergoing CN type coarsening. The turndown of the CSD curve at smallest sizes represents the disappearance of the small crystals, whereas the CSD flattening at largest sizes is indicative of the increase in growth rate to a maximum with grain size (Higgins, 1998). Thus, both the slope and intercept decrease during this process. In a closed system containing minerals of uniform composition, the total chemical composition is unaffected by coarsening.

Both coarsening and aggregation can lead to the loss of smaller crystals and gain of larger crystals. This occurs during coarsening by solution of smaller crystals and during aggregation by annexation and equilibration. Hence, the effects of aggregation on CSDs are extremely similar to textural coarsening, with maybe a lesser influence on smaller grains, and hence a smaller, or no turndown at smallest sizes. The similarity of effects on CSDs of these two magma processes is the reason why they are treated together. In both cases the slope of the CSDs becomes shallower. However, coarsening also produces a turndown for the smallest crystals, as is observed in many samples.

Group 'A' samples have steep CSDs, with a low maximum crystal size ($L_{max} < 1.5$ mm) and mostly with turndowns for the smallest crystals. Such CSDs preserve the initial CSDs produced by nucleation and growth slightly modified by coarsening, which has removed the smallest crystals. The CSDs do not record evidence for aggregation.

Group 'B' samples have strongly curved CSDs and extend out to greater maximum crystal size ($L_{max} > 2.3$ mm, except for MU73-1a and MU74-2a), again mostly with turndowns for the smallest crystals. The most extreme behaviour is that of sample MU69-1, which has amongst the lowest number of small crystals and extends out to the greatest maximum size. The gentler slopes at larger size fractions and turndowns are probably the result of coarsening, which is most developed in MU69-1. In these samples the curvature and rotation of the CSDs are signs of aggregation, as well. However, the additional effects of aggregation on the CSD are difficult to distinguish from the coarsening effects, since they are superimposed.

In other volcanoes, the early magmas in an eruption sequence are also the most coarsened (Tseax volcano, Canada, Higgins, 2009; Eldfell volcano, Iceland, Higgins & Roberge, 2007). This may occur during storage just before the eruption.

In all MU samples studied here aggregation textures (Fig. 1-2) are observed in varying but high proportions of crystals (~20-50%), with the glomerocrysts being composed of both deformed and undeformed crystals. We interpret some of these as formed by synneusis (Fig. 1-2a). The bulk driving force for aggregation, including synneusis, is the minimization of the surface energy of the crystal aggregate (Ikeda *et al.*, 2002). The synneusis aggregation mechanism typically occurs during crystal settling and accumulation in liquid-rich magma rather than a mush (Schwindinger & Anderson, 1989). Aggregation can also take place during cumulate formation via crystal settling. Unfortunately, it is unclear which process dominates.

1.6.5 Disruption of the cumulate and incorporation into the magma

The occurrence of deformed olivine crystals in all the MU lavas is consistent with a deep-seated cumulate origin, but how did this body become disrupted and how did parts of it enter the MU magmas? The cumulate olivines are mostly observed as individual crystals or small clusters of 2-4 grains on average. Even very small single crystals may be deformed, suggesting wholesale complete incorporation. Only one large dunitic xenolith has been recognized. Further, the CSDs of the bulk populations very often closely match those from the undeformed population (Figs. 1-16 and A1). Real divisions between both only appear at larger sizes, suggesting that though there may be small deformed crystals, the larger ones have a greater effect in the CSD plot shape. However, in the CSDs shown in Appendix A, even when there is a difference in the CSDs for a given sample, they still have approximately the same shape. NV-MU74-2a and NV-MU74-5a are good examples of that. This shape similarity could be seen as good evidence for early disruption and incorporation of cumulate (i.e., deformed olivines), which then underwent other subsequent magma processes that affected effectively a single population.

How did the cumulate body become so efficiently disrupted, that the deformed crystals became incorporated more or less individually into the magma? Clague & Denlinger (1994) argued that the production of large volumes of Hawaiian tholeiitic basalt coincides with the formation of olivine cumulates at magmatic temperatures ($>1100^{\circ}\text{C}$) containing a small amount of intercumulus liquid. This would prevent adjacent olivine crystals from adhering to one another. In turn it would also explain the efficiency of the disintegration of the cumulate when entrained by the magma. However, no precise estimate of the amount of liquid trapped in Kilauea cumulates is available in the literature, and no data in the present study can help to resolve this issue.

The incorporation of the disrupted cumulate olivines into new magma, which already must have contained olivine phenocrysts, is a case of crystal-magma mixing. This bulk magma process can be easily recognized using CSD analysis. The process could be chemically comparable to accumulation, because early, coarsened olivine crystals were added to late, less or not-coarsened magmas en-route to the surface. As a first approximation we can estimate this mixing to be in the minimum range of 0.1 to 3%, based on the volume fraction of deformed crystals measured in the rocks (Table 1-2).

The curvature of CSDs can be used as evidence for mixing: the mixing of two magmas with non-collinear straight CSDs produces a curved concave-up CSD (Higgins, 1996) (Fig. 1-10e). This is due to the logarithmic vertical scale on a classic CSD diagram. The global CSD consists of straight segments both at small and great sizes parallel to the original, individual CSDs and linked by a curved concave-up segment at intermediate sizes. The degree of curvature is dependent on the difference in original CSD slopes between the two magmas. On simple, linear, bivariate geochemical diagrams mixing will produce a compositional variation fitting a straight line, whereas fractionation will be indicated by a curved line. However, it is sometimes difficult to distinguish mixing from the curved patterns produced by fractional crystallisation, especially if there is little compositional variation.

Texturally, group 'A' samples mostly have straight CSDs and hence show no evidence of mixing. Most samples from group 'B' have moderately to strongly curved concave-up CSDs reflecting a multistage CSD history compatible with mixing. Furthermore, the CSDs of population B1 are parallel to the CSDs of group 'A', with slight displacement downwards. Thus, the magma inferred from group 'A' and population B1 is likely one of the magmatic components, that is, the late, less coarsened phenocrystic component. The straight CSDs have steep slopes and high

intercepts probably produced during the early stages of nucleation and growth of this phenocrystic population. The flatter CSDs of population B2 are thought to represent another mixing component, that is, the early, more coarsened cumulate component. This is confirmed by macroscopic evidence of mingling – that is abrupt crystallinity changes on the scale of centimetres – within the matrix of most lavas with curved CSDs from group ‘B’.

In summary, at least two distinct olivine populations are inferred from the textural and chemical analyses: (1) an antecrystic olivine population from a dunitic cumulate formed at greater depth (mostly at ~9-12 km deep, within the volcanic pile of the edifice) and then remobilized by the ascent of a juvenile magma, and (2) a phenocrystic olivine population that crystallized from the juvenile magma at shallower depth either in the summit reservoir or in the magma chamber(s) under Mauna Ulu area, or even both, and in the conduits. This type of solid-liquid mixing is mainly related to entrainment of cumulate olivine crystals by an ascending magma.

1.6.6 Movement to the summit and rift zones – Multiple magma mixing events?

Multiple parental magmas are common in many igneous systems. Therefore mixing is an important crustal process that modifies the bulk composition of lavas; as is the case for the Kilauea lavas (e.g. Garcia *et al.*, 2003). The term mixing is employed when the product is homogeneous. When the product still contains heterogeneities, as that described earlier, it is called mingling. We use the expression ‘multiple magmas’ if the magma components share a conduit.

Movement of the magma(s) through the crust to the surface appears to be complex, but several key points can be highlighted here, in relation to the prolonged MU 1969-1974 eruption. We propose that components of the MU magmas followed two different routes: one rose via the

main conduit to the summit reservoir and then moved out along the rift zone. The other moved along a deep low-angle decollement that lies underneath the volcanic shield, then rose under MU ridge, where the magmas mixed. Evidence for this decollement is provided by both the spatial distribution of hypocenters and high-resolution seismic imaging (Thurber *et al.*, 1989). The decollement plane has long been inferred to be of major importance in the bulk deformation of the edifice, i.e. seaward spreading of Kilauea's south flank (Borgia, 1994; Crosson & Endo, 1981; Delaney *et al.*, 1998; Delaney *et al.*, 1990; Dieterich, 1988; Lipman *et al.*, 1985; Okubo *et al.*, 1997; Owen *et al.*, 1995; Ryan, 1988; Wright & Klein, 2006; Wyss *et al.*, 2001), but never considered as a possible pathway for magma transfer, as we propose here. Yang *et al.* (1999) suggested that the oceanic crust-mantle boundary at ~15 km depth beneath the rift zones, was a possible transfer zone for magma to the rift zone, based on crystallisation pressures of 4-5 kbar inferred from augite and glass rind compositions. However, they did not mention the decollement and provide no further arguments for their idea. Details to support our suggested route along the decollement will follow.

The summit component is now mostly represented by the undeformed olivine phenocrysts, but some of the cumulates may also have formed here, on the conduit or chamber walls, and thus some antecrysts belong to this summit component, as well. The range in olivine composition suggests that these crystals grew from a range of different magmas, probably related by crystal fractionation. Such a process could occur principally in the main conduit and summit reservoir, but also in reservoirs in the upper rift system (e.g., Garcia *et al.*, 2003; Thornber, 2001). Here injections of new, undifferentiated magma could yield the Fo-rich crystals whereas Fo-poor crystals would have crystallized from more differentiated magmas. The ubiquitous presence of deformed crystals shows that a component must have originated by disruption of a deformed cumulate. It is

proposed that most antecryst-bearing magma transfer was along the basal decollement and then rose into the rift zone through near-vertical conduits. Transport through near-vertically dykes in the deep rift system to MU shallow region was hypothesized by several authors (e.g. Clague & Denlinger, 1994; Clague *et al.*, 1995; Delaney *et al.*, 1990; Dieterich, 1988; Ryan, 1988; Ryan *et al.*, 1981; Wright & Klein, 2006). In contrast to the seismically active shallow part of the rift zone (2-4 km), the deep part of the rift system is aseismic, maybe in part due to its partially molten nature (e.g. Clague & Denlinger, 1994; Ryan, 1988). Possibly partly made of a mush of picritic melt and olivine crystals (Ryan, 1988), it corresponds to a deforming, or dilating, zone of either a series of magmatic cumulates, an interconnected magmatic plumbing system, or a combination of both, extending down to the decollement, at ~9-12 km (e.g. Owen *et al.*, 1995).

The ratio of antecrysts to phenocrysts in a lava can be used to determine as a first approximation the proportion of the two components. The lava from the first year of eruption (MU69-1) contained 18% antecrysts among the total olivine population; in terms of bulk abundance relative to the whole section, it corresponds to 3% antecrysts in the sample. However, this ratio of antecrysts to phenocrysts dropped down to 3% for the 1970 lava, and then rose to 6%, 8% and finally 20% during the period 1970-1974. This suggests that the eruption started in 1969 with a large component of magma that had passed through the decollement and bypassed the summit reservoir. This source then diminished and was replaced by magma that flowed via the summit to the rift. With time the first magma source again became important.

The locations of the seismic hypocenters associated with the different phases of the eruption (Klein *et al.*, 1987, Fig. 43.98) are consistent with our model. During the period 1970-1972, the eruption initiation hypocenters were concentrated near summit of Kilauea, and propagated to Mauna Ulu (Klein *et al.*, 1987). This correlates well with our observation of low

ratios of antecrysts to phenocrysts for the same eruption time period. This correlation is indicative of magma movement through the rift zone from the summit to MU in 1970-1972; the main conduit and the summit reservoir are thus involved. In contrast, in 1969 and 1973-1974 the initiation points of swarms (hypocenters) were located in the vicinity of Pauahi or MU (Klein *et al.*, 1987), *aside from the rift zone main pathway passing through the summit*. This correlates with our observation of higher ratios of antecrysts to phenocrysts in the 1969 and 1973-1974 lavas. This correlation strongly suggests the importance of an alternative magma transport path from the primary conduit to the MU. Another route must be invoked, which bypassed the summit reservoir.

Transport of magma in pipe-like conduits from the base of the shield, that is the decollement, to the MU ridge has been suggested by some geophysical and numerical studies (Ryan, 1988; Ryan *et al.*, 1981). The pipe-like conduit was “formed in an area of weakness produced by the high-angle intersection of the rift zone with the Koae fracture system” (Ryan, 1988). Ryan (1988) considered that such pipe-like conduits transported magma vertically during the period April 1971 to December 1973. Wyss *et al.* (2001) found a strong b-value seismic anomaly in a structurally significant part of the rift zone, which they interpreted as the Hiiaka-Pauahi magma reservoir, located at the junction of the Koae Fault Zone and the rift eastward bend. The work of Klein *et al.* (1987) revealed that this reservoir was an aseismic zone within the rift, which is compatible with its intrinsic molten nature. Earthquake swarms showed that it was a starting point of intrusions that propagated both up- and down-rift.

It is proposed that the decollement is an ideal existing zone of weakness easily used for transport of magma, *especially when dilation occurred in the deep rift system just above it*, in response to spreading (on the decollement plane) of the south flank away from the rift zone.

Magma then moved up through pipe-like conduits en-route to MU, as they are thought to be relatively open, low-stress dilating pathways.

Unfortunately, there is no geophysical evidence from the MU eruption to suggest magma movement along the decollement. However, the zone from ~4-5 km deep to the decollement under the rift zone is mostly aseismic (Aki & Koyanagi, 1981; Koyanagi *et al.*, 1987; Wright & Klein, 2006). Here, magma can maintain a state of quasi-steady flow that does not generate measurable seismic signals. The decollement itself is not completely aseismic, as shown by location of the 1975 M7.2 Kalapana (e.g. Crosson & Endo, 1981) and other earthquakes.

1.7 MODEL OF MAUNA ULU MAGMA DYNAMICS IN THE CONTEXT OF KILAUEA DYNAMICS

To better understand and schematize how the plumbing of Kilauea Volcano works, in particular at crustal depths under the East Rift Zone, and to decipher what are the possible magma pathways and interconnections, we propose here a model of magma dynamics that focuses on the origin and history of the 1969-1974 MU magmas. Our model builds on earlier work, but is grounded in the textural and chemical data presented in this study that allow several magma solidification processes to be deciphered. Here we present the model as a semi-chronological series of events as illustrated in figure 1-20.

- (1) Olivine crystallized from primitive and fractionated magmas at various depths and locations in Kilauea. The crystals settled, aggregated and formed an olivine cumulate body mostly near the base of the summit reservoir and around the main crustal conduit from 4-5 to 10-12 km depth, and near the base of the volcanic pile beneath the East Rift

Zone at 9-12 km depth (e.g. Clague & Denlinger, 1994; Delaney *et al.*, 1990; Okubo *et al.*, 1997; Ryan, 1988; Wyss *et al.*, 2001). Crystallisation and accumulation ages based on CSD data range from 3 to 40 years. The cumulate never solidified completely and parts contained some intercumulus liquid.

- (2) The cumulate olivine crystals produced in event (1) underwent high-temperature plastic deformation induced by seaward flowing down of the hot (>1100°C), ductile dunite body (e.g. Clague & Denlinger, 1994; Clague *et al.*, 1995; Ryan, 1988; Wright & Klein, 2008).
- (3) Later olivine crystallized in the crustal summit magma reservoir and main conduit under Halema'uma'u Crater during the cooling of magnesian magmas (Wright & Klein, 2008). Crystallisation also occurred beneath the rift zone, at 4-5 km depth (e.g. Clague & Denlinger, 1994). The varying magma compositions were partly a result of crystal fractionation and olivine addition. Crystallisation ages based on CSD data range from 1 to 15 years. Olivine settled rapidly in the low viscosity primitive liquids and underwent synneusis-driven aggregation processes (Clague *et al.*, 1995).
- (4) A decollement fault at the base of the volcanic pile acted as a sub-horizontal conduit for magmas flowing from the base of the main conduit. The host rock comprised deformed olivine crystals, possibly with a small amount of interstitial liquid. The magma partly disrupted the host rock and entrained the deformed olivines, forming the antecrysts (e.g. Clague *et al.*, 1995; Yang *et al.*, 1994). The great variability of magma composition observed in the whole-rock data and olivine mineral chemistry of the MU lavas is mostly a result of this crystal-magma mixing. The magma migrated toward the surface underneath the upper East Rift Zone near Mauna Ulu vertically through secondary deep

pipe-like conduits (Ryan, 1988; Ryan *et al.*, 1981). Hence, one component of the MU magmas bypassed the summit reservoir.

- (5) Magma produced in event (3) was transported horizontally along the rift where it mixed with magma (4) rising vertically from the decollement. This may have occurred in small magma chambers under the Mauna Ulu area prior to eruption. At the start of the eruption in 1969 the deep magma source was important. However, this was almost completely replaced by the summit magma in 1970. During the next four years the amount of magma contributed by the deep source gradually returned to the initial value.

Here, we have proposed a model of magma dynamics under Mauna Ulu that can be applied at a larger scale to Kilauea Volcano, and possibly for other Hawaiian volcanoes. In particular, the conclusions of this study can help to reconstruct the magma dynamics and pathways of other long-lived rift eruptions, such as the on-going Pu'u 'O'o-Kupaianaha eruption.

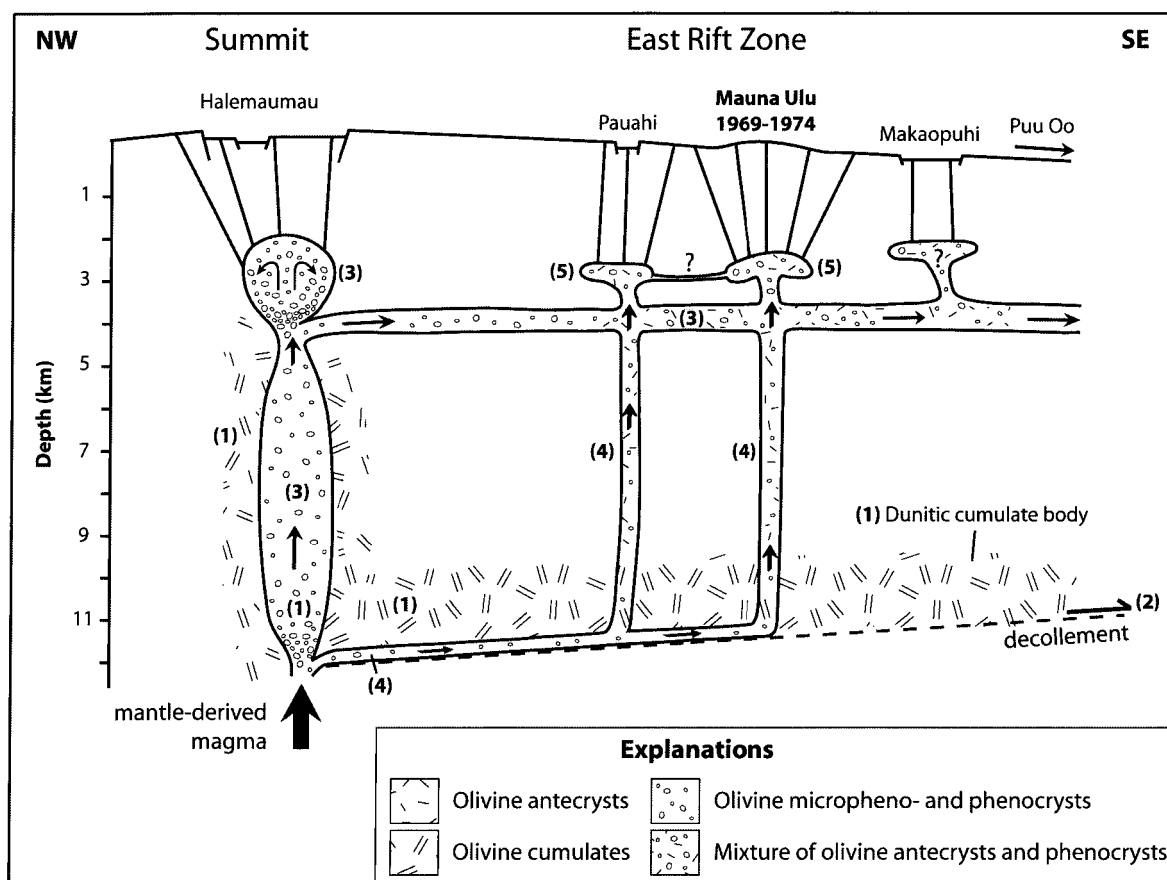


Figure 1-20. Schematic cross-section of Kilauea's magmatic plumbing system underneath the summit and upper East Rift Zone. The black arrows show magma transport through two main pathways. Horizontal movement of magma along the decollement disrupted and mobilised deformed olivines. These magmas were then transported through pipe-like conduits to reservoirs at depths of 3-4 km beneath Pauahi and Mauna Ulu areas. There also was crystallisation in the main conduit and the summit reservoir, possibly associated with olivine cumulates too, and lateral migration of magma through the conduits under the East Rift Zone where it mixed with the magma rising from the decollement. Numbers in parentheses refer to the stages of the 'Model of magma dynamics' proposed in the text. Not to scale horizontally.

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APPENDIX 1-1

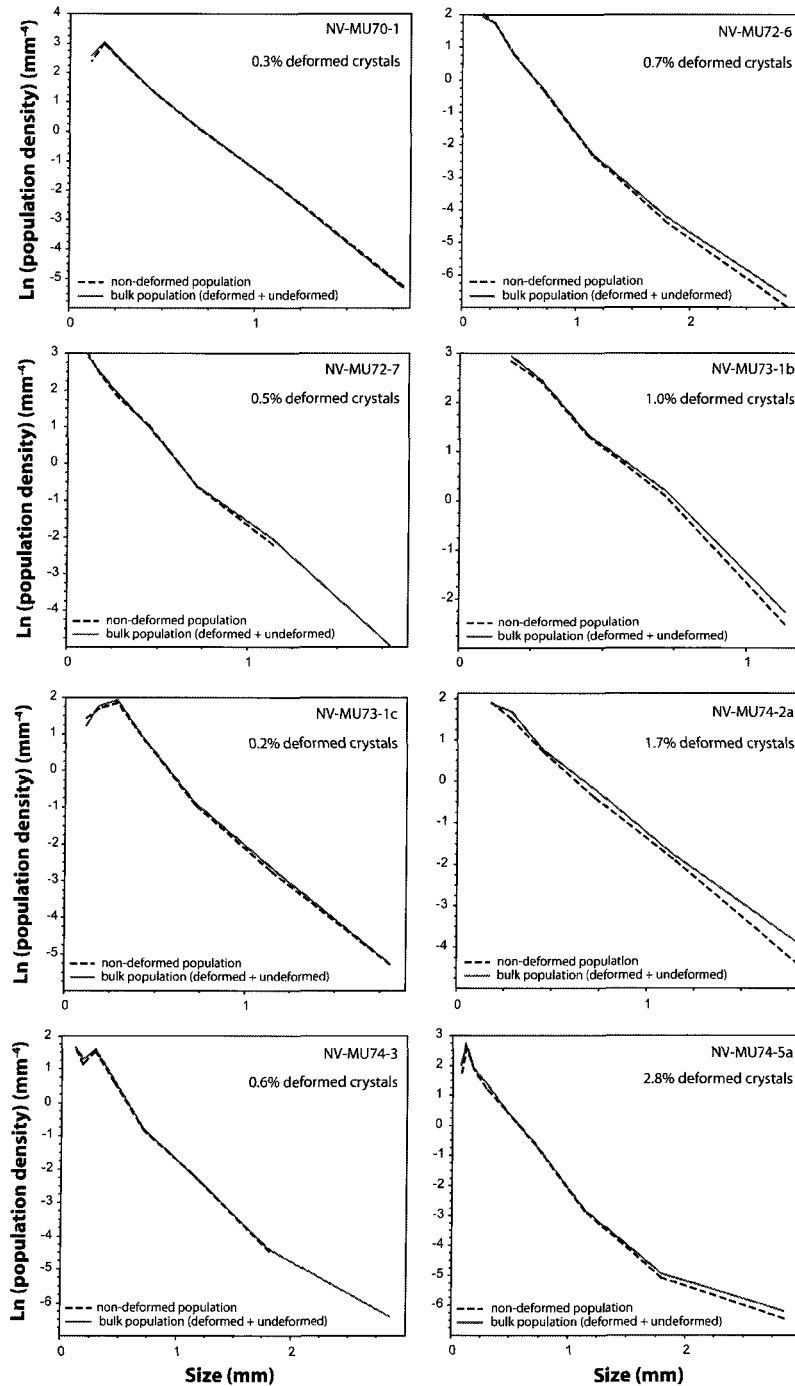


Figure 1-A1. CSD plots of the undeformed olivine population versus the total (deformed + undeformed) population for individual samples from Mauna Ulu, with the exception of the three samples in Figure 1-16 (NV-MU69-1, -MU72-2 and -MU73-1a). Quantitative estimate of the proportion of deformed crystals in each sample listed at the top right of the plot.

CHAPITRE 2

WHAT CAN CRYSTAL SIZE DISTRIBUTIONS AND OLIVINE COMPOSITIONS TELL US ABOUT MAGMA SOLIDIFICATION PROCESSES INSIDE KILAUEA IKI LAVA LAKE, HAWAII?

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2.1 ABSTRACT

Lava lakes offer the opportunity to investigate magma solidification and can be considered as a proxy for small magma chambers. Here we present olivine compositions and crystal size distributions (CSDs) from scoria and drill core samples from Kilauea Iki Lava Lake, which formed during the 1959 eruption of Kilauea Volcano, Hawaii. Three chemically distinct olivine populations were distinguished, on the basis of their forsterite (Fo) content: (1) a high-Fo population (Fo₈₅₋₈₈); (2) an intermediate-Fo population (Fo₇₇₋₈₁); and (3) a minor low-Fo population (Fo₇₂₋₇₆). Populations 1 and 2 both have deformed and undeformed crystals. The third population may be the result of rejuvenation. Olivine in the lower 60 m of lake has a less Fo-rich composition and more crystals are deformed. The CSD analysis yields estimates of the olivine residence time: 1-60 years. The shape of the olivine CSDs is fairly uniform with respect to depth. Curved CSDs are considered to be evidence of crystal or magma mixing. The turndown at smallest sizes of most lake CSDs may be the result of coarsening. Our CSD modeling does not support significant crystal settling and overall convection in the lava lake, although small advective currents are known to have occurred. The olivine vertical stratification could be an original feature. However, this is not consistent with supposed strong stirring of the lake magma due to intense activity over the 17 eruptive phases. Independent basal feeding of the lake during the eruption is another possibility, and may be seen as a likely mechanism to explain the stratification of the lake.

KEY WORDS: *Hawaii; Kilauea Iki; lava lake; olivine; crystal size distribution; solidification processes*

2.2 INTRODUCTION

A fundamental issue in igneous petrology is the understanding of magma solidification, with emphasis on processes at work in magma chambers. Since direct examination of mafic magma chambers is impossible, numerous experimental and numerical modelling studies have been conducted to gain insights into those processes, revealing that convection and crystal settling could play an important role, although it is much debated (e.g. Bowen, 1928; Brandeis and Jaupart, 1986; Davaille and Jaupart, 1993; Gibb and Henderson, 1992; Jarvis and Woods, 1994; Jaupart and Tait, 1995; Marsh, 1988a, 1989, 1996; Martin and Nokes, 1988, 1989; Rudman, 1992; Sparks et al., 1993; Weinstein et al., 1988). Although these studies have enabled valuable progress in the recognition and constraint of igneous mechanisms during solidification, the great complexity of natural environments necessitates the use of natural samples in conjunction with geological data acquired in the field. In this way, since they could be considered as “natural laboratories”, lava lakes offer the opportunity to investigate magma solidification via in-situ analyses and can be studied as a proxy for small magma chambers.

Volcanic products and lava lake samples are commonly studied by quantitative chemical and isotopic analysis. However, it is helpful to complement this approach by the quantitative analysis of rock textures. The texture of igneous rocks is one of the most direct means to investigate the physical and chemical conditions of solidification. Crystal size distributions (CSDs) are the most commonly studied aspect of textures. The CSD method provides valuable kinetic information for examining the link between crystal nucleation and growth, and the overall cooling and magma dynamics (Marsh, 2007). Mafic magmas have been the subject of many laboratory experiments and numerical models, but the understanding of solidification under natural conditions is of great

importance. The CSD approach offers this possibility of investigating solidification in natural dynamic systems, providing clues to both physical and chemical processes that have occurred.

Few CSD studies on natural basaltic (*sensu lato*) samples from lava lakes and sills have been published. The only published study of olivine CSDs in natural basaltic samples is that of Mangan (1990). She worked on scoria samples from the eruptive phases 1, 2, 4, 5, 7, 8, 9 and 16 of the same 1959 eruption of Kilauea Iki studied here. She measured few crystals in each sample, so her data has been amalgamated here into a single CSD. Cashman and Marsh (1988) worked mainly on plagioclase and oxides in tholeiitic basalt from the Makaopuhi lava lake, Kilauea. These early papers were exploratory in nature and time has come for more detailed work with a greater number of samples, combining textural and geochemical approaches. In this paper, our primary aim is to characterize texturally and geochemically the Kilauea Iki olivines in order to understand and quantify the main solidification processes of tholeiitic magmas at work within the lava lake. Our second purpose, closely related to the first one, is to clarify the active magma processes and possible pathways through the plumbing at depth underneath Kilauea Iki, by comparison with the dynamics of Mauna Ulu system.

This paper presents the results of textural and geochemical analyses from 3 eruption scoria samples, 34 drill core samples taken from 1967 to 1988 and spot analyses of over 430 individual olivine crystals. Here we consider the relationships among several early and late processes that are thought to have occurred within Kilauea Iki lava lake. We present new quantitative textural and in-situ geochemical information on the dynamics of these intra-lake processes and propose a model along with temporal constraints that explain the crustal magmatic origin of the distinct olivine populations.

2.3 THE 1959 KILAUEA IKI ERUPTION

Kilauea Iki lava lake formed during the 1959 eruption of Kilauea Volcano, an extensively studied near-summit eruption of picritic lava that lasted from November 14 to December 20 (Helz, 1987a, b; Helz, 2009; Macdonald and Katsura, 1961; Murata and Richter, 1966a; Richter and Eaton, 1960; Richter et al., 1970; Wright, 1973). It consisted of a sequence of 17 distinct eruptive phases, separated by periods of drainback of lava from the lake down the vent. The lava ponded in the pre-existing pit crater just next to the vent, and formed a small, undisturbed, self-roofed, picritic magma chamber (Richter *et al.*, 1970) with a volume of $\sim 38 \times 10^6 \text{ m}^3$ and an estimated mean MgO content of 15.4 wt %, corresponding to about 20 wt % olivine (Wright, 1973). Cycles of overturns of the surface of the lake also occurred during this period. With the exceptions of the first two days and the last eruptive phase, no significant changes in the overall mineralogy of the lavas during the entire eruption were noted (Richter and Murata, 1966). The erupted, tholeiitic olivine basalt lavas contained 4-30 vol. % of phenocryst olivine in a glassy or fine grained groundmass with abundant microlites of clinopyroxene, plagioclase and opaque minerals (Richter and Moore, 1966; Richter and Murata, 1966). Olivine occurred as varying sized crystals of up to 4 mm in diameter. Phenocrysts and microphenocrysts of clinopyroxene and plagioclase were only present in the early and extremely late lavas; hypersthene was absent or very minor.

Since the end of December 1959, when the crustal foundering ceased and the upper crust stabilised, the lava lake has been drilled repeatedly as it cooled and crystallised (Fig. 2-1a). 1960-62 drilling results are presented in detail in Richter and Moore (1966), a review of the results for the 1967-1981 drilling in Helz (1987a), and Helz (1993) for the final 1988 drilling. In 1981, it was possible to drill through the partially molten core and into the lower crust. Consequently, Helz and

Wright (1983) reported that the processes of large-scale internal differentiation occurring in the lake had stopped sometime between the 1979 drilling and 1981. Based on isotherm extrapolation, the lake is thought to have completely solidified in the mid-1990s, representing 35 years of cooling to reach the solidus throughout its maximum total thickness of 120-130 m (Barth *et al.*, 1994; Helz, 2009, and references therein).

In a recent review, Helz (2009, her Table 1) summarised the major differentiation processes within Kilauea Iki lava lake, and distinguished two different groups regarding the temperature range: a group of higher-temperature processes that occurred within the partially molten core, and a group of lower-temperature processes that were involved within the coherent crystal mush zones surrounding the molten core. Some of these processes have been described and discussed in earlier papers (Helz, 1980, 1987a; Helz, 1993; Helz *et al.*, 1989; Jellinek and Kerr, 2001). She also proposed the occurrence of double-layered lateral convection within the most olivine-poor part of the lake, between late 1962 and mid-1964. The variety and extent of all these processes document the complexity of the entire solidification of the lava body.

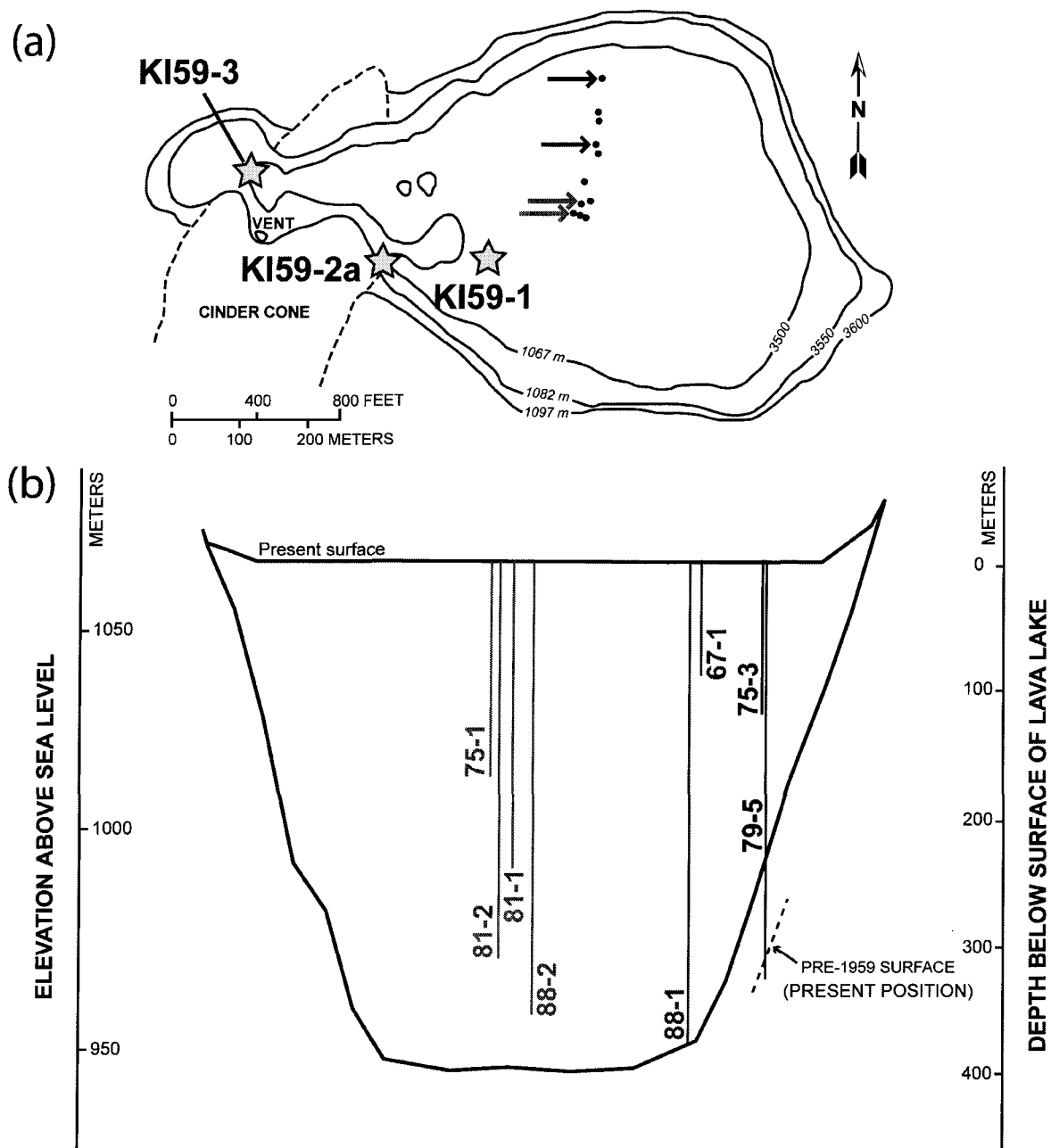


Figure 2-1. a) Plan view of the post-1959 surface of Kilauea Iki (modified after Helz, 1993). The larger dots indicate locations of holes drilled from 1967-1988. b) N-S cross-section of Kilauea Iki lava lake, with projected drill holes shown. Vertical exaggeration is 4:1. Modified after Barth et al. (1994).

2.4 MINERALOGY AND PETROGRAPHY OVERVIEW

2.4.1 Sampling

We sampled three eruptive scoria or spatter samples from the surface of Kilauea Iki lava lake, including one close to the main vent (Fig. 2-1b). All other samples were selected from the drill cores available at the National Museum of Natural History, Smithsonian Institution, Washington D.C. The holes KI67-1 (1967) and KI75-3 (1975) are located close to the northern edge of the lava lake, whereas KI75-1 (1975) lies in the centre of the lake (Fig. 2-1a). We also used thin sections of greater depths in holes KI79-5, KI81-1, KI88-1 and KI88-2. These samples are from the personal collection of R.T. Helz, USGS, and correspond to the drilling years 1979, 1981 and 1988, respectively. KI79-5 was drilled in the same location as KI75-3, KI81-1 as KI75-1 and KI88-1 as KI67-1.

This sampling covers a wide period in the solidification history of the lava lake, extending from 1967 to 1988. This ensures the recognition of a great variety of both early and late magma solidification processes regarding the depth within the solidifying body.

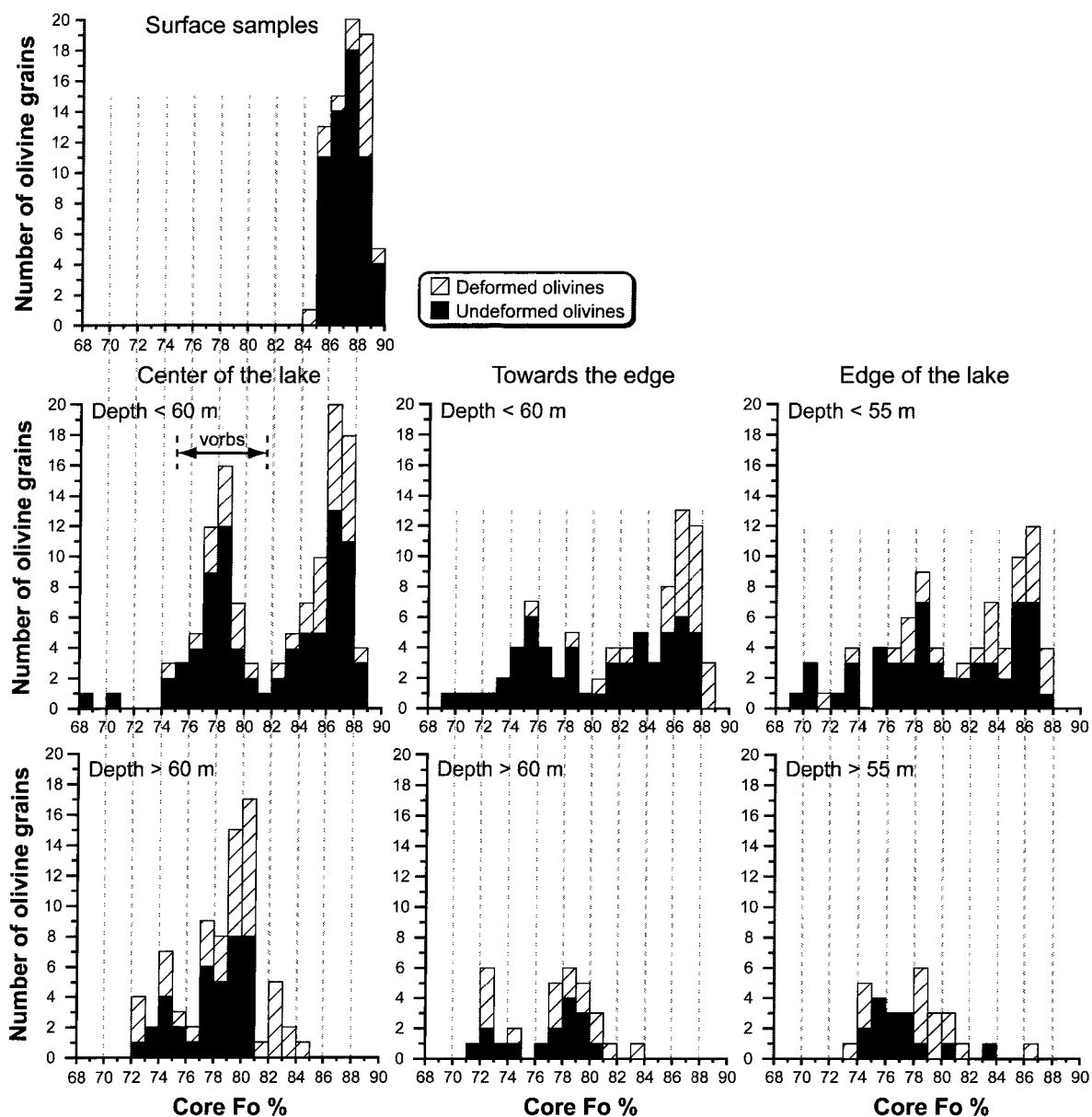


Figure 2-2. Histograms of olivine microphenocryst and phenocryst core compositions (in forsterite %) in lake and surface scoria samples, classified according to depth and location in the lake. Deformed and undeformed crystals are represented in different patterns. Hole numbers are: "Center of the lake", KI75-1, KI81-1, KI81-2 and KI88-2; "Towards the edge", KI67-1 and KI88-1; and "Edge of the lake", KI75-3 and KI79-5.

2.4.2 Macroscopic characteristics

2.4.2.1 *General remarks*

The samples consist of strongly to almost non-vesicular olivine tholeiites to picrites with no visible foliation. They contain 7-39 vol. % olivine in phenocrysts (>0.5 mm across) and microphenocrysts (>0.1 to <0.5 mm across) as the most common crystalline phase, with about 15-18 vol. % olivine in the surface samples. These volumes are vesicle-corrected. Detailed petrographic descriptions of Kilauea Iki lake samples, both eruptive pumice and drill core from the lake, are available in Richter and Eaton (1960), Murata and Richter (1966a), Richter and Moore (1966), Richter and Murata (1966), Helz (1980) and Helz (1987b).

2.4.2.2 *Olivines*

Helz (1987b) observed and defined five classes of olivine crystals in the magmas of the 1959 Kilauea eruption, from both lava lake and eruption pumice samples:

- (1) Irregular blocky crystals with deformation features;
- (2) Equant, euhedral, skeletal or strongly elongate grains;
- (3) Round or strongly resorbed grains;
- (4) Fragmental grains ('angular or conchoidal fragments');
- (5) Subhedral sulphide inclusion-bearing crystals.

As expected, all of these classes are also recognized in our samples, both in surface and drill core products. Class 1 consists here of large grains, 1-10 mm long, that are present in all the samples in varying proportions ranging from ~2 to 31% of the total crystal amount. Deformation is observed

optically as planar extinction discontinuities, undulose extinction, or subgrains in all the crystal sizes. Edge rounding or resorption are common features of this class, as well. Class 2 olivine is the most abundant. The grains commonly contain inclusions. We also noted rare evidence of deformation for this class, which is in slight contradiction with Helz (1987b). Class 3 olivine is fairly rare, typically a few percent of the total amount of olivines, as all the resorbed crystals with deformation features have been placed in class 1. Classes 4 and 5 are very rare and deformed grains are uncommon for both of those, even when the grains containing non-sulphide inclusion swarms are counted. The megacrysts noted by Helz (1987b) were not found in our samples. Although deformed crystals less than 0.5 mm in size are more difficult to recognize than larger ones, they were found in every sample. All deformed olivine crystals appear to be normally zoned.

2.4.2.3 Olivine aggregates

Most olivine crystals in every sample, and thus most of the Helz's classes appear to be part of larger olivine aggregates, as previously mentioned by Helz (1987b), Schwindinger and Anderson (1989), and Schwindinger (1999). Among the largest olivine aggregates, Helz (1987b) distinguished two groups: (1) open, glomerocrystic clusters of class 2 grains, and (2) dunitic or dense olivine aggregates. Group 2 is then subdivided into three subcategories: (1) dense aggregates of dominantly class 1 grains; (2) dense clusters of class 5 grains with sulphide-bearing inclusions; and (3) dunitic aggregates containing polygonal mosaics of unstrained olivine. In our samples olivine aggregates are omnipresent in varying but high proportions, with the glomerocrystic aggregates and the dense clusters of class 1 grains dominating. It is clearly the great majority of olivine crystals in all our samples. The clusters typically contain 2-6 individual crystals. This was also

observed in the 1969-74 Mauna Ulu lavas (Vinet and Higgins, 2010), but with overall smaller crystals and smaller aggregates. It is especially conspicuous in the Mauna Ulu lavas, and we note it as well here, that there is a great variability in the way in which the grains are connected in each aggregate. They range from adjacent, separate grains that may be connected in the third dimension, to clumps of crystals that are completely attached. Finally, some aggregates are re-equilibrated with originally separate grains becoming subgrains. In addition, we identified in KI67-1-05 one large dunitic aggregate of 230 grains containing rare deformed crystals. It is thought to resemble the last subcategory of dense aggregates of Helz (1987b), partly consisting in polygonal mosaics of undeformed olivines. It is the only example of this type of dense aggregate in our samples.

2.4.2.4 Other minerals

Kilauea Iki lavas also contain significant amounts of other minerals. Cr-spinel occurs within the olivine grains but mostly as separate crystals, commonly adjacent to the olivine grains. However, a few samples contain a lower amount of Cr-spinel that appears mostly included in olivine, as in the core KI81-1, for example. The Cr-spinel crystals are anhedral to euhedral and 10-100 μm in size. Clinopyroxene and plagioclase are rare as phenocrysts, but very common as microphenocrysts, 0.1 to 0.5 mm across, or microlites in the matrix. They may have grown during the eruption of their host lava and during the solidification period of the lava lake. Some microphenocrysts and phenocrysts of clinopyroxene show deformation features such as undulose or sector extinctions. Plagioclase, clinopyroxene and Cr-spinel were too small or not sufficiently abundant for their textures to be determined quantitatively.

2.4.2.5 *Vertical olivine-rich bodies*

Helz (1980, 1987a) described irregular, meter-scaled, pipe-like bodies that are mostly found at a depth range of 40-58 m within the lake, which she named vertical olivine-rich bodies (vorbs). Three lava lake samples in this study, KI75-1-141, KI75-1-145 and KI81-2-185.7, contain clear evidence of the presence of such vorbs. They are enriched in iron-rich olivine (Fo_{77-79}) and differentiated liquid, and have greater vesicle concentrations, relative to the adjacent host rock. Typically, the groundmass assemblage of olivine + augite + plagioclase in the vorb mostly formed adjacent to olivine, resulting in a related melt free of crystals. This textural observation is the key point in the recognition of such vorbs relative to their normal olivine-phyric matrix hosts. Helz et al. (1989, caption of Fig. 2b) further described the vorbs as containing “coarser, more abundant olivine, and more glass, in larger pools, than does the normal matrix rock”. They also noted that “the irregular distribution and grain size of groundmass phases [augite and plagioclase mostly] in the vorb contrast strongly with the uniform distribution and grain size of these phases in the normal matrix rock”. Helz (1987a) argued that the vorbs might be related, to a certain extent, to segregation veins. She also mentioned that their formation started by 1966, and ended in 1979.

2.4.2.6 *Olivine-depleted zone*

An olivine-depleted, low-MgO zone was identified to lie between ~18 to 40 m deep in the lake (Helz, 1980, 1987b; Helz, 2009; Helz et al., 1984; Helz and Wright, 1983). It contains 4.6 vol. % in average for the entire zone, of small, euhedral or rounded olivine phenocrysts (class 2) for the

great majority (Helz, 2009). Contrary to Helz (1980, 1987b) who noted the absence of class 1 olivines in this zone, here we observed some of those both in our samples and in other samples from National Rock and Ore Collections, National Museum of Natural History, Smithsonian Institution, Washington D.C. (not presented in this study). Helz (2009) also observed an olivine-rich, MgO-rich cumulate zone below ~55 m deep. She argued that the broadly S-shaped profile of MgO vs depth is consistent with redistribution of olivine phenocrysts within the lake in response to gravity, from the olivine-poor zone down to the olivine-rich zone. In this study, only a few samples from this olivine-depleted zone have been selected, near the upper and lower limits, because very most have insufficient olivine content to be included in the CSD analysis with reasonable uncertainties.

2.4.3 Chemical studies

2.4.3.1 Analytical methods

Olivine and glass major element compositions were measured in thin sections using the five-spectrometer, Cameca SX-100 electron-microprobe at the Université Laval Microanalysis Laboratory, Québec City, Canada. Both olivine and glass were analysed at 15 kV and 20 nA beam current, with a 5 μm spot size and using a wavelength dispersive spectra (WDS) mode. At least two spots, one in the core and one in the rim of each olivine, were analyzed to check for compositional zoning.

2.4.3.2 Olivine core and rim composition distribution

The cores and rims of over 430 olivine phenocrysts and microphenocrysts were analysed from 3 eruption scoria and 34 drill core samples. The forsterite (Fo) content of the olivine cores in all the Kilauea Iki lavas analysed here ranges from 68 to 90% (Fig. 2-2; Table 2-1). Within this wide range of compositions, olivine in the surface samples is distinguished by high core Fo (Fo₈₅₋₉₀) and limited zoning. This range contrasts with the generally lower Fo values of lava lake olivines, although there is a partial overlap. Amongst the lava lake samples, two distinct populations of olivines can be distinguished using their core Fo content: a high-Fo group (Fo₈₅₋₈₈) and a low-Fo group (Fo₇₇₋₈₁). This distribution seems to be mainly controlled by the olivines present at depths down to 60 m, and particularly in the centre of the lava lake. At depths of 60-90 m, the distribution of olivine composition changes. It consists of Fo₇₂₋₈₄ contents, with the range of the low-Fo group, Fo₇₇₋₈₁, dominating. This low-Fo group is clearly visible in the centre and toward the edge of the lake, whereas it is less evident at the edge of the lake. The high-Fo group is absent at these depths. Furthermore, at depths of 60-90 m some olivine grains are slightly Fo-depleted compared to the low-Fo group, with a bulk composition of Fo₇₂₋₇₆. Although minor in amount, those olivines may be a third population present within the lava lake. Finally, the composition of the olivine in the vorbs is in agreement with the Fo₇₇₋₇₉ range established by Helz (1987a).

In contrast to the Fo variation of the olivine cores, the overall composition of the olivine rims is much less variable and approximates a Gaussian distribution (Fig. 2-3). The major peak of composition is around Fo₇₂₋₇₆, which corresponds to the Fo range of the third olivine population defined by the crystal core composition. Also to be noted is the wider range of rim composition, Fo₅₇₋₈₈, as compared to the core compositions.

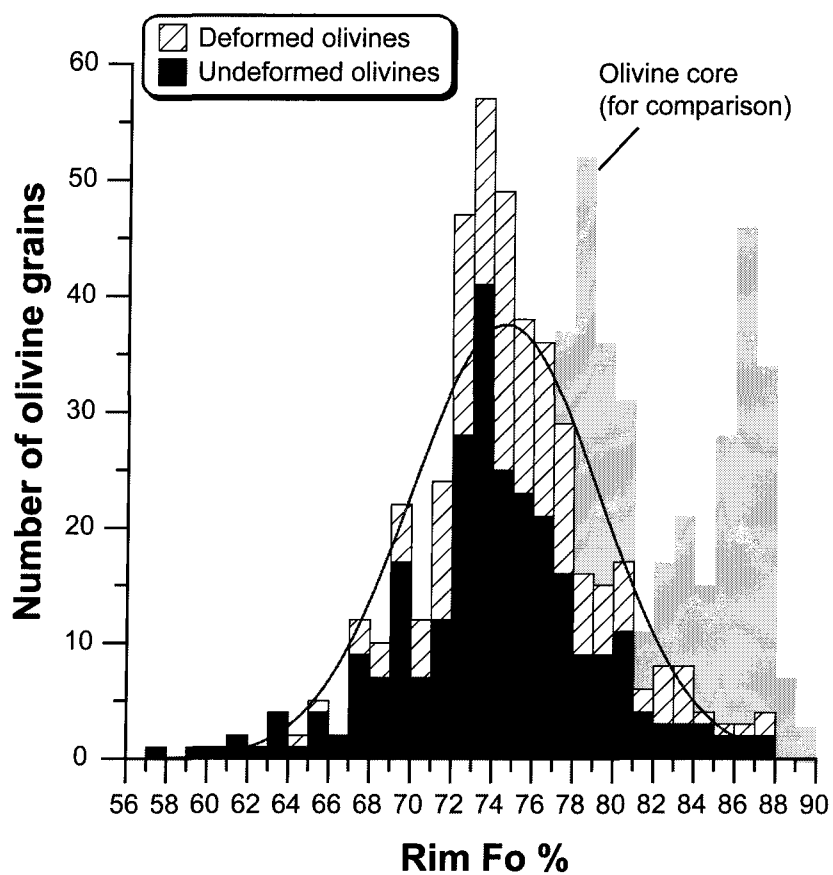


Figure 2-3. Histogram of olivine rim compositions (in Fo %) in lake samples. The overall composition of the olivine rims approximates a normal Gaussian distribution (in red). Deformed and undeformed crystals are represented in different patterns. Olivine core compositions, in light grey, are added for comparison

Table 2-1: Representative microprobe analyses of olivine cores and rims from Kilauea Iki surface and lake samples

Depth range (m):	-								0.5-3.5					
Sample:	KI-surface								KI67-1-05					
Size:	ph-c*	ph-r*	ph-c*	ph-r*	ph-c	ph-r	ph-c	ph-r	ph-c	ph-r	ph-c*	ph-r*		
SiO ₂	40.11	39.57	40.31	40.03	40.49	41.83	39.70	39.86	39.47	37.61	39.06	39.11		
MgO	47.20	44.62	47.55	47.47	47.33	46.99	44.91	44.85	46.63	38.48	45.90	44.13		
CaO	0.25	0.30	0.24	0.25	0.24	0.23	0.27	0.28	0.27	0.27	0.24	0.28		
MnO	0.16	0.21	0.19	0.13	0.10	0.19	0.20	0.23	0.22	0.40	0.14	0.22		
FeO	11.31	14.67	10.77	10.79	10.56	10.69	14.11	14.00	12.13	22.57	13.34	15.94		
NiO	0.32	0.21	0.36	0.35	0.37	0.41	0.29	0.29	0.40	0.30	0.38	0.38		
Total	99.52	99.68	99.53	99.22	99.37	100.56	99.56	99.63	99.25	99.74	99.18	100.42		
Fo%	88.2	84.4	88.7	88.7	88.9	88.7	85.0	85.1	87.3	75.3	86.0	83.1		
Depth range (m):	0.5-3.5													
Sample	KI75-3-05				KI75-1-02				KI75-1-05		KI75-1-11			
Size:	mph-c	mph-r	ph-c*	ph-r*	ph-c*	ph-r*	ph-c*	ph-r*	mph-c	mph-r	ph-c*	ph-r*	ph-c	ph-r
SiO ₂	39.78	35.79	39.75	36.46	39.28	38.78	40.09	39.25	39.22	38.18	39.78	37.12	37.76	36.89
MgO	44.95	27.15	44.23	31.18	44.66	44.12	48.33	46.72	46.29	40.25	46.59	35.99	39.42	35.42
CaO	0.28	0.22	0.29	0.30	0.28	0.29	0.23	0.24	0.27	0.29	0.26	0.23	0.27	0.22
MnO	0.24	0.51	0.23	0.34	0.18	0.21	0.07	0.20	0.25	0.30	0.13	0.31	0.28	0.34
FeO	13.91	36.15	15.38	31.15	15.22	15.92	10.21	12.59	12.92	20.39	12.55	25.75	21.69	26.58
NiO	0.33	0.18	0.33	0.14	0.32	0.34	0.41	0.40	0.31	0.24	0.41	0.22	0.27	0.25
Total	99.57	100.21	100.38	99.77	100.04	99.80	99.48	99.52	99.48	99.72	99.91	99.71	99.85	99.86
Fo%	85.2	57.2	83.7	64.1	83.9	83.2	89.4	86.9	86.5	77.9	86.9	71.4	76.4	70.4
Depth range (m):	5-14													
Sample:	KI67-1-20		KI67-1-35		KI75-3-18				KI75-3-46					
Size:	mph-c	mph-r	ph-c*	ph-r*	mph-c	mph-r	ph-c*	ph-r*	ph-c	ph-r	ph-c*	ph-r*		
SiO ₂	38.35	36.79	40.19	37.74	36.84	36.54	40.02	36.41	39.68	36.91	38.41	37.18		
MgO	40.85	35.05	48.61	38.43	35.45	35.02	45.58	32.56	45.02	33.31	40.03	34.95		
CaO	0.25	0.21	0.24	0.17	0.23	0.19	0.26	0.25	0.29	0.22	0.26	0.22		
MnO	0.25	0.34	0.12	0.31	0.34	0.51	0.17	0.28	0.20	0.32	0.36	0.37		
FeO	20.10	27.40	10.38	23.52	26.64	27.05	12.75	30.07	13.24	28.02	20.89	26.98		
NiO	0.30	0.27	0.49	0.24	0.28	0.33	0.41	0.19	0.36	0.22	0.27	0.28		
Total	100.24	100.16	100.07	100.46	99.90	99.88	99.25	99.92	98.98	99.09	100.35	100.05		
Fo%	78.4	69.5	89.3	74.4	70.4	69.8	86.4	65.9	85.9	67.9	77.4	69.8		

Table 2-1 (continued)

Depth range (m):	5-14								18-40					
Sample:	KI75-1-25				KI75-1-27				KI67-1-60		KI67-1-77			
Size:	ph-c	ph-r	ph-c*	ph-r*	ph-c	ph-r	ph-c*	ph-r*	ph-c	ph-r	ph-c*	ph-r*	ph-c	ph-r
SiO ₂	37.99	38.62	39.83	39.96	39.07	37.90	39.61	37.52	38.6	38.81	37.69	37.57	37.62	36.77
MgO	40.99	39.40	47.28	47.42	43.85	38.23	46.25	36.19	43.4	42.70	38.83	37.58	40.46	37.65
CaO	0.24	0.21	0.19	0.25	0.26	0.25	0.26	0.21	0.1	0.16	0.22	0.19	0.19	0.19
MnO	0.19	0.19	0.14	0.12	0.20	0.30	0.17	0.35	0.3	0.26	0.25	0.30	0.34	0.39
FeO	19.73	20.35	11.16	11.68	15.21	22.78	13.18	25.21	16.9	17.38	22.62	24.04	20.40	24.17
NiO	0.35	0.35	0.39	0.37	0.29	0.22	0.45	0.25	0.4	0.33	0.30	0.27	0.30	0.22
Total	99.50	99.27	99.21	99.91	99.06	99.76	100.28	99.91	99.8	99.71	100.02	100.07	99.44	99.56
Fo%	78.7	77.6	88.3	87.9	83.7	74.9	86.2	71.9	82.1	81.4	75.3	73.6	78.0	73.5
Depth range (m):	18-40								43-57					
Sample:	KI75-3-60				KI75-1-60				KI75-1-130		KI75-3-141			
Size:	ph-c*	ph-r*	ph-c	ph-r	ph-c	ph-r	ph-c*	ph-r*	ph-c	ph-r	ph-c*	ph-r*	ph-c	ph-r
SiO ₂	39.69	37.89	39.99	38.39	37.71	37.01	38.70	37.08	37.95	35.10	39.66	37.86	38.55	37.98
MgO	45.65	36.81	46.34	38.65	38.11	34.42	42.03	34.67	40.04	29.68	44.80	37.94	39.90	37.70
CaO	0.27	0.19	0.27	0.20	0.30	0.24	0.26	0.21	0.27	0.18	0.24	0.22	0.29	0.25
MnO	0.18	0.39	0.17	0.36	0.30	0.36	0.23	0.31	0.23	0.35	0.07	0.38	0.22	0.21
FeO	14.01	24.83	12.70	22.92	23.66	27.17	18.29	26.73	20.99	33.98	14.72	23.85	20.90	23.44
NiO	0.35	0.26	0.37	0.27	0.20	0.18	0.32	0.18	0.22	0.19	0.37	0.29	0.29	0.29
Total	100.31	100.43	99.98	100.88	100.36	99.47	99.91	99.32	99.71	99.71	100.03	100.67	100.25	99.93
Fo%	85.3	72.5	86.7	75.0	74.2	69.3	80.4	69.8	77.3	60.9	84.4	73.9	77.3	74.1
Depth range (m):	43-57								~60					
Sample:	KI79-5-182.4		KI75-1-141		KI81-2-185.7				KI79-5-193.3					
Size:	ph-c*	ph-r*	ph-c	ph-r	ph-c*	ph-r*	ph-c	ph-r	ph-c	ph-r	ph-c*	ph-r*		
SiO ₂	37.62	37.91	38.228	37.746	39.09	38.09	37.94	37.72	37.76	37.77	37.88	37.77		
MgO	38.16	37.51	38.425	37.353	43.17	39.57	40.19	38.44	38.29	37.25	38.11	37.57		
CaO	0.18	0.19	0.248	0.240	0.23	0.21	0.27	0.23	0.21	0.16	0.23	0.17		
MnO	0.26	0.35	0.263	0.347	0.16	0.27	0.24	0.37	0.35	0.23	0.31	0.23		
FeO	23.96	23.10	22.379	23.665	16.32	21.00	20.97	23.19	23.45	24.28	23.11	23.80		
NiO	0.24	0.28	0.212	0.266	0.34	0.31	0.30	0.23	0.27	0.26	0.26	0.26		
Total	100.58	99.50	99.80	99.75	99.46	99.86	99.99	100.38	100.49	100.07	99.92	99.86		
Fo%	74.0	74.3	75.4	73.8	82.5	77.1	77.4	74.7	74.4	73.2	74.6	73.8		

Table 2-1 (continued)

Depth range (m):	~60										75-90			
Sample:	KI88-1-205.2				KI81-1-200.2		KI88-2-197.7				KI79-5-250.0			
Size:	ph-c*	ph-r*	ph-c	ph-r	ph-c	ph-r	ph-c*	ph-r*	ph-c	ph-r	ph-c*	ph-r*	ph-c	ph-r
SiO ₂	39.20	37.78	38.38	37.84	38.36	38.20	39.37	37.80	38.09	37.58	40.02	37.74	37.98	37.48
MgO	44.22	38.10	40.37	38.14	40.78	40.69	43.70	38.81	38.32	37.15	46.96	38.47	38.66	36.38
CaO	0.24	0.19	0.26	0.17	0.26	0.27	0.23	0.21	0.22	0.15	0.29	0.17	0.26	0.15
MnO	0.14	0.20	0.21	0.31	0.24	0.30	0.15	0.34	0.36	0.33	0.22	0.19	0.25	0.31
FeO	16.07	23.17	19.64	22.92	19.37	19.68	15.96	21.46	23.43	23.89	12.55	22.65	22.76	24.62
NiO	0.33	0.31	0.29	0.28	0.30	0.27	0.31	0.23	0.31	0.22	0.37	0.25	0.27	0.23
Total	100.42	99.79	99.29	99.75	99.45	99.57	99.91	99.03	101.29	99.39	100.50	99.50	100.42	99.31
Fo%	83.1	74.6	78.6	74.8	79.0	78.7	83.0	76.3	74.5	73.5	87.0	75.2	75.2	72.5

Depth range (m):	75-90											
Sample:	KI88-1-298.1				KI81-1-249.7				KI81-1-299.9		KI88-2-296.1	
Size:	ph-c	ph-r	ph-c*	ph-r*	ph-c	ph-r	ph-c*	ph-r*	ph-c*	ph-r*	ph-c	ph-r
SiO ₂	37.57	37.45	37.39	37.36	38.61	38.57	38.41	38.69	39.03	37.83	37.44	37.63
MgO	36.37	36.41	36.78	36.62	42.18	41.91	41.92	42.19	44.14	38.43	36.98	36.96
CaO	0.20	0.20	0.22	0.18	0.26	0.27	0.25	0.26	0.25	0.23	0.22	0.21
MnO	0.34	0.26	0.23	0.30	0.28	0.29	0.23	0.25	0.18	0.26	0.31	0.41
FeO	25.65	25.61	24.47	25.17	18.38	18.29	18.64	18.66	15.80	22.14	24.64	25.05
NiO	0.24	0.30	0.28	0.26	0.31	0.30	0.24	0.26	0.31	0.28	0.26	0.24
Total	100.48	100.37	99.43	99.98	100.16	99.68	99.75	100.43	99.88	99.24	100.03	100.73
Fo%	71.6	71.7	72.8	72.2	80.4	80.3	80.0	80.1	83.3	75.6	72.8	72.4

Rock Mg-no. is $[\text{Mg}/(\text{Fe}^{2+} + \text{Mg})] \times 100$ assuming 90% of the total iron is Fe^{2+} ; ph, phenocryst; mph, microphenocrysts; r, rim; c, core; oxides are in wt %; Fo, forsterite; * deformed crystal.

2.4.3.3 *Fe-Mg zoning*

The great majority of the lava lake olivine crystals show moderate to strong normal zoning from core to rim, whereas olivines in surface samples are mostly unzoned or only marked by slight normal zoning, with a maximum Fo variation of ~4% (Table 2-1; Fig. 2-4). Almost no reverse zoning and no complex zoning are observed in any crystals. The Fo variation of normal zoning in the lava lake olivines is very variable, with a bulk maximum of 28% in KI75-3-05, at the edge of the lake. Overall, zoning increases with the increase of core Fo content, at every depth except 0.5-3.5 m, and for all of the drilling locations. It is somewhat more complex for samples from the centre of the lake; hence the centre and margin will be discussed separately.

Normal zoning in olivines from the centre of the lake (holes KI75-1, KI81-1, KI81-2 and KI88-2) varies markedly and extends from slightly negative values to ~17% Fo variation (Fig. 2-4). Normal zoning is strong in the upper 40 m and much weaker in the lower 50 m. Some unzoned olivines are present at very shallow depths (0.5 to 3.5 m), but they are much more abundant between 60 and 90 m.

Normal zoning in olivines from the margin of the lake (holes KI67-1, KI75-3, KI79-5 and KI88-1) varies from zero to ~28% Fo and decreases in magnitude downward. At shallow depths, 0.5-14 m, strong Fo variations are observed, which is expressed as a reduced range of higher core Fo content, Fo_{83-90} , (0.5-3.5 m), or for the entire range of core Fo contents, Fo_{70-90} , (5-14 m). Overall, the distribution of normal zoning is quite comparable for depths of 43-90 m. At the margin of the lake, it is thus at shallow levels that greatest zoning is developed in olivine. This is the region most strongly modified by foundering of crust.

Vorbs behave differently from other samples, in terms of Fo zoning. The vorb parts of KI75-1-141 and KI75-1-145 show a narrower range of zoning, with core and rim Fo content lying around Fo_{75-79} and Fo_{77-80} respectively. This tendency is less obvious for the third vorb-bearing (KI81-2-185.7).

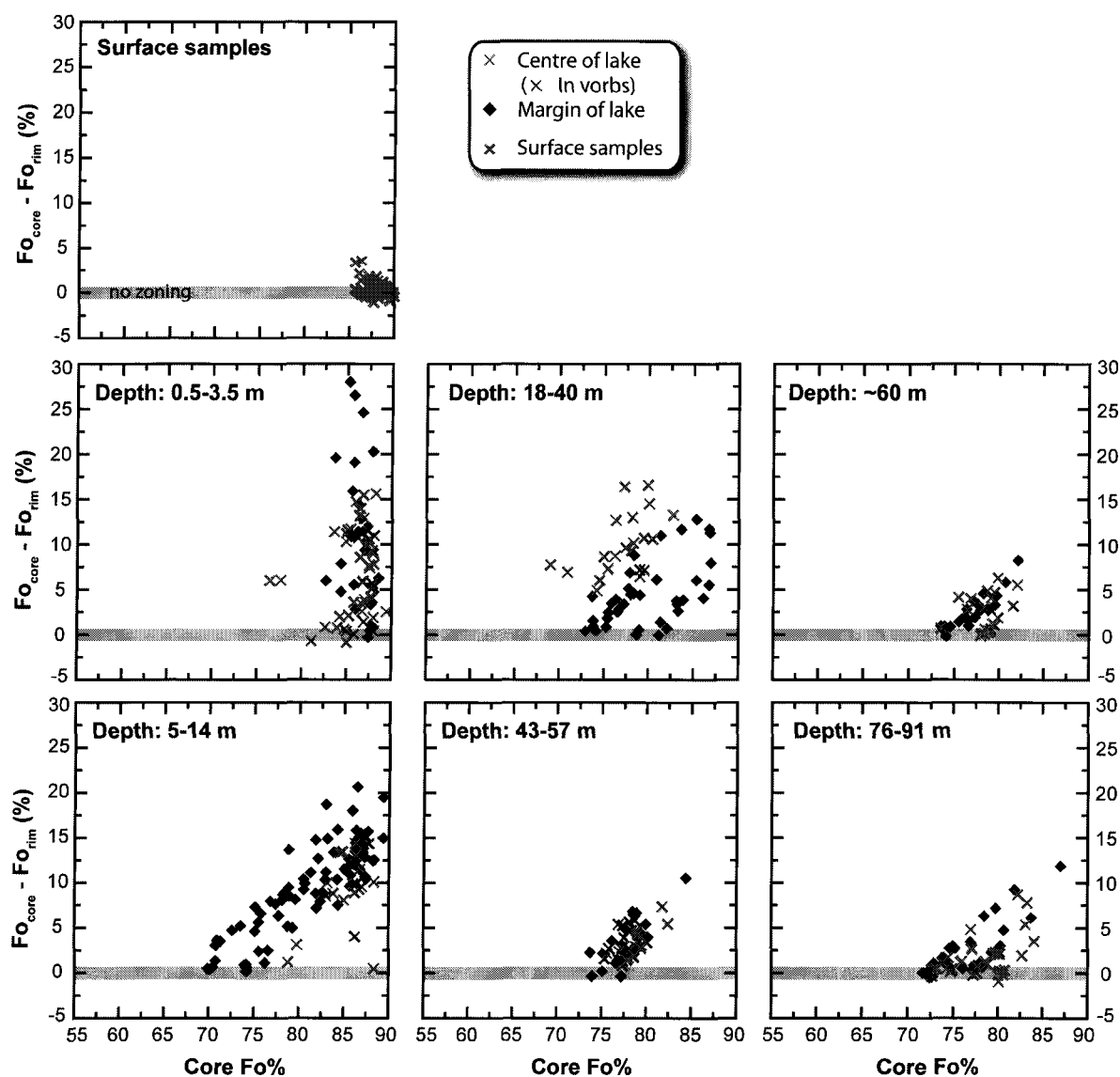


Figure 2-4. Variation of Fo content from core to rim versus Fo composition of the corresponding rim, for the olivine grains analyzed in Kilauea Iki samples, regarding the depth. The shaded area in each plot indicates the Fo composition range (around the variation value of 0) of no zoning. Grains normally zoned lie above this shaded area, and grains reversely zoned below it. Representative analyses given in Table 2-1

2.4.3.4 *CaO and NiO composition*

The CaO content of the cores of lava lake olivine is moderate and varies from 0.10 to 0.35 wt %, with our data fit well with the few previously published analyses (Helz, 1987b; Leeman and Scheidegger, 1977; Scowen et al., 1991) (Fig. 2-5a). The highest concentration of analyses lies between 0.2 and 0.3 wt % and comprises most of the deformed grains. The undeformed grains have a slightly wider compositional range. The CaO content of the rim covers about the same range of composition, for a larger range of Fo content. For comparison, the olivine from both our surface scoria samples and those from literature (Helz, 1987b) are also plotted. Scoria olivine has similar CaO composition than the lava lake olivine, with a narrower range of variation. All these CaO contents are typical of magmatic values and consistent with low pressure crystallisation in the crust (Jurewicz and Watson, 1988; Kamenetsky et al., 2006; Simkin and Smith, 1970; Stormer, 1973; Thompson and Gibson, 2000). Mantle values are significantly lower (Kamenetsky et al., 2006; Thompson and Gibson, 2000) (Fig. 2-5a).

The NiO content for the lava lake olivines ranges widely and shows a generally increasing trend from 0.13 to 0.49 wt % with increasing Fo content (Fig. 2-5b). The same kind of trend is also observed for the scoria olivines, with a narrower range of NiO composition of 0.24-0.43 wt %, and a steeper slope. The composition of the olivine rims is similar to that of the cores, but extends over a larger range of Fo content. Surprisingly, the core and rim composition of lava lake and scoria olivines from the previous published analyses of Helz (1987b) does not fit our data. In contrast, the microprobe analyses of five lava lake olivines from Leeman and Scheidegger (1977) match the highest NiO-Fo% values of our data. The lower NiO values found by Helz (1987b) may result from a lower analytical precision.

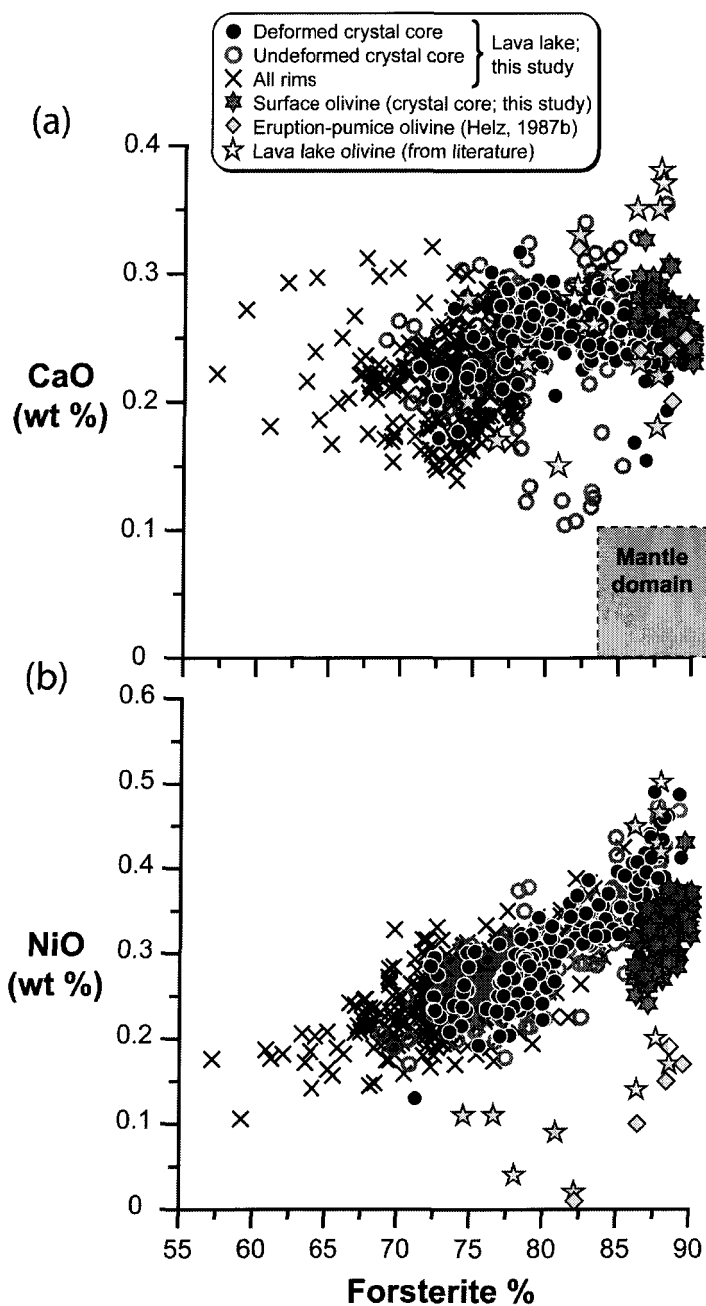


Figure 2-5. CaO (a) and NiO (b) content versus Fo content (%) of the core of both the undeformed and deformed olivine crystals analyzed in the lake samples. Also plotted are the rim compositions for the lake olivine, along with core compositions for the surface scoria olivine. Previous published analyses of eruption-pumice and lake olivines have been added for comparison. All olivines plot away up from the mantle domain indicated as the shaded area. Hence, no mantle xenocrysts are present here. Lava lake olivine: Helz (1987b), CaO and NiO; Leeman and Scheidegger (1977), CaO and NiO; Scowen et al. (1991), CaO. Mantle domain after Kamenetsky et al. (2006); Thompson and Gibson (2000)

2.4.4 Deformed olivine crystals

There are no systematic compositional differences between deformed and undeformed olivines in either microphenocrysts or phenocrysts (Fo content, Fig. 2-2; CaO and NiO contents, Fig. 2-5), and no differences in zoning amounts and distributions. In other words, only optical deformation features permit us to distinguish under the microscope the deformed crystals from the undeformed ones. This is in agreement with previous findings from lavas from the Hawaii Scientific Drilling Project (Garcia, 1996) and from some olivine-rich lavas taken amongst other Hawaiian picrites (Norman and Garcia, 1999; Wilkinson and Hensel, 1988).

The olivine crystals with deformation features are ubiquitous both in the scoria and lake samples. Deformed crystals have the same range of sizes of undeformed crystals. They are present as a variable proportion of total olivine crystals, ranging from 1.8-26.6 %, and up to 31.4 % in the vorb-only part of KI81-2-185.7 (Table 2-2). In terms of total volumetric abundance for each whole sample, it corresponds to 0.2-10.3 vol. % and up to 12.1 vol. % in the vorb-only part of KI81-2-185.7. In a plot of (ratio of deformed grains / total olivine number) versus depth (Fig. 2-6), one can observe that for depths <20 m up to three times more deformed grains occur at the centre (holes KI75-1, KI81-1, KI81-2 and KI88-2) and toward the edge (holes KI67-1 and KI88-1) of the lake relative to the edge (holes KI75-3 and KI79-5). In the olivine poor zone (20-40 m), there is a pronounced decrease in the proportion of deformed grains for samples at the centre of the lake. The same is noted as well for samples toward the edge of the lake, but it is less pronounced. At depths greater than 40 m deformed grains are again more abundant. This is observed for all the different drilling locations, but the proportion is more variable for the edge holes.

The vorbs of the hole KI75-1 have a similar proportion of deformed grains to the whole olivine population of the sample, at that depth. In contrast, the vorb part of KI81-2-185.7 has 31% deformed crystals, which is clearly much more abundant than that in any other samples.

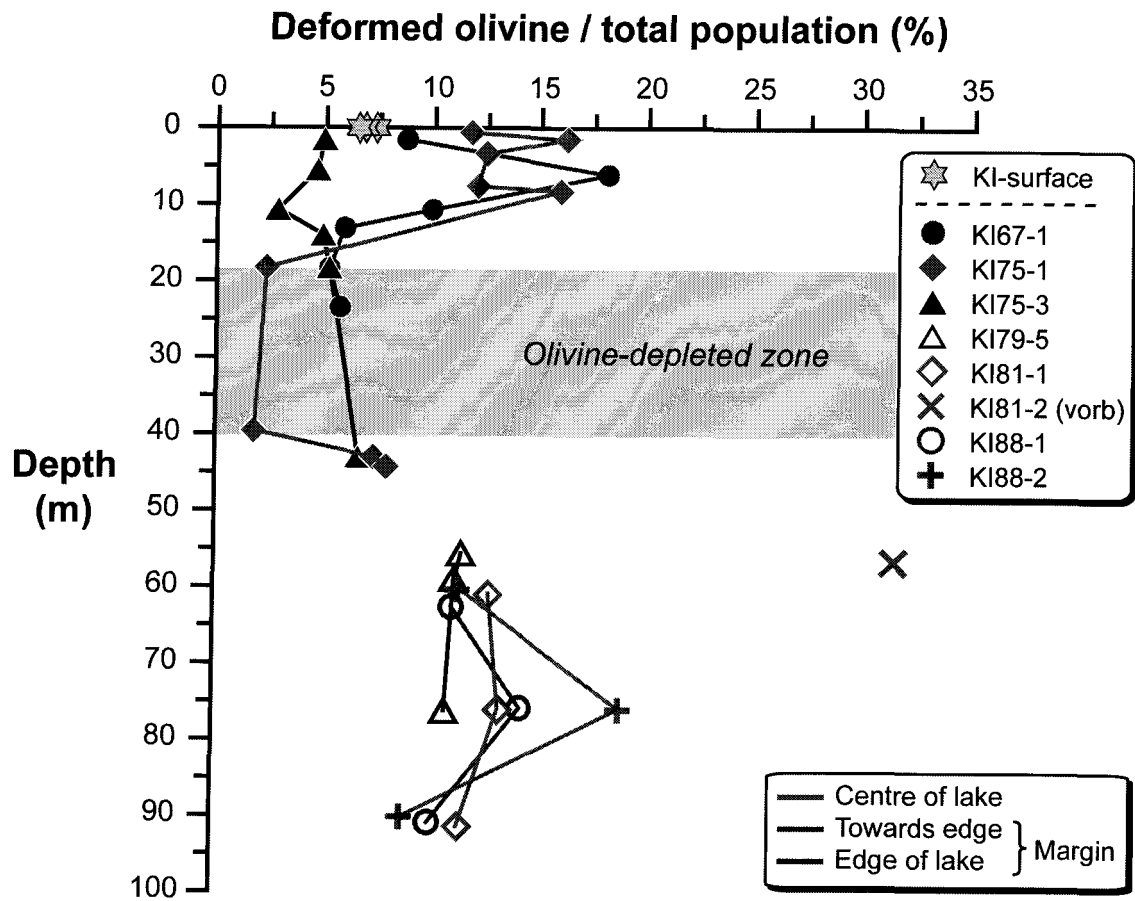


Figure 2-6. Ratio of deformed crystals to the total population plotted against the depth in the lava lake. Centre and margin of the lake are separated.

Table 2-2: Textural data (input and output) from *CSDCorrections*

Sample	Crystal morphology	R ² of shape best fit	Olivine vol. prop. (%)	Ratio deformed / total olivine population (%)	No. of crystals	L _{max} (mm)
NV-KI-59-1	rectangular prism	0.908	16.5	7.3	360	3.15
NV-KI-59-2a	rectangular prism	0.944	17.8	6.8	413	3.96
NV-KI-59-3	rectangular prism	0.922	14.8	6.5	370	3.53
KI67-1-05	rectangular prism	0.894	28.1	8.7	689	4.31
KI67-1-20	rectangular prism	0.843	30.3	18.1	265	5.54
KI67-1-35.1	rectangular prism	0.852	21.8	9.9	301	3.18
KI67-1-43	rectangular prism	0.924	23.6	5.9	539	3.38
KI67-1-60.4	rectangular prism	0.838	16.5	5.2	244	3.22
KI67-1-77	rectangular prism	0.924	15.5	5.7	280	3.12
KI75-1-02	rectangular prism	0.896	29.5	11.7	469	4.24
KI75-1-05	rectangular prism	0.934	34.7	16.2	473	5.64
KI75-1-11	rectangular prism	0.920	26.2	12.4	329	4.48
KI75-1-25	rectangular prism	0.881	29.7	12.0	356	4.48
KI75-1-27	rectangular prism	0.865	27.2	15.9	280	5.03
KI75-1-60	rectangular prism	0.850	7.2	2.3	117	2.16
KI75-1-130	rectangular prism	0.899	11.3	1.8	179	2.25
KI75-1-141	rectangular prism	0.894	24.2	6.3	423	2.79
<i>vorb-part</i>	-	-	27.7	7.3	401	-
KI75-1-145	rectangular prism	0.913	26.7	6.5	521	2.67
<i>vorb-part</i>	-	-	29.9	7.9	486	-
KI75-3-05	rectangular prism	0.766	14.2	4.9	223	3.30
KI75-3-18	rectangular prism	0.890	16.0	4.6	238	2.94
KI75-3-35	rectangular prism	0.888	12.5	2.8	359	2.71
KI75-3-46	rectangular prism	0.893	15.4	4.9	297	3.21
KI75-3-60	rectangular prism	0.911	17.6	5.2	277	3.93
KI75-3-141	rectangular prism	0.861	17.9	6.6	334	3.55
KI79-5-182.4	rectangular prism	0.912	25.3	11.4	279	3.00
KI79-5-193.3	rectangular prism	0.914	24.7	11.1	211	3.14
KI79-5-250.0	rectangular prism	0.882	20.9	10.7	286	3.47
KI81-1-200.2	rectangular prism	0.907	24.3	12.7	268	3.53
KI81-1-249.7	rectangular prism	0.926	28.1	13.2	407	3.28
KI81-1-299.9	rectangular prism	0.901	20.9	11.4	250	4.05
KI81-2-185.7	rectangular prism	0.953	38.6	26.6	285	4.35
<i>vorb-part</i>	-	-	43.3	31.4	156	-
KI88-1-205.4	rectangular prism	0.917	26.4	11.0	321	3.50
KI88-1-248.9	rectangular prism	0.858	30.0	14.2	361	3.06
KI88-1-298.1	rectangular prism	0.881	19.6	10.0	273	2.98
KI88-2-197.7	rectangular prism	0.873	26.1	11.3	268	3.77
KI88-2-249.5	rectangular prism	0.909	30.2	18.8	337	5.14
KI88-2-296.1	rectangular prism	0.902	15.5	8.7	268	3.59

Aspect ratios and R² calculated after Morgan and Jerram (2006). Olivine volume proportions are vesicle-corrected. L_{max} calculated from the three or four largest intersection lengths of each sample (after Marsh, 1998; O'Driscoll *et al.*, 2007).

2.5 QUANTITATIVE TEXTURAL MEASUREMENTS: CRYSTAL SIZE DISTRIBUTION (CSD)

2.5.1 Methods

To determine olivine CSDs, we stitched together high-resolution photomicrographs of each thin section to make one single image of each thin section. Subsequently, individual crystals were identified and outlined by using a polarizing microscope and a vector graphics editor. Effort was made to outline and measure separately each individual crystal in a cluster. In addition, all optically discernible deformed grains were outlined in a separate layer in order to compare the CSDs of undeformed versus total olivine populations. The digitized olivine texture images were then edited into simple greyscale compressed tiff format before importing into the software *ImageJ* (*Image Processing and Analysis in Java*). Area, orientation, and length of long and short axes of a best-fitted ellipse were determined.

We measured 120 to 940 olivine crystals in each thin section. Intersection data based on the major axis of the fitting ellipsoid are presented in Table 2-3. Intersection lengths were converted to 3-D CSDs using the program *CSDCorrections 1.3* (Higgins, 2000, 2002a; Higgins, 2006a). Logarithmic length intervals were used, with each bin $10^{0.2}$ times the size of the previous bin. There were no gaps in the CSDs; all bins contained crystals. Bins with less than three crystals were removed from the CSD analysis as not significant. The lower limit of the CSD was 0.1 mm, the smallest crystal that could be measured consistently and not necessarily the smallest crystal in the rock. However, for the glassy samples, we were able to measure all crystals.

To calculate CSDs from 2-D intersection measurements using *CSDCorrections 1.3*, we needed to specify the mean crystal shape as expressed by the crystal aspect ratio $S:I:L$ (short:intermediate:long dimensions). These aspect ratios were objectively estimated for each thin section, following Morgan & Jerram's (2006) *CSDSlice* method. A bulk fit value of the olivine shape, 1:1.15:1.6, was used for all the samples from Kilauea Iki, in order to plot and compare them adequately. This specific aspect ratio was chosen because, for every sample of this study, it appeared in every set of the five best fit values yielded as output results by *CSDSlice*. In contrast, all the individual CSD-derived textural parameters were obtained by using in the calculations the best fit individual aspect ratio (Table 2-2), among the set of the best five values estimated by *CSDSlice*. This gave greater accuracy in quantification. For the purposes of *CSDCorrections 1.3*, rock fabric was specified as massive; crystal roundness factor was estimated to be 0.5 in all cases. The complete list of all quantitative textural data from *CSDCorrections 1.3* is given in Table 2-2. In addition, vesicularity corrections were applied by estimating the proportion of vesicles in the sample, and then recalculating the area set in *CSDCorrections 1.3* by subtraction of the area of vesicles.

The CSD data were plotted on a $\ln(\text{population density})$ versus size diagram (Marsh, 1988b) (Fig. 2-7). For CSDs only slightly curved on a classical CSD diagram (S-type distribution of Higgins, 2006b), meaningful values for the slope and intercept of the CSD were determined by linear regression following Higgins (2006b). The 'Goodness of fit', Q , is a parameter that describes how well the data fit to a straight line (Table 2-4), taking into account the error in each point. A value of Q greater than 0.1 indicates a very significant fit, but values greater than 0.01 are also acceptable (Higgins, 2006b). The characteristic length (L_D) of the CSDs was calculated as $-1 / \text{slope}$. This

parameter is equal to the mean crystal length for a straight CSD that extends to all crystal sizes (Marsh, 1988b).

Strongly curved CSDs necessitated a different approach: the slope, intercept and L_D were calculated separately for the populations of smallest and largest crystals using the mean of several CSD bins in the corresponding domain. The volumetric olivine proportion is obtained by dividing the total area of the olivine crystals by the total measured area. A plot of phase volumetric proportion versus L_D (Fig. 2-8) is more useful than the commonly used graph of slope versus intercept, because of the correlation between these parameters produced by closure (pp 71-74 in Higgins, 2006a). In addition, the segmenting of the curved CSDs creates at least two different L_D values for one sample, and thus makes this L_D diagram more relevant, in particular in the discerning of distinct crystal populations. It also gives an idea of the curvature of the CSD.

The maximum 3-D crystal size is also an important (or useful) feature of the CSD as it indicates how far a CSD extends. However, no rigorous assessment is possible using two-dimensional data. An estimate of this characteristic is the maximum intersection size (L_{max} ; Fig. 2-9, Table 2-2). This is not very reliable measure because of the small number of large crystals present, which are not necessarily in the right orientation. Here, we used the average of the intersection length of the three or four largest crystals for each sample as suggested by Marsh (1998) and O'Driscoll *et al.* (2007).

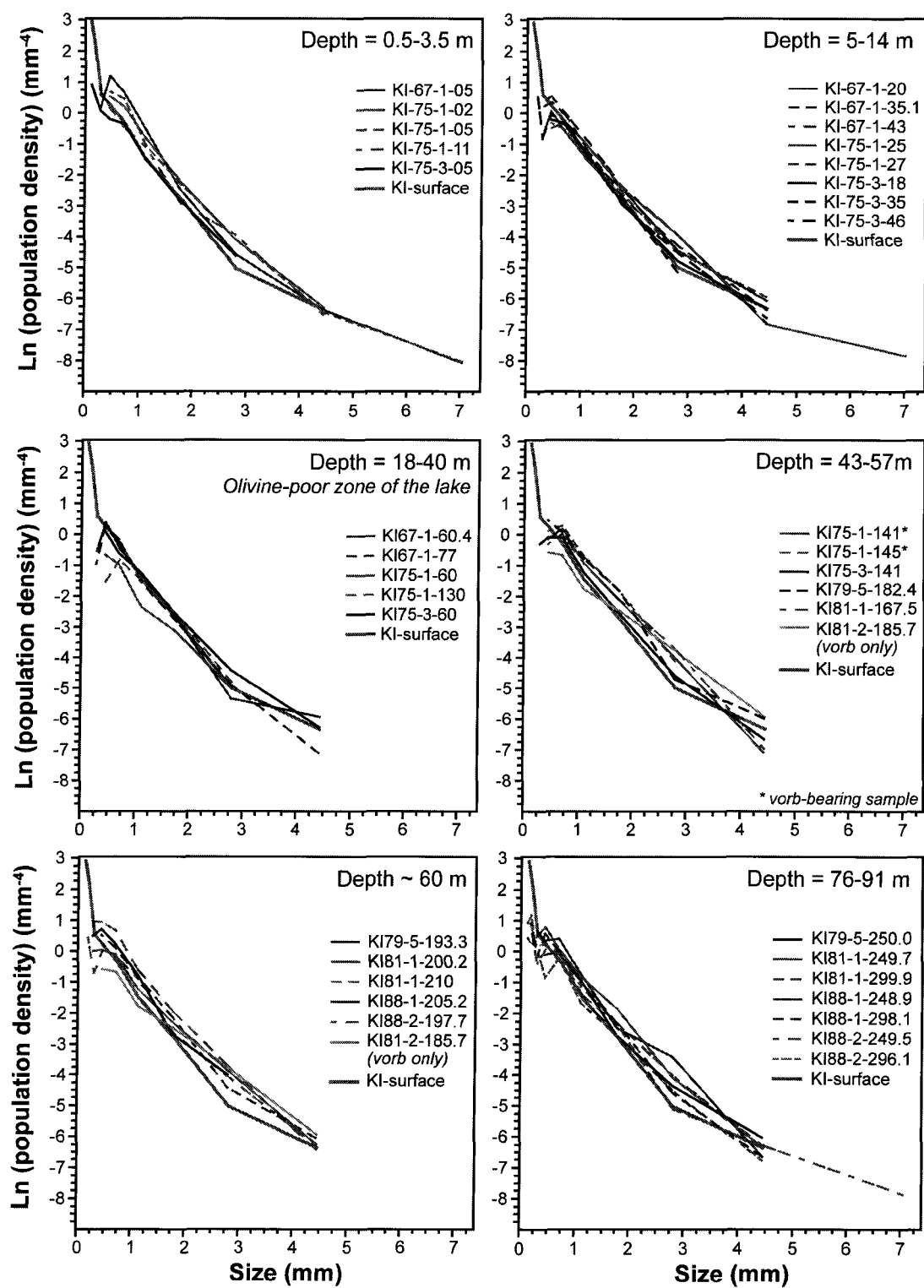


Figure 2-7. Olivine crystal size distributions (CSDs) of surface and lake samples, plotted at several fixed depth intervals that cover most of the thickness of the lake. The last part of the sample number is the depth in feet. The surface sample CSDs are shown in all frames as reference for variations. The mean best olivine shape for the CSD analysis was chosen to be 1:1.15:1.6 (ellipsoid).

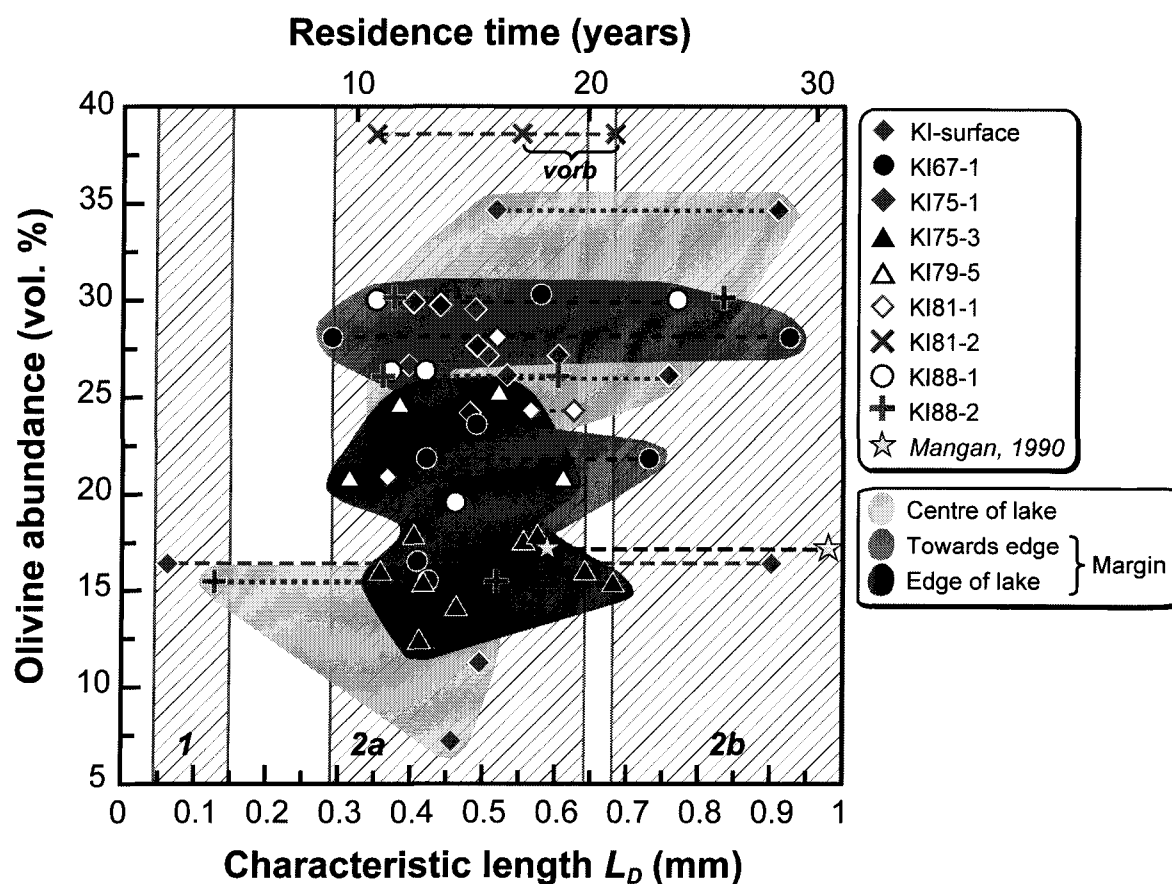


Figure 2-8. Characteristic length (L_D) versus olivine volumetric proportion in the surface and lake samples. L_D is equal to $-1 / \text{CSD slope}$. For the curved CSDs, two points per sample are plotted, which correspond to L_D of the extreme CSD segments. The volume-percent olivine axis applies to the whole sample. Also indicated on the X-axis is the residence time of crystal, assuming a mean growth rate of 10^{-9} mm/s (estimate after Cashman, 1993; Mangan, 1990). A single value of growth rate is used instead of a range of values for clarity.

Table 2-3: Crystal size ranges expressed as the number of crystals per unit area

Samples			Number of crystals per crystal size interval used in the CSD analysis										
Sample name	Vesicularity (%)	Total area measured (mm ²)	10.00-6.31 (mm)	6.31-3.98	3.98-2.51	2.51-1.58	1.58-1.00	1.00-0.631	0.631-0.398	0.398-0.251	0.251-0.158	0.158-0.100	0.100-0.0631
NV-KI-59-1	8.8	568.4			5	9	25	50	60	55	44	43	57
NV-KI-59-2a	11.4	866.3	2	6	16	40	55	79	54	28	82	49	
NV-KI-59-3	21.1	913.9		7	5	39	56	75	58	39	47	43	
KI67-1-05	16.8	929.6		3	7	17	54	152	215	169	60		
KI67-1-20	24.1	876.2		8	17	49	56	57	48	18			
KI67-1-35.1	5.6	587.6		5	17	33	61	79	58	29			
KI67-1-43	3.5	904.6		5	35	68	124	128	102	52			
KI67-1-60.4	23.2	787.4		8	7	32	57	59	58	21			
KI67-1-77	6.7	666.5		3	11	35	54	85	62	21			
KI75-1-02	25.9	933.3	2	7	25	65	111	133	91				
KI75-1-05	20.7	853.9	3	6	24	59	87	155	100				
KI75-1-11	10.1	849.4		6	32	39	92	82	69				
KI75-1-25	11.4	864.4		4	5	32	61	100	94	46			
KI75-1-27	10.2	878.9		11	20	52	65	73	45				
KI75-1-60	15.8	700.4				8	21	21	32	22			
KI75-1-130	12.3	778.1			11	41	53	47	19				
KI75-1-141	0	663.2		3	27	70	100	128	57				
<i>vorb-part</i>		569.2		3	27	70	97	119	55				
KI75-1-145	0	788.2			2	26	103	136	160	81			
<i>vorb-part</i>		648.6			2	25	101	129	149	70			
KI75-3-05	33.7	830.9			11	27	43	57	34	21	14	8	
KI75-3-18	20.4	786.0			6	12	31	62	71	41			
KI75-3-35	5.5	911.4			11	50	67	103	65	27	26	8	
KI75-3-46	4.3	795.5		5	17	34	66	77	52	21	19		
KI75-3-60	14.6	839.5	2	7	18	43	71	61	53				
KI75-3-141	4.9	775.7		5	16	47	75	98	54	27	10		
KI79-5-182.4	-	397.8		5	8	52	65	67	46				
KI79-5-193.3	-	311.1				12	19	48	57	42	20		
KI79-5-250.0	-	585.9			7	16	33	75	82	43			
KI81-1-200.2	-	519.4		5	22	44	55	68	41	22	9		
KI81-1-249.7	-	536.7		5	19	69	83	89	78	36			
KI81-1-299.9		516.3			5	7	39	45	67	40	18	20	7

Table 2-3 (continued)

KI81-2-185.7	-	492.0	2	11	26	45	56	74	48			
<i>vorb-part</i>		318.3	2	6	21	33	30	36	21			
KI88-1-205.4	-	501.6		6	13	48	77	84	60			
KI88-1-248.9	-	474.0	3	31	36	76	103	54	34			
KI88-1-298.1	-	457.1	3	10	38	58	68	55	21	16		
KI88-2-197.7	-	524.9		5	19	33	74	70	41	17	8	
KI88-2-249.5	-	586.6	3	6	22	51	68	83	55	29	14	5
KI88-2-296.1	-	677.1		4	16	38	47	78	30	27	22	

Table 2-4: CSD results including slope and intercept data calculated by CSDCorrections for each sample

Sample	Depth in lake (m)	Hole location on the lake	Drilling year	Slope		Intercept $\ln(n_0)$		Crystal growth time (years)		Nucleation rate ($\times 10^{-9} \text{ mm}^{-3}/\text{s}$)		Characteristic length (mm)	Goodness of fit (Q)
					error (1 σ)		error (1 σ)	G1	G2	G1	G2		
KI-surface	0	-	-	-1.11	0.18	-2.28	0.57	11.4	57.1	5.7	1.14	0.90	0.39
				-16.9	1.4	4.67	0.20	4.7	23.4	3.2	0.65	0.06	0.16
KI67-1-05	1.5	margin	1967	-3.46	0.15	2.67	0.12	3.7	18.3	6.7	1.34	0.29	0.19
				-1.08	0.20	-2.08	0.62	11.7	58.7	5.2	1.04	0.93	0.40
KI67-1-20	6.1	margin	1967	-1.73	0.09	0.51	0.14	7.3	36.7	1.3	0.26	0.58	0.08
KI67-1-35.1	10.7	margin	1967	-1.37	0.18	-0.33	0.44	9.3	46.3	0.8	0.17	0.73	0.70
				-2.38	0.18	1.50	0.19	5.3	26.6	3.8	0.75	0.42	0.68
KI67-1-43	13.1	margin	1967	-2.04	0.08	1.28	0.10	6.2	31.1	3.2	0.64	0.49	0.14
KI67-1-60.4	18.4	margin	1967	-2.45	0.16	1.35	0.18	5.2	25.9	3.4	0.68	0.41	0.77
KI67-1-77	23.5	margin	1967	-2.36	0.13	1.27	0.15	5.4	26.9	3.2	0.64	0.42	0.18
KI75-1-02	0.6	centre	1975	-2.05	0.09	1.56	0.12	6.2	30.9	3.9	0.78	0.49	0.31
KI75-1-05	1.5	centre	1975	-1.94	0.15	0.97	0.24	6.5	32.7	2.4	0.49	0.52	0.72
				-1.10	0.17	-1.28	0.57	11.5	57.7	3.2	0.64	0.91	0.14
KI75-1-11	3.4	centre	1975	-1.32	0.18	-0.60	0.43	9.6	48.0	1.5	0.30	0.76	0.16
				-1.88	0.32	0.89	0.27	6.7	33.7	2.2	0.45	0.53	0.52
KI75-1-25	7.6	centre	1975	-2.28	0.11	1.04	0.14	5.6	27.8	2.6	0.52	0.44	0.45
KI75-1-27	8.2	centre	1975	-1.66	0.08	0.34	0.13	7.6	38.2	0.9	0.17	0.60	0.02
				-1.98	0.13	0.70	0.19	6.4	32.0	1.8	0.35	0.51	0.53
KI75-1-60	18.3	centre	1975	-2.19	0.20	0.24	0.24	5.8	29.0	0.6	0.12	0.46	0.35
KI75-1-130	39.6	centre	1975	-2.02	0.17	0.55	0.24	6.3	31.4	1.4	0.28	0.50	0.60
KI75-1-141	43.0	centre	1975	-2.08	0.09	1.48	0.13	6.1	30.5	3.7	0.74	0.48	0.16
<i>vorb-part only</i>				-2.04	0.09	1.61	0.14	6.2	31.1	4.0	0.81	0.49	0.26
KI75-1-145	44.2	centre	1975	-2.53	0.11	1.79	0.13	5.0	25.1	4.5	0.90	0.40	0.21
<i>vorb-part only</i>				-2.49	0.12	1.89	0.14	5.1	25.5	4.7	0.95	0.40	0.20
KI75-3-05	1.5	margin	1975	-2.16	0.14	0.90	0.17	5.9	29.4	2.3	0.45	0.46	0.71
KI75-3-18	5.5	margin	1975	-2.80	0.27	1.46	0.27	4.5	22.6	3.7	0.73	0.36	0.89
				-1.56	0.20	-0.53	0.43	8.1	40.7	1.3	0.27	0.64	0.15
KI75-3-35	10.7	margin	1975	-2.43	0.14	1.22	0.17	5.2	26.1	3.1	0.61	0.41	0.06
KI75-3-46	14.0	margin	1975	-1.47	0.19	-0.51	0.44	8.6	43.1	1.3	0.26	0.68	0.66
				-2.40	0.19	1.15	0.20	5.3	26.4	2.9	0.58	0.42	0.53
KI75-3-60	18.3	margin	1975	-1.80	0.09	0.65	0.13	7.0	35.2	1.6	0.33	0.56	0.10
KI75-3-141	43.0	margin	1975	-2.48	0.20	1.49	0.22	5.1	25.6	3.7	0.75	0.40	0.25
				-1.74	0.19	0.26	0.40	7.3	36.4	0.7	0.13	0.57	0.23
KI79-5-182.4	55.6	margin	1979	-1.92	0.11	1.31	0.15	6.6	33.0	3.3	0.66	0.52	0.10

Table 2-4 (continued)

KI79-5-193.3	58.9	margin	1979	-2.61	0.17	1.71	0.18	4.9	24.3	4.3	0.86	0.38	0.12
KI79-5-250.0	76.2	margin	1979	-1.64	0.21	-0.39	0.42	7.7	38.7	1.0	0.20	0.61	0.31
				-3.19	0.29	1.87	0.25	4.0	19.9	4.7	0.94	0.31	0.48
KI81-1-200.2	61.0	centre	1981	-1.77	0.09	0.86	0.14	7.2	35.8	2.2	0.43	0.56	0.22
				-1.60	0.13	0.46	0.24	7.9	39.6	1.2	0.23	0.63	0.90
KI81-1-249.7	76.1	centre	1981	-1.93	0.09	1.42	0.12	6.6	32.9	3.6	0.71	0.52	0.20
KI81-1-299.9	91.4	centre	1981	-2.73	0.2	1.35	0.19	4.6	23.2	3.4	0.68	0.37	0.08
KI81-2-185.7	56.6	centre	1981	-1.47	0.16	-0.04	0.36	8.6	43.1	0.1	0.02	0.68	0.41
				-2.87	0.44	1.66	0.31	4.4	22.1	4.2	0.83	0.35	0.15
				-1.81	0.19	0.58	0.29	7.0	35.0	1.5	0.29	0.55	0.65
<i>vorb-part only</i>				-1.69	0.13	0.53	0.21	7.5	37.5	1.3	0.27	0.59	0.33
KI88-1-205.4	62.6	margin	1988	-2.39	0.12	1.42	0.13	5.3	26.5	3.6	0.71	0.42	0.06
				-2.7	0.15	1.67	0.16	4.7	23.5	4.2	0.84	0.37	0.93
KI88-1-248.9	75.9	margin	1988	-1.3	0.21	-0.13	0.5	9.8	48.8	0.3	0.07	0.77	0.04
				-2.86	0.22	2.34	0.23	4.4	22.2	5.9	1.17	0.35	0.72
KI88-1-298.1	90.9	margin	1988	-2.17	0.13	1.49	0.16	5.8	29.2	3.7	0.75	0.46	0.88
KI88-2-197.7	60.3	centre	1988	-1.66	0.22	-0.07	0.46	7.6	38.2	0.2	0.04	0.60	0.97
				-2.78	0.28	1.68	0.27	4.6	22.8	4.2	0.84	0.36	0.21
KI88-2-249.5	76.0	centre	1988	-1.20	0.19	-1.19	0.58	10.6	52.8	3.0	0.60	0.83	0.17
				-2.67	0.42	1.5	0.29	4.8	23.8	3.8	0.75	0.37	0.21
KI88-2-296.1	90.3	centre	1988	-1.93	0.17	0.22	0.27	6.6	32.9	0.6	0.11	0.52	0.86
				-7.9	2.47	2.17	0.63	1.6	8.0	5.4	1.09	0.13	0.95

Nucleation rates and crystal growth times (residence times) are calculated for olivine growth rates $G1 = 2.5 \times 10^{-9}$ mm/s and $G2 = 5 \times 10^{-10}$ mm/s (estimate after Cashman, 1993; Mangan, 1990; Marsh, 1988b, 1998; Wright and Fiske, 1971). The "Goodness of fit", Q , is a parameter that describes the quality of fit of the linear regression (see text for details).

2.5.2 CSD results

2.5.2.1 *General remarks*

Although all the Kilauea Iki drill core CSD patterns are broadly similar, they show significant differences in size, nucleation density and curvature, depending on the bulk drilling date, the drilling location and the proportions of deformed crystals (Table 2-4; Fig. 2-7). Samples have been classified by depth in figure 2-7, as a proxy for time of solidification. Most samples have CSDs that range from ~ 0.1 mm, the lower analytical limit after stereological conversion, to 4.5 mm; KI67-1-05, KI75-1-05, KI75-1-25 and KI88-2-249.5 extend to 7.0 mm; and KI75-3-05, KI75-3-35 and KI79-5-193.3 only extends to 2.8 mm in size (mid-interval intersection length). There is no simple link between CSDs and either drilling date, location or depth.

Many samples show a turndown of the CSD curves at smallest crystal sizes. However, we must verify whether this feature is a measurement artefact or a part of the CSD. In order to validate the true significance of this turndown in terms of magma processes, we plot the glassy samples separately and check for a turndown. If the turndown at the left side of the CSD exists for the glassy samples, then we can clearly say that it is not an artefact since all crystals are measured in glassy samples. The same holds true for some of the samples with a crystallized matrix that also have a turndown at smallest sizes where only crystals >0.1 mm were measured. Here four glassy samples were studied: KI81-2-185.7 and the three samples from the hole KI79-5. A clear turndown is observed for the KI79-5 samples (Fig. 2-7). In addition, no turndown occurs at the smallest sizes for surface samples that have been the most rapidly cooled; the slope is even much steeper and

still negative. This suggests real turndown for lava lake samples. Accordingly, the presence of a turndown is thus thought to reveal true magma processes there.

About half the CSDs have a significant curvature. The curvature is markedly variable and appears to be complex, in particular for the samples of the centre of the lake. It commonly consists of profiles with a steeper slope at small and middle sizes, sometimes both being almost linear, and a flatter slope at greater sizes, thus forming a clear change in the curve at the right side of the graph. Sometimes, the curvature is more continuous, or smooth. Most CSDs from the edge of the lake are curved. A convenient way of describing the characteristics of these plots is to divide the CSDs into two parts: (1) the main part, for crystal sizes >1.0 mm, and; (2) the part for the smallest sizes, i.e., <1.0 mm. The mean surface CSD is often used as a bench-mark for CSD variations. However, this does not imply that it is the most important unit.

2.5.2.2 CSD variations for sizes <1.0 mm

The left side of the CSD plot can be characterised in terms of slope and presence or absence of a turndown. The mean surface CSD has much steeper slope and greater intercept than the great majority of drill core CSDs, but no turndown. The turndown at sizes <0.75 mm is present for many CSDs of the margin of the lake, from shallow to deep levels in the lake. Certain ones display very steep slopes at sizes <0.5 mm. Turndown is also observed for some CSDs from the centre of lake, but mostly in the depth range 40-75 m. No turndown is observed for the CSD of the vorb part of KI81-2-185.7. At 75-90 m, one sample from the margin and two samples from the centre of the lake, exhibit steep slopes at smallest sizes that closely match the mean surface CSD. If this is not a stereological artefact, then it is thought to represent a common population of small crystals.

2.5.2.3 *CSD variations for sizes >1.0 mm*

The right side of the CSDs can be described in terms of slope and maximum size. The mean surface CSD has a relatively steep slope for sizes 1.0-2.8 mm (mid-sized interval), but a flatter slope for greater sizes than the great majority of lava lake samples. The latter is especially true at depths from 18 m down, both at the margin and centre of the lake. Significant variations of the drill core CSDs are commonly observed for crystals >1.0 mm. Sometimes these CSDs match with the mean surface CSD, either for the mid-sized interval, the greatest sizes, or both those parts of the CSD. This is observed for both the margin and centre of the lake. However, at depths >10 m, most drill core CSDs do not match the mean surface CSD for sizes >2.8 mm. At depths >60 m, both drill core CSDs of the margin and centre of lake often are displaced upward relative to the mean surface CSD. At 75-90 m, only KI81-1-299.9 has a CSD with the same pattern as the mean surface CSD; the other CSDs appear generally flatter for sizes 1.0-2.8 mm and steeper for sizes >2.8 mm.

In summary, there are only slight textural variations through the upper 100 meters of the lava lake. Although we note slightly more textural variations from 60-90 m, this graphical way of representing the evolution through time via multiple plots at constant depths definitely highlights the bulk homogeneity of the whole set of CSDs. However, several exceptions remain discernible, as demonstrated above.

2.5.2.4 Other CSD-derived results

A plot of characteristic length (segments and whole samples) versus olivine abundance shows great variations between samples, and may help for recognition of different crystal populations (Fig. 2-8). Two main groups regarding the characteristic length are recognised here: group 1 where $L_D < 0.15$ mm, typically ~ 0.1 ; and group 2 where L_D is 0.3-1.0 mm. It is convenient to further divide group 2 into two subcategories, which do not necessarily indicate different processes: subgroup 2a with L_D of 0.30-0.65 mm, and subgroup 2b with L_D of 0.7-1.0 mm. Group 1 is only composed of the lower end-member of both the surface samples and KI88-2-296.1. Subgroup 2b contains the upper end-member of KI67-1-05, KI67-1-35.1, KI75-1-05, KI75-1-11, KI75-3-46, KI88-1-248.9, KI88-2-249.5, KI81-2-185.7 (vorb) and of the scoria samples of Mangan (1990). In subgroup 2a, the most common one, lie all the remaining end-members or whole samples. These three groups or subgroups likely may be attributed to three distinct olivine populations. Furthermore, less variability is observed at the margin versus centre of the lake, as illustrated by the coloured areas in figure 2-8.

The total abundance of olivine is very variable (Fig. 2-8). Sample KI75-1-60, KI75-1-130 and KI88-2-296.1 are clearly poorer in olivine (7-16 vol. %) compared to the other samples from the same main hole(s) 1975-88 at the centre of the lake (21-33 vol. %). With a range of 13-30 vol. %, the samples studied from the main hole(s) 1967-88, towards the edge of the lake, show great variations in the olivine abundance, as well. In contrast, less variability in the olivine content is noted for the samples from the main hole(s) 1975-79 at the edge of the lake, with 13-25 volume %. It should be noted though that samples in hole KI75-3 and KI79-5 appear to define separate zones in the plot, the latter being depleted in olivine compared to the samples from greater

depths of KI79-5 hole. The olivine content of hole KI75-3 and KI79-5 samples shows little variation and range from 13 to 18 vol. % and 21 to 25 vol. %, respectively.

The vorb-bearing KI81-2-185.7 is the richest in olivine among all the samples we studied here, including our surface samples and the scoria samples from Mangan (1990). The distribution of L_D in this sample is comparable to the other 1981 lava lake samples.

There are significant differences in the maximum 3-D sizes (L_{max}) between the Kilauea Iki samples, with respect to depth (Fig. 2-9). The whole range of values is 2.1-5.6 mm, with a bigger concentration between 2.7 and 4.0 mm, including the surface samples. Overall, samples from the centre and toward the edge of the lake have higher L_{max} at depths 0-15 m, compared to deeper levels. In fact, for samples toward the edge of the lake L_{max} appears more or less constant, around 3.2 mm, from 15 m deep down. It is much more variable for samples from centre of the lake, with minimal values (~ 2 mm) at 20-40 m deep, the olivine-depleted zone defined by Helz (2009). It then increases almost continuously with depth up to 4 or 5 mm at ~ 80 -95 m deep. In contrast, the distribution of L_{max} for olivine in the edge of the lake begins decreasing with depth at shallower levels, and then reaches a maximum of ~ 4 mm at the top of the olivine-depleted zone. Finally L_{max} decreases again to a mean value of ~ 3 mm, although variations occur. Interestingly, L_{max} in the olivine-depleted zone seems to be controlled by the location in the lake: from lower values at the centre to higher ones at the edge, via intermediate values toward the edge.

Surprisingly, L_{max} of the pumice sample set of Mangan (1990) is comparable to our highest values, significantly higher than the L_{max} of our surface samples. In addition to the bulk variation of L_{max} over all the samples studied from 1975-88, internal variations among the samples from the same hole are also noted. The best example here is given by the clear dichotomy of the samples in

the hole KI75-1 where L_{\max} appears to range between 2.0 and 3.0 mm from 18 to 44 m deep, whereas it is about twice that value at shallower depths (1-8 m).

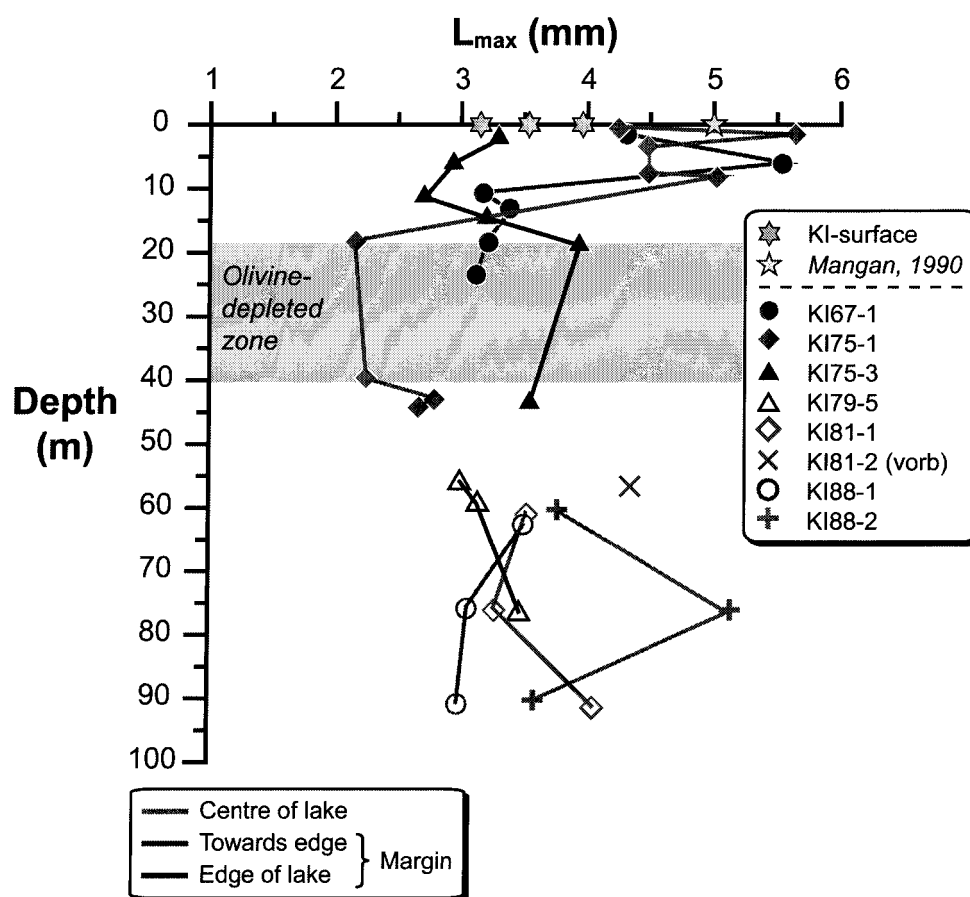


Figure 2-9. Maximum 3-D size (L_{max}) plotted against the depth in the lava lake. Centre and margin of the lake are separated and represented with different colours.

2.5.2.5 *Lateral and depth-related evolution: centre versus margin of the lake*

It is instructive to see if there are systematic textural differences between the centre and margin of the lake. We have therefore compiled characteristic length versus depth data for 1975-88 (0.5-91 m) at the centre of the lake, and 1967-88 (1.5-91 m) plus 1975-79 (1.5-76 m) at the northern margin of the lake (Fig. 2-10).

Characteristic length data for the samples from the center of the lake show the greatest variations compared to the drilling locations at the margin (Fig. 2-10a). Overall, greater values of L_D are found at both the shallowest and deepest parts of the lake. At depths <10 m, L_D varies widely from 0.45 to 0.90 mm. In contrast, less variation between 10-40 m deep is noted, but few samples have been studied in this depth interval that corresponds to the olivine-poor zone. A sudden change of L_D is noted both around 45 m and 60-65 m deep, with values varying from ~0.4 to ~0.6 mm, and from ~0.4 to >0.8 mm, respectively. From 65 m deep down, the range of L_D is large, and low values of 0.1-0.3 mm are also found. At depths 60-95 m, samples from 1988 drilling have L_D and residence time much more variable relative to the 1981 samples studied here. This is possibly the result of internal evolution through time. To a certain extent, the stronger variations observed at both shallower and deeper levels in the lake could be due to the effect of foundered crust at both upper and lower margins of the lake.

Characteristic length data for the samples from the margin of the lake are less variable than that observed for the centre (Fig. 2-10b). Until 15-20 m deep, the greatest variations are noted for the whole margin, with L_D varying from 0.3 to 0.9 mm. At depths 20-65 m, L_D appears more stable, with a limited range of 0.4-0.6 mm. However, no samples were studied between 25-60 m depth because olivine was not sufficiently abundant for a precise CSD to be determined. At depths >65

m, L_D varies from 0.35 to 0.75 mm. Overall, the textural variations observed for the whole lake margin are thought to reflect the presence of two distinct olivine populations throughout the entire thickness of Kilauea Iki lava lake: (1) a population of smaller crystals which estimated L_D is 0.30-0.65 mm; and (2) a population of bigger crystals with an estimated L_D of 0.7-0.9 mm. Those two populations correspond to the subgroups 2a and 2b previously defined.

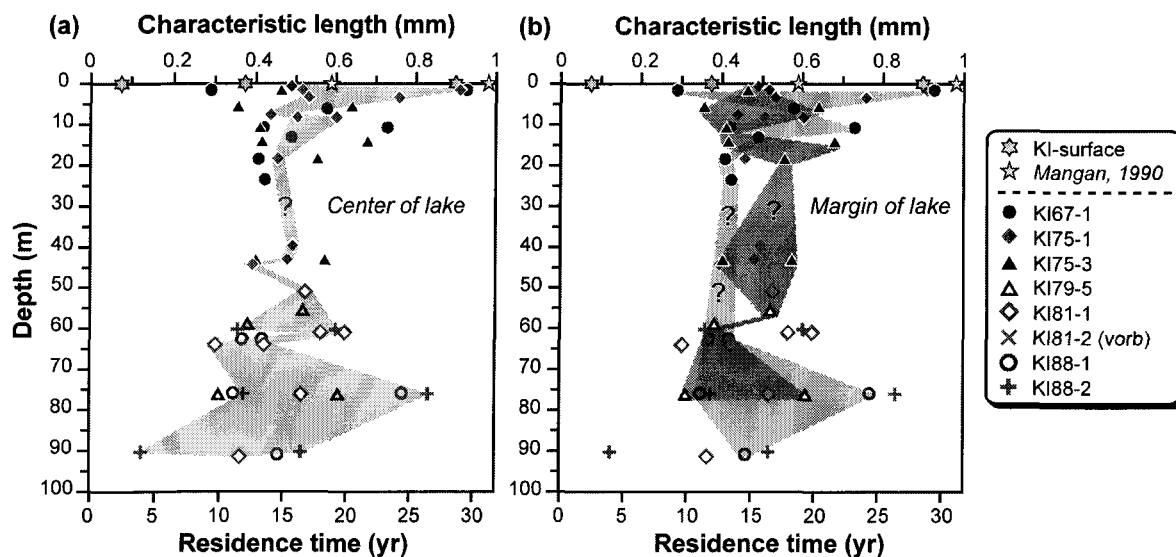


Figure 2-10. Characteristic length plotted against the depth in the lava lake. As in Figure 2-8, curved CSDs have two values for L_D . (a) The green-filled area shows the bulk variations for the centre of the lake. (b) The red-filled area shows the bulk variations for locations lying towards the edge, and the grey-filled one the bulk variations for the edge of the lake.

2.5.2.6 *Size distributions of deformed crystals*

The size distributions of deformed crystals may reveal aspects of how they were incorporated into the magmas. However, this is not easy to do as it is difficult to identify all deformed grains, especially the small ones. However, we feel we can consistently recognize deformed crystals larger than 0.15 mm, the size of the smallest, clearly deformed grains that were identified in this study.

In most samples the number of deformed crystals was so low that separate CSDs could not be constructed. Hence, we have compared the CSD of the total number of crystals with that of the undeformed crystals only (Fig. 2-11, and Appendix 2-1). In figure 2-11 three examples of the varying effects of the deformed crystals on the total CSDs are shown, for three distinct total amounts of deformed crystals. KI75-3-60 has only 0.9 vol. % deformed crystals and displays little differences in the CSD of undeformed versus total olivine populations (Fig. 2-11a). KI67-1-35.1 has an intermediate amount of deformed crystals (2.2 vol. %) and shows significant CSD changes for the larger crystal sizes (Fig. 2-11b). KI88-2-249.5 has a greater abundance of deformed crystals (5.7 vol. %) and thus exhibits marked contrasts between the two different CSDs (Fig. 2-11c); the undeformed CSD only extends to 2.4 mm and its slope is much steeper than that of the total CSD. Overall, we see that the difference between the total and undeformed CSDs is non-uniform throughout the entire CSD, being greater at larger sizes and almost non-existent at smallest sizes. In addition, no conspicuous link is noted with the depth, the location in the lake, or the vorb position.

In summary, all CSD plots show that deformed crystals are commonly larger than undeformed crystals. The deformed crystals hence appear to strongly affect the CSD, even where they are relatively few in number.

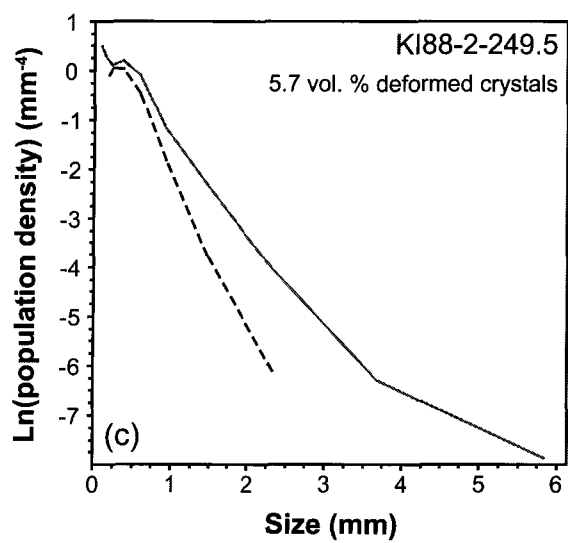
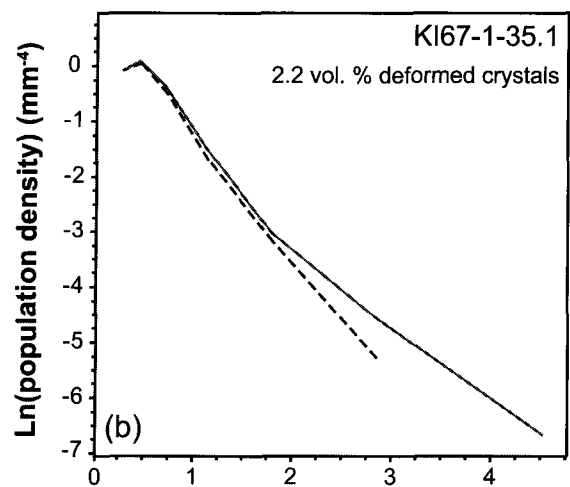
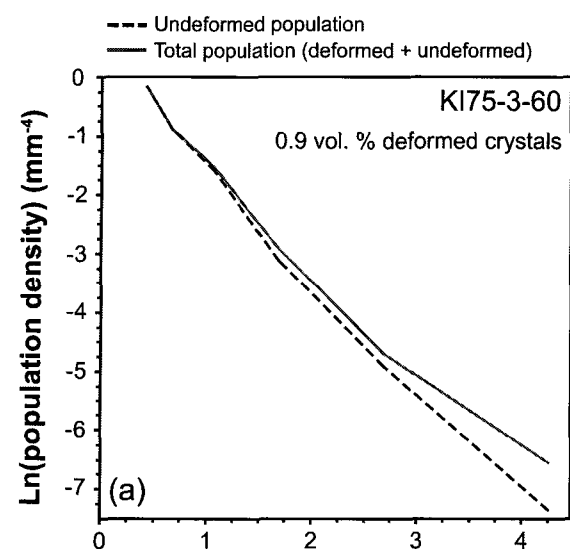


Figure 2-11. Representative CSDs of undeformed versus total olivine populations, showing the effects of deformed crystals on the total crystal size distribution. (a) KI75-3-60, with a low amount of deformed crystals (0.9 vol. %), displays little differences in the CSD of undeformed versus total olivine populations; (b) KI67-1-35.1, with an intermediate amount of deformed crystals (2.2 vol. %), shows significant CSD changes for the larger crystal domain; and (c) KI88-2-249.5, with a greater abundance of deformed crystals (5.7 vol. %), exhibits marked contrasts between the different CSDs, which are greater for large sizes. A quantitative estimate of the abundance of deformed crystals in each sample is shown at the top right of each plot.

2.5.2.7 *Olivine size distribution in the vorbs*

The original definition of vorbs included a crystal size aspect (Helz, 1980). It is therefore interesting to see if this is confirmed by quantitative textural measurements. The CSDs of the vorb part of KI75-1-141 and KI75-1-145 are very similar, except that KI75-1-145-vorb stops at 2.8 mm (Fig. 2-12a). In contrast, the CSD of KI81-2-185.7-vorb is rotated to a shallower slope with respect to the other vorbs (Fig 2-12a) and the CSDs of non-vorbs at similar depths (Fig. 2-7). It would be interesting to compare the vorb with non-vorb CSDs at those depths, but unfortunately, only KI81-2-185.7 has sufficient crystals in the non-vorb part of the sample to build an individual CSD (Fig. 2-12b). In this case, the vorb and non-vorb CSDs are clearly different. The vorb CSD has a bulk slope flatter than the non-vorb CSD, lower population densities at crystal sizes <1.5 mm for higher population densities at sizes >1.5 mm, and no change of the slope at sizes larger than 2.8 mm.

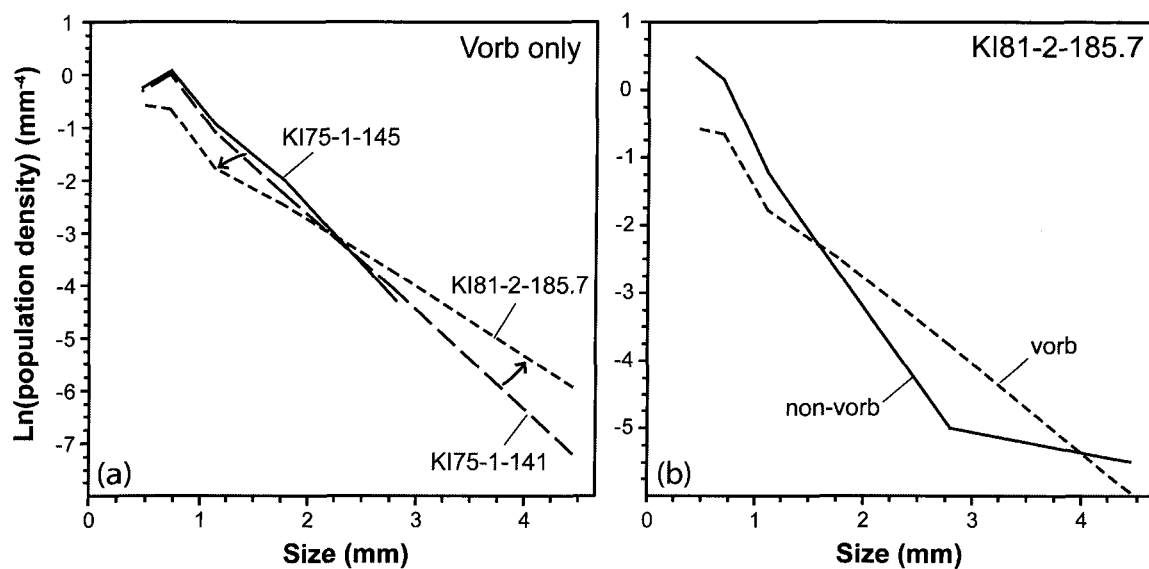


Figure 2-12. Crystal size distribution (CSD) of (a) the vorb-only part of selected samples; and (b) the vorb and non-vorb parts of KI81-2-185.7.

2.6 DISCUSSION

The first part of this section deals with olivine residence times in the whole Kilauea Iki system, that is in both the magmatic plumbing system and the lake itself. The next part is organised chronologically, as far as possible, from the source of the olivine crystals to the end of the cooling of the lake. It is divided into two parts: (1) processes that occurred before eruption, and; (2) processes that occurred in the lava lake. Then, geochemical and textural data are discussed to decipher how the variations are accounted for, and to draw a coherent petrogenetic model. Some aspects of a model, such as crystal nucleation, growth, and deformation, or coarsening, will only affect the rock texture. Other processes will affect both texture and geochemistry, such as mixing or crystal fractionation and accumulation.

2.6.1 Residence times of olivine in Kilauea Iki magmas

Crystal residence times can be estimated from S-type CSDs (Higgins, 2006b) if a CSD model is chosen, such as that of Marsh (1988b), and a growth rate defined. This model typically refers to simple nucleation and growth in a steady-state magma system. It is difficult to precisely define the growth rate; hence we use a range of values from 2.5×10^{-9} mm/s (G_1) to 5×10^{-10} mm/s (G_2), based on studies of similar rocks (Cashman, 1993, Mangan, 1990, Marsh, 1988, Marsh, 1998, Wright & Fiske, 1971). Residence time (τ) and L_D are related by the equation:

$$\tau = \frac{L_D}{\text{growth rate}}$$

Hence, the variations observed in this figure are qualitatively identical to that described for L_D in Section 4.2.4. The crystal populations from group 1 have mean crystal residence times of around 1-1.5 years when using G_1 , and 4-8 years when using G_2 (Table 2-4). Crystal populations from group 2 have calculated magma residence times significantly higher with values ranging from 4 to 12 years using G_1 in the calculations, and from 18 to 59 years using G_2 .

In summary, the shorter times of 1-8 years estimated from group 1 olivines are thought to represent the residence time of olivine crystallized both in the shallowest magma storage region before the eruption, and in the lava lake. Group 2 ages of 4-60 years reflect residence times of olivine crystallized deeper in the Kilauea Iki system before the eruption. These residence times agree with that determined by Mangan (1990) from olivine CSDs. She found an average olivine residence time of 11 ± 4 years for the 1959 Kilauea Iki eruption, with a growth rate of 10^{-9} mm/s. If we recalculate her data with our method, we find a range of residence times of ~18-30 years, using the same mean growth rate of 10^{-9} mm/s. Our results also accord well with the olivine residence times determined by Vinet and Higgins (2010) from olivine CSDs for Mauna Ulu lavas, Hawaii. They proposed olivine residence time ranges of 1-15 years and 3-40 years for the two texturally different populations revealed in the 1969-74 lavas, using the same growth rates as presented here.

Crystallisation timescales can also be discussed with regard to location and depth of the samples in the lake (Fig. 2-10). The residence times plotted in figure 10 were calculated with the use of a mean olivine growth rate of 10^{-9} mm/s, instead of a range of values presented above for clarity of the diagram. The emphasis is now given on the bulk quantitative changes of the residence times. Olivine residence times for magmas from the centre of the lake show greater variations than those from the margin (Fig. 2-10a). In addition, greater values are found at both

shallower and deeper levels for magmas from both the centre and margin. At depths <10 m, the residence times range from ~14 to 29 years, whereas it is more constant at 15-50 m depth, with values around 15 years. Deeper than 60-65 m, the values vary again more strongly from 4 to >25 years. For olivines from the margin of the lake, the residence time variations are less pronounced (Fig. 2-10b). At depths <10-15 m, the residence time varies from 10-30 years, with little variations at the edge of the lake. At depths 15-65 m, it appears more stable, centered around 12-18 years, although only few samples have been analysed. At depths >65 m, the residence time varies from 10-25 years

The variations of residence time (and other textural data) may reflect different crystal populations present throughout the entire Kilauea Iki lava lake. Overall, two distinct olivine populations are recognised both at the centre and margin of the lake, which correspond to the previously defined subgroups 2a and 2b: (1) subgroup 2a olivines, which are smaller crystals with estimated residence times of 10-20 years; and (2) subgroup 2b olivines, which are bigger crystals with an estimated residence time of ~24-30 years. To those, a third population of residence time <5 years, which corresponds to group 1 previously defined, may be added, but it is only present in a single sample, KI88-2-296.1. Our surface scoria samples contain the three olivine populations, whereas the eruptive pumice samples of Mangan (1990) does not include the group 1 olivines. This discrepancy is probably due to her lower CSD analytical precision.

2.6.2 Processes active in Kilauea Iki magma system before eruption

Although this paper focuses on processes active in the lava lake, it is necessary to start by drawing a synthetic portrait of the major processes at work in the magma system before the

eruption. Based on our geochemical and textural results, we consider that crystal or magma mixing, coarsening, crystal deformation and crystal aggregation played an important role in the formation of the 1959 Kilauea Iki magmas. Some of these processes were also active during the solidification of the lake. Consequently, a clear distinction and quantification of pre- and post-eruption processes is not always easy.

2.6.2.1 *Mixing*

Mixing is a common phenomenon in igneous systems and evidence for it can be found in field, textural and geochemical data. Mixing can involve distinct olivine populations from separate magmas or solid sources. We consider here both the role of magma and crystal mixing in the origin of the 1959 Kilauea Iki products.

The curvature of CSDs can be used as evidence for mixing: the mixing of two magmas with non-collinear straight CSDs produces a curved concave-up CSD (Higgins, 1996). This is due to the logarithmic vertical scale of a classical CSD diagram. This model can also be applied to a rock formed by two consecutive processes, each process providing straight CSDs. In either case, the origins of the larger and smaller crystal populations can be treated separately. Textural data has shown that mixing is important for the formation of some other basaltic rocks (e.g., Higgins, 2009; Vinet and Higgins, 2010).

Most of our samples are from the lava lake; hence the textural and chemical data acquired from these samples may have been affected by processes that occurred after the eruption in the lava lake. Consequently, data from these samples do not fully represent the original features of the magma system beneath Kilauea Iki, but instead, they likely are the result of a superposition of

processes of different time scales. The CSD and chemical data for the lake samples must thus be used with care here, and emphasis is given here to the surface samples.

Our surface CSDs are strongly curved concave-up (F-type), reflecting the presence of two or three olivine populations. The great similarity between our mean surface CSD and the recalculated scoria CSD from the data of Mangan (1990) is noteworthy (Fig. 2-13), and supports our results. The population of small crystals is proposed here to be the result of rejuvenation occurring just before or during the eruption, as the high value of the intercept and the steep CSD slope suggest a high nucleation rate and shorter residence time. The population of larger crystals must have been formed earlier than the small crystal population, and probably deeper in the plumbing as well, due to its relative higher amount of deformed grains. Due to its flatter CSD slope and lower intercept, this population was formed under lower nucleation rates during a longer time period, which is consistent with a smaller undercooling. Several locations within Kilauea plumbing may satisfy these characteristics, for example deeper levels in the summit chamber or even deeper cumulate layers at the base of the edifice. These latter possibilities have also been recently proposed by Vinet and Higgins (2010) in their textural and geochemical study of 1969-1974 Mauna Ulu eruption.

Magma mixing can be recognised using a series of chemical arguments, as well. Whole-rock and olivine compositions, along with petrographic data have been extensively discussed in previously published papers (e.g. Helz, 1987b; Murata and Richter, 1966a; Nicholls and Russell, 1991; Richter and Murata, 1966; Wright, 1973; Wright and Fiske, 1971). Based on major element data of eruptive samples from most phases, Murata and Richter (Murata and Richter, 1966a) and Wright (1973) recognised two different lava compositions, S-1 and S-2, and thus argued in favour of magma mixing during the eruption. Based on chemical and textural data of both eruptive scoria

and lake samples, Helz (1987b) also proposed a model in which there was mixing of two magmas with variable amounts of the magma components S-1 and S-2 regarding the eruptive phase. In several studies it is argued that a MgO-rich magma intruded into and mixed with several differentiated magmas during the Kilauea Iki eruption (Helz, 1987b; Schwindinger and Anderson, 1989; Wright and Fiske, 1971). The MgO-rich magma entrained olivines with gas and melt inclusions trapped at 5 kbars (e.g., Harris and Anderson, 1983). This MgO-rich magma is thus suggested to have risen from deep levels, i.e. the lower crust, within the Kilauea plumbing. Hence, it may have bypassed the main conduit and summit storage region, and mixed with the differentiated magmas at shallow depths (Anderson, 1995; Helz, 1987b). On the other hand, most of the olivines contain inclusions trapped at pressures <1 kbar, thus originating from shallow depths (Anderson and Brown, 1993). Alternatively, Murata and Richter (1966a, b) proposed a model in which the shallow magma chamber that fed the eruption was compositionally stratified: the differentiated magma at the top was erupted first, followed by the olivine-rich magma at the bottom.

In summary, the olivine populations found in our surface samples may have formed in different parts of the plumbing system, and undergone different crystallisation histories. Finally, they may have mixed some time before or maybe partly during the eruption.

The concave-up curvature of at least half of the lake CSDs is strong evidence for magma or crystal mixing. However, there is probably an overprinting of early (before eruption) and late (intra-lake) processes we should take into account in these CSDs. Although we feel that late mixing is of little importance and early mixing likely to be prominent because of the strong curvature of the mean surface CSD, for example, no estimate can be determined from the textural data, as it is difficult to distinguish them only using the final products. Furthermore, no strong petrological

evidence of mixing is noted among the lake olivine crystals studied here, which accords with the conclusions of earlier studies such as that of Helz (1987b).

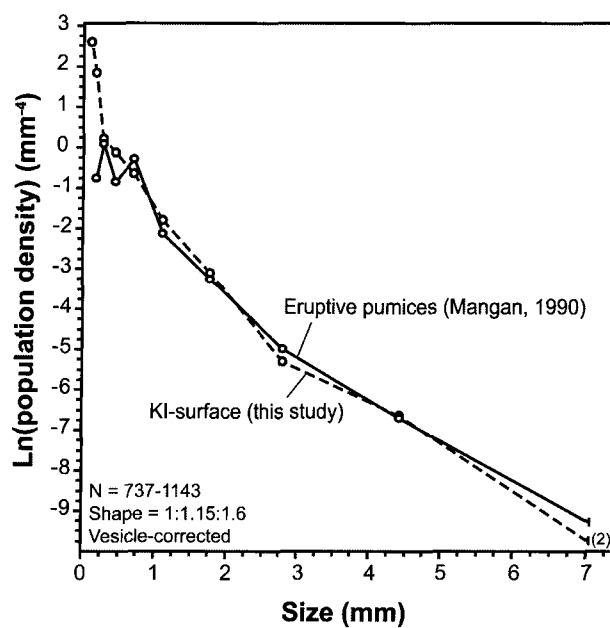


Figure 2-13. Comparison of our mean surface CSD versus recalculated pumice CSD from data of Mangan (1990). Number in parentheses at the right end of the KI-surface CSD refers to the number of grains analysed in the last CSD bin (largest crystal sizes).

2.6.2.2 *Coarsening*

In an igneous context, coarsening (Ostwald ripening, textural equilibration) is the solution of small crystals and the simultaneous growth of larger ones as a process of total surface energy minimization (see review by Voorhees, 1992). This process is most important at temperatures close to the mineral liquidus. Recently, Simakin and Bindeman (2008) showed that coarsening can be driven more effectively by thermal oscillations in the magma. The theoretical CSD evolution of magmas undergoing coarsening has been reviewed by Higgins (2006a). As an ideal result of coarsening, a straight CSD will be transformed into a hump-shaped CSD, lacking small crystals (Higgins, 1998; Higgins, 2006a). As coarsening progresses the slope of the right side of the CSD will decrease, as a result of the increase in growth rate to a maximum with grain size. In a closed system containing minerals of uniform composition, the total chemical composition is unaffected by coarsening.

The surface CSDs do not show turndowns at the left side of the plot. This could be considered as evidence against significant coarsening in the plumbing under Kilauea Iki before the eruption. However, we have only studied a restricted number of samples. Alternatively, rejuvenation after coarsening may have produced a new population of small crystals, hence overprinting the initial turndown at small sizes of the CSDs produced by coarsening. The implication of coarsening on the surface CSDs is difficult to rigorously test. One possibility may be that the population of larger crystals, which also contains many deformed grains, was partly produced by coarsening. In similar rocks from the 1969-74 Mauna Ulu eruption, Vinet and Higgins (2010) suggested that the main source for the population of large, coarse, and very often deformed olivine is a deep-seated cumulate lying at crustal levels at the base of the edifice. They

also argued in favour of some coarsening in the history of this specific olivine population that experienced variable degrees of deformation.

A number of lake samples do have a turndown at the left side of the CSDs. These samples have thus been coarsened. It might have happened in the lava lake during solidification, or earlier in the magma plumbing system of Kilauea Iki.

2.6.2.3 *Deformation of crystals*

Deformed olivine crystals are ubiquitous both in the scoria and lake samples, with an abundance of 0.2 to 12 vol. %, or 2-31 % of the total crystal population (Fig. 2-6; Table 2-2). They are of great interest here for the evaluation of the origin of the different crystal populations, and subsequently for the study of olivine distribution or redistribution in the lava lake. It is proposed here that the deformed crystals are not mantle xenocrysts, but instead are antecrysts, in the sense of that term as used by Jerram and Martin (2008). This is similar to the ideas of Helz (1987b) for these specific Kilauea Iki olivines. There are also antecrysts of deformed clinopyroxene.

The assertion that some olivine crystals are antecrysts is in accord with the popular idea that the deformed crystals in other volcanic rocks in Hawaii are unlikely to represent mantle xenocrysts or unrelated cumulates from different volcanoes (Baker et al., 1996; Garcia, 1996; Norman and Garcia, 1999; Wilkinson and Hensel, 1988). This idea is consistent with:

- (1) the magmatic origin shown by the fairly high CaO values, around 0.20-0.30 wt. %, of the scoria olivines (e.g., Norman and Garcia, 1999; Simkin and Smith, 1970; Thompson and Gibson, 2000);

- (2) the persistence of melt and Cr-spinel inclusions in both deformed and undeformed crystals;
- (3) the very slight or absence of compositional differences between deformed and undeformed olivines, even in minor components such as CaO, NiO and MnO, that suggests that the deformed olivines have a cognate origin with the undeformed ones.

Strictly speaking, these arguments apply only for the scoria olivines since the composition of the lava lake samples and their olivines must have been modified by intra-lake internal differentiation and reequilibration throughout the entire cooling period. We have thus avoided a comparison of lava lake olivine core contents versus olivine composition of the mantle domain, which may be unreliable in this case.

Sakyi *et al.* (2008) studied the dislocation microstructures in olivine grains from Kilauea. They found high densities of dislocations ($>10^6 \text{ cm}^{-2}$) and thus suggest that the olivines have experienced high-temperature plastic deformation. This interpretation is in agreement with the kink-banded olivines observed in our samples, given that kink-bands normally form upon migration of dislocations during thermal restoration at high temperature. The appearance of those kink-bands attests to a strong deformation regime (Nicolas and Poirier, 1976), in the solid state (e.g., Green II and Radcliffe, 1972).

Helz (1987b) proposed that the deformed olivines of the 1959 Kilauea Iki eruption were derived originally from conduit walls, lined by deep crustal cumulates from earlier Kilauea tholeiites. Wilkinson & Hensel (1988) proposed that the deformed olivines may be cognate with their hosts and only reflect phenocryst deformation during magma flow through narrow feeders and conduits, at elevated (but unknown) pressures. In their review of geochemical and geophysical data on Piton de la Fournaise volcano, La Réunion Island, Peltier *et al.* (2009) suggested that

deformed olivine antecrysts were derived from the walls of the summit magma chamber or from olivine-enriched layer stored at the base of the reservoir. Others argued in favour of a dunitic cumulate body formed at the base of, and partly under the shallow reservoirs, at a depth of 4-10 km within the edifice (Clague and Denlinger, 1994; Okubo et al., 1997; Wright and Klein, 2006; Wyss et al., 2001). Vinet and Higgins (2010) reintroduced this idea of cumulate-derived olivines by associating it with the influence of the active decollement lying at the base of the edifice, at 9-12 km deep.

Although the deformed crystals have a different history from the undeformed crystals, they have very similar compositions. Hence, some undeformed crystals may have the same origin as the deformed ones in the decollement zone. Perhaps there were domains isolated from stress where olivine grains remained undeformed.

2.6.3 Processes active in Kilauea Iki lava lake

2.6.3.1 Crystal settling

The solidification of a thick magma body such as the Kilauea Iki lava lake is commonly assumed to be dominated by crystal settling and accumulation, although debate is still active. In particular, olivine settling is thought to be important in the solidification and differentiation of the Kilauea Iki lake products, on the basis of their chemical and petrological effects (e.g. Hagerty et al., 2006; Helz, 1980, 1987a; Helz et al., 1989; Jellinek and Kerr, 2001; Richter and Moore, 1966). Settling of coarser olivine crystals in the lava lake was proposed to have mostly controlled the formation of an olivine-depleted zone between 18 and 40 m depth (Helz, 1987a; Mangan and Helz,

1985). This process is thought to have been efficient only from 1963 to 1972 (Helz, 1980), mostly resulting in the current distribution of olivine in the lake. Olivine settling is also argued to be a major process involved in the differentiation that occurred in the magma chamber before the 1959 Kilauea Iki eruption (Murata and Richter, 1966b; Richter and Murata, 1966).

The abundance of olivine determined in our study can also be examined to see if it supports olivine settling. The distribution of olivine in the centre of the lava lake follows the generally settling-related S-shaped pattern with respect to depth noted by Helz (1980; 2009) (Fig. 2-14; Table 2-2): an olivine-rich zone at upper levels (<18 m), with up to 35 vol. % olivine; an olivine-depleted zone at intermediate levels (18-40 m), with olivine contents of 7 vol. % in KI75-1-60 (~18 m) and 11 vol. % in KI75-1-130 (~40 m); and an olivine accumulation zone at deeper levels (>45 m), with values up to 30 vol. %. This pattern is less pronounced for the margin of the lake, and even totally absent for the very edge that has only 13-18 vol. % olivine over the upper 40 m. Although only a few samples from the depths corresponding to the olivine-poor zone are studied here, the strong decrease in olivine abundance seems significant. To a lesser extent, this decrease in total percent olivine is noted as well at the margin of the lake, with 15-17 vol. % in KI67-1-60.4 (18 m) and KI67-1-77 (24 m). This tendency is also observed in a plot of L_{\max} versus depth (Fig. 2-9). In addition, we have examined several other thin sections of different drill cores from the depth range 18-40 m in which the olivine content was very low, <5-10 vol. % typically, and olivines >2 mm in size essentially absent. The olivine distribution pattern of the accumulation zone is similar for samples from both the centre and margin of the lake, with slightly lower values for the very edge. In a first view, our data seem thus to support the idea of crystal settling.

Crystal settling is commonly proposed based on chemical data or total olivine abundance, but recently a few studies have used textural data such as the CSD analysis (e.g. Higgins, 2002b;

Hoshide *et al.*, 2006; Marsh, 1998). In order to constrain how olivine settling is likely to affect the lake CSDs, we have applied a simple numerical model (Fig. 2-15) developed by Higgins (2002b). It deals with the separation of crystals and liquid in response to differences in density, following the Stokes' law. Crystals are initially distributed throughout a magma column and are allowed to descend to an accumulation zone at the base. Drag or interactions between particles was not considered. Because of the simplicity of this model, it should only be viewed as a point of departure, which is probably more accurate for the initial stages of the process. It is also useful as a proxy for constraining time limits of the settling process.

The model begins with a population of 1 million spherical olivine crystals that have a straight CSD. The initial CSD was set using the parameters of the middle segment of our mean surface CSD (size range 0.5-3.0 mm), but with only 1% of the crystal concentration (see Fig. 12 in Higgins, 2002b). This model is more dilute than reality so that crystals would not 'overflow' from the accumulation zone. Other input parameters were the height of the column (120 m), the thickness of the accumulation zone (1 m), the time interval (variable), the density of crystals and magma (3500 and 2700 kg/m³, respectively), and the viscosity of the magma (100 Pa.s). Density and viscosity of magma are that of Jellinek and Kerr (2001). In Figure 2-15 are represented the initial CSD used in the modeling versus the modeled CSDs after settling along different time scales. Some lake CSDs at depths around 60 m and greater have also been plotted for scale and result validation. The modeled CSDs are those of the bottom part of the magma column, which includes both the accumulated crystals and those still suspended in the liquid (see Higgins, 2002b).

Initially, the modeled CSDs remain linear and rotate around the intercept with the slope being shallower. By increasing the time intervals, the right ends of the CSDs reach a limit as all the crystals are precipitated, and the intercept begins to increase. The late stages coincide with the

appearance of a turndown for small crystals. Our simple model indicates that, for small amounts of olivine settling, the only effect on the CSD of the 'cumulate' zone will be that the slope becomes shallower.

The differences in slope of the modeled CSDs can be used to estimate the settling time necessary to produce the slope variations of our lake CSDs. They are reproduced after less than 100 hours of settling, which is actually too short in reality. Also, with such a model we can say that settling time is unreasonably long (>15 years) to produce turndowns for small crystals that we see and hence that the progression of increasing turndown with time is more likely due to coarsening. Thus, the CSDs may have been modified and overprinted by intra-lake coarsening. This hypothesis has previously been proposed by Higgins (2002b), for example, for the Kiglapait layered mafic intrusion, Labrador.

In summary, settling does not appear to be of significant importance here. This assertion is similar to that proposed by Cashman and Marsh (1988) for the 1965 Makaopuhi lava lake, based on the straightness of the plagioclase and opaque phase CSDs. Furthermore, the characteristic length of the Kilauea Iki lake samples does not vary much throughout the entire thickness of the lake, whereas it should because of the contrast of density between olivine and magma. In contrast, the volume proportion of olivine varies by a factor of up to six or seven.

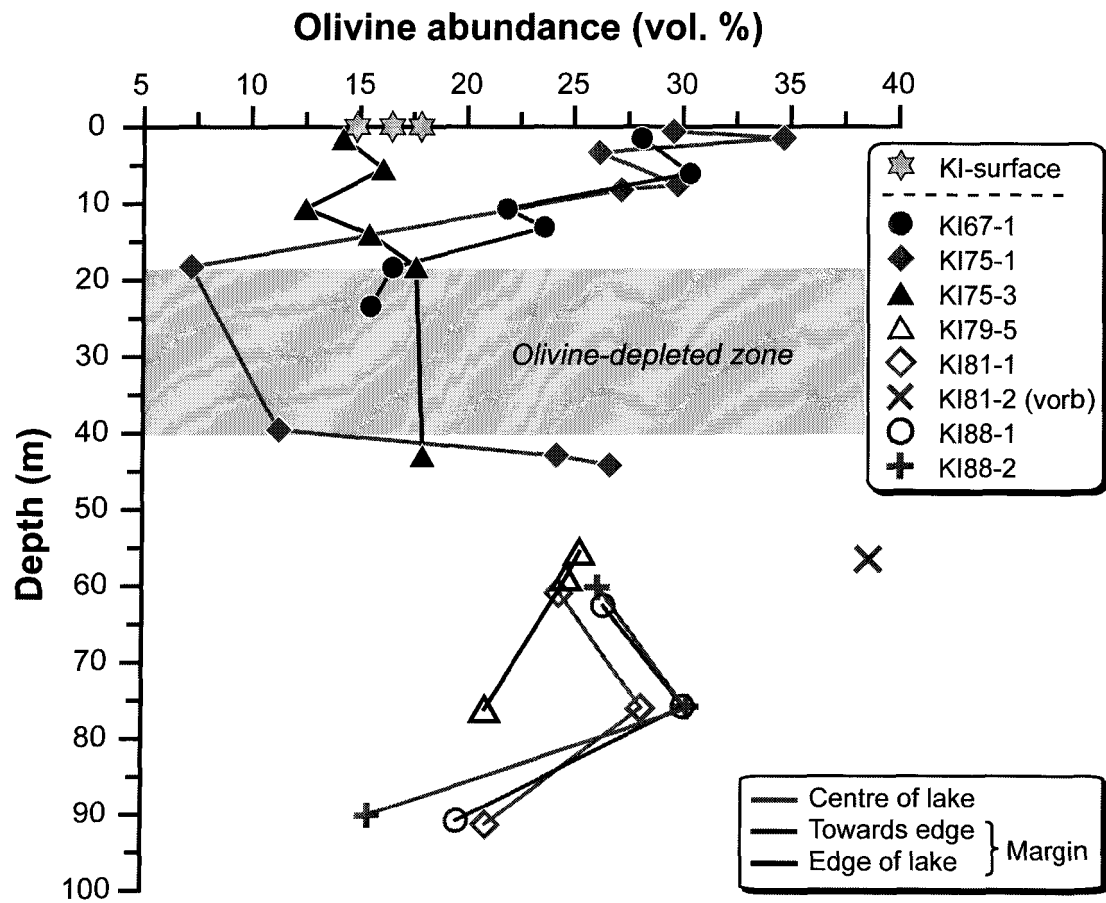


Figure 2-14. Olivine abundance (in volume %) plotted against the depth in the lava lake. Centre and margin of the lake are separated and represented in different colours.

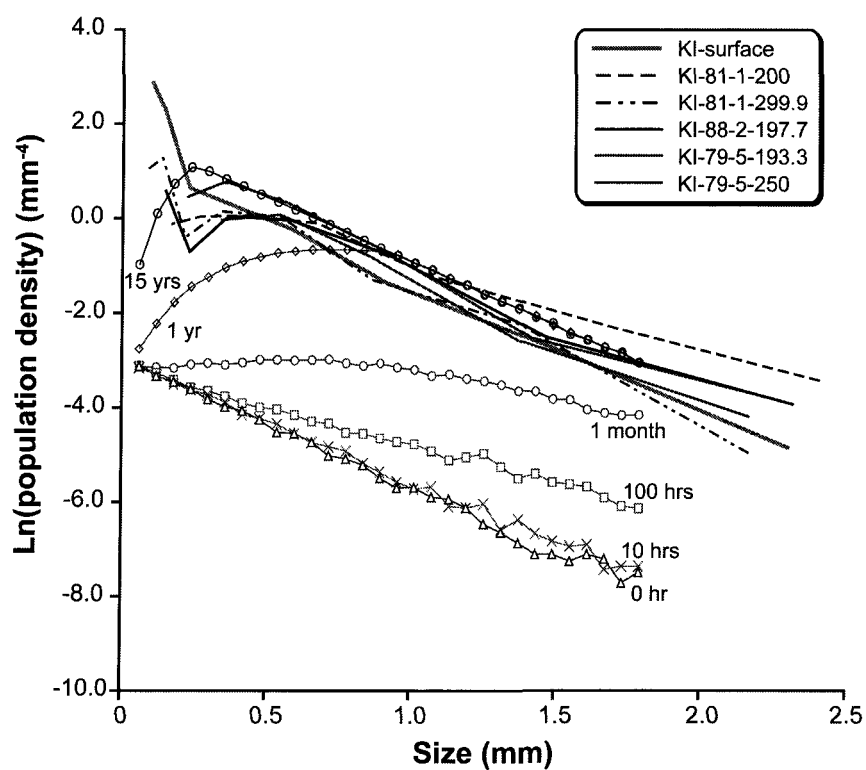


Figure 2-15. Modeled CSDs after olivine settling. Some lake CSDs at depths around 60 m and greater have also been plotted for scale and result validation. The initial CSD was taken to be the straight middle segment of our mean surface CSD, at size range 0.5-3.0 mm. The model used here follows the procedure of Higgins (2002b).

2.6.3.2 *Convective or advective redistribution of crystals*

In igneous geology, natural convection motions of magmas or melts are generated by density differences due to thermal or compositional gradients. Advection is the relative transport of matter, such as crystals, or heat, by the larger-scale motion of currents in magma or melt. This may be of thermal, compositional or other origin. It is actually difficult in natural environments to distinguish which process really occurs relative to the other, which is why they are treated here together. Both processes are commonly involved in the redistribution of crystals and other materials; thus, the process of differentiation in many thick magma bodies generally occurs both at depth (in magma chambers) and at surface levels (in active lava lakes) (e.g. Harris *et al.*, 2005; Witham and Llewellyn, 2006). The occurrence of significant convection or advection in passive, or inactive lava lakes (see definition in Harris *et al.*, 2005; Swanson *et al.*, 1979) is heavily more debated (e.g. Davaille and Jaupart, 1993; Evans and Moore, 1968; Hermance and Colp, 1982; Hort *et al.*, 1999; Jellinek and Kerr, 2001; Moore and Evans, 1967; Rupke and Hort, 2004; Shaw *et al.*, 1968; Worster *et al.*, 1993; Wright and Okamura, 1977). Below, we will only use the term 'convection' for the purpose of simplification.

In natural systems such as the Kilauea Iki lava lake, the timing, extent and location of convection is difficult to evaluate. However, this process may have a significant control on the way the crystals are redistributed in the magma, and thus, may lead to some textural record. Hence, the CSD approach may help in constraining the possible effect of convection on the redistribution patterns of olivine crystals in the Kilauea Iki lava lake. To evaluate this possibility we have chosen to use the same CSD numerical model and its parameters as for settling (Higgins, 2002b), but with the addition of convection velocity as another input parameter. Upward convection competes

with settling whereas downward convection enhances it. One should expect that, if convection occurred in a significant way both in terms of time and space, it would result in a significant textural response possibly seen in the CSDs. However, a problem arises here. If we use realistic advection velocities, such as $\sim 1.4 \times 10^{-3}$ m/s, which corresponds to one overturn per day, then we see no settling at all or complete settling, depending on the source of the vertical motion. Hence, CSDs cannot be used to determine if there was convection.

Absence of significant large-scale convection in the lava lake was proposed in previous studies (Barth et al., 1994; Gibb and Henderson, 1992; Helz, 1980; Helz et al., 1989). It accords with the conclusions of Marsh (1989) that convection is only possible when the crystal fraction is low, although no precise values can be easily estimated. However, it is in marked contrast with the view of Jellinek and Kerr (2001) of a “vigorous stirring in the magma”, driven by descending thermal convection and ascending compositional convection. Also, it somewhat contrasts with the convective flow differentiation of crystals proposed for the 1965 Makaopuhi lava lake (Cashman and Marsh, 1988; Wright and Okamura, 1977), which was thought to be particularly active in the deeper, non-vesiculated lava of this lake.

Convection movements can be associated with diapiric transfers, which are considered here as the mechanisms responsible for the vorb formation. Segregation veins and vorbs are widely distributed in the Kilauea Iki lava lake (Helz, 1979; Helz, 1980, 1987a; Helz et al., 1989; Mangan and Helz, 1985, 1986; Richter and Moore, 1966). The focus is only given here to the vorbs, and not to segregation veins, since the latter are not present in our samples. Vorbs were interpreted to play an important role in the evolution of the lake magma, but only in the depth range 18-58 m, with the 40-58 m interval being more intensely affected (Helz *et al.*, 1989). In this study, most textural criteria of Helz have been recognised. We also observed that all vorbs are from the centre

of the lake, which is consistent with the observations of Helz (1980) and Helz et al. (1989) that few or no vorb structures have been recognised at the margin of the lake. Slow convective movement can produce the vorb CSDs. The vorb part of KI81-2-185.7 is a good example for that. Relative to the mean surface CSD, this vorb has a CSD rotated clockwise around 1.5 mm size (Fig. 2-7). This CSD pattern, much more conspicuous for this vorb than for the vorbs in KI75-1-141 and KI75-1-145, is likely a result of the advective currents upward induced by the diapiric transfer responsible for the vorb formation. In summary, the vorb-related diapiric transfer is thought here to have participated in the olivine redistribution through the lava lake during its cooling, but only for the low-Fo group and in a narrow-scale manner.

2.6.3.3 *Final crystal growth*

The presence of several olivine populations, with zoned or resorbed crystals, along with a significant proportion of deformed olivines, suggest that crystallisation of the Kilauea Iki magma(s) occurred at different times and locations or depths in Kilauea's magma system. Several authors have argued that post-emplacement crystallisation has occurred in thick basaltic magma bodies, such as Kilauea Iki lava lake (e.g. Jellinek and Kerr, 2001; Richter and Moore, 1966; Scowen et al., 1991), as well as the prehistoric and modern Makaopuhi lava lakes (Evans and Moore, 1968; Hagerty et al., 2006; Moore and Evans, 1967; Wright and Okamura, 1977). But how can we distinguish crystals that formed before the eruption from crystals that nucleated and grew later within the lava lake? In the Kilauea Iki lava lake, we noted that the rim Fo content follows a normal Gaussian distribution throughout the entire thickness of the lake, around a mean of Fo₇₄ (Fig. 2-3). We also see a minor population of crystals with this as a core composition, particularly towards

the base (>60 m), and edge of the lake. Undercooling was strong here, which produced high nucleation rates that allowed rejuvenation. However, the volume proportion of newly formed (rejuvenated) crystals in the lake was not quantified for our samples.

Some of our straight lake CSDs, with no apparent turndowns, may result from post-emplacement growth. This is also applicable for the steep part at the left side of the CSD for samples from 1988 drillings (mostly at the centre and a little at the margin of the lake), or samples from the first 15 m of the lake, at the northern edge. This minor feature is possibly attributable to rejuvenation.

2.6.3.4 Origin of the chemical and mineralogical vertical stratification

One of the major issues of this study lies in the understanding of the mechanisms that originated and maybe preserved the vertical stratification of the Kilauea Iki lava lake, as defined by the olivine characteristics. There is no simple explanation to account for the presence at depths >60 m of olivine crystals with core Fo content of 72-81%. Similarly, all Fo-rich olivines are constrained to the first 60 m of the lake. This cannot be ascribed to full crystal reequilibration with the residual, more evolved melt in the lake. This view is in marked contrast with the work of Scowen et al. (1991) that argued in favour of reequilibration by cationic diffusion, with Mg, Al, Cr moving outwards from the olivine, and Fe and Ti inwards. We feel that Fe-Mg diffusion between olivine and the interstitial melt would not be efficient enough, even at magmatic temperatures around 1100°C, to allow such reequilibration to extend to every crystal core. Our analytical data show that only the rim has been affected. Also, there is divergence between the composition of surface olivine and that of lake materials, the former being more Fo-rich.

Another question arises as how to explain the higher ratio of deformed crystals to the total population at depths >60 m, compared to shallower levels in the lake? One possibility is that deformed crystals were generally larger, and hence sank faster. However, crystal settling was shown above to be inefficient and hence no simple explanation can be found for this data either.

Several early studies suggested that the shallow magma chamber that fed the 1959 Kilauea Iki eruption was chemically and mineralogically stratified (Murata and Richter, 1966b; Richter and Murata, 1966): the upper zone was composed of an assemblage olivine (Fo_{82}) + cpx + plg; the lower zone was composed of coarse, euhedral to rounded olivine Fo_{86-88} ; and the base was formed by an olivine cumulate that was subsequently mobilised during episodes of high rates of lava discharge. To a certain extent, this idea is consistent with our observations of the presence of olivines Fo_{85-88} in the upper 55-60 m, and of olivines having compositions around Fo_{80} at depths >60 m. The lake stratification may thus be representative of this initial chamber stratification. In support of this idea is the variation of whole-rock MgO through the lake that is broadly similar to that observed in eruption samples (data from Murata and Richter, 1966a), if we take into account the volume of the eruptive products. Such an original feature of the filling process could have survived more or less unchanged since emplacement if there was no crystal settling and convection. Gibb and Henderson (1992) concurred with this idea. They concluded that “the magmas have not undergone turbulent convection and that gravity settling has usually played only a minor modifying role” since the lake emplacement. Our CSD data discussed above are consistent with this possibility as well, in the sense they suggest that crystal settling, and probably convection also, are processes of little importance in the evolution of the lake. We assume here that diapiric transfer of material (e.g. vorbs) was not large-scale enough to induce wholesale crystal

redistribution. Instead, the vorb-related transport may be viewed only for redistributing low-Fo olivine to the upper parts of the lava lake.

In contrast, a number of later works suggested that the lake was well-mixed (e.g. Richter *et al.*, 1970). This idea is based on the many cycles of rapid influx and drainback of lava into the lava lake, together with the repeated sub-crustal injections of lava and crust foundering; all of which have occurred over the 17 eruptive episodes of the eruption. Eaton *et al.* (1987) also proposed that the lake was well-mixed on the basis of volcano summit deformation data. They suggested that large-scale mixing in the summit reservoir occurred under Kilauea Iki before and during the eruption. Mixing occurred in response to repeated refilling events by the injection of new, hotter, more primitive and gas-rich magma from deeper levels in the crust, along with repeated drainback of heavier, degassed lake lava into the reservoir. Helz (1987b) also argued in favour of a well-mixed summit reservoir after phase 1 of the 1959 eruption, based mostly on olivine compositions and textures. Hence, although the case of non-homogenised magma that preserved the stratification seems convincing, it does not accord with these arguments of undoubtedly well-stirred lake magma.

An overall alternative could be the involvement during the eruption of another magma source that was injected into the lake through the base, or the deep part of the lateral walls (Richter *et al.*, 1970). This idea of basal feeding of the lake has been proposed for most active lava lakes. Here we feel it is not necessarily incompatible with Hawaiian lava lakes formed in a pre-existing pit-crater, taking into account the spatial configuration of the plumbing system beneath Kilauea summit and East Rift Zone. This is fairly consistent with the model of cycling of magma between the summit reservoir and Kilauea Iki lava lake of Eaton *et al.* (1987), along with the magma budget of the eruption they drew. The only concrete evidence for this basal replenishment

of the lava lake during eruption lies in the chronological narrative of the eruption by Richter et al. (1970). They repeatedly described sub-crustal, or cryptic, draining of the lake lava, with no visual active backflow to the vent, and sometimes what we could interpret as addition of magma under the lava lake. In this case, the implications on the way the lava lakes, previously thought to be 'inactive', are interpreted in terms of magma differentiation during cooling would be considerable.

2.7 CONCLUSIONS

This study of the CSDs and olivine compositions of the 1959 Kilauea Iki products and lava lake has yielded the following conclusions:

- (1) Both the geochemistry and the CSDs reveal two major and one minor populations of olivine crystals, which are not necessarily directly related;
- (2) There are no systematic compositional differences between deformed and undeformed olivine crystals;
- (3) Significant percentage of olivine crystals are antecrysts; but no mantle xenocrysts are present;
- (4) The textural data show evidence of coarsening, mostly occurring before the eruption;
- (5) Crystal or magma mixing was prominent before the eruption in the Kilauea Iki plumbing system, but became less active or inactive during the cooling of the lake;
- (6) Olivine in the lower part of the lake (>60 m deep) has a less Fo-rich composition and more crystals are deformed, but there are no simple explanations to account for this stratification;

- (7) The CSDs show no significant evidence of both crystal settling and convection, although local advective currents are known to have formed numerous segregation veins and vorbs;
- (8) The vertical stratification, mainly in terms of olivine distribution and chemistry, could be an original feature, probably derived from the stratification of the summit reservoir at the time of the eruption. However, this hypothesis is not consistent with supposed strong stirring of the lake magma due to sub-crustal injections of lava over the 17 eruptive phases, along with repeated draining through the vent and crust foundering. Basal feeding of the lake during the eruption is another possibility, and may be seen as a likely mechanism to explain the stratification of the lake.

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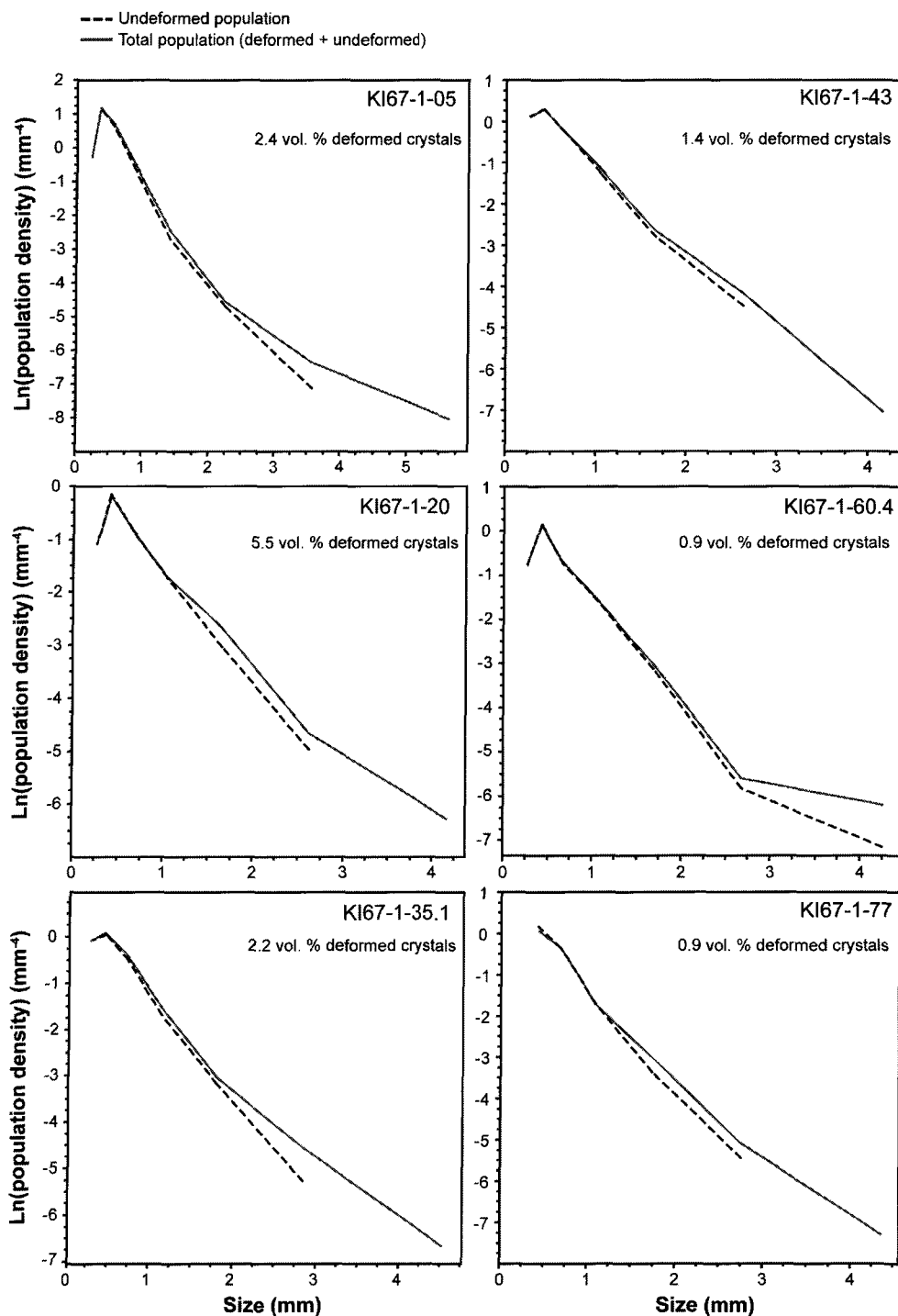
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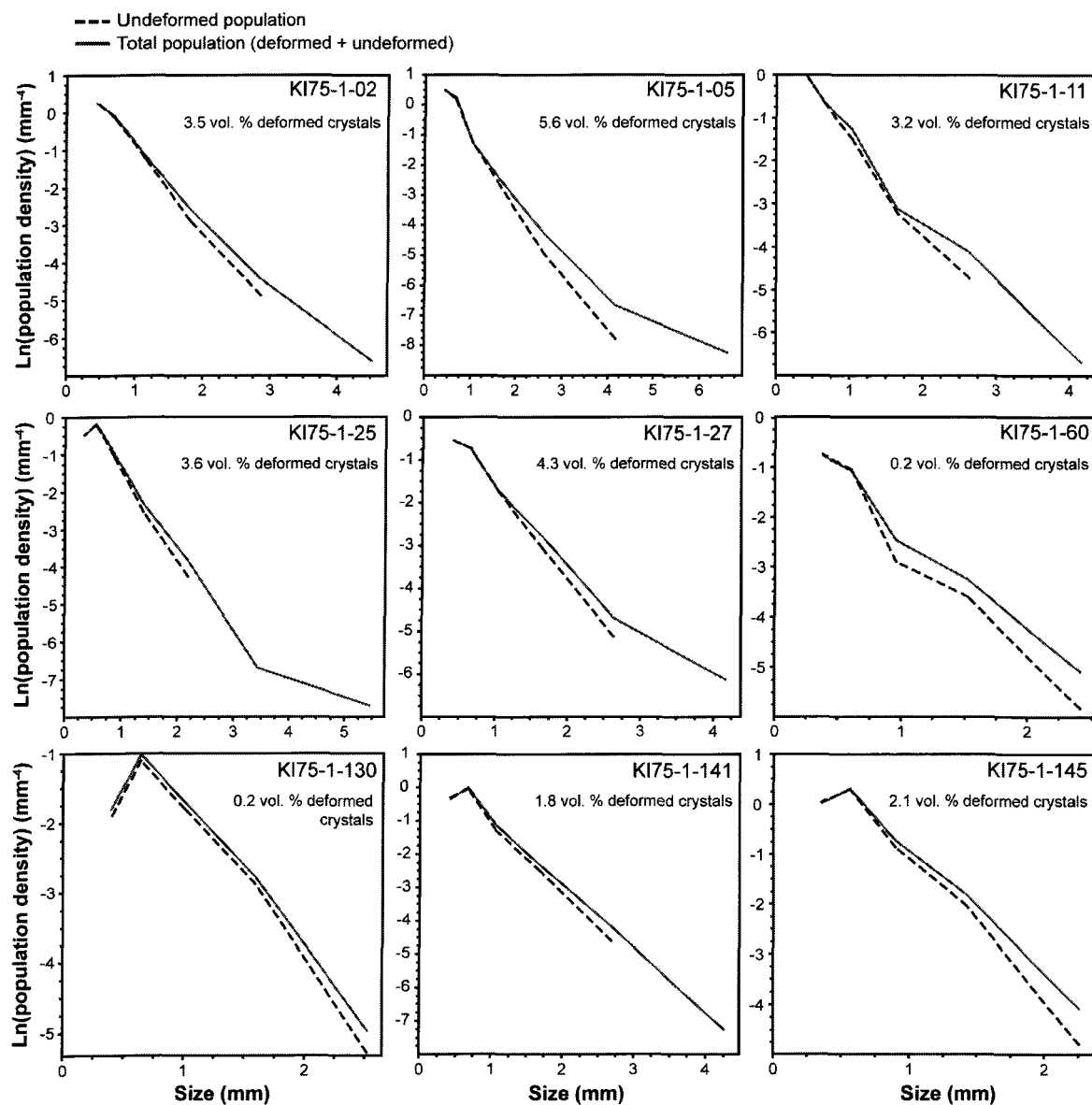
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APPENDIX 2-1

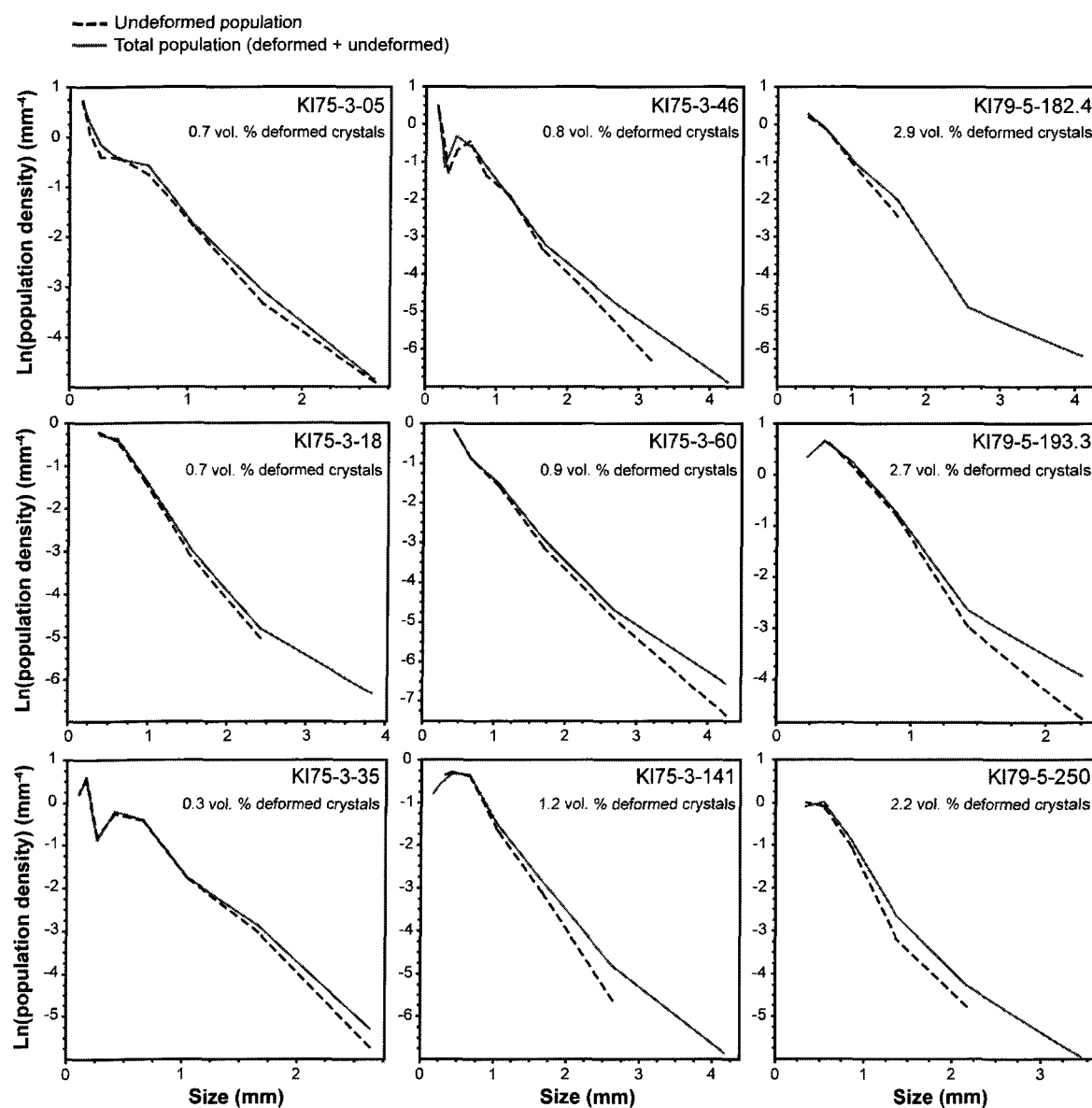
All CSDs of undeformed versus total olivine populations, showing the effects of deformed crystals on the total crystal size distribution. A quantitative estimate of the abundance of deformed crystals in each sample is shown at the top right of each plot.



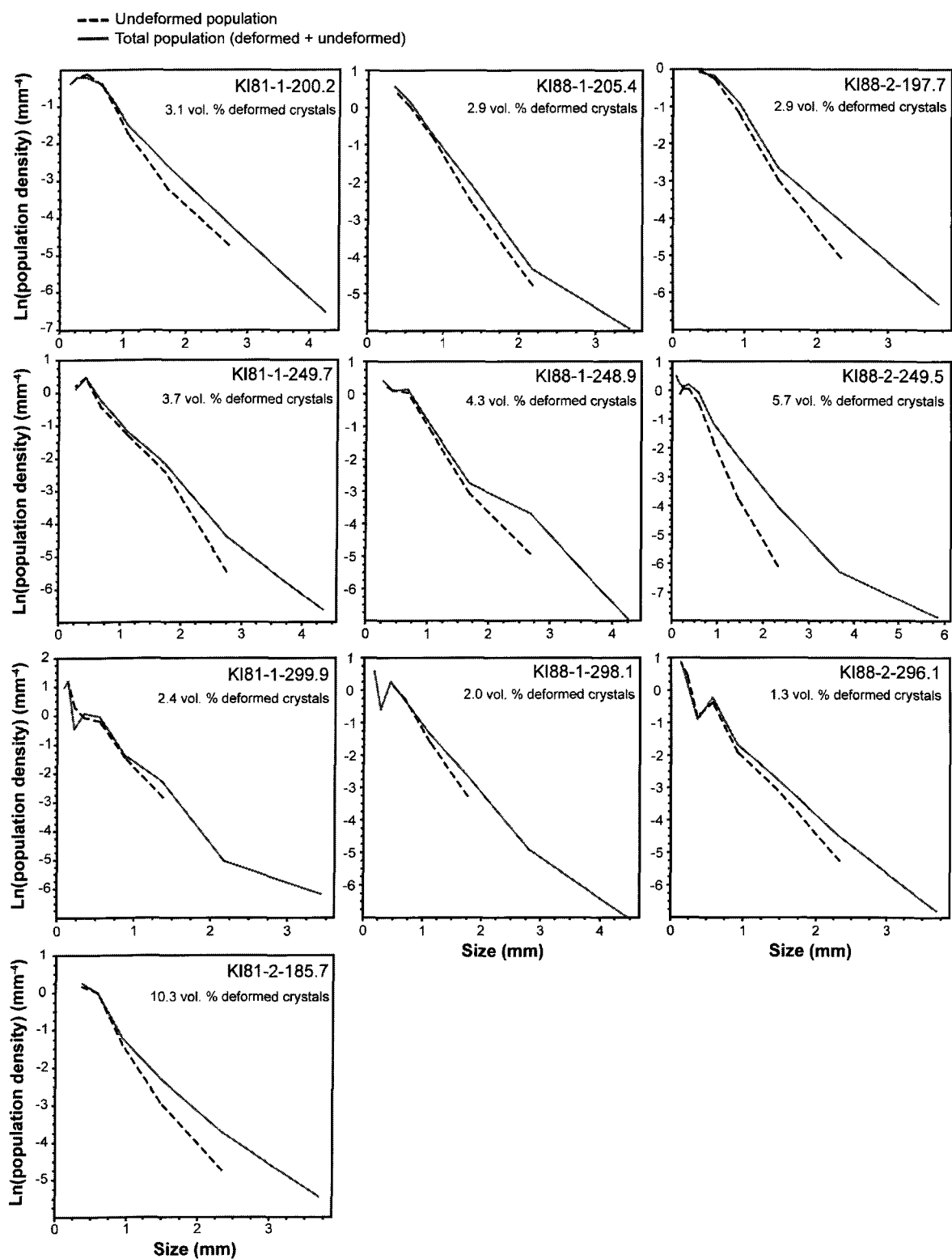
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AVANT-PROPOS AU CHAPITRE 3

CONTRIBUTION DES AUTEURS

Le Chapitre 3 est le résultat d'une collaboration avec Dr. Roberta Flemming, responsable du Laboratoire de Microanalyse par Rayons X, *The University of Western Ontario, London, Canada*. L'UQAC n'avait pas l'appareillage nécessaire à l'analyse par diffraction des rayons X de mes cristaux d'olivine. J'ai ainsi pu recueillir les données indispensables à la rédaction de l'article qui constitue ce troisième chapitre de thèse. Dr. Flemming m'a aidé à correctement utiliser l'appareillage et les divers logiciels de traitement de données ; elle a par ailleurs rédigé la partie « Methods » de l'article.

CHAPITRE 3

CRYSTAL STRUCTURE, MOSAICITY AND STRAIN ANALYSIS OF HAWAIIAN OLIVINES USING IN SITU MICRO X-RAY DIFFRACTION (μ XRD)

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3.1 ABSTRACT

Deformation of olivine in a volcanic context is poorly constrained, although deformed olivine is abundant in some volcanic rocks, and its presence is important for the definition of the magmatic history of volcanic edifices such as Kilauea Volcano, Hawaii. Deformed olivines at Kilauea originate in the lower crust, therefore, the classic approaches and interpretations applied to mantle-derived olivine are not applicable here. In this paper we present data on deformed olivine crystals from Kilauea lava samples using an *in-situ* μ XRD technique. Our results allow the validation, and refinement, of optical observations of olivine deformation. We also confirm the presence of deformation for every olivine crystal size, and quantify it. There are significant correlations between deformation intensity (strain-related mosaicity) and the olivine composition and crystal size. Although this technique does not allow the simple estimation of the P-T conditions of deformation and crystal formation or magmatic history, some constraints have been provided here, in particular the estimation of a threshold over which deformation undoubtedly occurred. Micro XRD is shown here to be an easy-to-use, fast, low-cost, non-destructive and less-ambiguous technique to determine the presence of strain in the crystal structure of magmatic olivines, and quantify it, even for very small crystals.

Keywords: olivine, deformation, crystal strain, microstructure, mosaicity, μ XRD, Hawaii, Kilauea, Mauna Ulu, Kilauea Iki

3.2 INTRODUCTION

In Earth Sciences, the mechanisms responsible for the deformation of a rock, and hence associated plastic deformation of its constitutive minerals, have been debated for decades. Most emphasis has been given on plastic deformation of mantle-forming silicates, with predominance of olivine and pyroxenes. The main reason for this is that the understanding of plastic flow in the mantle is important in the creation and refining of models explaining the geodynamic processes and geophysical or rheological properties of the mantle, in particular the upper mantle (e.g. Mercier, 1985). Rock deformation is influenced by a number of intrinsic and extrinsic factors (Hobbs, 1985). Some control the migration rate of dislocations, whereas others control the shape, orientation and distribution of grains, and thus leave an imprint on the microfabric. Temperature is one of the most important factors influencing the deformation. Many studies have also been carried out on the dislocation mechanisms themselves. For example, olivine deforms by dislocation creep in the shallow mantle (Raterron et al., 2009). Constitutive equations have been proposed to describe dislocation mechanisms in single olivine crystals, taking into account a wide range of thermodynamic, thermomechanical and microstructural conditions based on compressional creep experiments at various pressures and temperatures (Bai and Kohlstedt, 1992; Bai et al., 1991; Demouchy et al., 2009; Durham and Goetze, 1977; Evans and Goetze, 1979; Kohlstedt and Goetze, 1974; Li et al., 2004; Raterron et al., 2004).

In contrast, deformation of olivine now found in a volcanic context has remained poorly constrained, although deformed olivine is abundant in some volcanic rocks, and its presence is important for the definition of the magmatic history of such volcanic edifices (e.g. Kilauea Volcano, Hawaii; Piton de la Fournaise Volcano, La Réunion Island). In such a volcanic context, volcano-

related microfabrics are rare. There is no direct relation, or even no relation at all with mantle flow issues, and thus no need to get lattice-preferred orientation data, for example, for magmatic olivines. Therefore, classic approaches, techniques and interpretations applied to mantle-related issues are not applicable, and few data are available.

In this paper we investigate the deformation of olivine crystals from Kilauea Iki and Mauna Ulu lava samples, using an innovative, *in-situ* μ XRD technique which measures streak length along the Debye ring (chi angle) in two-dimensional X-ray images (Flemming, 2007). Since it is a recently developed technique, there are very few quantitative data available for comparison with our data. Previous studies have examined quartz (Flemming, 2007), zircon (Moser et al., 2009), orthopyroxene (Izawa et al., 2009; Izawa et al., *in review*), and diamonds (Smith et al., 2010), but none have examined olivine. Consequently, our aim is to quantify deformation using such parameters as the full width at half maximum (FWHM) of peak intensity along the Debye rings. The focus is also given to the validation of this simple μ XRD approach that allows: (1) the clear recognition of the presence of significant strain in olivine single crystals, even for very small grains; and, (2) the quantification of the strain-related mosaicity for many different crystallographic planes. This approach is intended to be easy to apply for a large range of olivine-bearing rock samples. In turn, reference values, in particular FWHM data, recovered on olivines with well known origin would be available for the scientific community.

3.3 MATERIALS AND METHODS

3.3.1 Materials

All olivines are from 1969-1974 Mauna Ulu (MU) lava samples and 1959 Kilauea Iki (KI) eruption products. Polished thin sections were examined so that olivine crystals for analysis by μ XRD. All the samples were previously studied using both quantitative textural and geochemical methods (Vinet and Higgins, 2010; Vinet and Higgins, *in revision*). These previous studies indicated the presence of multiple olivine populations that underwent different solidification histories, including a deformed component that is thought to originate from a cumulate body at the base of the spreading volcano. A range of both optically deformed (that is, crystals with optical evidence for deformation) and undeformed olivine crystals of varying sizes and shapes were chosen for study. The optical deformation features observed in olivine are kink-bands, undulose extinction and rectangular subgrains. We also included some olivines for which the deformed or undeformed character is ambiguous.

3.3.2 Micro XRD methods

X-ray diffraction data were collected using a Bruker D8 Discover micro X-ray diffractometer at the University of Western Ontario. The diffractometer was operated using Cu-K α radiation generated at 40 kV and 40 mA. The instrument was equipped with a Gobel mirror parallel optics system and pinhole collimation to produce a nominal beam diameter of 500 μ m. Uncoated polished thin sections were mounted on an XYZ stage, and precise sample positioning was accomplished using an optical microscope monitor and laser system (reproducible to within 12.5

μm). Diffracted X-rays were detected using a Hi-Star Detector and General Area Detector Diffraction System (GADDS) software. The two-dimensional detector was positioned at a distance of ~ 15 cm from the sample and intersected a 35° segment of diffraction space. The θ - θ geometry of the diffractometer enabled the sample to remain horizontal and stationary, while the source and the detector moved independently. The angle between the sample and the source is θ_1 while the angle between the sample and the detector is θ_2 , where $\theta_1 + \theta_2 = 2\theta$, as measured at the centre of the detector.

Samples were analyzed in omega scan mode, where the X-ray optics (source = θ_1 ; detector = θ_2) were rotated through an omega angle ω , at fixed 2θ . At the start of the omega scan, the source and detector were positioned at $\theta_1 = x^\circ$ and $\theta_2 = y^\circ$, respectively, such that the starting $2\theta = \theta_1 + \theta_2 = x + y$. Both source and detector were rotated clockwise, by an omega angle of ω , stopping at $\theta_1 = x + \omega$ and $\theta_2 = y - \omega$, such that the final $2\theta = \theta_1 + \theta_2 = x + \omega + y - \omega = x + y$. At all points during omega rotation, 2θ remained constant, where $\theta_1 + \theta_2 = 2\theta = x + y$. Because 2θ has remained constant from the frame of reference of the detector it appears that the sample has rotated by $-\omega$. This enabled a greater number of lattice planes in a single crystal to satisfy the Bragg's Law diffraction condition. Omega Scan conditions were as follows: Frame 1: $\theta_1=15^\circ$, $\theta_2=18.5^\circ$, $\omega=8^\circ$, time=20 minutes; Frame 2: $\theta_1=18^\circ$, $\theta_2=45^\circ$, $\omega=30^\circ$, time=60 minutes.

Two-dimensional GADDS images yield information in both the 2θ and chi (χ) dimensions. When using two-dimensional detection methods, polycrystalline samples give powder rings, or Debye rings, indicative of crystallites having all orientations simultaneously, whereas single crystals give diffraction spots, indicative of a single orientation. Each diffraction spot having lattice plane $\{hkl\}$ lies along an arc of radius $2\theta_{hkl}$. Strained crystals exhibit strain-related mosaicity, visible

as elongation or streaking along the Debye ring, as measured by chi angle, χ . For each GADDs image, peak intensity was integrated in two different ways. First, intensity was integrated along chi (i.e. integrated along each Debye ring) and plotted as a function of 2θ , to create a one-dimensional plot of intensity versus 2θ , reminiscent of a conventional powder diffraction pattern. Such a plot is shown in Figure 1d. These plots are used to assign the Miller index to the observed olivine diffraction lines, using olivines in the International Centre for Diffraction Data (ICDD) database. (e.g. olivine was indexed using card # 79-1192). This is typically the only method of integration performed on two-dimensional data, as it is necessary for phase identification. Additionally, we performed an integration of narrow regions of 2θ as a function of chi angle. Such integration is shown in Figure 1e. This enabled examination of the peak shape along the Debye ring as well as quantitative analysis of peak width along the Debye ring.

3.4 RESULTS AND DISCUSSION

Microstructure and strain information for single olivine crystals can be obtained on 2-D GADDs images and on the resulting μ XRD patterns (Fig. 1). A single homogeneous grain analysed by μ XRD will have a single diffracted X-ray for each hkl plane, producing a single spot on the GADDs detector (Flemming, 2007). In contrast, an inhomogeneously strained grain will produce “streaking” or asterism of the diffraction spots in the GADDs image (Fig. 1c) (Flemming, 2007). These fundamental characteristics are well observed for the olivine grains studied here, with variability in the brightness of the diffraction spot and length of the streak.

An objective of this study is to validate both the optical petrographic observations of olivine deformation and the μ XRD technique recently developed for *in-situ* analysis purposes (Flemming,

2007; Izawa et al., 2009; Izawa et al., *in review*; Moser et al., 2009; Smith et al., 2010). A comparison between the petrographic and XRD evidence of deformation for each crystal is presented in Table 1. Overall, most optical observations of deformation in olivine crystals (Vinet and Higgins, 2010; Vinet and Higgins, *in revision*) were validated or refined using μ XRD. With two exceptions, all the optically deformed crystals of any size also show, as expected, streaked diffraction patterns that are attributed to strain. Also, with the exceptions of three grains, all optically undeformed crystals of any size exhibit single diffraction spots in the GADDS images, which are attributed to the absence of strain. The three exceptions are grains with streaked diffraction patterns. Hence, the μ XRD technique has highlighted the presence of strain in these grains whereas it was not seen optically. Rare deformed or undeformed grains have diffraction spots that are difficult to interpret. Therefore, for these grains we were unable to clearly determine whether there is presence of strain. Finally, for most optically ambiguous grains (with respect to their deformation nature), we have been able to determine if they were deformed or undeformed using μ XRD.

Olivine deformation was quantified using the full width at half maximum (FWHM) of the line shape along the direction of Debye ring (chi angle χ). It is a useful parameter of quantification as it represents a simple way of estimating the quantity of strain-related mosaicity recorded in a single crystal, and hence the deformation intensity. Intensity of the diffracted rays was integrated along the Debye rings to produce plots of intensity vs. χ (Fig. 1e). In such plots, the FWHM values of several plane reflections were determined for 52 single olivine crystals from two Mauna Ulu and five Kilauea Iki lava samples. Deformed crystals cover the whole range of FWHM values (Table 2; Fig. 2), 0.3-3.3°, with most analyses ranging from 0.3 to 1.8°. In contrast, undeformed crystals are constrained to FWHM < 1.0°, with a peak around 0.5° (Fig. 2). It is demonstrated in this study that

every single crystal with FWHM > 1.0° clearly shows both optical and XRD evidence of strain, with very rare exceptions. We propose that this be used as a threshold value, above which deformation has occurred. However, we cannot determine information on deformation for crystals with FWHM < 1.0°, as these values comprise both deformed and undeformed crystals.

The FWHM parameter can also be useful for the recognition of correlations with respect to the *hkl* planes (Miller indexes) or other independent parameters such as the olivine composition or crystal size. In a given lava sample, i.e. thin section, FWHM is uncorrelated to the *hkl* index of the reflecting lattice planes (Table 2). In contrast, there is a broad positive correlation between FWHM and the forsterite content (Fig. 3). It is especially true for samples from Kilauea Iki, although an exception could be seen around Fo₉₀. In a plot of all reflecting lattice planes (all Miller indexes), we also found a somewhat weak, but significant positive correlation between FWHM and the crystal size (Fig. 4), which was chosen here to be the diameter of a circle of equivalent area to the ellipsoid fitted to the crystal outline (Table 2). This correlation becomes even stronger for individual lattice planes such as: (024) / (330), (041), (061), (062), (222), and (251) (Fig. 5). In contrast, there is no simple correlation between FWHM and crystal size when data are plotted individually for each lava sample (Fig. 6). We notice here that the μ XRD data are consistent with previous petrographic findings (Vinet and Higgins, 2010; Vinet and Higgins, *in revision*), that is, deformation has affected all crystal sizes and, to a lesser extent, compositions, with a slight predominance for larger crystals but not for Fo-rich olivines.

The quantitative parameter FWHM can be related to mosaicity in the crystal lattice, as demonstrated by Flemming (2007) using quartz in quartzite from La Malbaie, Quebec. Mosaicity (or mosaicism) refers to the presence of numerous subgrains with slightly to significantly misoriented crystal lattices relative to each other, in a uniform single crystal (French and Koeberl,

2010). It is seen in X-ray diffraction patterns as a broadening of the normally sharp diffraction peaks of lattice planes (Hanss et al., 1978). Also, Flemming (2007) found a correlation between inhomogeneous strain in minerals and “streaking” of the diffraction spots in the 2-D GADDS image. As for her quartz grains, in our polished thin sections of Mauna Ulu and Kilauea Iki lavas, optically unstrained grains of olivine (Fig. 7a) exhibited single diffraction spots in the corresponding GADDS image (Fig. 7b). In contrast, the strained, or deformed, olivine grains optically exhibiting kink-bands, undulose extinction or rectangular subgrains (Fig. 7c), show distorted diffraction spots, or streaks, in the corresponding GADDS image (Fig. 7d). Streaked diffraction spots may be interpreted to be caused by strain-related mosaicity or non-ideality of the crystal lattice that is imposed by inhomogeneous strain (Flemming, 2007). As argued by Flemming, these observations are consistent with streaking or asterism, as observed by Hörz and Quaide (1973) after shock experiments on a variety of minerals, and recently treated in the meteorite impact review of French and Koeberl (2010). Although these observations have been applied to minerals shocked by a meteorite impact, they are clearly similar to what we have observed here for magmatic olivines, and may help in our interpretations. Furthermore, for many of our strained olivines exhibiting streaks (strain-related mosaicity) in the diffraction pattern, the GADDS image (Fig. 7d) reveals that each streak is resolved into a row of diffraction spots. This was also observed in a quartz sample by Flemming (2007). She measured the GADDS image and found a consistent angular separation of 2° between spots. She clearly showed that GADDS images can be used for quantification of strain-related textures. She also interpreted the angular separation between spots to be a small angle of misorientation between subgrains, which may represent “edge dislocations climbing into alignment to form planar arrays of dislocations”. This is also applicable to our deformed olivines

showing the same type of strain-related mosaicity, and is suggestive of power law creep (Langdon, 1985).

MU69-1b - Grain 15 (spot 17)

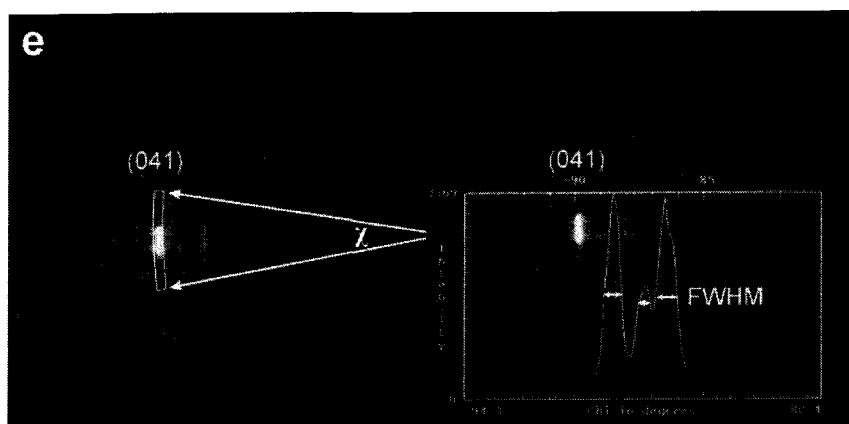
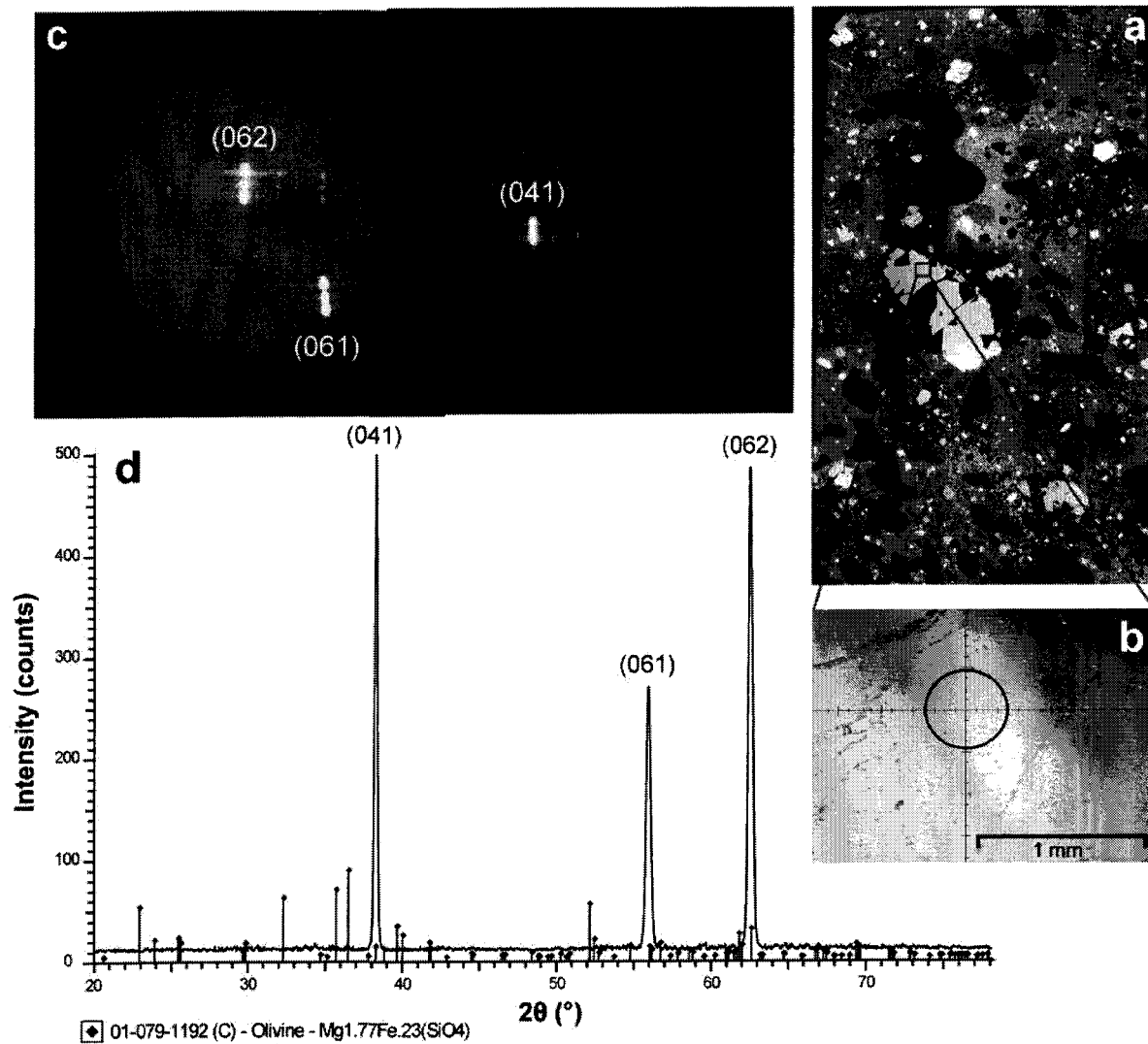


Figure 3-1. Micro X-ray diffraction data from MU69-1b-grain 15.

(a) Photomicrograph of the whole thin section under crossed polarized light. **(b)** Image of spot 17 as seen under the optical microscope monitor at maximum magnification; black circle represents approximate beam diameter using 500 μm beam. **(c)** GADDs images from spot 17 showing streaked diffraction patterns, attributed to strain (omega scan, 2 frames, frame 1 (right) = 20 min, frame 2 (left) = 40 min, 500 μm beam). **(d)** Plot of intensity versus 2θ for spot 17 (after background subtraction) showing a good match with an ideal olivine composition: $\text{Mg}_{1.77}\text{Fe}_{0.23}(\text{SiO}_4)$; hkl planes indicated on top of peaks. **(e)** GADDs image from spot 17 (frame 1) showing integrated box along Debye ring (left), and plot of intensity vs. χ (right), the angle subtended by the Debye ring; white arrows in right frame indicate full width at half maximum (FWHM), the representative FWHM value being the sum of the three distances in this case where there are subgrains.

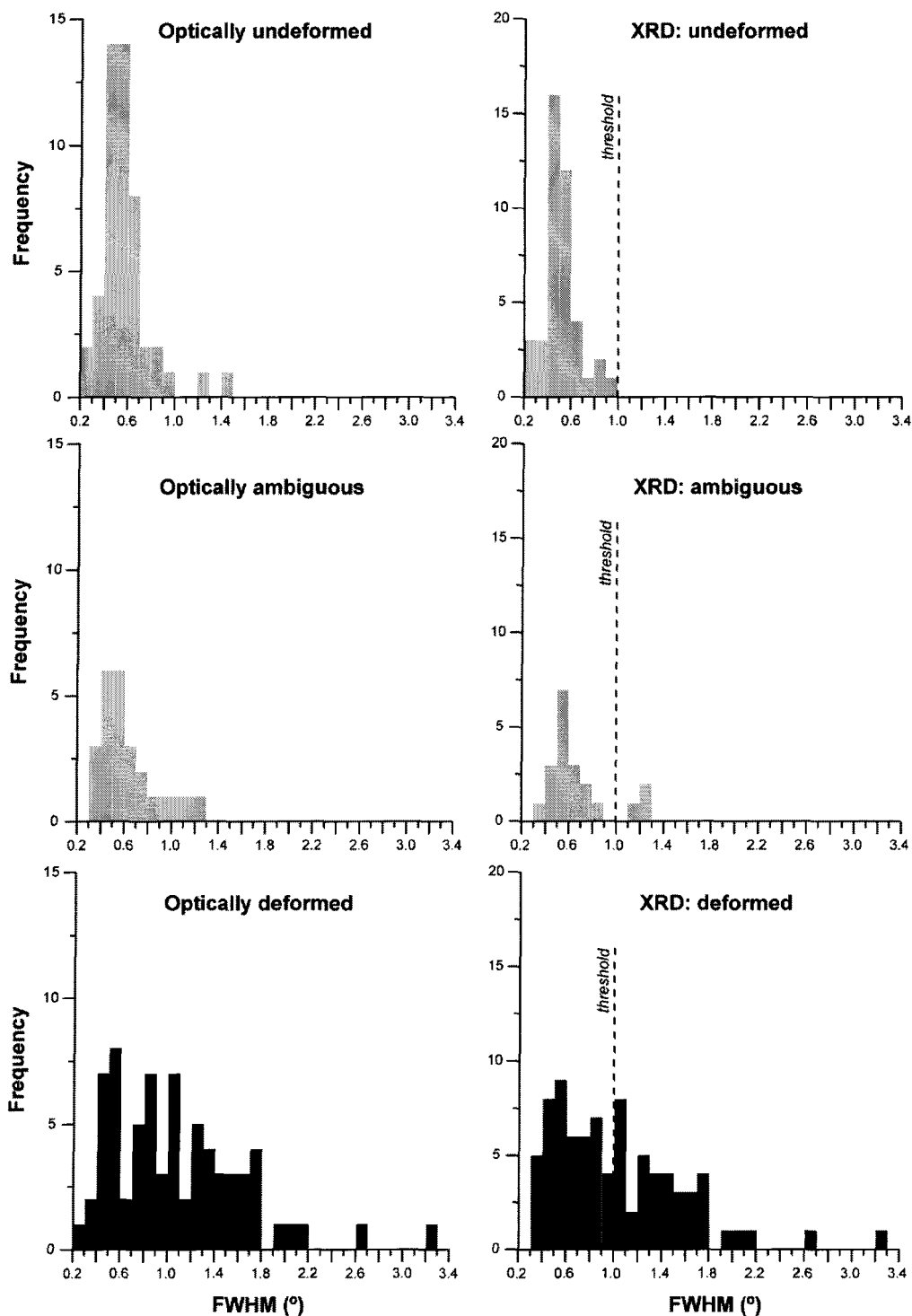


Figure 3-2. Full width at half maximum (FWHM) values for undeformed, deformed and ambiguous olivine grains, with optical (left) and XRD (right) distinction. Vertical dashed line indicates a threshold value of FWHM (1.0°) over which there is clear evidence of strain.

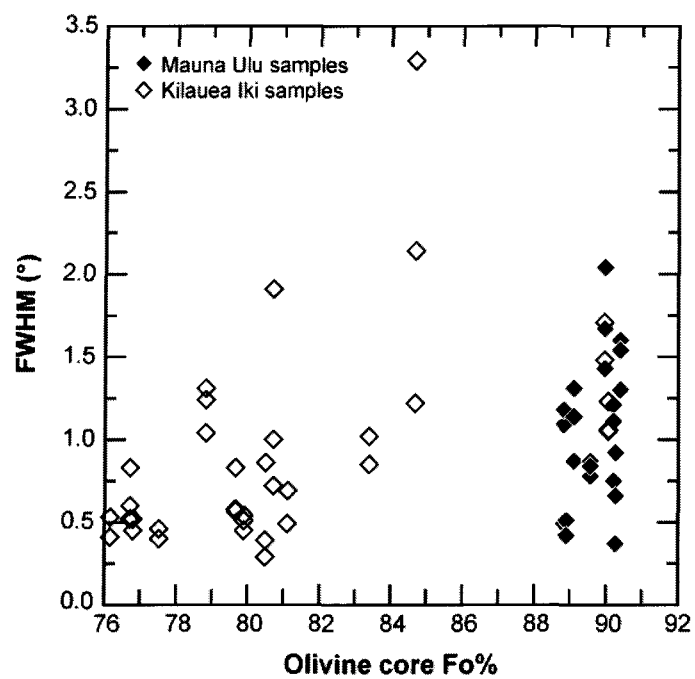


Figure 3-3. Full width at half maximum (FWHM) versus olivine core forsterite (Fo) content for all samples analysed by μ XRD.

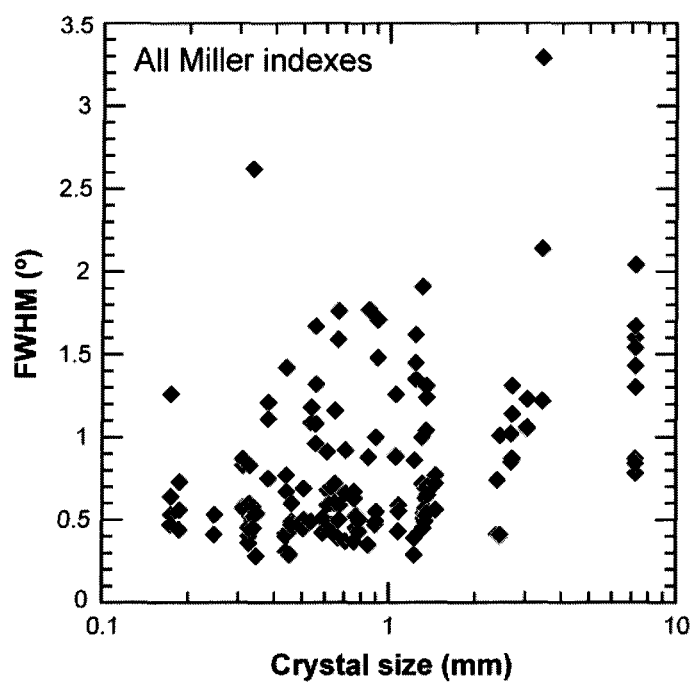


Figure 3-4. Full width at half maximum (FWHM) versus crystal size (chosen to be the diameter of circle of equivalent area to an ellipsoid fitted to outline), for all hkl indexes at a time.

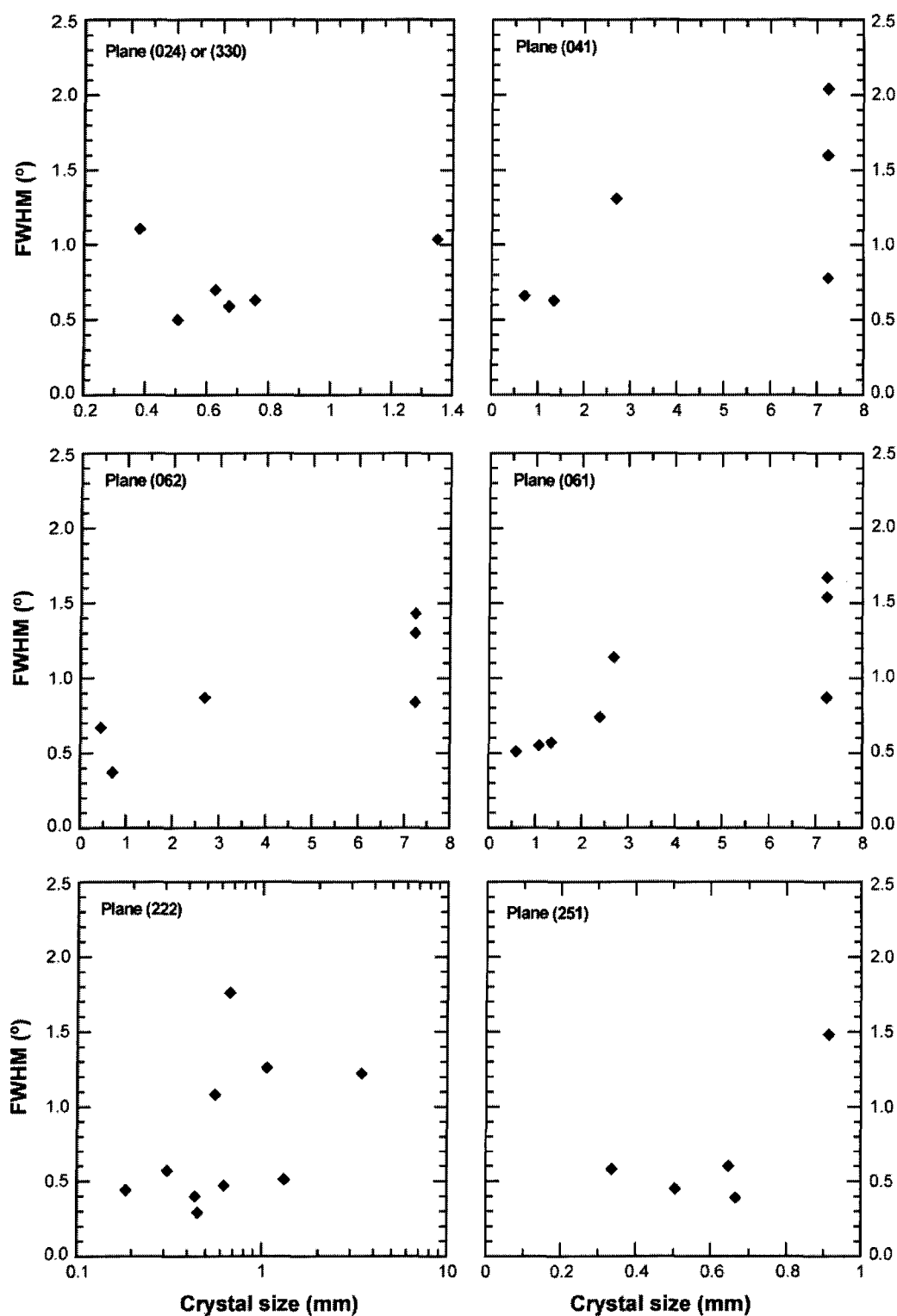


Figure 3-5. Full width at half maximum (FWHM) versus crystal size (chosen to be the diameter of circle of equivalent area to an ellipsoid fitted to outline), for selected individual hkl indexes.

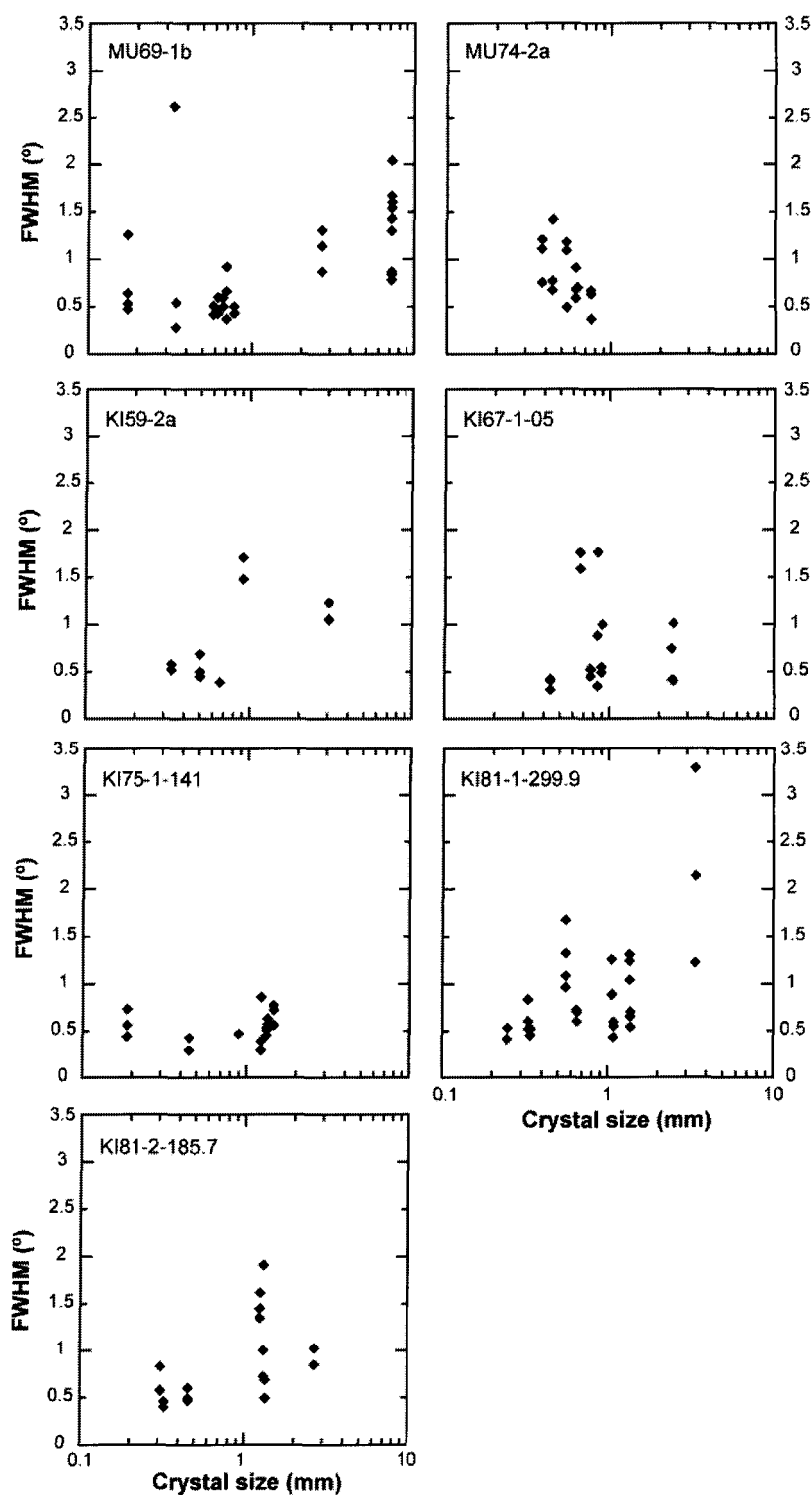


Figure 3-6. Full width at half maximum (FWHM) versus crystal size (chosen to be the diameter of circle of equivalent area to an ellipsoid fitted to outline), for each analysed sample. All analysed hkl indexes are plotted per sample.

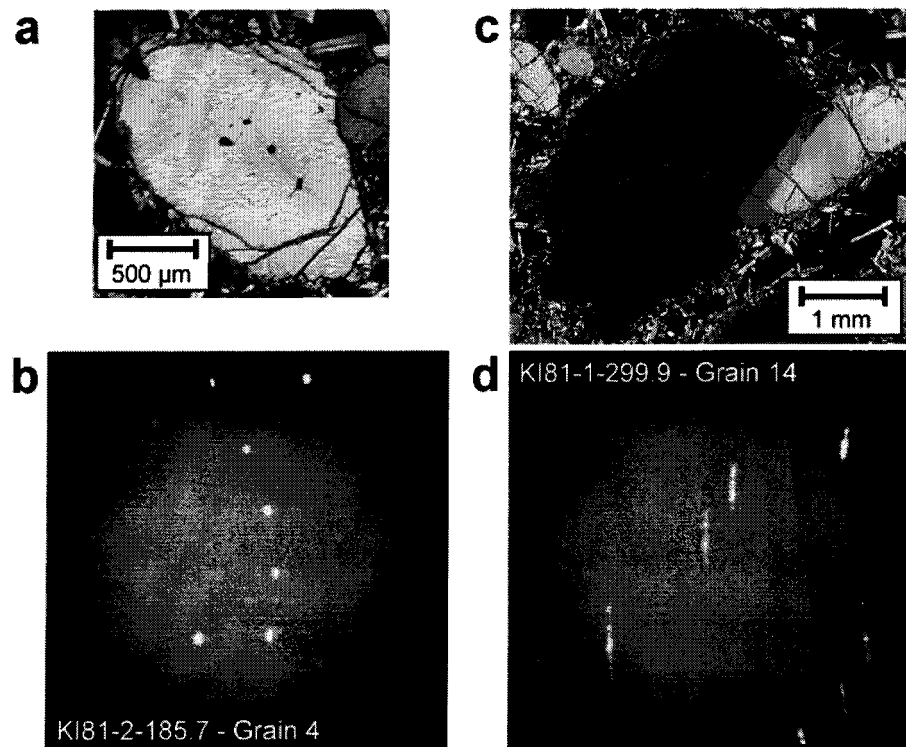


Figure 3-7. Polarizing light microscope images and μ XRD data for strained versus unstrained Kilauean olivine.

(a) Photomicrograph showing one undeformed olivine grain under crossed polarized light. **(b)** General area detector diffraction system (GADDS) image corresponding to the grain in (a), exhibiting circular diffraction spots (omega scan, 40 min, 500 μ m beam). **(c)** Photomicrograph showing kink-bands and undulose extinction of a deformed olivine grain under crossed polarized light. **(d)** GADDS image of olivine grain from the photomicrograph in (c) (omega scan, 40 min, 500 μ m beam).

Table 3-1: Comparison between petrographic and μ XRD evidence of deformation

Sample	Grain or spot number	Petrography	μ XRD	Similarity of observations
KI59-2a	1	undeformed	undeformed	yes
	2	deformed	slightly deformed?	ambiguous
	3	undeformed	deformed (streak = mosaicity)	no
	4	deformed	deformed (mosaicity + asterism)	yes
	5	deformed	deformed (streak)	yes
KI67-1-05	1	deformed?	undeformed	refined
	2	deformed	deformed (streak)	yes
	3	deformed?	slightly deformed; plane (140) in particular	refined
	4	deformed	deformed (mosaicity + asterism)	yes
	5	undeformed?	undeformed	refined
	6	deformed	deformed (mosaicity + asterism)	yes
	7	undeformed	undeformed	yes
KI75-1-141	6	deformed	undeformed	no
	7	deformed	deformed (but, 2 grains?)	yes
	8	deformed	deformed (but, 2 grains?)	yes
	9	undeformed	undeformed	yes
	10	deformed	deformed	yes
	11	deformed	undeformed	no
KI81-1-299.9	12	undeformed	undeformed (2 grains?)	yes
	8	deformed	deformed (mosaicity + asterism)	yes
	9	deformed?	undeformed	refined
	10	deformed?	slightly deformed? (satellite grains?)	ambiguous
	11	deformed	deformed (mosaicity + asterism)	yes
	12	deformed	deformed	yes
	13	undeformed	slightly deformed; plane (331) in particular	no
	14	deformed	deformed (mosaicity + asterism)	yes
	15	deformed?	slightly deformed? (satellite grains?)	ambiguous
	16	undeformed	undeformed	yes
KI81-2-185.7	17	undeformed?	undeformed	refined
	1	deformed	deformed (streak)	yes
	2	undeformed	undeformed (satellite grains)	yes
	3	deformed	deformed (streak)	yes
	4	undeformed	undeformed	yes
	5	deformed	deformed (streak)	yes
	6	deformed	slightly deformed	yes
MU69-1b	7	deformed?	slightly deformed? (satellite grains?); plane (222) in particular	ambiguous
	7	deformed	deformed (mosaicity + asterism)	yes
	8	deformed	deformed (streak)	yes
	9	undeformed?	slightly deformed?; plane (112) in particular	ambiguous
	10	undeformed	undeformed	yes
	11	undeformed	undeformed	yes
	12	deformed?	slightly def	refined
	13	undeformed	undeformed	yes
	14	deformed	deformed (mosaicity)	yes
	15	deformed	deformed (mosaicity + asterism)	yes
	16	deformed	deformed (mosaicity + asterism)	yes
	17	deformed	deformed (mosaicity + asterism)	yes
	18	undeformed?	undeformed	refined

Table 3-1 (continued)

MU74-2a	1	undeformed	slightly deformed (asterism)	no
	2	undeformed	slightly deformed?	ambiguous
	3	deformed	deformed (mosaicity + asterism)	yes
	4	deformed?	deformed? (asterism or satellite grain?)	ambiguous
	5	deformed?	deformed (streak)	refined
	6	undeformed?	slightly deformed	refined

Table 3-2: Full width at half maximum (FWHM) of plane reflections integrated along the direction of Debye ring (chi angle χ), for single deformed and undeformed olivine crystals from Mauna Ulu (MU) and Kilauea Iki (KI) volcanic rocks

Sample	Grain/spot number	<i>hkl</i>	Dimensions of ellipsoid fitted to outline		Diameter of circle (mm)*	Crystal size class	FWHM (°)	Note
			Major axis (mm)	Minor axis (mm)				
KI59-2a	1	002	0.76	0.33	0.50	small	0.69	
	1	251	0.76	0.33	0.50	small	0.45	
	1	330/024	0.76	0.33	0.50	small	0.50	
	2	002/121	0.39	0.29	0.34	small	0.52	
	2	251	0.39	0.29	0.34	small	0.58	
	3	251	0.88	0.50	0.67	small	0.39	
	4	121	1.37	0.61	0.91	small	1.71	
	4	251	1.37	0.61	0.91	small	1.48	
	5	301	4.28	2.16	3.04	large	1.05	
	5	311	4.28	2.16	3.04	large	1.23	
KI67-1-05	5	312	4.28	2.16	3.04	large	1.06	
	1	120	0.93	0.87	0.90	small	1.00	
	1	242/251	0.93	0.87	0.90	small	0.49	
	1	322	0.93	0.87	0.90	small	0.55	
	2 (spot 3)	061	2.68	2.12	2.38	large	0.74	
	2 (spot 3)	331	2.68	2.12	2.38	large	0.41	
	3 (spot 4)	140	2.98	2.01	2.44	large	1.01	
	3 (spot 4)	241	2.98	2.01	2.44	large	0.40	
	3 (spot 4)	340	2.98	2.01	2.44	large	0.41	
	4 (spot 5)	111	1.01	0.44	0.66	small	1.59	
	4 (spot 5)	112	1.01	0.44	0.66	small	1.77	
	4 (spot 5)	222	1.01	0.44	0.66	small	1.76	
	5 (spot 6)	112	0.47	0.41	0.44	small	0.31	
	5 (spot 6)	222	0.47	0.41	0.44	small	0.40	
	5 (spot 6)	241	0.47	0.41	0.44	small	0.42	
	6 (spot 7)	130	0.93	0.77	0.85	small	0.35	
	6 (spot 7)	240	0.93	0.77	0.85	small	1.77	2-3 2 individuals?
	6 (spot 7)	241	0.93	0.77	0.85	small	0.88	
	7 (spot 9)	043	0.93	0.63	0.77	small	0.52	
	7 (spot 9)	133	0.93	0.63	0.77	small	0.53	
KI75-1-141	7 (spot 9)	134	0.93	0.63	0.77	small	0.45	
	6	122	0.63	0.33	0.45	small	0.43	
	6	222	0.63	0.33	0.45	small	0.29	
	7	133	2.25	0.94	1.45	medium	0.56	
	7	134	2.25	0.94	1.45	medium	0.77	
	7	350	2.25	0.94	1.45	medium	0.72	
	8	041	1.74	1.03	1.34	medium	0.63	
	8	061	1.74	1.03	1.34	medium	0.57	
	8	331	1.74	1.03	1.34	medium	0.53	
	9	021	1.25	1.20	1.23	medium	0.86	
	9	143/062	1.25	1.20	1.23	medium	0.29	
	9	332/063	1.25	1.20	1.23	medium	0.39	
	10	213/242	1.54	1.13	1.32	medium	0.54	
	10	222	1.54	1.13	1.32	medium	0.51	
	10	322	1.54	1.13	1.32	medium	0.45	
	11	242/251	1.01	0.80	0.90	small	0.47	
	12	122	0.20	0.17	0.19	small	0.56	
	12	222	0.20	0.17	0.19	small	0.44	
	12	242	0.20	0.17	0.19	small	0.73	2 individuals?

Table 3-2 (continued)

KI81-1-299	8	222	1.32	0.85	1.06	small	1.26
	8	312	1.32	0.85	1.06	small	0.89
	8	322	1.32	0.85	1.06	small	0.88
	9	142	0.42	0.27	0.34	small	0.52
	9	152	0.42	0.27	0.34	small	0.45
	10	111	0.52	0.21	0.33	small	0.83
	10	112	0.52	0.21	0.33	small	0.60
	10	134	0.52	0.21	0.33	small	0.52
	11	024/330	2.07	0.88	1.35	medium	1.04
	11	111	2.07	0.88	1.35	medium	1.24
	11	311	2.07	0.88	1.35	medium	1.31
	12	111	0.66	0.47	0.56	small	1.67
	12	122	0.66	0.47	0.56	small	1.32
	12	222	0.66	0.47	0.56	small	1.08
	12	322	0.66	0.47	0.56	small	0.96
	13	061	1.44	0.81	1.08	small	0.55
	13	142	1.44	0.81	1.08	small	0.59
	13	331	1.44	0.81	1.08	small	0.43
	14	112	3.82	3.05	3.42	large	2.14
	14	122	3.82	3.05	3.42	large	3.29
	14	222	3.82	3.05	3.42	large	1.22
	15	121/002	0.73	0.57	0.65	small	0.70
	15	134	0.73	0.57	0.65	small	0.72
	15	251	0.73	0.57	0.65	small	0.60
	16	043	0.28	0.22	0.25	small	0.53
	16	114/162	0.28	0.22	0.25	small	0.41
	17	130	1.70	1.09	1.36	medium	0.70
	17	150	1.70	1.09	1.36	medium	0.65
	17	242	1.70	1.09	1.36	medium	0.54
KI81-2-185.7	1	113	3.27	2.18	2.67	large	1.02
	1	123	3.27	2.18	2.67	large	1.02
	1	214	3.27	2.18	2.67	large	0.85
	2	111	0.32	0.30	0.31	small	0.83
	2	211	0.32	0.30	0.31	small	0.58
	2	222	0.32	0.30	0.31	small	0.57
	3	112	1.38	1.11	1.24	medium	1.62
	3	204	1.38	1.11	1.24	medium	1.45
	3	311	1.38	1.11	1.24	medium	1.35
	4	210	1.71	1.05	1.34	medium	0.69
	4	312	1.71	1.05	1.34	medium	0.49
	5	132	1.64	1.04	1.30	medium	1.91
	5	133	1.64	1.04	1.30	medium	0.72
	5	240/123	1.64	1.04	1.30	medium	1.00
	6	312	0.39	0.28	0.33	small	0.36
	6	322	0.39	0.28	0.33	small	0.40
	7	112	0.58	0.37	0.46	small	0.60
	7	133	0.58	0.37	0.46	small	0.49
	7	134	0.58	0.37	0.46	small	0.47

Table 3-2 (continued)

MU69-1b	7	041	3.11	2.33	2.69	large	1.31	2 individuals?
	7	061	3.11	2.33	2.69	large	1.14	
	7	062	3.11	2.33	2.69	large	0.87	
	8	112	0.45	0.25	0.34	small	2.62	
	8	112	0.45	0.25	0.34	small	1.16	
	9	002	0.20	0.15	0.17	small	0.64	
	9	112	0.20	0.15	0.17	small	1.26	
	9	130	0.20	0.15	0.17	small	0.53	
	9	131	0.20	0.15	0.17	small	0.47	
	10	024/330	0.75	0.60	0.67	small	0.59	
	10	134	0.75	0.60	0.67	small	0.50	
	11	002	0.40	0.30	0.35	small	0.54	
	11	004	0.40	0.30	0.35	small	0.28	
	12	222	0.68	0.58	0.63	small	0.47	
	12	311	0.68	0.58	0.63	small	0.60	
	12	322	0.68	0.58	0.63	small	0.43	
	13	061	0.66	0.53	0.59	small	0.51	
	13	331	0.66	0.53	0.59	small	0.42	
	14	021	0.79	0.64	0.71	small	0.92	
	14	041	0.79	0.64	0.71	small	0.66	
	14	062	0.79	0.64	0.71	small	0.37	
	15 (spot 15)	061	10.06	5.20	7.23	megacryst	0.87	
	15 (spot 15)	062	10.06	5.20	7.23	megacryst	0.84	
	15 (spot 15)	041	10.06	5.20	7.23	megacryst	0.78	
	15 (spot 16)	061	10.06	5.20	7.23	megacryst	1.54	
	15 (spot 16)	062	10.06	5.20	7.23	megacryst	1.30	
	15 (spot 16)	041	10.06	5.20	7.23	megacryst	1.60	
	15 (spot 17)	061	10.06	5.20	7.23	megacryst	1.67	
	15 (spot 17)	062	10.06	5.20	7.23	megacryst	1.43	
	15 (spot 17)	041	10.06	5.20	7.23	megacryst	2.04	
	16 (spot 18)	140	1.14	0.54	0.79	small	0.50	
	16 (spot 18)	241	1.14	0.54	0.79	small	0.43	
	16 (spot 18)	261	1.14	0.54	0.79	small	0.50	
MU74-2a	1	021	0.48	0.41	0.44	small	1.42	2 individuals? 2 individuals?
	1	042	0.48	0.41	0.44	small	0.77	
	1	062	0.48	0.41	0.44	small	0.67	
	2	024/330	0.69	0.57	0.63	small	0.70	
	3	112	0.57	0.51	0.54	small	0.49	
	3	123	0.57	0.51	0.54	small	1.09	
	3	134	0.57	0.51	0.54	small	1.18	
	4	002	0.39	0.37	0.38	small	1.21	
	4	004	0.39	0.37	0.38	small	0.75	
	4	024/330	0.39	0.37	0.38	small	1.11	
	5	004	0.81	0.71	0.76	small	0.37	
	5	024/330	0.81	0.71	0.76	small	0.63	
	5	113	0.81	0.71	0.76	small	0.67	
	6	140	0.80	0.46	0.61	small	0.91	
	6	241	0.80	0.46	0.61	small	0.68	
	6	242	0.80	0.46	0.61	small	0.59	

* Diameter of circle of equivalent area than fitted-ellipsoid

3.5 CONCLUDING REMARKS

This μ XRD study of some Kilauea Iki and Mauna Ulu olivines has yielded the following conclusions and implications:

- The optical petrographic observations of olivine deformation were validated and refined using the μ XRD technique;
- Micro XRD applied to single olivine crystals in Hawaiian tholeiitic basalts is shown to be an easy-to-use, fast, low-cost, non-destructive and less-ambiguous technique to determine the presence of strain in the crystal structure, and possibly quantify it, even for very small crystals;
- Although this technique does not allow the simple estimation of the P-T conditions of deformation and crystal formation or magmatic history, some constraints have been provided here, in particular the estimation of a threshold over which deformation undoubtedly occurred;
- In the future, it would be interesting to carry out such μ XRD analyses for a wide range of single, deformed and undeformed olivine crystals with precise constraints on their formation and magmatic environments, such as upper, lower, and sub-continental lithospheric mantle, shallow magma chambers, volcano cumulates, etc. A complementary approach would be to analyse experimental olivines by μ XRD, olivines being formed under well-constrained temperature, pressure and other thermodynamic parameters. Overall, this data collecting approach using the μ XRD technique will allow the formation of a complete reference data set of qualitative and quantitative parameters useful to document deformation of magmatic olivines;

- Although mosaicity is often a shock effect, it can also be produced by endogenic processes (French and Koeberl, 2010, and references therein). French and Koeberl noted that there are no quantitative estimations of intensity of mosaicity in the literature, along with a lack of data about characteristics of mosaicity versus mineral composition or grain size, for example. In the present study, we gain insights into the relationship between the intensity of mosaicity, as estimated using FWHM, and olivine composition along with grain size. We also yield constraints on the level of mosaicity in basaltic rocks not affected by impact events, so that it could be used either in meteorite studies as a background level of non-shock-produced mosaicity, or in other studies of igneous rocks as a useful level range for comparison purposes.

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SYNTHÈSE ET CONCLUSION

Dans cette thèse, l'étude de la solidification des magmas basaltiques, et donc les articles qui constituent ce travail, s'articulent autour du même composant central : l'olivine. Ce minéral est certes très commun et généralement abondant dans les basaltes, mais il se retrouve ici aussi sous la forme de cristaux déformés, d'origine magmatique. Ce point constitue clairement un aspect crucial de l'étude, la présence d'olivines déformées étant généralement une particularité des roches mantelliques et plutoniques, mais beaucoup plus rare en contexte volcanique. Les cristaux déformés présents dans nos laves du Kilauea ont pour source commune majoritaire un cumulat de base d'édifice, situé au niveau du décollement majeur, à ~10-12 km de profondeur. L'analyse structurale de ces cristaux d'olivine déformés s'insère dans l'optique générale d'une meilleure compréhension de la déformation et du système magmatique hôte.

Dans le premier article (Chapitre 1), les principaux processus de solidification magmatique ont été examinés et quantifiés à des profondeurs de 0 à 12 km à l'intérieur du Kilauea. Un modèle pétrogénétique et semi-chronologique simple expliquant la dynamique du système magmatique ("*plumbing*") à ces profondeurs est également proposé. Ce modèle traite de l'origine et de l'histoire de solidification des magmas de l'éruption de 1969-1974 du Mauna Ulu, mais rend aussi compte de la dynamique plus globale qui affecte les "entrailles" du Kilauea. En résumé, cet article met en évidence l'existence de deux populations majeures d'olivines magmatiques, cogénétiques par ailleurs. Parmi ces olivines, de nombreux cristaux déformés (présence de bandes de déformation, extinction ondulante, etc.) sont présents. Aucune olivine mantellique n'est reconnue

ici ; les cristaux déformés sont en fait d'origine magmatique. Les résultats montrent que les magmas à l'origine de l'éruption du Mauna Ulu ont suivi deux trajets différents : (1) transit du magma transportant la plupart des olivines déformées le long du décollement majeur sous l'édifice, puis remontée à travers des sortes de conduits verticaux jusqu'à la région sous Mauna Ulu ; et (2) remontée du magma transportant les olivines non déformées par le conduit magmatique principal jusqu'au réservoir sommital, puis transfert sous la *East Rift Zone* et mélange des magmas dans des réservoirs secondaires.

Dans le deuxième article (Chapitre 2) l'approche combinée de la chimie *in situ* et des CSDs de l'olivine appliquée aux échantillons de surface du Kilauea Iki et aux échantillons du lac de lave (profondeurs : 0-90 m), a permis de déterminer et quantifier toute une série de processus de solidification pré-éruption et intra-lac. La géochimie et les CSDs révèlent la présence de deux populations majeures et une population mineure d'olivines ; ces populations ne sont pas nécessairement directement reliées. À l'image du Mauna Ulu, nombre de cristaux présentent des évidences pétrologiques de déformation, mais aucun xénocrystal mantellique n'est présent. Nos résultats suggèrent l'influence majeure du mélange de magmas (ou cristaux) et du mûrissement textural dans le système magmatique avant éruption. Ces processus deviennent peu actifs lors du refroidissement du lac. La modélisation CSD n'a fourni aucune évidence de sédimentation gravitaire des cristaux, ni de convection. Cet article met aussi l'accent sur l'existence dans le lac de lave d'une stratification verticale minéralogique et chimique dont l'origine reste quelque peu ambiguë. Il se peut que ce soit le résultat d'une stratification originelle, dérivée de la stratification du réservoir magmatique au moment de l'éruption. Cependant, cette hypothèse est en désaccord avec les observations faites en cours d'éruption qui démontrent clairement que le magma du lac a

subi un fort brassage tout au long des 17 phases éruptives. Le remplissage du lac de lave par sa base lors de l'éruption constitue une autre possibilité proposée ici.

Le troisième article (Chapitre 3) présente l'analyse microstructurale *in situ* par micro-diffraction des rayons X (μ XRD) d'olivines déformées et non déformées provenant de certains échantillons déjà étudiés dans les articles 1 et 2. Cette étude utilise une technique innovante, non destructive, peu coûteuse et rapide à mettre en œuvre permettant de recueillir des informations sur la structure interne des cristaux, ainsi que le mode et l'intensité de déformation. Dr. Roberta Flemming, *University of Western Ontario, London, Canada*, m'a appris comment utiliser le diffractomètre à rayons X et m'a aidé dans l'analyse des données brutes. Elle a également participé à la rédaction de la méthodologie de l'article. Les résultats ont permis de valider les observations pétrographiques de déformation faites à l'aide du microscope. Cette analyse μ XRD a aussi permis de confirmer la présence de déformation pour toutes les tailles de grains d'olivine, sans corrélation simple avec leur chimie, et de quantifier cette déformation. Cette technique ne permet cependant pas une estimation simple des conditions pression-température de déformation ou de formation des cristaux, ni d'apporter d'informations sur l'histoire magmatique. Il a cependant été possible de fixer un seuil quantitatif au-delà duquel toute olivine est déformée de façon significative.

Globalement, cette thèse a permis de mieux comprendre et de quantifier les principaux processus de solidification des magmas basaltiques, via l'étude des produits volcaniques de l'éruption latérale de 1969-1974 du Mauna Ulu (système ouvert), et l'éruption sommitale de 1959 du Kilauea Iki (système fermé). L'olivine a été le minéral de choix puisque très abondant, précoce dans l'histoire de la solidification, et adapté à l'étude combinée des compositions chimiques et des

textures magmatiques. La chromite aurait été un minéral complémentaire intéressant dans ce type d'approche, puisqu'il est lui aussi abondant, précoce, et souvent associé à l'olivine dans nos échantillons. Cependant, l'étude de la chromite fera uniquement partie de projets futurs.

Deux modèles de solidification magmatique sont proposés (Chapitres 1 et 2), l'un appliqué aux systèmes ouverts et l'autre aux systèmes fermés. Nos modèles se révèlent être partiellement en accord avec les modèles présentés dans la problématique. Ils amènent également une part significative de nouveauté ou de raffinement, en particulier grâce à la quantification des mécanismes en général, incluant une estimation temporelle via l'analyse CSD. Par exemple, les mouvements convectifs et la sédimentation gravitaire à grande échelle dans le lac de lave sont ici considérés comme négligeables, ce qui est en contradiction avec la plupart des travaux antérieurs.

Un modèle de transfert des magmas à l'intérieur du système magmatique du Kilauea est également proposé (Chapitre 1). Ce modèle s'applique sur toute l'épaisseur de l'édifice (0-12 km de profondeur).

Cette thèse traite aussi plus en détail l'aspect déformation magmatique de l'olivine, et amène une quantification appliquée au milieu naturel. Une réponse positive est apportée à la question de la quantification et reconnaissance de ce type de déformation magmatique. La combinaison de la pétrographie classique et de l'analyse structurale *in situ* par μ XRD se révèle être une approche adaptée dans ce cas. Cependant, la question du mode de déformation à l'œuvre dans la structure même de l'olivine ne trouve pas de réponse précise ici ; la μ XRD n'apporte pas toutes les informations requises. Plus précisément, le lien entre les résultats μ XRD et le mode de déformation cristalline n'est pas direct, si lien simple il y a.

D'un point de vue méthodologique global, cette étude démontre clairement l'intérêt de combiner les approches texturale et géochimique, notamment dans le cas de systèmes naturels en domaine volcanique. Cette conclusion est vraie autant en système ouvert que fermé, même si la plupart des modèles CSD sont par défaut adaptés aux systèmes ouverts ; une approximation en système fermé est alors requise, mais tout à fait valable. Dans les Chapitres 1 et 2 la double approche basée sur la combinaison distribution de la taille des cristaux et géochimie, a permis d'apporter davantage d'informations quantitatives sur les processus de solidification des magmas basaltiques. En effet, certains de ces processus n'affectent que la texture des roches (i.e. nucléation et croissance cristallines, déformation, mûrissement textural) ; ils ne sont donc quantifiables que par l'approche texturale. D'autres processus affectent à la fois la texture et la chimie des roches (i.e. mélange, fractionnement et accumulation) ; ils peuvent donc être étudiés par la combinaison des deux approches, texturale et géochimique. Cependant, les effets texturaux de plusieurs processus (e.g. agrégation vs. mûrissement textural) sur les CSDs peuvent se superposer (partiellement ou totalement), et donc engendrer une certaine ambiguïté dans l'interprétation des résultats. Ce cas de figure a régulièrement posé problème tout au long du doctorat, et l'aspect géochimique s'est alors révélé indispensable pour aller plus de l'avant.

Concernant l'analyse microstructurale de la déformation des olivines, il serait intéressant à terme de mener ce genre d'études μ XRD sur une large gamme de cristaux, olivine et autres silicates naturels, déformés et non déformés, dont les conditions de formation et l'environnement magmatique sont parfaitement connus. L'apport de l'analyse μ XRD d'olivines (ou autres silicates) expérimentales serait également significatif. Cette démarche permettrait de créer une base de données qualitatives et quantitatives complète, utile pour documenter la déformation magmatique des minéraux silicatés, trop peu contrainte à l'heure actuelle. Notre étude μ XRD met

en évidence la relation entre l'intensité de déformation (ou de "mosaïcité"), exprimée par le paramètre FWHM, et la composition de l'olivine ou la taille de grain. Le niveau de mosaïcité, très utile dans les études sur les météorites, est ainsi contraint ici dans le cas de roches basaltiques non affectées par un impact météorique, et pourrait servir de base de comparaison pour de futures études sur les météorites (comme seuil de référence) ou sur les roches ignées.

Finalement, même si la signification des résultats est difficile à prédire dans un domaine d'étude en cours de développement telle que la quantification texturale appliquée aux roches magmatiques, cette thèse apporte l'évidence que l'application d'une dynamique de recherche exploratoire peut aboutir à la formation de modèles cohérents. De plus, le développement de *l'outil textural pour l'étude des magmas pourrait aboutir à de nouvelles conclusions (ou théories)* quant aux processus magmatiques associés. Par exemple, si le mûrissement textural est un processus pétrologique comparable en importance à la décantation/sédimentation des cristaux, il pourrait avoir des implications importantes sur la façon dont les compositions chimiques et isotopiques des roches et minéraux sont interprétées. De plus, la quantification texturale des roches magmatiques est un domaine de recherche en pleine croissance. Il est donc nécessaire de profiter de toutes les opportunités pour un avancement rapide des connaissances en la matière. En résumé, ce travail participe pleinement à l'essor global de la pétrologie vers l'aspect textural et microstructural.

ANNEXE 1

Compositions en éléments majeurs et mineurs d'olivines des laves de l'éruption de 1969-1974 du Mauna Ulu. Ces données sont utilisées dans l'article 1 (i.e. pour les graphiques et dans le texte), mais seule une sélection a été publiée.

Sample	Spot	Deformed	(wt%)					CaO	MnO	FeO*	NiO	Total	(mol)		
			SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO						MgO	FeO	Fo%
MU69-1A 1b1	rim 1a		40,20	0,00	0,05	0,09	47,61	0,24	0,11	11,47	0,45	100,22	1,18	0,14	89,2
MU69-1A 1b2	rim 2		39,49	0,02	0,06	0,04	46,05	0,25	0,16	13,32	0,30	99,70	1,14	0,17	87,3
MU69-1A 1b3	rim 1b		39,26	0,02	0,03	0,05	45,07	0,29	0,14	14,92	0,31	100,13	1,12	0,19	85,7
MU69-1A 1c1	core 1		39,87	0,01	0,00	0,04	47,50	0,26	0,16	11,41	0,34	99,63	1,18	0,14	89,2
MU69-1A 1c2	core 2		39,98	0,01	0,02	0,06	47,99	0,25	0,08	11,33	0,38	100,08	1,19	0,14	89,3
MU69-1A 2b1	rim a	X	39,44	0,02	0,07	0,04	46,42	0,28	0,21	13,24	0,28	100,00	1,15	0,17	87,4
MU69-1A 2b2	rim b	X	39,49	0,00	0,00	0,14	46,78	0,26	0,17	12,48	0,39	99,79	1,16	0,16	88,1
MU69-1A 2c1	core	X	39,72	0,00	0,03	0,01	47,38	0,24	0,11	11,42	0,39	99,31	1,18	0,14	89,2
MU69-1A 3b1	rim 1		39,26	0,02	0,06	0,02	44,88	0,27	0,22	14,78	0,30	99,85	1,11	0,19	85,7
MU69-1A 3b3	rim 3		39,40	0,00	0,00	0,07	45,72	0,27	0,15	13,82	0,30	99,74	1,13	0,17	86,8
MU69-1A 3c1	core 1		39,79	0,00	0,07	0,09	46,72	0,27	0,16	12,37	0,37	99,85	1,16	0,15	88,2
MU69-1A 3c2	core 2		39,55	0,00	0,04	0,03	46,71	0,25	0,11	12,37	0,32	99,38	1,16	0,15	88,2
MU69-1A 3c3	core 3		39,49	0,00	0,07	0,13	46,94	0,25	0,18	11,94	0,32	99,32	1,16	0,15	88,6
MU69-1A 4b1	rim 1		39,51	0,01	0,06	0,08	46,57	0,25	0,17	12,51	0,33	99,53	1,16	0,16	88,1
MU69-1A 4b2	rim 2		39,77	0,02	0,05	0,10	46,75	0,28	0,14	12,64	0,38	100,13	1,16	0,16	88,0
MU69-1A 4b3	rim 3		39,30	0,05	0,07	0,10	44,41	0,35	0,22	15,32	0,21	100,03	1,10	0,19	85,2
MU69-1A 4c1	core 1		39,70	0,01	0,00	0,07	47,39	0,25	0,15	11,41	0,35	99,34	1,18	0,14	89,2
MU69-1A 4c2	core 2		39,88	0,01	0,07	0,08	47,52	0,24	0,14	11,59	0,40	99,96	1,18	0,15	89,0
MU69-1A 4c3	core 3		39,77	0,00	0,02	0,13	47,20	0,25	0,13	12,42	0,37	100,31	1,17	0,16	88,3
MU69-1A 5b1	rim	X	39,45	0,02	0,00	0,02	45,19	0,26	0,21	14,29	0,35	99,84	1,12	0,18	86,2
MU69-1A 5c1	core	X	40,06	0,00	0,05	0,12	47,61	0,23	0,17	11,91	0,44	100,58	1,18	0,15	88,8
MU69-1A 6b1	rim 1		39,28	0,00	0,05	0,02	45,52	0,27	0,20	13,72	0,30	99,38	1,13	0,17	86,8
MU69-1A 6b2	rim 2		39,02	0,03	0,06	0,06	44,76	0,28	0,10	14,66	0,33	99,31	1,11	0,18	85,8
MU69-1A 6b3	rim 3		39,45	0,03	0,06	0,04	46,22	0,26	0,16	13,10	0,31	99,63	1,15	0,16	87,5
MU69-1A 6c1	core 1		39,80	0,01	0,20	0,06	47,63	0,23	0,21	11,40	0,37	99,90	1,18	0,14	89,2
MU69-1A 6c2	core 2		39,71	0,01	0,10	0,10	47,11	0,24	0,20	11,97	0,38	99,81	1,17	0,15	88,6
MU69-1A 6c3	core 3		39,66	0,00	0,09	0,05	47,19	0,25	0,14	11,76	0,41	99,54	1,17	0,15	88,8
MU69-1A 7b1	rim 1	X	39,19	0,02	0,06	0,05	44,74	0,26	0,15	14,88	0,33	99,71	1,11	0,19	85,6
MU69-1A 7c1	core 1	X	39,75	0,00	0,00	0,06	47,62	0,24	0,15	11,55	0,42	99,79	1,18	0,14	89,1
MU69-1A 7c2	core 2		39,67	0,01	0,06	0,10	47,11	0,24	0,03	12,18	0,35	99,75	1,17	0,15	88,5
MU69-1A 8b1	rim		39,59	0,04	0,01	0,03	46,30	0,28	0,17	12,96	0,38	99,78	1,15	0,16	87,6
MU69-1A 8c1	core		39,80	0,01	0,07	0,02	47,40	0,24	0,16	11,80	0,40	99,92	1,18	0,15	88,8
MU69-1A 9b1	rim a	X	38,84	0,01	0,07	0,00	43,73	0,22	0,24	16,25	0,38	99,75	1,09	0,20	84,2
MU69-1A 9b2	rim b	X	39,48	0,01	0,05	0,13	46,16	0,26	0,22	13,32	0,33	99,97	1,15	0,17	87,3

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU69-1A 9b3	rim c	X	39,40	0,00	0,06	0,06	46,34	0,26	0,11	13,03	0,33	99,59	1,15	0,16	87,6
MU69-1A 9b4	rim d	X	39,63	0,03	0,05	0,03	46,14	0,27	0,23	13,02	0,34	99,77	1,14	0,16	87,5
MU69-1A 9c1	core a	X	37,91	0,01	0,05	0,00	39,56	0,19	0,29	21,62	0,25	99,89	0,98	0,27	78,4
MU69-1A 9c2	core b	X	37,98	0,00	0,00	0,00	39,42	0,21	0,30	21,71	0,24	99,86	0,98	0,27	78,2
MU69-1A 10b1	rim 1		39,65	0,00	0,07	0,11	46,30	0,26	0,16	12,94	0,35	99,85	1,15	0,16	87,6
MU69-1A 10b2	rim 2		39,51	0,03	0,05	0,09	45,90	0,28	0,14	13,37	0,34	99,69	1,14	0,17	87,2
MU69-1A 10c1	core 1		40,09	0,00	0,05	0,13	48,10	0,22	0,18	11,32	0,37	100,49	1,19	0,14	89,4
MU69-1A 10c2	core 2		40,11	0,02	0,06	0,06	47,42	0,23	0,16	11,48	0,35	99,91	1,18	0,14	89,1
MU69-1A 10c3	core 3		40,06	0,02	0,04	0,08	47,74	0,26	0,12	11,34	0,39	100,07	1,18	0,14	89,3
MU69-1A 10c4	core 4		40,00	0,01	0,00	0,09	47,76	0,25	0,18	11,11	0,37	99,79	1,18	0,14	89,5
MU69-1A 11b1	rim a	X	38,63	0,02	0,05	0,07	43,68	0,31	0,14	15,89	0,29	99,09	1,08	0,20	84,5
MU69-1A 11b2	rim b	X	38,91	0,01	0,05	0,01	42,71	0,27	0,23	17,70	0,31	100,22	1,06	0,22	82,7
MU69-1A 11c1	core a	X	40,20	0,01	0,05	0,08	48,00	0,24	0,16	11,65	0,39	100,79	1,19	0,15	89,1
MU69-1A 11c2	core b	X	39,51	0,01	0,06	0,16	46,85	0,24	0,22	12,21	0,38	99,65	1,16	0,15	88,4
MU69-1A 12b1	rim 1		39,25	0,00	0,00	0,08	44,05	0,28	0,20	15,90	0,29	100,06	1,09	0,20	84,6
MU69-1A 12c1	core 1		39,91	0,00	0,05	0,12	46,91	0,24	0,22	12,67	0,39	100,56	1,16	0,16	88,0
MU69-1A 12c2	core 2		39,56	0,01	0,05	0,01	46,41	0,24	0,19	12,81	0,28	99,56	1,15	0,16	87,8
MU69-1A 12c3	core 3		39,78	0,00	0,00	0,15	47,15	0,23	0,22	12,67	0,35	100,55	1,17	0,16	88,1
MU69-1A 13b1	rim a	X	39,77	0,05	0,00	0,14	45,79	0,28	0,14	13,96	0,39	100,54	1,14	0,17	86,7
MU69-1A 13b2	rim b	X	38,79	0,03	0,08	0,09	42,71	0,27	0,20	17,43	0,26	99,91	1,06	0,22	82,9
MU69-1A 13c1	core	X	39,55	0,00	0,00	0,01	47,04	0,25	0,04	11,82	0,34	99,08	1,17	0,15	88,7
MU69-1A 14b1	rim 1a	X	39,17	0,04	0,04	0,08	44,52	0,31	0,19	14,90	0,31	99,60	1,10	0,19	85,5
MU69-1A 14b3	rim 2	X	39,29	0,02	0,03	0,05	45,63	0,28	0,29	13,85	0,28	99,77	1,13	0,17	86,7
MU69-1A 14c1	core 1	X	40,15	0,02	0,00	0,03	46,91	0,20	0,18	12,53	0,32	100,38	1,16	0,16	88,1
MU69-1A 14c2	~rim 1b	X	39,68	0,02	0,05	0,09	46,43	0,25	0,21	12,70	0,36	99,82	1,15	0,16	87,9
MU69-1A 14c3	core 2	X	39,50	0,01	0,00	0,03	46,06	0,21	0,15	13,24	0,35	99,60	1,14	0,17	87,3
MU69-1B 1b1	rim a	X	38,71	0,04	0,06	0,03	43,07	0,30	0,17	17,08	0,22	99,72	1,07	0,21	83,3
MU69-1B 1b2	rim b	X	37,74	0,05	0,04	0,04	39,27	0,34	0,20	21,17	0,21	99,07	0,97	0,27	78,6
MU69-1B 1b3	rim c	X	38,59	0,03	0,04	0,08	42,73	0,31	0,22	17,87	0,21	100,12	1,06	0,22	82,6
MU69-1B 1b4	rim d	X	39,74	0,03	0,06	0,01	46,43	0,27	0,22	12,95	0,38	100,10	1,15	0,16	87,7
MU69-1B 1c1	core a	X	39,84	0,01	0,04	0,08	47,39	0,25	0,18	11,48	0,41	99,69	1,18	0,14	89,1
MU69-1B 1c2	core b	X	39,84	0,01	0,00	0,13	47,39	0,22	0,04	11,34	0,39	99,38	1,18	0,14	89,2
MU69-1B 2b1	rim c	X	39,38	0,01	0,03	0,07	46,71	0,26	0,18	12,55	0,37	99,57	1,16	0,16	88,1
MU69-1B 2b4	rim d	X	39,18	0,01	0,05	0,05	44,28	0,27	0,23	14,95	0,24	99,31	1,10	0,19	85,4
MU69-1B 2c1	core a	X	39,97	0,00	0,06	0,12	47,32	0,24	0,23	12,07	0,36	100,37	1,17	0,15	88,6
MU69-1B 2c2	core b	X	39,63	0,03	0,08	0,11	47,31	0,27	0,19	11,57	0,33	99,54	1,17	0,14	89,0
MU69-1B 2c3	core c	X	39,83	0,01	0,06	0,10	47,56	0,28	0,18	11,24	0,35	99,62	1,18	0,14	89,3
MU69-1B 2c4	core d	X	40,10	0,01	0,06	0,14	47,63	0,27	0,13	11,30	0,37	100,07	1,18	0,14	89,3
MU69-1B 3b1	rim a		39,44	0,04	0,06	0,12	44,94	0,30	0,18	14,95	0,31	100,33	1,12	0,19	85,6
MU69-1B 3b2	rim b		39,68	0,00	0,04	0,07	46,00	0,25	0,12	14,19	0,37	100,73	1,14	0,18	86,5

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU69-1B 3b3	rim c		39,30	0,00	0,03	0,00	44,90	0,27	0,14	14,90	0,29	99,85	1,11	0,19	85,7
MU69-1B 3c1	core		39,61	0,00	0,06	0,02	46,35	0,27	0,22	12,90	0,35	99,79	1,15	0,16	87,7
MU69-1B 4b1	rim		38,63	0,04	0,08	0,07	43,74	0,30	0,17	15,78	0,24	99,05	1,09	0,20	84,6
MU69-1B 4c1	core		39,51	0,00	0,04	0,08	46,07	0,21	0,21	13,10	0,30	99,50	1,14	0,16	87,4
MU69-1B 5b1	rim a		39,33	0,01	0,04	0,02	46,52	0,25	0,13	12,81	0,32	99,43	1,15	0,16	87,8
MU69-1B 5b2	rim b		38,94	0,02	0,03	0,05	44,56	0,26	0,23	15,25	0,32	99,66	1,11	0,19	85,3
MU69-1B 5b3	rim c		36,78	0,08	0,04	0,04	34,36	0,29	0,31	28,24	0,17	100,35	0,85	0,35	70,7
MU69-1B 5b4	rim d		39,02	0,01	0,00	0,04	44,44	0,26	0,24	15,12	0,30	99,45	1,10	0,19	85,3
MU69-1B 5b5	rim e		38,62	0,03	0,05	0,04	43,31	0,27	0,24	16,57	0,30	99,47	1,07	0,21	83,8
MU69-1B 5c1	core		39,45	0,00	0,00	0,06	46,38	0,27	0,19	12,90	0,30	99,58	1,15	0,16	87,7
MU69-1B 6b1	rim a	X	39,58	0,01	0,03	0,08	47,07	0,25	0,19	13,00	0,33	100,54	1,17	0,16	87,8
MU69-1B 6b2	rim b	X	39,43	0,01	0,05	0,05	45,39	0,26	0,22	14,46	0,30	100,24	1,13	0,18	86,1
MU69-1B 6b3	rim c	X	39,87	0,03	0,05	0,08	47,19	0,27	0,05	11,56	0,37	99,48	1,17	0,14	89,0
MU69-1B 6b4	rim d	X	39,66	0,02	0,04	0,02	46,59	0,25	0,12	12,54	0,38	99,66	1,16	0,16	88,0
MU69-1B 6c1	core a	X	39,83	0,00	0,06	0,08	47,30	0,27	0,10	11,80	0,34	99,82	1,17	0,15	88,8
MU69-1B 6c2	core b	X	39,64	0,00	0,05	0,05	46,73	0,27	0,20	11,99	0,32	99,26	1,16	0,15	88,5
MU69-1B 6c3	core c	X	39,88	0,02	0,00	0,06	47,63	0,24	0,14	11,14	0,46	99,55	1,18	0,14	89,4
MU69-1B 6c4	core d	X	39,71	0,00	0,08	0,14	47,23	0,21	0,12	11,60	0,43	99,54	1,17	0,15	89,0
MU69-1B 7b1	rim 1	X	39,15	0,04	0,02	0,08	43,74	0,31	0,21	16,09	0,32	99,97	1,09	0,20	84,3
MU69-1B 7b2	rim 2		36,12	0,10	0,00	0,01	31,73	0,29	0,22	30,53	0,15	99,17	0,79	0,38	67,3
MU69-1B 7b5	rim 4		39,30	0,04	0,06	0,05	44,70	0,29	0,23	14,76	0,30	99,75	1,11	0,18	85,7
MU69-1B 7c1	core 1	X	40,08	0,00	0,04	0,05	47,46	0,24	0,17	11,27	0,38	99,69	1,18	0,14	89,3
MU69-1B 7c2	core 2a		40,08	0,02	0,02	0,01	47,64	0,24	0,08	11,10	0,44	99,66	1,18	0,14	89,5
MU69-1B 7c3	core 2b		39,99	0,00	0,07	0,08	47,73	0,24	0,14	11,18	0,33	99,75	1,18	0,14	89,4
MU69-1B 7c4	core 3		39,80	0,00	0,00	0,04	47,65	0,25	0,15	11,30	0,41	99,64	1,18	0,14	89,3
MU69-1B 7c5	core 4		40,19	0,00	0,00	0,06	47,60	0,26	0,20	11,07	0,40	99,79	1,18	0,14	89,5
MU69-1B 7c6	core 5		39,74	0,01	0,10	0,02	47,71	0,23	0,10	11,35	0,41	99,67	1,18	0,14	89,3
MU69-1B 8b1	rim a		37,70	0,03	0,05	0,04	38,85	0,34	0,19	22,30	0,19	99,70	0,96	0,28	77,5
MU69-1B 8b2	rim b		39,20	0,00	0,05	0,03	45,35	0,28	0,08	14,14	0,34	99,47	1,13	0,18	86,4
MU69-1B 8c1	core		39,58	0,03	0,01	0,05	46,33	0,26	0,20	13,13	0,28	99,87	1,15	0,16	87,5
MU69-1B 9b1	rim a		38,88	0,04	0,17	0,07	42,52	0,41	0,24	17,27	0,19	99,82	1,05	0,22	83,0
MU69-1B 9b2	rim b		39,10	0,01	0,01	0,08	43,08	0,32	0,19	16,99	0,27	100,06	1,07	0,21	83,4
MU69-1B 9b3	rim c		39,04	0,02	0,07	0,01	43,25	0,31	0,18	16,44	0,29	99,64	1,07	0,21	83,9
MU69-1B 9c1	core		39,47	0,01	0,06	0,06	46,26	0,25	0,20	12,71	0,30	99,32	1,15	0,16	87,8
MU69-1B 10b1	rim 1		38,89	0,03	0,07	0,08	43,55	0,28	0,17	16,63	0,31	100,02	1,08	0,21	83,8
MU69-1B 10b2	rim 2		38,50	0,01	0,01	0,07	42,85	0,29	0,19	16,78	0,30	99,03	1,06	0,21	83,5
MU69-1B 10b3	rim 3		38,59	0,09	0,18	0,12	43,07	0,31	0,22	17,61	0,26	100,47	1,07	0,22	82,9
MU69-1B 10c1	core 1		39,60	0,00	0,05	0,04	46,39	0,28	0,16	12,99	0,31	99,86	1,15	0,16	87,6
MU69-1B 10c2	core 2		39,54	0,01	0,09	0,09	46,57	0,27	0,13	12,67	0,34	99,71	1,16	0,16	87,9
MU69-1B 10c3	core 3		39,85	0,00	0,00	0,08	46,95	0,27	0,25	12,41	0,33	100,14	1,16	0,16	88,2
MU69-1B 11b1	rim a		38,86	0,01	0,00	0,00	43,53	0,15	0,26	16,30	0,29	99,42	1,08	0,20	84,1

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU69-1B 11b2	rim b		39,41	0,05	0,00	0,01	44,88	0,30	0,11	14,76	0,33	99,85	1,11	0,18	85,8
MU69-1B 11b3	rim c		38,81	0,05	0,00	0,12	44,15	0,30	0,18	15,20	0,32	99,13	1,10	0,19	85,2
MU69-1B 11b4	rim d		39,30	0,02	0,02	0,05	45,11	0,26	0,21	14,38	0,28	99,62	1,12	0,18	86,1
MU69-1B 11c1	core a		38,50	0,00	0,05	0,05	42,03	0,21	0,31	18,95	0,29	100,41	1,04	0,24	81,5
MU69-1B 11c2	core b		38,60	0,04	0,07	0,11	42,22	0,16	0,24	18,08	0,28	99,81	1,05	0,23	82,2
MU69-1B 12b1	rim a		39,17	0,01	0,01	0,06	44,10	0,30	0,12	15,56	0,32	99,65	1,09	0,19	84,9
MU69-1B 12b2	rim b		38,89	0,04	0,07	0,07	43,58	0,29	0,22	16,34	0,25	99,75	1,08	0,20	84,1
MU69-1B 12c1	core		39,58	0,00	0,04	0,00	46,29	0,25	0,16	12,88	0,37	99,61	1,15	0,16	87,7
MU69-1B 13b1	rim		39,29	0,04	0,05	0,17	46,01	0,29	0,17	14,06	0,29	100,38	1,14	0,18	86,6
MU69-1B 13c1	core		39,57	0,01	0,03	0,09	46,98	0,26	0,14	13,14	0,34	100,57	1,17	0,16	87,6
MU70-1 1b1	rim a		38,99	0,01	0,06	0,04	42,99	0,32	0,28	17,05	0,21	99,95	1,07	0,21	83,3
MU70-1 1b2	rim b		39,71	0,01	0,03	0,02	44,45	0,31	0,19	16,18		100,90	1,10	0,20	84,5
MU70-1 1c1	core		39,35	0,00	0,08	0,02	45,40	0,27	0,20	14,48	0,26	100,09	1,13	0,18	86,1
MU70-1 2b1	rim 2		38,66	0,01	0,06	0,00	42,62	0,28	0,24	16,90	0,16	98,99	1,06	0,21	83,3
MU70-1 2c1	core 1		39,46	0,01	0,05	0,04	45,45	0,25	0,09	14,26	0,30	99,97	1,13	0,18	86,3
MU70-1 2c2	core 2		39,51	0,02	0,04	0,03	45,40	0,24	0,24	14,63	0,29	100,44	1,13	0,18	86,0
MU70-1 2c3	core 3		39,26	0,00	0,05	0,06	45,56	0,28	0,22	14,06	0,32	99,81	1,13	0,18	86,5
MU70-1 3b1	rim 1	X	38,64	0,00	0,03	0,04	42,93	0,31	0,15	17,11	0,24	99,46	1,07	0,21	83,3
MU70-1 3c1	core 1	X	39,74	0,04	0,04	0,04	46,39	0,12	0,22	13,29	0,38	100,26	1,15	0,17	87,4
MU70-1 3c2	core 2	X	39,12	0,03	0,03	0,00	44,11	0,16	0,19	15,92	0,28	99,85	1,09	0,20	84,6
MU70-1 4b1	rim 1		39,32	0,02	0,05	0,04	44,96	0,26	0,26	14,85	0,27	100,03	1,12	0,19	85,7
MU70-1 4b2	rim 2		39,39	0,01	0,03	0,03	44,32	0,25	0,16	16,43	0,27	100,92	1,10	0,21	84,2
MU70-1 4c1	core 1		39,71	0,00	0,08	0,12	46,62	0,26	0,16	12,93	0,37	100,30	1,16	0,16	87,7
MU70-1 4c2	core 2		39,54	0,02	0,05	0,06	45,70	0,23	0,25	13,98	0,36	100,19	1,13	0,18	86,6
MU70-1 5b1	rim 1	X	39,21	0,00	0,05	0,07	43,69	0,31	0,24	17,05	0,26	100,91	1,08	0,21	83,5
MU70-1 5b2	rim 2	X	38,58	0,01	0,05	0,03	42,43	0,32	0,26	17,52	0,23	99,42	1,05	0,22	82,7
MU70-1 5c1	core 1	X	39,53	0,00	0,05	0,03	46,30	0,27	0,19	13,46	0,31	100,17	1,15	0,17	87,2
MU70-1 5c2	core 2	X	39,68	0,00	0,06	0,03	46,30	0,25	0,23	13,81	0,33	100,68	1,15	0,17	86,9
MU70-1 6b1	rim 1		39,01	0,02	0,05	0,03	44,04	0,30	0,19	16,47	0,25	100,36	1,09	0,21	84,1
MU70-1 6b2	rim 3		38,80	0,00	0,04	0,04	42,89	0,30	0,25	17,09	0,22	99,65	1,06	0,21	83,2
MU70-1 6c1	core 1		39,35	0,00	0,06	0,00	44,35	0,27	0,29	16,25	0,26	100,83	1,10	0,20	84,4
MU70-1 6c2	core 2		39,55	0,01	0,03	0,09	45,50	0,24	0,16	14,87	0,32	100,76	1,13	0,19	85,8
MU70-1 6c3	core 3		39,45	0,02	0,04	0,05	44,26	0,25	0,17	15,91	0,23	100,41	1,10	0,20	84,6
MU70-1 7b1	rim	X	38,75	0,01	0,03	0,05	43,05	0,30	0,20	17,42	0,21	100,05	1,07	0,22	83,0
MU70-1 7c1	core	X	39,51	0,02	0,03	0,00	45,62	0,27	0,18	14,43	0,26	100,34	1,13	0,18	86,2
MU70-1 8b1	rim		38,78	0,00	0,06	0,07	43,06	0,31	0,24	16,58	0,23	99,38	1,07	0,21	83,7
MU70-1 8c1	core		39,58	0,00	0,04	0,00	45,05	0,28	0,20	14,94	0,31	100,42	1,12	0,19	85,7
MU70-1 9b1	rim		38,99	0,01	0,03	0,03	44,00	0,29	0,28	16,19	0,23	100,08	1,09	0,20	84,3
MU70-1 9c1	core		39,40	0,00	0,07	0,04	45,33	0,26	0,19	14,36	0,22	99,88	1,12	0,18	86,2

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU70-1 10b1	rim		39,30	0,01	0,05	0,09	45,62	0,25	0,24	14,07	0,28	99,90	1,13	0,18	86,5
MU70-1 10c1	core		39,19	0,01	0,05	0,05	45,22	0,24	0,18	14,56	0,33	99,87	1,12	0,18	86,0
MU70-1 11b1	rim 1		38,81	0,02	0,06	0,07	42,84	0,33	0,26	17,58	0,24	100,25	1,06	0,22	82,8
MU70-1 11b2	rim 2		38,73	0,01	0,07	0,10	42,89	0,30	0,19	17,34	0,23	99,88	1,06	0,22	83,0
MU70-1 11c1	core 1		39,36	0,00	0,05	0,04	44,84	0,23	0,15	15,40	0,29	100,37	1,11	0,19	85,2
MU70-1 11c2	core 2		39,28	0,02	0,03	0,04	44,21	0,25	0,19	15,96	0,21	100,21	1,10	0,20	84,6
MU70-1 12b1	rim 1		39,02	0,01	0,05	0,04	43,84	0,27	0,21	16,08	0,28	99,82	1,09	0,20	84,4
MU70-1 12b2	rim 2	X	39,19	0,01	0,04	0,07	44,12	0,27	0,23	16,36	0,29	100,57	1,09	0,20	84,2
MU70-1 12c1	core 1		39,18	0,00	0,05	0,09	44,73	0,26	0,19	15,28	0,28	100,05	1,11	0,19	85,3
MU70-1 12c2	core 2	X	39,80	0,00	0,04	0,00	45,40	0,28	0,14	14,52	0,29	100,52	1,13	0,18	86,1
MU70-1 13b1	rim		39,83	0,00	0,07	0,05	45,83	0,24	0,21	14,15	0,31	100,73	1,14	0,18	86,5
MU70-1 13c1	core		39,86	0,01	0,02	0,01	45,79	0,27	0,23	13,95	0,32	100,49	1,14	0,17	86,7
MU70-1 14b1	rim		39,36	0,00	0,03	0,06	45,19	0,27	0,24	14,49	0,22	99,91	1,12	0,18	86,1
MU70-1 14c1	core		39,58	0,00	0,08	0,00	45,30	0,28	0,28	14,83	0,28	100,63	1,12	0,19	85,8
MU70-1 15b1	rim 1		39,12	0,00	0,00	0,12	44,65	0,27	0,22	15,68	0,26	100,34	1,11	0,20	84,9
MU70-1 15b2	rim 2	X	38,46	0,01	0,04	0,09	42,71	0,31	0,28	17,59	0,30	99,81	1,06	0,22	82,8
MU70-1 15c1	core 1		39,48	0,00	0,03	0,00	44,99	0,28	0,18	15,48	0,23	100,71	1,12	0,19	85,2
MU70-1 15c2	core 2	X	39,59	0,00	0,02	0,04	44,90	0,28	0,26	14,80	0,22	100,16	1,11	0,19	85,7
MU70-1 16b1	rim 1		38,34	0,00	0,05	0,02	41,08	0,26	0,16	19,51	0,25	99,69	1,02	0,24	80,7
MU70-1 16b2	rim 2		38,83	0,04	0,03	0,03	42,68	0,30	0,19	17,66	0,28	100,05	1,06	0,22	82,7
MU70-1 16c1	core 1		39,33	0,01	0,02	0,09	43,42	0,24	0,23	16,98	0,34	100,68	1,08	0,21	83,5
MU70-1 16c2	~core 2		38,51	0,02	0,04	0,00	41,23	0,23	0,26	19,85	0,28	100,44	1,02	0,25	80,4
MU70-1 16c3	~core 3		38,49	0,02	0,06	0,07	40,53	0,23	0,24	20,67	0,24	100,58	1,01	0,26	79,5
MU70-1 16c4	~core 4		38,42	0,01	0,07	0,02	40,07	0,23	0,28	21,39	0,33	100,86	0,99	0,27	78,8
MU70-1 17b1	rim a		39,14	0,01	0,03	0,04	43,29	0,28	0,19	16,98	0,22	100,21	1,07	0,21	83,5
MU70-1 17b2	rim b		39,14	0,02	0,06	0,05	43,63	0,29	0,20	17,01	0,22	100,61	1,08	0,21	83,6
MU70-1 17c1	core		39,80	0,01	0,04	0,12	46,01	0,24	0,18	14,09	0,31	100,80	1,14	0,18	86,6
MU70-1 18b1	rim		39,70	0,01	0,06	0,18	45,76	0,26	0,20	14,58	0,34	101,09	1,14	0,18	86,1
MU70-1 18c1	core		39,98	0,00	0,05	0,11	47,72	0,24	0,14	11,75	0,37	100,35	1,18	0,15	88,9
MU70-1 19b1	rim a		39,11	0,02	0,08	0,01	43,21	0,28	0,24	17,07	0,23	100,29	1,07	0,21	83,4
MU70-1 19b2	rim b		39,04	0,01	0,03	0,05	43,44	0,30	0,09	17,24	0,26	100,45	1,08	0,22	83,3
MU70-1 19c1	core		39,09	0,01	0,03	0,00	44,62	0,19	0,28	15,97	0,29	100,47	1,11	0,20	84,7
MU72-2 1b1	rim 1		39,01	0,00	0,09	0,00	42,74	0,29	0,20	17,48	0,22	100,05	1,06	0,22	82,9
MU72-2 1b2	rim 2		39,17	0,01	0,05	0,02	43,68	0,29	0,26	16,68	0,23	100,42	1,08	0,21	83,8
MU72-2 1c1	core 1		39,19	0,00	0,05	0,01	43,90	0,28	0,23	16,35	0,22	100,27	1,09	0,20	84,2
MU72-2 1c2	core 2		39,16	0,02	0,05	0,06	43,99	0,31	0,22	16,11	0,20	100,15	1,09	0,20	84,4
MU72-2 2b1	rim		39,03	0,00	0,03	0,10	42,94	0,33	0,24	17,82	0,18	100,67	1,07	0,22	82,7
MU72-2 2c1	core		39,74	0,00	0,02	0,05	45,97	0,29	0,20	13,94	0,30	100,51	1,14	0,17	86,7
MU72-2 3b1	rim a		38,60	0,00	0,05	0,00	42,54	0,31	0,29	17,26	0,22	99,26	1,06	0,22	83,0
MU72-2 3b2	rim b		38,87	0,01	0,06	0,04	42,56	0,31	0,29	17,33	0,20	99,69	1,06	0,22	82,9

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU72-2 3c1	core		39,08	0,00	0,04	0,05	43,81	0,28	0,30	16,53	0,29	100,38	1,09	0,21	84,0
MU72-2 4b1	rim		39,12	0,03	0,04	0,00	43,11	0,31	0,31	17,72	0,14	100,81	1,07	0,22	82,8
MU72-2 4c1	core		39,50	0,00	0,01	0,07	44,56	0,28	0,21	15,89	0,21	100,72	1,11	0,20	84,7
MU72-2 5b1	rim a		38,81	0,01	0,04	0,01	42,59	0,33	0,20	17,40	0,19	99,63	1,06	0,22	82,9
MU72-2 5b2	rim b		39,44	0,01	0,02	0,01	43,42	0,32	0,22	17,51	0,26	101,19	1,08	0,22	83,1
MU72-2 5c1	core		38,85	0,00	0,03	0,02	42,15	0,29	0,25	18,63	0,21	100,45	1,05	0,23	81,8
MU72-2 6b1	rim 1a		39,48	0,01	0,07	0,02	44,91	0,27	0,18	14,86	0,33	100,16	1,11	0,19	85,7
MU72-2 6b2	rim 1b		38,65	0,01	0,03	0,08	42,94	0,28	0,21	16,85	0,24	99,28	1,07	0,21	83,5
MU72-2 6c1	core 1		39,50	0,00	0,00	0,05	45,38	0,28	0,22	14,57	0,29	100,35	1,13	0,18	86,1
MU72-2 6c2	~core 2		39,48	0,00	0,05	0,04	45,64	0,25	0,14	14,22	0,31	100,14	1,13	0,18	86,4
MU72-2 7b1	rim 1		38,82	0,00	0,03	0,10	42,66	0,31	0,27	17,85	0,22	100,30	1,06	0,22	82,6
MU72-2 7b2	rim 2		38,95	0,00	0,05	0,10	43,17	0,29	0,19	17,38	0,25	100,38	1,07	0,22	83,1
MU72-2 7c1	core 1		39,39	0,03	0,09	0,10	44,99	0,27	0,25	14,77	0,23	100,18	1,12	0,19	85,8
MU72-2 7c2	core 2		39,42	0,01	0,04	0,04	44,70	0,25	0,23	15,27	0,30	100,25	1,11	0,19	85,3
MU72-2 8b1	rim		38,76	0,03	0,05	0,04	43,09	0,30	0,23	16,91	0,28	99,68	1,07	0,21	83,5
MU72-2 8c1	core		39,29	0,00	0,04	0,07	45,03	0,27	0,28	14,98	0,24	100,22	1,12	0,19	85,6
MU72-2 9b1	rim 1		38,43	0,02	0,05	0,02	42,01	0,33	0,26	18,44	0,27	99,86	1,04	0,23	81,9
MU72-2 9b2	rim 2		38,81	0,02	0,02	0,08	42,64	0,30	0,21	17,91	0,23	100,25	1,06	0,22	82,5
MU72-2 9c1	core 1		39,74	0,00	0,04	0,06	46,24	0,25	0,30	13,54	0,34	100,56	1,15	0,17	87,1
MU72-2 9c2	core 2		39,26	0,01	0,05	0,07	44,88	0,26	0,19	15,26	0,29	100,27	1,11	0,19	85,3
MU72-2 10b1	rim 1a		38,75	0,01	0,04	0,00	43,13	0,34	0,28	16,94	0,21	99,72	1,07	0,21	83,5
MU72-2 10b2	rim 1b		38,95	0,03	0,08	0,13	43,21	0,34	0,23	17,14	0,28	100,40	1,07	0,21	83,3
MU72-2 10c1	core 1		39,46	0,00	0,02	0,04	44,99	0,26	0,25	15,00	0,33	100,35	1,12	0,19	85,6
MU72-2 10c2	core 2		39,05	0,00	0,05	0,09	43,46	0,28	0,23	17,03	0,22	100,46	1,08	0,21	83,5
MU72-2 11b1	rim a		38,75	0,00	0,04	0,05	42,52	0,32	0,18	17,88	0,23	99,98	1,06	0,22	82,5
MU72-2 11b2	rim b		38,73	0,04	0,03	0,05	42,48	0,37	0,22	18,18	0,17	100,27	1,05	0,23	82,2
MU72-2 11c1	core		38,91	0,00	0,04	0,06	43,34	0,26	0,31	17,54	0,25	100,73	1,08	0,22	83,0
MU72-2 12b1	rim 1		38,79	0,00	0,05	0,08	43,17	0,31	0,16	16,95	0,21	99,72	1,07	0,21	83,5
MU72-2 12b2	rim 2		38,77	0,01	0,04	0,10	43,44	0,31	0,29	16,83	0,22	100,02	1,08	0,21	83,6
MU72-2 12c1	core 1		39,43	0,02	0,04	0,10	45,17	0,28	0,32	14,97	0,26	100,60	1,12	0,19	85,7
MU72-2 12c2	core 2		39,01	0,00	0,07	0,17	43,77	0,25	0,24	16,78	0,26	100,56	1,09	0,21	83,8
MU72-2 13b1	rim 1	X	39,19	0,01	0,07	0,01	43,47	0,31	0,20	16,90	0,20	100,40	1,08	0,21	83,6
MU72-2 13c1	core 1		39,81	0,01	0,07	0,08	47,28	0,25	0,16	12,07	0,39	100,14	1,17	0,15	88,6
MU72-2 13c2	core 2		39,40	0,00	0,03	0,00	46,32	0,26	0,22	13,34	0,36	99,98	1,15	0,17	87,3
MU72-2 14b1	rim 1		39,06	0,04	0,05	0,07	43,12	0,32	0,22	17,06	0,23	100,17	1,07	0,21	83,4
MU72-2 14b2	rim 2		38,86	0,01	0,05	0,02	43,37	0,30	0,26	16,45	0,17	99,48	1,08	0,21	83,9
MU72-2 14c1	core 1	X	39,64	0,01	0,04	0,06	45,28	0,25	0,16	14,21	0,35	100,02	1,12	0,18	86,3
MU72-2 14c2	core 2		39,51	0,00	0,13	0,05	45,35	0,26	0,25	14,58	0,26	100,36	1,13	0,18	86,0
MU72-2 15b1	rim		38,65	0,02	0,05	0,09	42,78	0,33	0,23	17,95	0,24	100,36	1,06	0,22	82,5
MU72-2 15c1	core		39,05	0,00	0,03	0,04	43,30	0,30	0,25	17,18	0,22	100,40	1,07	0,22	83,3
MU72-2 16b1	rim 1		38,91	0,01	0,05	0,05	43,13	0,30	0,28	17,23	0,23	100,21	1,07	0,22	83,2

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU72-2 16b2	rim 2		38,94	0,01	0,05	0,14	43,15	0,32	0,22	17,16	0,19	100,19	1,07	0,21	83,3
MU72-2 16c1	core 1		39,03	0,01	0,03	0,04	43,39	0,29	0,22	16,84	0,24	100,12	1,08	0,21	83,6
MU72-2 16c2	core 2		38,90	0,01	0,05	0,04	43,32	0,28	0,15	16,77	0,25	99,81	1,07	0,21	83,7
MU72-2 17b1	rim 1		39,16	0,03	0,06	0,00	43,21	0,31	0,25	16,96	0,25	100,23	1,07	0,21	83,5
MU72-2 17c1	core 1		39,19	0,01	0,04	0,06	44,28	0,24	0,09	15,40	0,26	99,59	1,10	0,19	85,1
MU72-2 17c2	core 2		39,20	0,02	0,11	0,05	43,85	0,26	0,28	16,11	0,24	100,17	1,09	0,20	84,4
MU72-6-1 1b1	rim 1		39,76	0,01	0,06	0,08	45,80	0,24	0,14	13,99	0,34	100,43	1,14	0,18	86,6
MU72-6-1 1c1	core 1		40,13	0,00	0,07	0,14	48,23	0,22	0,17	11,16	0,43	100,57	1,20	0,14	89,5
MU72-6-1 1c2	core 2		39,86	0,01	0,07	0,19	48,30	0,22	0,11	10,89	0,44	100,12	1,20	0,14	89,8
MU72-6-1 2b1	rim 1a		38,85	0,00	0,04	0,11	41,67	0,27	0,24	19,05	0,16	100,44	1,03	0,24	81,2
MU72-6-1 2b2	rim 1b		38,51	0,01	0,07	0,09	41,75	0,32	0,27	18,83	0,21	100,09	1,04	0,24	81,5
MU72-6-1 2c1	core 1		38,54	0,01	0,06	0,06	41,66	0,28	0,27	19,37	0,22	100,48	1,03	0,24	81,0
MU72-6-1 2c2	~core 2		38,51	0,01	0,03	0,06	41,94	0,27	0,32	19,02	0,22	100,40	1,04	0,24	81,4
MU72-6-1 3b1	rim a		39,03	0,02	0,04	0,02	42,84	0,29	0,23	18,17	0,20	100,84	1,06	0,23	82,4
MU72-6-1 3b2	rim b		38,93	0,01	0,03	0,08	43,14	0,29	0,23	17,65	0,24	100,59	1,07	0,22	82,9
MU72-6-1 3c1	core		38,45	0,00	0,01	0,04	41,68	0,24	0,29	19,32	0,20	100,25	1,03	0,24	81,0
MU72-6-1 4b1	rim 1		39,29	0,01	0,07	0,09	43,81	0,25	0,17	16,71	0,30	100,71	1,09	0,21	83,9
MU72-6-1 4b2	rim 2		39,26	0,01	0,03	0,06	44,51	0,23	0,24	15,94	0,30	100,58	1,10	0,20	84,7
MU72-6-1 4c1	core 1		39,45	0,02	0,05	0,08	45,17	0,21	0,21	14,99	0,34	100,55	1,12	0,19	85,7
MU72-6-1 4c2	core 2		38,49	0,01	0,04	0,11	41,56	0,22	0,23	19,58	0,25	100,50	1,03	0,25	80,8
MU72-6-1 5b1	rim a	X	39,78	0,01	0,04	0,05	45,51	0,26	0,21	14,48	0,31	100,65	1,13	0,18	86,2
MU72-6-1 5b2	rim b	X	39,52	0,01	0,06	0,06	45,75	0,24	0,20	13,70	0,35	99,91	1,14	0,17	86,9
MU72-6-1 5c1	core	X	40,41	0,00	0,06	0,12	48,52	0,24	0,15	11,05	0,51	101,10	1,20	0,14	89,7
MU72-6-1 6b1	rim a		38,73	0,00	0,05	0,04	42,60	0,30	0,33	18,02	0,21	100,31	1,06	0,23	82,4
MU72-6-1 6b2	rim b		39,23	0,03	0,04	0,00	43,15	0,32	0,22	17,92	0,28	101,19	1,07	0,22	82,7
MU72-6-1 6c1	core		38,70	0,00	0,05	0,01	41,40	0,27	0,26	19,62	0,20	100,51	1,03	0,25	80,7
MU72-6-1 7b1	rim 1		39,66	0,01	0,09	0,09	45,59	0,26	0,20	14,35	0,28	100,56	1,13	0,18	86,3
MU72-6-1 7b2	rim 2		39,57	0,02	0,05	0,04	45,51	0,28	0,18	14,50	0,25	100,41	1,13	0,18	86,1
MU72-6-1 7c1	core 1		39,92	0,00	0,05	0,06	47,25	0,26	0,08	12,52	0,43	100,56	1,17	0,16	88,2
MU72-6-1 7c2	core 2		39,62	0,00	0,07	0,02	46,50	0,24	0,15	13,40	0,42	100,42	1,15	0,17	87,3
MU72-6-1 8b1	rim a		38,90	0,01	0,05	0,04	42,80	0,29	0,27	17,71	0,22	100,33	1,06	0,22	82,7
MU72-6-1 8b2	rim b		39,16	0,02	0,04	0,13	42,77	0,33	0,29	18,41	0,15	101,32	1,06	0,23	82,2
MU72-6-1 8c1	core		38,89	0,01	0,03	0,06	41,98	0,26	0,13	19,22	0,24	100,84	1,04	0,24	81,2
MU72-6-1 9b1	rim		38,93	0,00	0,05	0,03	42,27	0,27	0,23	18,80	0,22	100,86	1,05	0,24	81,7
MU72-6-1 9b2	rim 2		39,40	0,02	0,09	0,06	43,02	0,29	0,35	17,86	0,28	101,42	1,07	0,22	82,7
MU72-6-1 9c1	core 1		39,15	0,01	0,05	0,07	43,83	0,23	0,19	16,86	0,32	100,73	1,09	0,21	83,7
MU72-6-1 9c2	core 2		38,81	0,01	0,07	0,00	41,51	0,25	0,35	19,32	0,23	100,56	1,03	0,24	81,0
MU72-6-1 9c3	core 3		38,84	0,01	0,05	0,02	42,07	0,26	0,20	19,47	0,24	101,15	1,04	0,24	81,1
MU72-6-2 1b1	rim a		38,30	0,02	0,07	0,01	42,64	0,31	0,26	16,99	0,24	98,84	1,06	0,21	83,3
MU72-6-2 1b2	rim b		38,69	0,01	0,04	0,00	42,66	0,31	0,23	17,50	0,25	99,70	1,06	0,22	82,8

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU72-6-2 1b3	rim c		38,53	0,02	0,07	0,06	42,14	0,32	0,27	17,00	0,25	98,67	1,05	0,21	83,1
MU72-6-2 1c1	core		38,74	0,00	0,05	0,08	41,87	0,26	0,25	18,60	0,29	100,15	1,04	0,23	81,7
MU72-6-2 2b1	rim 1a		39,53	0,03	0,06	0,16	45,62	0,27	0,21	13,57	0,28	99,74	1,13	0,17	86,9
MU72-6-2 2b2	rim 1b		38,83	0,02	0,05	0,09	41,81	0,31	0,27	19,01	0,23	100,65	1,04	0,24	81,3
MU72-6-2 2b3	rim 2		38,95	0,03	0,04	0,05	44,34	0,28	0,22	15,17	0,26	99,39	1,10	0,19	85,3
MU72-6-2 2c1	core 1		40,08	0,00	0,05	0,05	47,69	0,23	0,16	11,50	0,39	100,14	1,18	0,14	89,1
MU72-6-2 2c2	core 2		39,87	0,01	0,04	0,08	47,48	0,23	0,24	11,71	0,36	100,04	1,18	0,15	88,9
MU72-6-2 3b1	rim 1a		38,15	0,01	0,04	0,00	41,76	0,30	0,26	18,36	0,21	99,09	1,04	0,23	81,8
MU72-6-2 3b2	rim 1b		38,61	0,01	0,04	0,04	42,08	0,29	0,12	18,04	0,27	99,54	1,04	0,23	82,2
MU72-6-2 3c1	core 1		38,28	0,02	0,07	0,01	41,77	0,24	0,26	18,71	0,22	99,63	1,04	0,23	81,6
MU72-6-2 3c2	core 2		38,54	0,00	0,03	0,00	41,50	0,26	0,24	19,13	0,21	99,92	1,03	0,24	81,1
MU72-6-2 4b1	rim a		38,63	0,02	0,04	0,02	42,29	0,29	0,27	18,14	0,21	99,95	1,05	0,23	82,2
MU72-6-2 4b2	rim b		38,60	0,03	0,09	0,06	42,55	0,30	0,23	17,81	0,23	99,95	1,06	0,22	82,6
MU72-6-2 4c1	core		39,41	0,01	0,04	0,05	45,77	0,20	0,22	14,37	0,43	100,52	1,14	0,18	86,3
MU72-6-2 5b1	rim 1a		38,50	0,01	0,05	0,06	41,68	0,27	0,18	18,95	0,22	99,96	1,03	0,24	81,3
MU72-6-2 5b2	rim 1b		38,46	0,01	0,13	0,04	42,09	0,30	0,19	18,32	0,23	99,79	1,04	0,23	82,0
MU72-6-2 5c1	core 1a		39,38	0,00	0,06	0,09	45,59	0,28	0,17	14,11	0,35	100,05	1,13	0,18	86,5
MU72-6-2 5c2	core 1b		39,54	0,00	0,05	0,05	45,42	0,27	0,14	14,27	0,29	100,04	1,13	0,18	86,3
MU72-6-2 5c3	core 2		38,50	0,02	0,10	0,06	41,28	0,25	0,25	19,65	0,27	100,38	1,02	0,25	80,6
MU72-6-2 6b1	rim a	X	39,27	0,01	0,05	0,01	45,39	0,24	0,24	13,84	0,36	99,41	1,13	0,17	86,7
MU72-6-2 6b2	rim b	X	39,71	0,00	0,09	0,07	46,05	0,25	0,23	14,30	0,26	100,99	1,14	0,18	86,4
MU72-6-2 6c1	core	X	40,05	0,01	0,08	0,06	48,07	0,24	0,24	10,90	0,39	100,06	1,19	0,14	89,7
MU72-6-2 7b1	rim 1		38,60	0,03	0,06	0,00	42,51	0,31	0,30	17,59	0,18	99,57	1,05	0,22	82,7
MU72-6-2 7b2	rim 2		38,68	0,02	0,05	0,06	42,87	0,29	0,26	17,59	0,21	100,06	1,06	0,22	82,8
MU72-6-2 7c1	core 1		38,53	0,01	0,08	0,05	41,68	0,27	0,24	19,07	0,30	100,28	1,03	0,24	81,2
MU72-6-2 7c2	core 2		38,73	0,00	0,04	0,00	42,13	0,29	0,30	18,39	0,21	100,10	1,05	0,23	81,9
MU72-6-2 7c3	core 3		38,39	0,01	0,07	0,03	41,35	0,29	0,18	19,18	0,25	99,78	1,03	0,24	81,0
MU72-6-2 7c4	core 4		38,54	0,00	0,06	0,09	41,57	0,28	0,14	18,98	0,23	99,96	1,03	0,24	81,3
MU72-6-2 7c5	core 5		38,60	0,00	0,06	0,00	42,00	0,28	0,25	18,80	0,19	100,17	1,04	0,24	81,6
MU72-6-2 8b1	rim 1a		38,28	0,02	0,10	0,08	42,51	0,29	0,26	17,53	0,29	99,40	1,05	0,22	82,8
MU72-6-2 8b2	rim 1b		38,92	0,02	0,05	0,05	42,94	0,31	0,27	18,07	0,27	100,92	1,07	0,23	82,5
MU72-6-2 8c1	core 1		38,27	0,03	0,06	0,00	41,42	0,29	0,27	19,45	0,16	99,97	1,03	0,24	80,8
MU72-6-2 8c2	core 2		38,65	0,00	0,06	0,00	41,72	0,28	0,25	19,20	0,21	100,37	1,04	0,24	81,2
MU72-6-2 9b1	rim 1		38,70	0,04	0,04	0,04	42,78	0,31	0,41	17,75	0,23	100,32	1,06	0,22	82,7
MU72-6-2 9b2	rim 3		38,80	0,04	0,04	0,05	43,10	0,31	0,22	17,39	0,26	100,23	1,07	0,22	83,1
MU72-6-2 9c1	core 1		39,10	0,02	0,06	0,08	44,02	0,28	0,22	15,82	0,25	99,86	1,09	0,20	84,6
MU72-6-2 9c2	core 2		39,20	0,02	0,06	0,06	44,89	0,26	0,21	15,14	0,26	100,10	1,11	0,19	85,4
MU72-6-2 9c3	core 3		38,53	0,01	0,04	0,00	41,58	0,28	0,22	19,18	0,25	100,09	1,03	0,24	81,1
MU72-6-2 10b1	rim a		39,04	0,03	0,06	0,00	43,39	0,30	0,21	17,04	0,20	100,26	1,08	0,21	83,5
MU72-6-2 10b2	rim b		38,91	0,00	0,04	0,07	43,48	0,27	0,23	16,51	0,24	99,76	1,08	0,21	83,9

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU72-6-2 10b3	rim c		38,83	0,02	0,05	0,05	43,26	0,30	0,25	17,24	0,22	100,21	1,07	0,22	83,3
MU72-6-2 10c1	core		39,50	0,00	0,05	0,04	45,67	0,26	0,13	13,56	0,32	99,53	1,13	0,17	87,0
MU72-6-2 11b1	rim 1		38,88	0,01	0,04	0,07	43,28	0,27	0,23	16,02	0,28	99,08	1,07	0,20	84,3
MU72-6-2 11b2	rim 2		38,98	0,01	0,05	0,05	43,73	0,27	0,27	15,72	0,28	99,39	1,09	0,20	84,6
MU72-6-2 11b3	~rim 3		38,97	0,01	0,04	0,08	43,78	0,28	0,17	16,55	0,27	100,17	1,09	0,21	84,0
MU72-6-2 11c1	core 1		39,14	0,01	0,07	0,03	44,71	0,26	0,27	14,90	0,30	99,67	1,11	0,19	85,6
MU72-6-2 11c2	core 2		39,78	0,02	0,06	0,00	45,65	0,28	0,21	14,16	0,30	100,50	1,13	0,18	86,5
MU72-6-2 11c3	~core 3		39,53	0,00	0,05	0,04	45,62	0,27	0,18	14,04	0,24	100,00	1,13	0,18	86,6
MU72-6-2 12b1	rim a		38,95	0,00	0,03	0,09	43,68	0,28	0,21	17,16	0,28	100,70	1,08	0,21	83,4
MU72-6-2 12b2	rim b		39,05	0,02	0,06	0,00	43,50	0,28	0,12	16,85	0,34	100,25	1,08	0,21	83,6
MU72-6-2 12c1	core a		40,08	0,00	0,05	0,16	46,60	0,26	0,18	13,02	0,38	100,74	1,16	0,16	87,6
MU72-6-2 12c2	core b		39,65	0,00	0,06	0,06	45,86	0,26	0,18	13,48	0,39	99,98	1,14	0,17	87,1
MU72-6-2 13b1	rim 1	X	38,48	0,02	0,04	0,03	42,00	0,30	0,26	18,77	0,26	100,16	1,04	0,24	81,6
MU72-6-2 13b2	rim 2a	X	38,39	0,00	0,05	0,00	41,16	0,29	0,11	18,95	0,20	99,15	1,02	0,24	81,1
MU72-6-2 13b3	rim 2b	X	38,83	0,02	0,06	0,00	42,54	0,29	0,30	17,91	0,24	100,21	1,06	0,22	82,5
MU72-6-2 13c1	core 1	X	38,35	0,01	0,04	0,07	41,18	0,29	0,29	19,23	0,22	99,71	1,02	0,24	80,9
MU72-6-2 13c2	core 2	X	38,51	0,03	0,01	0,03	41,06	0,28	0,25	19,09	0,20	99,48	1,02	0,24	81,0
MU72-7 1b1	rim a		38,48	0,04	0,04	0,08	42,18	0,32	0,14	19,34	0,13	100,77	1,05	0,24	81,2
MU72-7 1b2	rim b		38,75	0,02	0,06	0,00	42,82	0,28	0,25	18,28	0,22	100,67	1,06	0,23	82,3
MU72-7 1b3	rim c		38,70	0,04	0,01	0,00	42,08	0,34	0,21	18,51	0,21	100,14	1,04	0,23	81,8
MU72-7 1c1	core a		39,42	0,02	0,08	0,07	45,05	0,31	0,23	15,83	0,25	101,26	1,12	0,20	84,9
MU72-7 1c2	core b		39,33	0,02	0,04	0,10	44,48	0,28	0,24	15,42	0,26	100,17	1,10	0,19	85,1
MU72-7 2b1	rim a		38,59	0,03	0,04	0,00	41,30	0,31	0,22	19,06	0,25	99,82	1,02	0,24	81,1
MU72-7 2b2	rim b		38,27	0,06	0,04	0,06	40,82	0,35	0,21	19,71	0,21	99,74	1,01	0,25	80,4
MU72-7 2b3	rim c		38,84	0,03	0,05	0,00	42,49	0,37	0,27	19,61	0,19	101,89	1,05	0,25	81,1
MU72-7 2c1	core		39,44	0,01	0,01	0,09	45,49	0,25	0,26	14,34	0,40	100,32	1,13	0,18	86,3
MU72-7 3b1	rim		39,03	0,02	0,08	0,00	43,28	0,30	0,22	16,89	0,20	100,03	1,07	0,21	83,5
MU72-7 4b1	rim 1		39,15	0,04	0,03	0,04	43,87	0,34	0,23	17,56	0,18	101,48	1,09	0,22	83,2
MU72-7 4b2	rim 2		37,72	0,07	0,00	0,03	39,31	0,30	0,22	21,48	0,15	99,34	0,98	0,27	78,4
MU72-7 4c1	core 1		39,48	0,01	0,00	0,00	45,53	0,24	0,12	14,25	0,28	99,91	1,13	0,18	86,4
MU72-7 4c2	core 2		39,25	0,00	0,00	0,02	44,47	0,28	0,19	15,58	0,22	100,04	1,10	0,20	85,0
MU72-7 4c3	core 3		39,12	0,01	0,05	0,05	43,91	0,29	0,27	16,90	0,27	100,88	1,09	0,21	83,7
MU72-7 5b1	rim 1		38,48	0,03	0,05	0,11	41,70	0,31	0,21	18,84	0,22	99,97	1,03	0,24	81,4
MU72-7 5b3	rim 3		38,91	0,01	0,06	0,00	44,00	0,31	0,23	17,81	0,26	101,62	1,09	0,22	83,0
MU72-7 5c1	core 1		38,99	0,02	0,04	0,07	43,06	0,29	0,25	16,72	0,20	99,64	1,07	0,21	83,6
MU72-7 5c2	core 2		38,90	0,03	0,07	0,01	42,78	0,30	0,24	17,53	0,20	100,08	1,06	0,22	82,9
MU72-7 5c3	core 3		38,81	0,01	0,03	0,04	43,58	0,30	0,28	17,23	0,26	100,54	1,08	0,22	83,4
MU72-7 6b1	rim 1a		38,51	0,00	0,49	0,11	43,71	0,28	0,28	16,81	0,22	100,44	1,08	0,21	83,7
MU72-7 6b2	rim 1b		38,88	0,01	0,04	0,08	43,03	0,29	0,28	17,01	0,24	99,90	1,07	0,21	83,4

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fe%
MU72-7 6b4	rim 4		38,36	0,02	0,05	0,04	41,61	0,33	0,25	19,35	0,20	100,22	1,03	0,24	81,0
MU72-7 6c1	core 1a		39,33	0,00	0,01	0,01	45,37	0,27	0,25	14,86	0,31	100,40	1,13	0,19	85,8
MU72-7 6c2	core 1b		39,41	0,00	0,04	0,05	45,39	0,26	0,14	14,61	0,29	100,21	1,13	0,18	86,0
MU72-7 6c3	core 2		39,51	0,00	0,00	0,06	45,48	0,27	0,16	14,42	0,26	100,17	1,13	0,18	86,2
MU72-7 6c4	core 3	X	39,49	0,01	0,04	0,00	45,21	0,27	0,30	14,59	0,25	100,17	1,12	0,18	86,0
MU72-7 7b1	rim 1a	X	38,42	0,01	0,06	0,06	44,26	0,34	0,17	17,85	0,16	101,37	1,10	0,22	83,1
MU72-7 7b2	rim 1b	X	38,68	0,01	0,04	0,07	42,12	0,31	0,25	18,54	0,17	100,21	1,05	0,23	81,8
MU72-7 7c1	core 1	X	38,99	0,01	0,00	0,00	44,17	0,28	0,20	16,89	0,26	100,81	1,10	0,21	83,8
MU72-7 7c2	core 2		38,69	0,00	0,09	0,04	42,94	0,30	0,29	17,18	0,20	99,75	1,07	0,22	83,2
MU72-7 8b1	rim 1		39,05	0,01	0,04	0,04	43,31	0,27	0,19	16,77	0,21	99,88	1,07	0,21	83,6
MU72-7 8b2	rim 2		38,93	0,04	0,05	0,05	42,44	0,33	0,11	18,19	0,18	100,31	1,05	0,23	82,2
MU72-7 8c1	core 1		39,47	0,00	0,06	0,03	45,18	0,28	0,16	15,81	0,20	101,19	1,12	0,20	85,0
MU72-7 8c2	core 2		38,87	0,00	0,00	0,06	44,36	0,29	0,22	15,62	0,27	99,71	1,10	0,20	84,9
MU72-7 9b1	rim a	X	37,84	0,02	0,04	0,00	38,37	0,26	0,31	24,29	0,17	101,35	0,95	0,30	75,8
MU72-7 9b2	rim b	X	38,22	0,03	0,04	0,00	42,85	0,30	0,30	19,95	0,22	101,92	1,06	0,25	81,0
MU72-7 9b4	rim d	X	37,84	0,05	0,03	0,02	39,23	0,33	0,26	22,68	0,20	100,66	0,97	0,28	77,4
MU72-7 9c1	core	X	39,21	0,01	0,05	0,05	44,28	0,26	0,24	15,79	0,25	100,16	1,10	0,20	84,7
MU72-7 10b2	rim 1b	X	38,63	0,06	0,06	0,09	43,41	0,32	0,29	18,62	0,17	101,66	1,08	0,23	82,2
MU72-7 10b3	rim 2	X	38,86	0,02	0,04	0,06	42,37	0,32	0,27	18,17	0,19	100,34	1,05	0,23	82,2
MU72-7 10c1	core 1	X	39,18	0,00	0,04	0,07	45,36	0,26	0,20	14,77	0,36	100,23	1,13	0,18	85,9
MU72-7 10c2	core 2a	X	39,04	0,02	0,05	0,04	43,63	0,27	0,24	16,26	0,32	99,87	1,08	0,20	84,2
MU72-7 10c3	core 2b	X	39,25	0,02	0,04	0,05	44,94	0,25	0,19	15,60	0,31	100,70	1,12	0,20	85,1
MU72-7 11b1	rim 1a	X	38,20	0,01	0,04	0,02	41,06	0,33	0,23	18,86	0,19	98,93	1,02	0,24	81,2
MU72-7 11b2	rim 1b	X	38,87	0,03	0,05	0,07	43,45	0,31	0,28	16,97	0,22	100,24	1,08	0,21	83,5
MU72-7 11b3	rim 2		38,39	0,06	0,09	0,01	40,93	0,32	0,18	20,08	0,19	100,28	1,02	0,25	80,1
MU72-7 11c1	core 1	X	39,33	0,00	0,03	0,06	44,78	0,27	0,17	15,03	0,29	99,99	1,11	0,19	85,5
MU72-7 11c2	core 2		39,55	0,02	0,01	0,10	44,96	0,27	0,25	15,07	0,22	100,52	1,12	0,19	85,5
MU72-7 12b1	rim 1a		38,74	0,03	0,03	0,05	42,66	0,32	0,37	17,53	0,21	99,95	1,06	0,22	82,8
MU72-7 12b2	rim 1b		38,77	0,02	0,03	0,07	42,43	0,32	0,25	17,40	0,21	99,48	1,05	0,22	82,8
MU72-7 12c1	core 1		38,82	0,01	0,06	0,07	43,34	0,29	0,33	16,48	0,22	99,67	1,08	0,21	83,9
MU72-7 12c2	core 2		38,97	0,00	0,00	0,04	43,62	0,27	0,21	16,74	0,22	100,10	1,08	0,21	83,8
MU72-7 13b1	rim 1		39,07	0,01	0,06	0,06	42,98	0,30	0,26	17,29	0,23	100,26	1,07	0,22	83,1
MU72-7 13b3	rim 3		38,25	0,03	0,04	0,01	41,68	0,32	0,34	18,97	0,20	99,85	1,03	0,24	81,3
MU72-7 13c1	core 1		39,26	0,01	0,00	0,06	44,30	0,28	0,24	16,21	0,29	100,66	1,10	0,20	84,4
MU72-7 13c2	core 2		38,98	0,02	0,06	0,07	43,54	0,29	0,26	17,12	0,17	100,51	1,08	0,21	83,4
MU72-7 13c3	core 3		39,22	0,00	0,01	0,05	44,81	0,29	0,17	15,10	0,23	99,89	1,11	0,19	85,5
MU72-7 13c4	core 4		39,23	0,00	0,00	0,02	44,61	0,28	0,20	15,07	0,29	99,71	1,11	0,19	85,4
MU72-7 14b1	rim a	X	39,19	0,03	0,06	0,02	44,53	0,28	0,17	15,40	0,31	100,00	1,10	0,19	85,1
MU72-7 14b2	rim b	X	39,23	0,01	0,06	0,11	45,66	0,26	0,21	13,95	0,37	99,86	1,13	0,17	86,6
MU72-7 14c1	core a	X	39,56	0,01	0,08	0,08	47,55	0,25	0,14	11,35	0,38	99,39	1,18	0,14	89,2
MU72-7 14c2	core b	X	39,72	0,02	0,03	0,09	47,27	0,24	0,11	11,85	0,32	99,65	1,17	0,15	88,8

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU73-1a 1b1	rim 1	X	38,75	0,00	0,05	0,07	42,87	0,28	0,16	17,28	0,23	99,73	1,06	0,22	83,1
MU73-1a 1c1	core 1	X	39,31	0,01	0,05	0,05	44,68	0,28	0,24	15,75	0,24	100,64	1,11	0,20	84,9
MU73-1a 1c2	core 2	X	39,21	0,01	0,09	0,07	44,87	0,28	0,23	15,12	0,29	100,19	1,11	0,19	85,5
MU73-1a 2b1	rim 1		38,89	0,00	0,05	0,04	43,25	0,31	0,28	17,16	0,24	100,22	1,07	0,21	83,3
MU73-1a 2c1	core 1		38,49	0,00	0,06	0,07	43,54	0,28	0,25	17,33	0,26	100,28	1,08	0,22	83,3
MU73-1a 2c2	~core 2		39,07	0,02	0,04	0,00	43,49	0,30	0,24	17,04	0,17	100,36	1,08	0,21	83,5
MU73-1a 3b1	rim 2		38,50	0,01	0,04	0,03	42,27	0,32	0,23	18,21	0,27	99,88	1,05	0,23	82,1
MU73-1a 3c1	core 1	X	38,95	0,03	0,05	0,00	43,81	0,17	0,13	16,63	0,29	100,10	1,09	0,21	83,9
MU73-1a 3c2	core 2		38,84	0,01	0,05	0,10	43,39	0,29	0,29	17,12	0,25	100,34	1,08	0,21	83,4
MU73-1a 4b1	rim	X	39,05	0,00	0,06	0,09	44,22	0,27	0,28	15,17	0,27	99,46	1,10	0,19	85,2
MU73-1a 4c1	core	X	39,52	0,00	0,05	0,03	45,98	0,24	0,11	13,50	0,36	99,78	1,14	0,17	87,1
MU73-1a 5b1	rim	X	38,69	0,00	0,04	0,08	42,51	0,32	0,34	17,71	0,23	99,96	1,05	0,22	82,6
MU73-1a 5c1	core	X	39,00	0,01	0,00	0,03	43,63	0,28	0,22	16,21	0,24	99,66	1,08	0,20	84,2
MU73-1a 6b1	rim 3		38,70	0,01	0,05	0,09	43,34	0,27	0,30	16,32	0,23	99,31	1,08	0,20	84,0
MU73-1a 6b2	rim 1		38,71	0,02	0,05	0,07	42,56	0,28	0,26	17,23	0,17	99,35	1,06	0,22	83,0
MU73-1a 6c1	core 1		39,87	0,00	0,05	0,08	46,85	0,24	0,21	12,44	0,43	100,19	1,16	0,16	88,2
MU73-1a 6c2	core 2		39,87	0,02	0,05	0,15	47,30	0,23	0,26	11,69	0,39	99,98	1,17	0,15	88,9
MU73-1a 6c3	core 3		39,79	0,00	0,06	0,04	46,79	0,23	0,21	12,54	0,35	100,01	1,16	0,16	88,1
MU73-1a 6c4	core 4		39,30	0,00	0,05	0,06	45,84	0,24	0,17	13,48	0,34	99,50	1,14	0,17	87,1
MU73-1a 7b1	rim 2		38,65	0,01	0,06	0,08	42,49	0,28	0,22	17,51	0,22	99,53	1,05	0,22	82,8
MU73-1a 7b2	rim 1		38,43	0,02	0,03	0,12	41,46	0,32	0,22	19,39	0,17	100,17	1,03	0,24	80,9
MU73-1a 7c1	core 1		39,02	0,00	0,06	0,07	43,54	0,27	0,27	16,94	0,26	100,45	1,08	0,21	83,6
MU73-1a 7c2	core 2		38,78	0,01	0,11	0,02	45,09	0,26	0,22	14,65	0,28	99,45	1,12	0,18	85,9
MU73-1a 8b1	rim a	X	38,74	0,02	0,10	0,00	42,68	0,32	0,27	18,00	0,20	100,36	1,06	0,23	82,4
MU73-1a 8b2	rim b	X	39,00	0,00	0,04	0,03	43,11	0,32	0,24	17,79	0,17	100,77	1,07	0,22	82,8
MU73-1a 8c1	core	X	39,63	0,01	0,04	0,11	45,40	0,26	0,15	14,55	0,26	100,45	1,13	0,18	86,1
MU73-1a 9b1	rim 2		38,87	0,02	0,04	0,04	42,32	0,31	0,28	18,02	0,20	100,12	1,05	0,23	82,3
MU73-1a 9b2	rim 3		39,15	0,02	0,05	0,08	42,73	0,31	0,13	17,77	0,21	100,46	1,06	0,22	82,6
MU73-1a 9c1	core 1		38,94	0,02	0,04	0,11	42,46	0,31	0,22	18,09	0,21	100,38	1,05	0,23	82,3
MU73-1a 9c2	core 2		39,07	0,00	0,05	0,09	42,83	0,27	0,21	17,56	0,23	100,31	1,06	0,22	82,9
MU73-1a 9c3	core 3		38,92	0,01	0,04	0,12	42,99	0,28	0,23	17,21	0,23	100,02	1,07	0,22	83,2
MU73-1a 10b1	rim 1	X	38,89	0,03	0,04	0,01	42,88	0,32	0,25	17,39	0,26	100,09	1,06	0,22	83,0
MU73-1a 10b2	rim 2		38,89	0,02	0,08	0,06	42,25	0,32	0,20	17,81	0,27	99,91	1,05	0,22	82,5
MU73-1a 10c1	core 1	X	39,33	0,01	0,06	0,01	43,97	0,29	0,24	16,44	0,24	100,62	1,09	0,21	84,1
MU73-1a 10c2	core 2		39,01	0,00	0,04	0,00	43,17	0,29	0,21	17,22	0,22	100,16	1,07	0,22	83,2
MU73-1a 11b1	rim 1		38,95	0,00	0,03	0,03	42,94	0,31	0,33	17,64	0,21	100,47	1,07	0,22	82,8
MU73-1a 11c1	core 1		39,18	0,01	0,07	0,05	43,32	0,27	0,27	17,05	0,25	100,47	1,07	0,21	83,4
MU73-1a 11c2	core 2		39,03	0,01	0,05	0,10	42,92	0,33	0,19	17,40	0,24	100,29	1,06	0,22	83,0
MU73-1a 12b1	rim 1		38,93	0,00	0,05	0,11	43,41	0,28	0,27	17,03	0,23	100,31	1,08	0,21	83,5
MU73-1a 12b2	rim 2		39,00	0,02	0,04	0,06	42,89	0,31	0,17	17,68	0,19	100,36	1,06	0,22	82,8

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU73-1a 12c1	core 1		39,21	0,01	0,05	0,00	44,38	0,29	0,31	15,61	0,22	100,08	1,10	0,20	84,9
MU73-1a 12c2	core 2		39,17	0,01	0,04	0,02	44,40	0,27	0,16	16,06	0,21	100,36	1,10	0,20	84,6
MU73-1a 13b1	rim 1		38,97	0,02	0,06	0,06	43,59	0,30	0,26	17,25	0,20	100,71	1,08	0,22	83,4
MU73-1a 13b2	rim 2		38,84	0,00	0,07	0,08	43,50	0,28	0,19	16,57	0,22	99,79	1,08	0,21	83,9
MU73-1a 13c1	core 1		39,24	0,02	0,06	0,04	44,11	0,26	0,27	16,34	0,29	100,62	1,09	0,20	84,2
MU73-1a 13c2	core 2		39,05	0,02	0,07	0,11	43,86	0,26	0,30	16,59	0,24	100,49	1,09	0,21	84,0
MU73-1a 14b1	rim	X	38,78	0,02	0,04	0,06	42,36	0,34	0,26	18,20	0,17	100,22	1,05	0,23	82,2
MU73-1a 14c1	core	X	39,06	0,01	0,07	0,03	43,81	0,25	0,23	16,68	0,25	100,42	1,09	0,21	83,9
MU73-1a 15b1	rim		38,75	0,01	0,05	0,00	42,43	0,33	0,26	18,00	0,22	100,03	1,05	0,23	82,4
MU73-1a 15c1	core		39,31	0,02	0,04	0,06	44,68	0,31	0,17	15,66	0,27	100,51	1,11	0,20	85,0
MU73-1a 16b1	rim 1		39,31	0,01	0,04	0,00	43,66	0,29	0,24	17,01	0,20	100,79	1,08	0,21	83,6
MU73-1a 16c1	core 1		39,34	0,01	0,02	0,03	44,46	0,27	0,19	15,93	0,26	100,49	1,10	0,20	84,7
MU73-1a 16c2	core 2		39,29	0,00	0,07	0,08	43,84	0,27	0,15	16,59	0,25	100,59	1,09	0,21	84,0
MU73-1a 17b1	rim 1		39,09	0,00	0,03	0,03	43,91	0,26	0,24	16,41	0,29	100,26	1,09	0,21	84,1
MU73-1a 17c1	core 1		40,16	0,00	0,07	0,06	47,58	0,25	0,09	11,31	0,43	99,97	1,18	0,14	89,3
MU73-1a 17c2	core 2		40,03	0,00	0,07	0,13	46,27	0,22	0,19	13,30	0,34	100,54	1,15	0,17	87,3
MU73-1b 1b1	rim 1		38,67	0,02	0,06	0,10	42,89	0,30	0,30	17,60	0,25	100,17	1,06	0,22	82,8
MU73-1b 1b2	rim 2		38,96	0,04	0,05	0,03	42,53	0,32	0,26	17,60	0,22	100,02	1,06	0,22	82,7
MU73-1b 1c1	core 1		39,16	0,03	0,05	0,02	44,24	0,24	0,24	16,05	0,27	100,29	1,10	0,20	84,5
MU73-1b 1c2	core 2		39,51	0,03	0,05	0,04	44,72	0,27	0,25	15,70	0,26	100,85	1,11	0,20	84,9
MU73-1b 2b1	rim		38,61	0,01	0,05	0,06	42,48	0,30	0,33	17,93	0,20	100,00	1,05	0,22	82,4
MU73-1b 2c1	core		38,99	0,02	0,03	0,06	43,96	0,29	0,24	16,28	0,20	100,07	1,09	0,20	84,2
MU73-1b 3b1	rim		38,94	0,03	0,05	0,08	42,68	0,32	0,19	17,71	0,18	100,20	1,06	0,22	82,7
MU73-1b 3c1	core		39,32	0,02	0,05	0,05	44,57	0,24	0,14	15,45	0,29	100,12	1,11	0,19	85,1
MU73-1b 4b1	rim 1		38,70	0,03	0,04	0,08	41,81	0,33	0,29	18,73	0,20	100,21	1,04	0,23	81,6
MU73-1b 4c1	core 1		39,00	0,01	0,04	0,07	42,63	0,31	0,13	17,44	0,24	99,87	1,06	0,22	82,9
MU73-1b 4c3	~core 3		38,72	0,00	0,04	0,05	42,43	0,30	0,26	17,89	0,16	99,85	1,05	0,22	82,5
MU73-1b 5b1	rim 1		38,80	0,01	0,03	0,06	42,76	0,32	0,27	17,76	0,25	100,28	1,06	0,22	82,7
MU73-1b 5b2	rim 2		38,53	0,06	0,06	0,00	41,77	0,36	0,20	18,65	0,20	99,82	1,04	0,23	81,6
MU73-1b 5c1	core 1		40,35	0,00	0,04	0,00	48,32	0,26	0,22	10,82	0,31	100,31	1,20	0,14	89,8
MU73-1b 5c2	~core 2		38,87	0,01	0,05	0,06	42,85	0,31	0,25	17,17	0,24	99,79	1,06	0,22	83,2
MU73-1b 6b1	rim 1		38,50	0,02	0,05	0,05	42,17	0,32	0,28	18,66	0,18	100,27	1,05	0,23	81,7
MU73-1b 6c1	core 1		38,54	0,00	0,05	0,03	42,27	0,29	0,17	18,19	0,20	99,78	1,05	0,23	82,2
MU73-1b 6c2	core 2		38,78	0,00	0,05	0,03	43,13	0,30	0,25	17,42	0,23	100,24	1,07	0,22	83,1
MU73-1b 6c3	core 3		39,56	0,01	0,03	0,05	45,23	0,25	0,25	14,84	0,31	100,58	1,12	0,19	85,8
MU73-1b 7b1	rim 2		38,57	0,05	0,07	0,08	41,88	0,35	0,31	18,83	0,20	100,37	1,04	0,24	81,5
MU73-1b 7c1	core 1		38,82	0,02	0,59	0,12	42,73	0,32	0,24	17,11	0,26	100,19	1,06	0,21	83,2
MU73-1b 7c2	core 2		38,94	0,02	0,05	0,06	43,00	0,29	0,19	17,76	0,23	100,55	1,07	0,22	82,7
MU73-1b 8b1	rim 2		38,57	0,03	0,03	0,04	42,74	0,31	0,26	17,57	0,26	99,81	1,06	0,22	82,8

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU73-1b 8b2	rim 1		39,10	0,00	0,06	0,07	43,72	0,27	0,23	16,75	0,23	100,43	1,08	0,21	83,8
MU73-1b 8c1	core 1		39,14	0,01	0,03	0,10	43,31	0,27	0,22	16,97	0,25	100,34	1,07	0,21	83,5
MU73-1b 8c2	core 2		38,96	0,01	0,03	0,10	43,54	0,28	0,23	16,64	0,23	100,03	1,08	0,21	83,8
MU73-1b 9b1	rim a		38,52	0,03	0,05	0,07	41,60	0,34	0,25	19,13	0,19	100,19	1,03	0,24	81,2
MU73-1b 9b2	rim b		38,65	0,02	0,10	0,12	41,72	0,32	0,23	18,60	0,21	99,99	1,04	0,23	81,6
MU73-1b 9c1	core		39,12	0,01	0,04	0,05	42,95	0,31	0,23	17,38	0,28	100,39	1,07	0,22	83,0
MU73-1b 10b1	rim 1	X	39,01	0,02	0,01	0,04	43,21	0,29	0,27	17,37	0,26	100,48	1,07	0,22	83,1
MU73-1b 10c1	core 1	X	39,16	0,01	0,01	0,09	44,28	0,28	0,11	16,04	0,20	100,19	1,10	0,20	84,5
MU73-1b 10c2	core 2		39,40	0,00	0,07	0,13	44,85	0,30	0,18	15,02	0,23	100,15	1,11	0,19	85,5
MU73-1b 10c3	core 3		38,92	0,00	0,04	0,03	43,17	0,31	0,23	17,42	0,22	100,36	1,07	0,22	83,1
MU73-1b 10c4	core 4		39,09	0,00	0,04	0,00	44,24	0,27	0,16	15,73	0,27	99,83	1,10	0,20	84,8
MU73-1b 11b1	rim 1		38,52	0,02	0,06	0,03	42,01	0,33	0,16	18,33	0,22	99,70	1,04	0,23	81,9
MU73-1b 11b2	rim 2		38,50	0,02	0,06	0,06	41,40	0,35	0,35	18,96	0,21	99,91	1,03	0,24	81,2
MU73-1b 11c1	core 1		39,01	0,00	0,04	0,06	42,67	0,29	0,22	17,81	0,25	100,40	1,06	0,22	82,6
MU73-1b 11c2	core 2		39,45	0,00	0,03	0,13	44,34	0,28	0,24	16,05	0,27	100,82	1,10	0,20	84,5
MU73-1b 11c3	core 3		39,04	0,02	0,06	0,04	43,17	0,30	0,26	17,16	0,20	100,25	1,07	0,22	83,3
MU73-1b 12b1	rim		38,79	0,02	0,03	0,00	42,38	0,31	0,35	17,90	0,22	100,02	1,05	0,22	82,4
MU73-1b 12c1	core		39,16	0,01	0,02	0,04	44,49	0,33	0,20	15,42	0,30	99,98	1,10	0,19	85,1
MU73-1c 1b1	rim a		39,02	0,02	0,04	0,08	43,00	0,30	0,23	18,01	0,25	100,96	1,07	0,23	82,5
MU73-1c 1b2	rim b		38,79	0,02	0,18	0,05	42,82	0,29	0,19	17,54	0,22	100,12	1,06	0,22	82,9
MU73-1c 1c1	core		39,27	0,00	0,03	0,00	44,18	0,28	0,27	16,37	0,25	100,68	1,10	0,21	84,2
MU73-1c 2b1	rim		38,84	0,00	0,06	0,02	42,87	0,29	0,23	17,65	0,25	100,24	1,06	0,22	82,8
MU73-1c 2c1	core		39,16	0,01	0,09	0,05	43,57	0,27	0,15	17,33	0,20	100,84	1,08	0,22	83,3
MU73-1c 3b1	rim		38,81	0,00	0,04	0,03	43,55	0,30	0,22	17,17	0,22	100,38	1,08	0,22	83,4
MU73-1c 3c1	core		39,18	0,00	0,02	0,07	44,56	0,26	0,20	15,58	0,24	100,12	1,11	0,20	85,0
MU73-1c 4b1	rim 1	X	38,96	0,03	0,06	0,11	43,00	0,30	0,22	17,70	0,22	100,58	1,07	0,22	82,8
MU73-1c 4c1	core 1	X	38,94	0,00	0,03	0,00	43,35	0,29	0,23	17,08	0,21	100,17	1,08	0,21	83,4
MU73-1c 4c2	core 2	X	39,14	0,02	0,06	0,07	43,29	0,30	0,23	17,49	0,20	100,81	1,07	0,22	83,1
MU73-1c 5b1	rim 1		38,58	0,02	0,06	0,00	42,49	0,31	0,28	18,04	0,22	100,04	1,05	0,23	82,3
MU73-1c 5b2	rim 2		38,94	0,01	0,05	0,08	42,87	0,31	0,28	18,16	0,25	100,98	1,06	0,23	82,4
MU73-1c 5c1	core 1		39,52	0,00	0,08	0,14	45,88	0,25	0,24	13,75	0,31	100,17	1,14	0,17	86,9
MU73-1c 5c2	core 2		39,22	0,00	0,04	0,06	44,21	0,26	0,22	16,36	0,28	100,66	1,10	0,20	84,3
MU73-1c 6b1	rim 1a		39,30	0,00	0,04	0,07	44,81	0,27	0,28	15,27	0,33	100,39	1,11	0,19	85,3
MU73-1c 6b2	rim 1b		39,40	0,00	0,06	0,12	45,18	0,26	0,25	15,25	0,32	100,86	1,12	0,19	85,4
MU73-1c 6b3	rim 2		39,09	0,00	0,05	0,05	43,32	0,28	0,16	17,37	0,23	100,55	1,07	0,22	83,2
MU73-1c 6c1	core 1		39,13	0,01	0,08	0,04	44,49	0,29	0,21	16,07	0,24	100,58	1,10	0,20	84,6
MU73-1c 6c2	core 2		39,00	0,01	0,08	0,05	43,34	0,29	0,19	17,06	0,18	100,24	1,08	0,21	83,4
MU73-1c 7b1	rim		38,77	0,01	0,06	0,03	42,74	0,31	0,29	17,56	0,24	100,01	1,06	0,22	82,8
MU73-1c 7c1	core		38,97	0,01	0,07	0,03	43,43	0,29	0,26	17,14	0,24	100,43	1,08	0,21	83,4

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU73-1c 8b1	rim		38,85	0,01	0,03	0,07	43,19	0,27	0,23	16,79	0,25	99,70	1,07	0,21	83,6
MU73-1c 8c1	core		39,50	0,00	0,04	0,00	45,18	0,27	0,15	14,87	0,30	100,36	1,12	0,19	85,7
MU73-1c 9b1	rim 1	X	38,64	0,02	0,03	0,05	41,75	0,33	0,33	18,94	0,20	100,30	1,04	0,24	81,4
MU73-1c 9b2	rim 2	X	38,40	0,03	0,04	0,01	39,72	0,31	0,31	21,90	0,14	100,88	0,99	0,27	78,2
MU73-1c 9c1	core 1	X	39,26	0,01	0,04	0,02	43,76	0,27	0,21	16,74	0,23	100,57	1,09	0,21	83,8
MU73-1c 9c2	core 2	X	39,09	0,00	0,08	0,03	43,68	0,29	0,21	17,35	0,25	100,99	1,08	0,22	83,3
MU73-1c 10b1	rim a		38,80	0,02	0,05	0,06	42,85	0,29	0,20	17,55	0,27	100,10	1,06	0,22	82,9
MU73-1c 10b2	rim b		39,03	0,01	0,03	0,07	42,67	0,32	0,15	18,25	0,22	100,77	1,06	0,23	82,2
MU73-1c 10c1	core		38,83	0,01	0,05	0,00	43,02	0,17	0,32	17,90	0,39	100,68	1,07	0,22	82,6
MU73-1c 11b1	rim 1a		38,72	0,00	0,06	0,09	43,17	0,30	0,16	17,52	0,28	100,33	1,07	0,22	83,0
MU73-1c 11b2	rim 1b		39,05	0,03	0,04	0,03	42,87	0,31	0,24	17,64	0,23	100,49	1,06	0,22	82,8
MU73-1c 11c1	core 1		39,41	0,02	0,05	0,11	44,97	0,26	0,28	14,97	0,29	100,37	1,12	0,19	85,6
MU73-1c 11c2	~core 2		39,09	0,02	0,05	0,01	44,01	0,29	0,28	16,21	0,25	100,21	1,09	0,20	84,3
MU73-1c 11c3	~core 3		39,14	0,02	0,02	0,04	43,33	0,31	0,26	17,35	0,29	100,79	1,08	0,22	83,2
MU73-1c 12b1	rim a		39,10	0,00	0,03	0,02	43,74	0,27	0,18	15,97	0,35	99,68	1,09	0,20	84,4
MU73-1c 12b2	rim b		39,13	0,00	0,07	0,09	43,98	0,28	0,26	16,69	0,28	100,83	1,09	0,21	83,9
MU73-1c 12c1	core		39,37	0,01	0,06	0,06	44,94	0,26	0,23	15,42	0,32	100,69	1,12	0,19	85,2
MU73-1c 13b1	rim 2		39,27	0,00	0,05	0,10	44,07	0,27	0,28	16,10	0,23	100,41	1,09	0,20	84,4
MU73-1c 13b2	rim 5		39,09	0,00	0,05	0,02	43,07	0,29	0,24	17,41	0,31	100,52	1,07	0,22	83,1
MU73-1c 13c1	core 1		39,47	0,00	0,06	0,04	44,76	0,26	0,17	15,34	0,26	100,34	1,11	0,19	85,2
MU73-1c 13c2	core 2		39,23	0,01	0,04	0,03	44,42	0,20	0,19	15,41	0,24	99,78	1,10	0,19	85,1
MU73-1c 13c3	core 3		39,24	0,00	0,05	0,05	44,72	0,27	0,27	15,25	0,26	100,11	1,11	0,19	85,3
MU73-1c 13c4	core 4		39,31	0,00	0,06	0,07	44,83	0,26	0,20	15,41	0,29	100,43	1,11	0,19	85,2
MU73-1c 13c5	core 5		39,51	0,00	0,03	0,00	45,02	0,26	0,30	15,12	0,23	100,47	1,12	0,19	85,5
MU73-1c 14b1	rim		38,95	0,03	0,08	0,06	43,06	0,30	0,33	17,16	0,20	100,24	1,07	0,21	83,2
MU73-1c 14c1	core		39,19	0,01	0,04	0,02	44,00	0,24	0,15	16,58	0,23	100,48	1,09	0,21	84,0
MU73-1c 15b1	rim 2	X	38,81	0,00	0,07	0,06	42,21	0,30	0,27	18,33	0,20	100,25	1,05	0,23	82,0
MU73-1c 15c1	core 1		39,76	0,00	0,05	0,07	47,07	0,24	0,19	12,25	0,39	100,03	1,17	0,15	88,4
MU73-1c 15c2	core 2	X	39,66	0,00	0,07	0,07	46,59	0,23	0,12	12,67	0,39	99,84	1,16	0,16	87,9
MU73-1c 16b1	rim a		38,99	0,01	0,05	0,04	42,84	0,31	0,23	17,17	0,22	99,88	1,06	0,22	83,2
MU73-1c 16b2	core b		38,82	0,02	0,07	0,06	42,86	0,32	0,24	17,43	0,19	100,04	1,06	0,22	83,0
MU73-1c 16c1	core a		39,01	0,00	0,04	0,07	43,17	0,28	0,19	17,80	0,32	100,89	1,07	0,22	82,8
MU73-1c 16c2	rim b		38,85	0,00	0,05	0,00	43,02	0,28	0,26	17,52	0,28	100,25	1,07	0,22	82,9
MU74 2a a1	rim 1a	X	39,80		0,04	0,01	44,73	0,28	0,15	14,30	0,24	99,65	1,11	0,18	86,1
MU74 2a a2	rim 2a		39,56		0,06	0,00	44,72	0,25	0,16	14,25	0,27	99,26	1,11	0,18	86,1
MU74 2a a3	core 2		39,87		0,06	0,11	44,82	0,25	0,11	14,03	0,27	99,66	1,11	0,18	86,4
MU74 2a a4	rim 2b		39,63		0,05	0,06	44,20	0,30	0,17	14,85	0,22	99,49	1,10	0,19	85,5
MU74 2a a5	rim 1b	X	39,69		0,04	0,00	44,99	0,24	0,23	13,94	0,27	99,40	1,12	0,17	86,5
MU74 2a a6	rim 1c	X	39,76		0,02	0,05	44,37	0,29	0,19	14,71	0,23	99,62	1,10	0,18	85,7

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU74 2a b1	core		40,25		0,05	0,06	47,70	0,23	0,13	10,55	0,39	99,36	1,18	0,13	90,0
MU74 2a b2	rim a		40,35		0,05	0,08	45,78	0,23	0,19	13,50	0,30	100,60	1,14	0,17	87,0
MU74 2a b3	rim b		39,66		0,06	0,08	45,08	0,25	0,21	13,10	0,32	99,07	1,12	0,16	87,2
MU74 2a c1	core 1	X	40,26		0,06	0,08	46,80	0,23	0,11	11,15	0,38	99,07	1,16	0,14	89,3
MU74 2a c2	rim 1	X	39,65		0,08	0,03	45,30	0,23	0,19	13,32	0,29	99,08	1,12	0,17	87,1
MU74 2a c3	~core 2	X	39,93		0,07	0,07	46,49	0,21	0,16	11,77	0,34	99,07	1,15	0,15	88,7
MU74 2a c4	~rim 2	X	39,90		0,05	0,10	45,87	0,23	0,16	12,55	0,30	99,15	1,14	0,16	87,9
MU74 2a d1	core		39,71		0,04	0,08	44,56	0,26	0,22	14,74	0,24	99,86	1,11	0,18	85,7
MU74 2a d2	rim		39,67		0,03	0,09	44,37	0,28	0,13	14,84	0,21	99,63	1,10	0,19	85,5
MU74 2a e1	core 1		39,42		0,05	0,01	44,18	0,28	0,21	14,76	0,27	99,22	1,10	0,18	85,6
MU74 2a e2	rim 2a		39,65		0,05	0,04	44,15	0,29	0,19	14,65	0,23	99,25	1,10	0,18	85,6
MU74 2a e3	rim 2b		39,64		0,06	0,04	44,47	0,28	0,19	14,76	0,21	99,64	1,10	0,18	85,6
MU74 2a f1	core		40,02		0,02	0,09	44,87	0,28	0,15	14,40	0,22	100,07	1,11	0,18	86,1
MU74 2a f2	rim a		39,83		0,02	0,04	44,40	0,28	0,14	14,97	0,20	99,89	1,10	0,19	85,4
MU74 2a f3	rim b		39,81		0,05	0,08	44,33	0,27	0,16	14,68	0,28	99,83	1,10	0,18	85,7
MU74 2a g1	core 1	X	40,54		0,04	0,14	47,92	0,22	0,19	10,42	0,39	99,86	1,19	0,13	90,1
MU74 2a g2	rim 1	X	39,73		0,07	0,09	44,45	0,26	0,17	14,35	0,20	99,43	1,10	0,18	86,0
MU74 2a g3	core 2	X	40,64		0,07	0,11	48,17	0,23	0,11	10,36	0,31	100,17	1,20	0,13	90,2
MU74 2a h1	core 1	X	40,42		0,06	0,07	47,16	0,24	0,10	11,76	0,33	100,15	1,17	0,15	88,8
MU74 2a h2	rim 1	X	39,86		0,02	0,00	45,18	0,26	0,19	13,62	0,29	99,69	1,12	0,17	86,8
MU74 2a h3	core 2	X	40,26		0,05	0,07	46,86	0,25	0,19	11,77	0,37	99,89	1,16	0,15	88,7
MU74 2a i1	core		40,37		0,04	0,07	46,91	0,24	0,16	11,72	0,38	100,08	1,16	0,15	88,8
MU74 2a i2	rim a		40,42		0,05	0,09	46,86	0,23	0,21	11,90	0,34	100,12	1,16	0,15	88,6
MU74 2a i3	rim b		39,85		0,05	0,05	44,21	0,30	0,24	14,99	0,25	99,94	1,10	0,19	85,4
MU74 2a j1	core	X	40,30		0,05	0,07	47,34	0,24	0,15	11,13	0,35	99,63	1,17	0,14	89,4
MU74 2a j2	rim a	X	39,97		0,05	0,09	45,67	0,23	0,20	13,20	0,33	99,90	1,13	0,17	87,3
MU74 2a j3	rim b	X	40,15		0,04	0,03	44,77	0,29	0,24	14,45	0,23	100,21	1,11	0,18	86,0
MU74 2a k1	rim a		39,43		0,05	0,10	45,23	0,24	0,15	13,63	0,35	99,17	1,12	0,17	86,8
MU74 2a k2	core		39,95		0,05	0,05	46,02	0,21	0,16	12,84	0,35	99,64	1,14	0,16	87,7
MU74 2a k3	rim b		39,45		0,05	0,01	43,94	0,28	0,15	14,80	0,30	98,99	1,09	0,19	85,5
MU74 2a l1	rim a		39,96		0,05	0,05	45,76	0,26	0,15	13,16	0,32	99,75	1,14	0,16	87,3
MU74 2a l2	core		39,81		0,05	0,00	45,22	0,27	0,20	13,58	0,27	99,40	1,12	0,17	86,8
MU74 2a l3	rim b		39,59		0,03	0,10	43,47	0,32	0,18	15,42	0,21	99,32	1,08	0,19	84,8
MU74 2a m1	~core	X	39,90		0,04	0,06	45,68	0,27	0,19	13,44	0,27	99,86	1,13	0,17	87,1
MU74 2a m2	~rim a	X	39,70		0,05	0,00	45,13	0,27	0,18	13,97	0,20	99,69	1,12	0,18	86,5
MU74 2a m3	rim b	X	39,76		0,04	0,06	44,57	0,26	0,21	14,94	0,25	100,10	1,11	0,19	85,5
MU74 2a n1	core	X	40,12		0,05	0,10	46,20	0,25	0,20	12,39	0,32	99,63	1,15	0,16	88,1
MU74 2a n2	rim a	X	39,71		0,04	0,04	45,02	0,24	0,17	13,41	0,23	98,85	1,12	0,17	86,9
MU74 2a n3	rim b	X	39,73		0,05	0,08	44,29	0,28	0,21	14,83	0,23	99,70	1,10	0,19	85,5
MU74 2a o1	core a		39,16		0,05	0,00	41,32	0,27	0,20	18,29	0,22	99,52	1,03	0,23	81,7

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU74 2a o2	rim a		39,74		0,05	0,07	44,09	0,29	0,18	15,23	0,19	99,83	1,09	0,19	85,1
MU74 2a o3	core b		38,83		0,05	0,04	40,63	0,25	0,22	19,52	0,22	99,84	1,01	0,24	80,5
MU74 2a o4	rim b		39,58		0,03	0,02	43,68	0,30	0,22	15,39	0,25	99,47	1,08	0,19	84,9
MU74 2a p1	core		39,77		0,06	0,14	45,65	0,28	0,22	13,68	0,24	100,04	1,13	0,17	86,9
MU74 2a p2	rim a		39,57		0,05	0,11	44,70	0,28	0,10	14,72	0,23	99,78	1,11	0,18	85,7
MU74 2a p3	rim b		39,57		0,04	0,06	44,47	0,28	0,15	14,32	0,27	99,27	1,10	0,18	86,0
MU74 2a q1	core		39,21		0,05	0,05	44,09	0,25	0,25	14,59	0,26	98,76	1,09	0,18	85,7
MU74 2a q2	rim		39,22		0,05	0,00	43,72	0,28	0,19	15,19	0,24	98,89	1,08	0,19	85,1
MU74 2a r1	core	X	39,75		0,05	0,08	45,16	0,25	0,24	13,70	0,31	99,53	1,12	0,17	86,7
MU74 2a r2	rim	X	39,84		0,04	0,06	45,71	0,25	0,08	13,31	0,29	99,69	1,13	0,17	87,2
MU74 2a s1	core		39,75		0,04	0,08	45,01	0,25	0,20	14,06	0,26	99,64	1,12	0,18	86,4
MU74 2a s2	~rim		39,79		0,06	0,08	44,52	0,26	0,16	14,54	0,26	99,66	1,10	0,18	85,8
MU74 2a t1	core 1		40,00		0,06	0,14	45,88	0,25	0,12	12,94	0,28	99,67	1,14	0,16	87,5
MU74 2a t2	core 2		39,72		0,03	0,09	44,93	0,27	0,23	13,90	0,28	99,45	1,11	0,17	86,5
MU74 2a t3	~core 3		39,77		0,02	0,01	43,82	0,27	0,22	15,13	0,23	99,47	1,09	0,19	85,2
MU74 2a v1	core	X	38,71		0,04	0,07	40,98	0,26	0,25	18,47	0,16	98,94	1,02	0,23	81,5
MU74 2a v2	rim a	X	36,92		0,03	0,00	33,24	0,22	0,31	27,25	0,15	98,11	0,82	0,34	70,7
MU74 2a v3	rim b	X	39,00		0,04	0,05	41,60	0,31	0,23	17,58	0,16	99,10	1,03	0,22	82,4
MU74 2a w1	core		38,97		0,05	0,00	41,20	0,27	0,24	18,68	0,20	99,80	1,02	0,23	81,4
MU74 2a w2	rim a		39,82		0,02	0,09	44,27	0,28	0,12	14,87	0,21	99,77	1,10	0,19	85,5
MU74 2a w3	rim b		39,83		0,05	0,00	44,23	0,29	0,14	14,97	0,27	99,79	1,10	0,19	85,4
MU74-3 1c1	core a	X	39,99	0,00	0,05	0,11	47,74	0,23	0,17	11,59	0,38	100,31	1,18	0,15	89,1
MU74-3 1c2	core b	X	39,84	0,00	0,05	0,04	47,20	0,24	0,16	12,57	0,39	100,50	1,17	0,16	88,1
MU74-3 1c3	core c	X	40,01	0,02	0,08	0,09	47,87	0,24	0,21	12,18	0,41	101,12	1,19	0,15	88,6
MU74-3 2b1	rim a		38,44	0,04	0,04	0,04	39,99	0,32	0,35	21,80	0,22	101,24	0,99	0,27	78,4
MU74-3 2b2	rim b		38,12	0,03	0,05	0,03	39,93	0,31	0,36	21,27	0,24	100,33	0,99	0,27	78,8
MU74-3 2c1	core		38,84	0,00	0,03	0,00	43,79	0,27	0,27	16,42	0,19	99,83	1,09	0,21	84,1
MU74-3 3b1	rim 1		39,09	0,01	0,06	0,09	42,30	0,26	0,27	18,35	0,27	100,72	1,05	0,23	82,0
MU74-3 3b2	rim 2		39,04	0,04	0,03	0,06	42,32	0,27	0,24	18,29	0,27	100,56	1,05	0,23	82,1
MU74-3 3c1	core 1	X	40,05	0,01	0,07	0,07	47,80	0,24	0,03	11,26	0,40	99,95	1,19	0,14	89,4
MU74-3 3c2	core 2	X	40,32	0,00	0,04	0,19	48,20	0,25	0,09	11,25	0,45	100,80	1,20	0,14	89,5
MU74-3 3c3	core 3	X	40,26	0,00	0,06	0,08	47,79	0,23	0,20	11,25	0,44	100,30	1,19	0,14	89,4
MU74-3 3c4	core 4		40,27	0,00	0,08	0,08	47,84	0,25	0,09	11,24	0,46	100,32	1,19	0,14	89,4
MU74-3 4b1	rim 1	X	38,34	0,00	0,05	0,07	40,65	0,29	0,33	20,45	0,23	100,40	1,01	0,26	79,7
MU74-3 4b2	rim 3		38,75	0,02	0,06	0,07	42,54	0,28	0,31	18,30	0,21	100,55	1,06	0,23	82,2
MU74-3 4c1	core 1	X	40,05	0,00	0,09	0,14	48,10	0,24	0,13	10,90	0,39	100,03	1,19	0,14	89,7
MU74-3 4c2	core 2		40,16	0,00	0,03	0,13	48,04	0,23	0,17	11,14	0,45	100,38	1,19	0,14	89,5
MU74-3 4c3	core 3		40,30	0,01	0,13	0,05	48,02	0,23	0,20	11,31	0,48	100,74	1,19	0,14	89,4
MU74-3 5b1	rim a		38,31	0,03	0,14	0,01	39,92	0,32	0,29	21,41	0,18	100,66	0,99	0,27	78,7

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU74-3 5b2	rim b		37,91	0,05	0,03	0,01	38,74	0,33	0,40	22,97	0,20	100,68	0,96	0,29	77,0
MU74-3 5b3	rim c		38,66	0,03	0,05	0,12	41,97	0,31	0,19	18,93	0,22	100,52	1,04	0,24	81,5
MU74-3 5c1	core		38,92	0,01	0,04	0,02	42,35	0,29	0,16	18,35	0,22	100,38	1,05	0,23	82,0
MU74-3 6b1	rim 1a	X	38,53	0,02	0,07	0,01	42,04	0,30	0,24	17,88	0,26	99,34	1,04	0,22	82,3
MU74-3 6b2	rim 1b	X	39,20	0,03	0,08	0,06	42,69	0,29	0,33	18,26	0,20	101,17	1,06	0,23	82,2
MU74-3 6b3	rim 2	X	39,01	0,03	0,04	0,02	42,52	0,30	0,23	18,22	0,24	100,62	1,06	0,23	82,2
MU74-3 6b4	rim 1c	X	39,20	0,01	0,05	0,02	42,80	0,31	0,34	18,13	0,20	101,05	1,06	0,23	82,4
MU74-3 6c1	core 1	X	38,86	0,00	0,03	0,02	41,67	0,22	0,25	19,44	0,22	100,74	1,03	0,24	80,9
MU74-3 6c2	~core 2	X	38,74	0,02	0,06	0,07	42,01	0,29	0,29	18,61	0,23	100,36	1,04	0,23	81,7
MU74-3 7b1	rim a		38,55	0,01	0,06	0,03	41,78	0,30	0,30	19,23	0,23	100,50	1,04	0,24	81,1
MU74-3 7b2	rim b		38,75	0,01	0,05	0,00	41,71	0,31	0,22	18,86	0,19	100,12	1,03	0,24	81,4
MU74-3 7b3	rim c		37,92	0,04	0,04	0,01	39,24	0,33	0,28	22,06	0,25	100,20	0,97	0,28	77,9
MU74-3 7c1	core a		38,60	0,02	0,05	0,06	41,68	0,26	0,29	19,48	0,24	100,71	1,03	0,24	80,9
MU74-3 7c2	core b		38,70	0,00	0,04	0,06	42,14	0,29	0,25	18,57	0,21	100,26	1,05	0,23	81,8
MU74-3 7c3	core c		38,72	0,02	0,04	0,06	41,99	0,29	0,25	18,54	0,22	100,15	1,04	0,23	81,8
MU74-3 7c4	core d		38,83	0,02	0,05	0,06	41,64	0,28	0,25	19,24	0,23	100,62	1,03	0,24	81,1
MU74-3 9b1	rim 1	X	39,07	0,02	0,05	0,04	41,79	0,28	0,32	19,11	0,25	100,99	1,04	0,24	81,2
MU74-3 9b2	rim 2	X	39,29	0,02	0,05	0,06	44,12	0,26	0,25	16,41	0,31	100,83	1,09	0,21	84,2
MU74-3 9b3	rim 3	X	38,04	0,03	0,06	0,00	39,85	0,29	0,19	20,92	0,20	99,59	0,99	0,26	79,1
MU74-3 9c1	core 1	X	40,24	0,00	0,05	0,05	47,95	0,23	0,23	11,52	0,39	100,66	1,19	0,14	89,2
MU74-3 9c2	core 2	X	39,79	0,02	0,08	0,11	46,78	0,23	0,20	13,02	0,40	100,62	1,16	0,16	87,7
MU74-3 9c3	core 3	X	40,07	0,00	0,06	0,10	47,35	0,23	0,20	12,01	0,37	100,39	1,17	0,15	88,6
MU74-3 9c4	core 4	X	40,24	0,00	0,07	0,06	47,31	0,21	0,16	12,56	0,45	101,09	1,17	0,16	88,2
MU74-3 11b1	rim a		38,61	0,01	0,05	0,00	40,75	0,31	0,25	20,65	0,16	100,80	1,01	0,26	79,6
MU74-3 11b2	rim b		39,35	0,03	0,05	0,00	44,20	0,27	0,25	16,35	0,31	100,85	1,10	0,20	84,3
MU74-3 11c1	core		39,55	0,00	0,04	0,05	45,21	0,25	0,29	14,97	0,28	100,67	1,12	0,19	85,7
MU74-3 12b1	rim 1		38,79	0,03	0,05	0,08	40,51	0,29	0,21	20,47	0,24	100,68	1,01	0,26	79,7
MU74-3 12b2	rim 2a	X	38,24	0,05	0,05	0,01	40,62	0,27	0,31	20,55	0,21	100,34	1,01	0,26	79,7
MU74-3 12b3	rim 2b	X	38,87	0,02	0,04	0,03	41,90	0,29	0,25	19,19	0,19	100,79	1,04	0,24	81,2
MU74-3 12c1	core 1		39,37	0,00	0,05	0,00	45,18	0,33	0,28	15,08	0,31	100,62	1,12	0,19	85,6
MU74-3 12c2	core 2	X	39,61	0,00	0,07	0,05	45,18	0,27	0,22	15,14	0,28	100,86	1,12	0,19	85,5
MU74-3 13b1	rim a	X	38,48	0,00	0,03	0,01	41,16	0,28	0,15	19,84	0,26	100,22	1,02	0,25	80,4
MU74-3 13b2	rim b	X	39,51	0,00	0,04	0,00	44,64	0,28	0,18	16,01	0,29	100,95	1,11	0,20	84,7
MU74-3 13b3	rim c	X	38,10	0,04	0,04	0,00	40,10	0,29	0,31	21,58	0,21	100,68	0,99	0,27	78,6
MU74-3 13c1	core	X	39,91	0,00	0,05	0,05	46,55	0,25	0,14	13,58	0,29	100,85	1,15	0,17	87,2
MU74-3 14b1	rim a		39,30	0,02	0,06	0,04	43,46	0,27	0,16	16,78	0,28	100,40	1,08	0,21	83,7
MU74-3 14b2	rim b		38,60	0,02	0,05	0,03	40,52	0,27	0,24	20,98	0,24	100,98	1,01	0,26	79,3
MU74-3 14b3	rim c		38,81	0,04	0,05	0,06	41,85	0,28	0,24	18,65	0,23	100,20	1,04	0,23	81,6
MU74-3 14c1	core		40,17	0,00	0,06	0,09	47,24	0,24	0,20	12,43	0,39	100,85	1,17	0,16	88,3
MU74-3 15b1	rim	X	38,82	0,00	0,05	0,02	41,97	0,29	0,26	18,63	0,20	100,24	1,04	0,23	81,7

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU74-3 15c1	core	X	40,44	0,00	0,05	0,12	47,97	0,22	0,18	11,01	0,46	100,47	1,19	0,14	89,6
MU74-3 16b1	rim a	X	38,39	0,04	0,07	0,03	41,03	0,32	0,22	19,32	0,21	99,69	1,02	0,24	80,8
MU74-3 16b2	rim b	X	38,52	0,04	0,06	0,00	41,46	0,32	0,28	18,75	0,27	99,71	1,03	0,23	81,4
MU74-3 16c1	core	X	39,30	0,01	0,06	0,00	44,22	0,27	0,19	16,27	0,30	100,66	1,10	0,20	84,3
MU74-3 17b1	rim a	X	38,29	0,05	0,05	0,09	40,24	0,33	0,29	20,38	0,24	100,00	1,00	0,26	79,6
MU74-3 17b2	rim b	X	38,76	0,01	0,06	0,07	40,89	0,32	0,28	20,51	0,23	101,14	1,01	0,26	79,8
MU74-3 17c1	core	X	38,62	0,00	0,05	0,06	41,51	0,27	0,28	19,20	0,22	100,23	1,03	0,24	81,1
MU74-3 18b1	rim a		38,39	0,03	0,06	0,04	41,12	0,30	0,35	20,04	0,22	100,60	1,02	0,25	80,2
MU74-3 18b2	rim b		38,92	0,02	0,04	0,02	41,52	0,30	0,33	19,71	0,20	101,10	1,03	0,25	80,7
MU74-3 18c1	core		39,12	0,01	0,04	0,00	43,71	0,29	0,31	16,97	0,22	100,69	1,08	0,21	83,6
MU74-5a 1b1	rim		37,84	0,04	0,13	0,04	40,71	0,37	0,22	19,17	0,17	98,71	1,01	0,24	80,8
MU74-5a 1c1	core		39,24	0,00	0,02	0,09	45,18	0,24	0,19	14,75	0,24	99,96	1,12	0,18	85,8
MU74-5a 2b1	rim 1		38,91	0,01	0,04	0,00	43,81	0,30	0,21	15,52	0,21	99,03	1,09	0,19	84,8
MU74-5a 2b2	rim 2		39,02	0,02	0,05	0,07	45,05	0,30	0,13	15,05	0,24	99,94	1,12	0,19	85,6
MU74-5a 2c1	core 1		39,42	0,00	0,05	0,13	46,22	0,24	0,23	13,15	0,30	99,74	1,15	0,16	87,4
MU74-5a 2c2	core 2		39,41	0,00	0,05	0,07	46,70	0,24	0,08	12,71	0,36	99,65	1,16	0,16	87,9
MU74-5a 3b1	rim a		38,81	0,00	0,05	0,09	43,55	0,30	0,27	15,99	0,24	99,32	1,08	0,20	84,4
MU74-5a 3b2	rim b		39,09	0,04	0,04	0,08	44,57	0,30	0,25	16,06	0,25	100,71	1,11	0,20	84,6
MU74-5a 3c1	core a		39,39	0,01	0,00	0,07	44,68	0,27	0,25	15,29	0,33	100,31	1,11	0,19	85,3
MU74-5a 3c2	core b		38,96	0,02	0,03	0,08	44,14	0,33	0,18	15,55	0,28	99,56	1,10	0,19	84,9
MU74-5a 4b1	rim 1a		38,95	0,03	0,03	0,04	44,30	0,31	0,23	15,74	0,25	99,92	1,10	0,20	84,8
MU74-5a 4b2	rim 1b		39,06	0,02	0,04	0,03	44,40	0,29	0,23	15,33	0,28	99,73	1,10	0,19	85,2
MU74-5a 4b3	rim 2		39,08	0,03	0,07	0,05	44,99	0,32	0,20	16,76	0,27	101,76	1,12	0,21	84,2
MU74-5a 4b4	rim 3		38,81	0,04	0,04	0,02	42,38	0,31	0,32	17,39	0,24	99,54	1,05	0,22	82,8
MU74-5a 4c1	core 1		39,24	0,01	0,05	0,07	45,35	0,27	0,17	14,20	0,25	99,61	1,13	0,18	86,3
MU74-5a 4c2	core 2		39,12	0,00	0,07	0,01	45,13	0,26	0,08	14,29	0,26	99,28	1,12	0,18	86,2
MU74-5a 4c3	core 3		39,67	0,00	0,03	0,02	45,28	0,25	0,16	14,53	0,24	100,19	1,12	0,18	86,1
MU74-5a 5b1	rim a		38,84	0,02	0,05	0,02	43,36	0,31	0,24	16,50	0,26	99,60	1,08	0,21	83,9
MU74-5a 5b2	rim b		39,19	0,00	0,00	0,10	44,34	0,33	0,23	15,50	0,23	99,93	1,10	0,19	85,0
MU74-5a 5c1	core		39,37	0,00	0,04	0,03	45,56	0,26	0,19	14,03	0,28	99,76	1,13	0,18	86,5
MU74-5a 6b1	rim		39,80	0,00	0,04	0,08	44,33	0,29	0,23	16,56	0,24	101,59	1,10	0,21	84,1
MU74-5a 6c1	core		38,70	0,00	0,07	0,01	44,16	0,26	0,26	15,82	0,24	99,53	1,10	0,20	84,7
MU74-5a 7b1	rim a		38,88	0,03	0,03	0,10	43,08	0,32	0,27	16,64	0,29	99,65	1,07	0,21	83,7
MU74-5a 7b2	rim b		38,81	0,02	0,08	0,05	42,72	0,29	0,28	17,40	0,30	99,96	1,06	0,22	82,9
MU74-5a 7c1	core a		39,17	0,00	0,04	0,01	45,64	0,27	0,07	14,26	0,30	99,75	1,13	0,18	86,4
MU74-5a 7c2	core b		38,99	0,00	0,08	0,07	44,89	0,25	0,15	14,59	0,25	99,29	1,11	0,18	85,9
MU74-5a 8b3	rim 3	X	39,11	0,00	0,06	0,03	44,65	0,30	0,26	14,80	0,33	99,58	1,11	0,19	85,7
MU74-5a 8b5	rim 5	X	39,01	0,06	0,03	0,03	43,43	0,33	0,23	16,40	0,26	99,81	1,08	0,21	84,0
MU74-5a 8b7	rim 7	X	39,63	0,01	0,04	0,07	45,19	0,27	0,24	14,14	0,32	99,92	1,12	0,18	86,4

Annexe 1 (suite)

Sample	Spot	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU74-5a 8b8	rim 8	X	39,05	0,00	0,09	0,01	43,98	0,32	0,24	15,86	0,27	99,82	1,09	0,20	84,6
MU74-5a 8c1	core 1	X	39,70	0,00	0,03	0,07	46,95	0,24	0,18	12,13	0,38	99,67	1,16	0,15	88,5
MU74-5a 8c2	core 2	X	39,67	0,00	0,06	0,00	46,26	0,23	0,12	12,76	0,36	99,51	1,15	0,16	87,8
MU74-5a 8c3	core 3	X	39,64	0,00	0,05	0,10	47,13	0,23	0,17	11,65	0,41	99,40	1,17	0,15	88,9
MU74-5a 8c4	core 4	X	39,84	0,01	0,05	0,05	47,52	0,28	0,14	11,35	0,42	99,69	1,18	0,14	89,2
MU74-5a 8c5	core 5	X	39,42	0,00	0,03	0,07	45,66	0,22	0,13	13,75	0,32	99,60	1,13	0,17	86,8
MU74-5a 8c6	core 6	X	39,92	0,01	0,07	0,05	47,64	0,26	0,05	11,21	0,36	99,58	1,18	0,14	89,4
MU74-5a 8c7	core 7	X	39,92	0,00	0,06	0,08	47,75	0,25	0,13	11,30	0,40	99,88	1,18	0,14	89,3
MU74-5a 8c8	core 8	X	39,67	0,00	0,00	0,07	47,05	0,24	0,15	11,58	0,36	99,15	1,17	0,15	88,9
MU74-5a 9b1	rim a	X	39,40	0,04	0,00	0,00	44,84	0,29	0,17	14,78	0,36	99,89	1,11	0,19	85,7
MU74-5a 9b2	rim c	X	39,07	0,04	0,04	0,04	44,25	0,30	0,29	15,21	0,26	99,50	1,10	0,19	85,2
MU74-5a 9b3	rim d	X	38,93	0,04	0,05	0,02	43,83	0,32	0,26	15,60	0,21	99,27	1,09	0,20	84,8
MU74-5a 9c1	core	X	39,86	0,00	0,08	0,10	47,61	0,22	0,13	11,36	0,37	99,78	1,18	0,14	89,2
MU74-5a 9c2	rim b	X	39,40	0,00	0,05	0,10	45,19	0,24	0,19	14,16	0,31	99,66	1,12	0,18	86,3
MU74-5a 10b1	rim 1	X	38,50	0,03	0,00	0,08	42,97	0,34	0,23	16,63	0,28	99,07	1,07	0,21	83,7
MU74-5a 10b4	rim 4		38,68	0,02	0,06	0,02	42,83	0,31	0,23	17,17	0,20	99,54	1,06	0,22	83,2
MU74-5a 10c1	core 1	X	39,68	0,00	0,08	0,03	46,65	0,25	0,13	12,16	0,37	99,40	1,16	0,15	88,4
MU74-5a 10c2	core 2	X	40,01	0,01	0,10	0,06	47,52	0,22	0,18	11,65	0,42	100,18	1,18	0,15	89,0
MU74-5a 10c3	core 3	X	39,83	0,00	0,03	0,04	46,99	0,21	0,08	12,02	0,37	99,60	1,17	0,15	88,6
MU74-5a 10c4	core 4		39,33	0,00	0,00	0,08	44,67	0,25	0,14	14,79	0,34	99,61	1,11	0,19	85,7
MU74-5a 11b1	rim 1a	X	37,27	0,01	0,03	0,00	36,48	0,26	0,25	25,13	0,26	99,72	0,91	0,31	74,2
MU74-5a 11b2	rim 1b	X	37,91	0,05	0,00	0,00	39,44	0,34	0,30	21,05	0,15	99,26	0,98	0,26	78,8
MU74-5a 11b3	rim 2		37,62	0,04	0,04	0,06	38,73	0,34	0,32	22,58	0,22	99,97	0,96	0,28	77,3
MU74-5a 11b4	rim 1c		38,06	0,05	0,00	0,03	38,03	0,34	0,24	23,78	0,19	100,78	0,94	0,30	76,0
MU74-5a 11c1	core 1	X	39,18	0,00	0,06	0,08	44,51	0,27	0,20	15,69	0,23	100,24	1,10	0,20	84,9
MU74-5a 11c2	core 2		38,24	0,00	0,05	0,00	40,27	0,27	0,18	20,80	0,24	100,05	1,00	0,26	79,3
MU74-5a 11c3	core 3		38,45	0,02	0,04	0,03	41,86	0,27	0,18	18,72	0,26	99,86	1,04	0,23	81,6
MU74-5a 12b1	rim a	X	38,62	0,02	0,05	0,04	42,67	0,31	0,20	16,71	0,23	98,89	1,06	0,21	83,5
MU74-5a 12b2	rim b	X	38,62	0,01	0,02	0,10	42,66	0,35	0,21	16,62	0,23	98,87	1,06	0,21	83,6
MU74-5a 12c1	core	X	39,04	0,02	0,00	0,03	45,40	0,24	0,11	14,09	0,28	99,21	1,13	0,18	86,5
MU74-5a 13b1	rim 1		38,54	0,02	0,07	0,02	43,23	0,35	0,29	15,96	0,26	98,78	1,07	0,20	84,3
MU74-5a 13c1	core 1		40,08	0,00	0,04	0,05	47,95	0,23	0,10	11,07	0,41	99,96	1,19	0,14	89,6
MU74-5a 13c2	core 2		39,69	0,00	0,00	0,17	47,26	0,25	0,17	11,52	0,39	99,44	1,17	0,14	89,0
MU74-5a 14b1	rim 1		37,83	0,01	0,00	0,02	39,48	0,26	0,24	21,47	0,24	99,56	0,98	0,27	78,5
MU74-5a 14b2	rim 2		37,88	0,01	0,03	0,06	39,58	0,26	0,26	20,96	0,22	99,26	0,98	0,26	78,9
MU74-5a 14c1	core 1		40,13	0,01	0,00	0,03	47,86	0,23	0,17	10,88	0,42	99,74	1,19	0,14	89,7
MU74-5a 14c2	core 2		39,86	0,00	0,08	0,09	47,63	0,20	0,07	10,98	0,36	99,26	1,18	0,14	89,6
MU74-5a 15b1	rim a		39,12	0,01	0,04	0,06	44,09	0,31	0,25	15,93	0,20	100,02	1,09	0,20	84,6
MU74-5a 15b2	rim b		39,16	0,03	0,16	0,04	44,06	0,32	0,25	15,83	0,26	100,12	1,09	0,20	84,6
MU74-5a 15c1	core		38,83	0,00	0,02	0,00	42,91	0,31	0,23	17,26	0,18	99,77	1,06	0,22	83,1

Annexe 1 (suite)

Sample	Spot	Deformed	SiO₂	TiO₂	Al₂O₃	Cr₂O₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
MU74-5a 16b1	rim a	X	38,90	0,02	0,06	0,03	44,31	0,29	0,25	15,33	0,28	99,46	1,10	0,19	85,1
MU74-5a 16b2	rim b	X	38,73	0,02	0,04	0,05	43,77	0,31	0,13	16,74	0,20	100,00	1,09	0,21	83,8
MU74-5a 16c1	core a	X	39,75	0,02	0,09	0,04	47,59	0,25	0,16	10,95	0,37	99,24	1,18	0,14	89,6
MU74-5a 16c2	core b	X	39,22	0,00	0,05	0,01	44,98	0,27	0,13	14,26	0,28	99,20	1,12	0,18	86,2

FeO*: total Fe; **Fo%** = $100 \times [\text{MgO} / (\text{MgO} + \text{FeO})]_{\text{mol}}$

ANNEXE 2

Compositions en éléments majeurs et mineurs d'olivines du Kilauea Iki provenant des scories et d'échantillons de forage du lac de lave. Ces données sont utilisées dans l'article 2, mais seule une sélection a été soumise pour publication.

Sample	Spot	Depth (m)	Deformed	(wt%)				MgO	CaO	MnO	FeO*	NiO	Total	(mol)		
				SiO2	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃							MgO	FeO	Fo%
Surface																
KI59 1 c o1	core	-	X	40,13	-	0,05	0,12	47,28	0,26	0,13	11,43	0,33	99,90	1,75	0,24	89,1
KI59 1 c o2	~rim 1a	-	X	39,92	-	0,07	0,10	46,49	0,25	0,15	11,45	0,35	98,79	1,74	0,24	88,9
KI59 1 c o3	rim 1b	-	X	40,00	-	0,05	0,06	46,54	0,26	0,16	11,50	0,35	98,94	1,74	0,24	88,9
KI59 1 c o4	~rim 1c	-	X	40,20	-	0,05	0,10	46,94	0,27	0,14	11,74	0,31	99,74	1,74	0,24	88,8
KI59 1 d o1	core 1a	-	X	40,31	-	0,09	0,03	47,55	0,24	0,19	10,77	0,36	99,53	1,76	0,22	89,7
KI59 1 d o2	rim	-	X	40,03	-	0,06	0,06	47,47	0,25	0,13	10,79	0,35	99,22	1,76	0,23	89,7
KI59 1 d o3	core 1b	-	X	39,97	-	0,07	0,07	47,20	0,26	0,12	10,70	0,33	98,72	1,76	0,22	89,8
KI59 1 h o1	core	-	X	39,67	-	0,07	0,04	46,06	0,24	0,16	12,24	0,33	98,98	1,73	0,26	88,2
KI59 1 h o3	rim 1b	-	X	39,95	-	0,06	0,11	46,30	0,27	0,21	12,06	0,27	99,36	1,73	0,25	88,4
KI59 1 h o4	~rim 1c	-	X	40,14	-	0,07	0,06	46,35	0,27	0,14	12,37	0,31	99,77	1,72	0,26	88,1
KI59 1 j o1	core	-	X	40,11	-	0,05	0,12	47,20	0,25	0,16	11,31	0,32	99,52	1,75	0,24	89,2
KI59 1 j o2	rim 1a	-	X	39,57	-	0,07	0,04	44,62	0,30	0,21	14,67	0,21	99,68	1,68	0,31	85,8
KI59 1 j o3	rim 1b	-	X	40,21	-	0,06	0,08	47,07	0,26	0,18	11,49	0,35	99,70	1,74	0,24	89,0
KI59 1 j o4	rim 1c	-	X	40,35	-	0,06	0,07	47,27	0,26	0,16	11,24	0,31	99,72	1,75	0,23	89,3
KI59 1 l o1	core 1	-	X	39,61	-	0,05	0,02	44,89	0,28	0,17	14,09	0,27	99,37	1,69	0,30	86,3
KI59 1 l o2	~rim 2a	-	X	39,46	-	0,05	0,02	44,76	0,27	0,21	14,02	0,27	99,10	1,69	0,30	86,3
KI59 1 l o3	core 2	-	X	39,20	-	0,05	0,05	44,43	0,27	0,23	13,88	0,25	98,49	1,69	0,30	86,4
KI59 1 l o4	rim 2b	-	X	39,40	-	0,05	0,04	44,54	0,28	0,19	14,09	0,26	98,90	1,68	0,30	86,2
KI59 1 l o5	rim 2c	-	X	39,27	-	0,05	0,05	44,66	0,27	0,18	13,95	0,25	98,70	1,69	0,30	86,4
KI59 1 p o1	core	-	X	40,07	-	0,05	0,07	47,35	0,26	0,12	10,78	0,43	99,17	1,76	0,22	89,7
KI59 1 p o2	~rim 1a	-	X	39,99	-	0,06	0,10	47,24	0,23	0,22	11,00	0,38	99,22	1,75	0,23	89,5
KI59 1 p o3	rim 1b	-	X	39,79	-	0,07	0,10	46,22	0,25	0,10	12,14	0,39	99,05	1,73	0,26	88,3
KI59 1 a o1	core 1	-		39,95	-	0,03	0,04	46,85	0,25	0,16	11,67	0,36	99,35	1,74	0,24	88,8
KI59 1 a o2	core 2	-		40,22	-	0,06	0,11	46,93	0,25	0,13	11,41	0,29	99,39	1,74	0,24	89,1
KI59 1 a o3	core 3	-		40,48	-	0,06	0,01	47,15	0,26	0,17	11,38	0,37	99,88	1,74	0,24	89,1
KI59 1 a o4	rim 1	-		41,21	-	0,05	0,03	46,35	0,26	0,20	12,02	0,33	100,45	1,70	0,25	88,4
KI59 1 b o1	core 1	-		40,99	-	0,05	0,06	46,95	0,26	0,17	11,14	0,30	99,90	1,73	0,23	89,3
KI59 1 b o2	rim 1	-		39,61	-	0,05	0,07	46,72	0,27	0,12	11,33	0,31	98,59	1,75	0,24	89,1
KI59 1 b o3	rim 2a	-		40,09	-	0,03	0,00	46,80	0,26	0,17	11,72	0,36	99,55	1,74	0,24	88,8
KI59 1 b o4	core 2	-		40,28	-	0,06	0,08	46,85	0,25	0,16	11,29	0,30	99,28	1,74	0,24	89,2
KI59 1 b o5	rim 2b	-		39,90	-	0,06	0,07	46,72	0,27	0,19	11,99	0,30	99,61	1,74	0,25	88,5
KI59 1 e o1	core	-		39,87	-	0,05	0,07	46,20	0,29	0,17	12,78	0,31	99,74	1,72	0,27	87,7

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI59 1 e o3	rim 1b	-		40,08	-	0,04	0,04	45,97	0,27	0,19	12,44	0,30	99,37	1,71	0,26	88,0
KI59 1 f o1	core 1	-		40,47	-	0,05	0,05	47,57	0,26	0,10	11,41	0,34	100,25	1,75	0,24	89,2
KI59 1 f o2	core 2	-		39,99	-	0,04	0,14	46,30	0,27	0,17	12,76	0,36	100,02	1,72	0,27	87,8
KI59 1 f o3	core 3	-		39,86	-	0,03	0,02	46,50	0,27	0,14	12,17	0,33	99,31	1,73	0,26	88,3
KI59 1 f o4	rim 3	-		39,74	-	0,05	0,06	45,53	0,29	0,19	13,05	0,31	99,22	1,71	0,27	87,4
KI59 1 f o5	rim 1	-		39,58	-	0,04	0,03	45,79	0,29	0,20	13,17	0,30	99,38	1,72	0,28	87,3
KI59 1 g o1	core	-		39,61	-	0,05	0,09	44,85	0,27	0,20	13,84	0,31	99,36	1,69	0,29	86,5
KI59 1 g o2	rim 1a	-		39,99	-	0,03	0,10	45,24	0,26	0,13	13,74	0,31	99,78	1,69	0,29	86,7
KI59 1 g o3	rim 1b	-		39,74	-	0,05	0,06	44,99	0,28	0,18	13,74	0,33	99,47	1,69	0,29	86,6
KI59 1 i o1	core	-		39,94	-	0,04	0,04	46,65	0,27	0,14	12,06	0,32	99,46	1,74	0,25	88,4
KI59 1 i o2	rim 1a	-		39,62	-	0,04	0,07	44,79	0,27	0,21	14,17	0,31	99,48	1,68	0,30	86,2
KI59 1 i o3	rim 1b	-		39,92	-	0,06	0,04	45,96	0,26	0,14	12,58	0,30	99,33	1,72	0,26	87,9
KI59 1 k o1	core 1	-		39,75	-	0,05	0,02	46,62	0,27	0,13	11,78	0,34	98,95	1,74	0,25	88,7
KI59 1 k o2	rim 1	-		39,88	-	0,06	0,09	45,66	0,29	0,18	13,02	0,34	99,52	1,71	0,27	87,4
KI59 1 k o3	core 2	-		39,92	-	0,05	0,07	46,12	0,27	0,18	12,22	0,27	99,09	1,72	0,26	88,2
KI59 1 k o4	rim 2	-		40,10	-	0,06	0,05	46,63	0,27	0,24	12,03	0,32	99,70	1,73	0,25	88,5
KI59 1 k o5	~rim 3	-		39,84	-	0,07	0,04	45,83	0,28	0,21	12,89	0,32	99,56	1,71	0,27	87,6
KI59 1 m o1	~rim 1	-		40,67	-	0,06	0,05	48,10	0,24	0,16	10,62	0,35	100,25	1,76	0,22	90,0
KI59 1 m o2	~core 1	-		40,41	-	0,06	0,11	47,88	0,26	0,14	10,40	0,36	99,61	1,77	0,22	90,1
KI59 1 m o3	~core 2	-		40,65	-	0,05	0,09	47,77	0,25	0,18	10,39	0,35	99,74	1,76	0,21	90,1
KI59 1 n o1	core	-		39,61	-	0,05	0,05	45,41	0,29	0,15	13,36	0,28	99,26	1,70	0,28	87,1
KI59 1 n o2	rim 1a	-		39,71	-	0,03	0,04	44,99	0,28	0,26	13,38	0,30	99,01	1,69	0,28	87,0
KI59 1 n o3	rim 1b	-		39,80	-	0,05	0,08	45,13	0,28	0,21	13,08	0,33	98,98	1,70	0,28	87,2
KI59 1 o o1	core	-		40,49	-	0,07	0,03	47,80	0,26	0,13	10,99	0,36	100,14	1,76	0,23	89,6
KI59 1 o o2	rim 1a	-		40,45	-	0,07	0,04	46,94	0,25	0,14	10,89	0,38	99,38	1,74	0,23	89,5
KI59 1 o o3	rim 1b	-		40,16	-	0,07	0,06	47,19	0,24	0,17	10,91	0,38	99,33	1,75	0,23	89,5
KI59 1 q o1	core	-		39,93	-	0,04	0,10	46,04	0,28	0,18	12,68	0,27	99,51	1,72	0,27	87,8
KI59 1 q o2	rim 1a	-		39,93	-	0,06	0,02	44,63	0,30	0,14	13,95	0,30	99,34	1,68	0,29	86,4
KI59 1 q o3	rim 1b	-		39,99	-	0,03	0,07	46,13	0,26	0,11	12,70	0,34	99,73	1,72	0,27	87,8
KI59 1 r o1	core 1	-		40,10	-	0,05	0,11	46,80	0,24	0,19	11,75	0,33	99,57	1,74	0,25	88,7
KI59 1 r o2	~core 2	-		39,93	-	0,05	0,11	46,66	0,25	0,18	11,59	0,29	99,05	1,74	0,24	88,9
KI59 1 r o3	rim 2a	-		39,78	-	0,05	0,08	46,34	0,27	0,11	12,07	0,32	99,06	1,73	0,25	88,4
KI59 1 r o4	rim 2b	-		40,28	-	0,07	0,07	46,90	0,25	0,14	11,48	0,37	99,69	1,74	0,24	89,0
KI59 1 r o5	rim 1	-		40,34	-	0,06	0,10	46,77	0,26	0,14	11,60	0,35	99,64	1,73	0,24	88,9

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	FeO
KI59 2a c o1	core 1	-	X	40,42	-	0,05	0,09	44,67	0,30	0,11	13,85	0,28	99,23	1,68	0,29	86,5
KI59 2a c o2	rim 1	-	X	39,97	-	0,04	0,05	44,99	0,30	0,24	13,42	0,24	99,15	1,69	0,28	86,9
KI59 2a c o3	core 2	-	X	41,21	-	0,05	0,06	44,82	0,30	0,17	13,71	0,21	99,53	1,68	0,29	86,6
KI59 2a c o4	rim 2	-	X	40,31	-	0,08	0,09	47,13	0,24	0,12	10,45	0,34	98,87	1,75	0,22	90,0
KI59 2a d o1	core	-	X	40,49	-	0,07	0,10	45,87	0,25	0,21	12,31	0,30	99,09	1,72	0,26	88,1
KI59 2a d o2	rim 1a	-	X	40,34	-	0,08	0,05	46,30	0,24	0,20	11,53	0,32	100,03	1,70	0,24	88,8
KI59 2a g o1	core	-	X	40,07	-	0,06	0,09	46,88	0,24	0,16	10,82	0,31	98,90	1,74	0,23	89,5
KI59 2a g o2	rim 1a	-	X	40,41	-	0,07	0,08	47,39	0,24	0,18	10,35	0,32	99,12	1,75	0,22	90,1
KI59 2a g o3	rim 1b	-	X	40,48	-	0,05	0,14	46,60	0,25	0,07	11,17	0,25	98,86	1,73	0,23	89,2
KI59 2a l o1	core	-	X	40,47	-	0,05	0,05	44,57	0,33	0,17	13,44	0,28	98,58	1,68	0,29	86,8
KI59 2a l o2	rim 1a	-	X	40,77	-	0,05	0,10	45,52	0,29	0,17	12,53	0,33	99,00	1,71	0,26	87,8
KI59 2a l o3	rim 1b	-	X	40,27	-	0,05	0,02	45,72	0,30	0,20	12,70	0,29	99,15	1,77	0,22	90,1
KI59 2a l o4	rim 1c	-	X	40,71	-	0,08	0,04	47,79	0,23	0,14	10,39	0,35	99,00	1,72	0,27	87,7
KI59 2a b o1	core	-	X	39,89	-	0,08	0,06	44,59	0,27	0,15	13,82	0,26	98,46	1,69	0,29	86,5
KI59 2a b o2	rim 1a	-	X	39,88	-	0,07	0,03	47,21	0,24	0,13	10,86	0,31	98,81	1,76	0,23	89,6
KI59 2a b o3	rim 1b	-	X	40,15	-	0,07	0,05	46,50	0,25	0,11	11,68	0,32	98,95	1,74	0,25	88,7
KI59 2a f o1	core	-	X	39,65	-	0,05	0,07	45,55	0,26	0,16	12,63	0,32	99,43	1,70	0,26	87,7
KI59 2a f o2	rim 1a	-	X	39,88	-	0,08	0,13	46,99	0,23	0,19	10,69	0,41	100,56	1,71	0,22	89,7
KI59 2a f o3	rim 1b	-	X	39,67	-	0,05	0,10	47,33	0,24	0,10	10,56	0,37	99,37	1,75	0,22	89,9
KI59 2a e o1	core 1	-	X	40,13	-	0,03	0,08	47,38	0,28	0,18	10,63	0,34	98,98	1,76	0,22	89,8
KI59 2a e o2	rim 1	-	X	39,11	-	0,04	0,07	47,51	0,29	0,15	10,61	0,37	99,44	1,76	0,22	89,9
KI59 2a e o3	rim 2a	-	X	39,80	-	0,05	0,05	47,07	0,28	0,12	10,87	0,36	99,27	1,74	0,23	89,5
KI59 2a e o4	core 2	-	X	39,93	-	0,06	0,09	45,84	0,28	0,21	12,66	0,31	99,15	1,72	0,27	87,8
KI59 2a e o5	rim 2b	-	X	40,38	-	0,05	0,08	46,39	0,30	0,16	12,10	0,28	99,42	1,73	0,25	88,3
KI59 2a e o6	rim 3	-	X	41,83	-	0,05	0,00	46,38	0,28	0,24	12,35	0,34	99,37	1,73	0,26	88,1
KI59 2a e o7	core 3	-	X	40,49	-	0,05	0,09	45,29	0,27	0,18	12,66	0,29	98,74	1,70	0,27	87,6
KI59 2a h o1	core	-	X	39,70	-	0,05	0,08	46,01	0,25	0,18	12,99	0,28	100,00	1,71	0,27	87,5
KI59 2a h o2	rim 1a	-	X	40,05	-	0,04	0,05	45,60	0,27	0,13	12,70	0,28	99,43	1,70	0,27	87,7
KI59 2a h o3	rim 1b	-	X	39,64	-	0,04	0,06	45,95	0,29	0,18	12,64	0,32	99,36	1,72	0,27	87,8
KI59 2a i o1	core 1	-	X	39,70	-	0,06	0,06	45,84	0,29	0,23	12,50	0,34	99,58	1,71	0,26	87,9
KI59 2a i o2	rim 1	-	X	40,16	-	0,04	0,05	45,18	0,28	0,17	13,15	0,24	98,79	1,70	0,28	87,2
KI59 2a i o3	core 2	-	X	40,09	-	0,03	0,03	45,15	0,28	0,15	13,08	0,27	98,86	1,70	0,28	87,2
KI59 2a i o4	core 3	-	X	39,86	-	0,07	0,10	45,07	0,29	0,23	13,07	0,24	98,60	1,70	0,28	87,2
KI59 2a i o5	rim 3	-	X	40,19	-	0,06	0,07	47,03	0,27	0,20	11,46	0,36	99,76	1,74	0,24	89,0
KI59 2a j o1	core	-	X	39,65	-	0,02	0,05	45,59	0,26	0,19	12,98	0,31	99,87	1,69	0,27	87,4
KI59 2a j o2	rim 1a	-	X	39,84	-	0,05	0,05	45,78	0,27	0,17	12,72	0,29	99,30	1,71	0,27	87,7
KI59 2a j o3	rim 1b	-	X	39,47	-	0,06	0,09	47,30	0,26	0,12	10,67	0,34	99,30	1,75	0,22	89,8
KI59 2a k o1	core	-	X	40,29	-	0,05	0,08	46,96	0,28	0,16	11,63	0,32	100,24	1,73	0,24	88,9
KI59 2a k o2	rim 1a	-	X	40,44	-	0,07	0,06	47,24	0,25	0,21	10,87	0,39	99,37	1,75	0,23	89,6
KI59 2a k o3	rim 1b	-	X	39,97	-	0,07	0,13	48,18	0,25	0,09	10,44	0,38	100,23	1,76	0,21	90,2
KI59 2a m o1	core	-	X	39,96	-	0,05	0,09	47,26	0,25	0,15	11,32	0,28	99,54	1,75	0,24	89,2

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI59 2a m o2	rim 1a	-		40,15	-	0,03	0,10	46,29	0,25	0,22	12,50	0,29	99,95	1,72	0,26	88,0
KI59 2a m o3	~rim 1b	-		40,21	-	0,03	0,09	47,27	0,27	0,09	11,05	0,32	99,51	1,75	0,23	89,4
KI59 2a n o1	~core	-		40,26	-	0,07	0,03	46,62	0,27	0,19	11,78	0,31	99,53	1,73	0,25	88,7
KI59 2a n o2	rim 1a	-		42,91	-	0,06	0,05	46,37	0,29	0,17	12,06	0,31	102,22	1,67	0,24	88,4
KI59 2a n o3	rim 1b	-		40,56	-	0,03	0,09	46,66	0,28	0,25	11,97	0,32	100,15	1,72	0,25	88,5
KI59 2a o o1	~rim 1a	-		40,61	-	0,06	0,10	47,26	0,25	0,11	11,01	0,35	99,75	1,74	0,23	89,5
KI59 2a o o2	core	-		40,27	-	0,06	0,10	47,74	0,24	0,14	10,36	0,37	99,27	1,77	0,22	90,1
KI59 2a o o3	~rim 1b	-		40,25	-	0,06	0,10	47,16	0,26	0,18	11,69	0,33	100,05	1,74	0,24	88,9
KI59 2a p o1	core	-		39,39	-	0,05	0,07	45,66	0,28	0,12	12,54	0,30	98,41	1,72	0,27	87,8
KI59 2a p o2	rim 1a	-		39,64	-	0,05	0,04	45,38	0,29	0,23	12,99	0,30	99,03	1,70	0,27	87,4
KI59 2a p o3	rim 1b	-		40,07	-	0,05	0,16	45,98	0,27	0,15	12,42	0,36	99,56	1,71	0,26	88,0
KI59 3 a o1	~core 1	-	X	39,89	-	0,03	0,05	45,44	0,29	0,22	13,21	0,26	99,38	1,70	0,28	86,0
KI59 3 a o2	~rim 1a	-	X	41,43	-	0,05	0,08	45,64	0,28	0,12	13,15	0,28	101,03	1,67	0,27	86,1
KI59 3 a o3	rim 1b	-	X	39,82	-	0,05	0,05	45,64	0,28	0,14	13,35	0,31	99,64	1,71	0,28	85,9
KI59 3 a o6	~core 3	-	X	39,55	-	0,03	0,06	45,28	0,29	0,12	13,17	0,32	100,57	1,67	0,27	86,0
KI59 3 a o7	core 4	-	X	39,32	-	0,04	0,07	45,33	0,28	0,16	13,07	0,32	99,12	1,70	0,28	86,1
KI59 3 a o8	rim 4a	-	X	40,02	-	0,06	0,07	45,25	0,30	0,17	13,00	0,31	98,75	1,70	0,28	86,1
KI59 3 a o9	rim 4b	-	X	39,21	-	0,05	0,06	44,34	0,26	0,23	14,07	0,32	98,74	1,68	0,30	84,9
KI59 3 e o1	core	-	X	40,56	-	0,06	0,05	44,38	0,28	0,17	14,63	0,24	99,90	1,66	0,31	84,4
KI59 3 e o2	~rim 1a	-	X	40,32	-	0,06	0,05	44,44	0,25	0,20	14,41	0,25	98,97	1,68	0,31	84,6
KI59 3 e o3	rim 1b	-	X	40,55	-	0,03	0,00	45,50	0,28	0,21	13,39	0,28	99,65	1,70	0,28	85,8
KI59 3 k o1	subgrain	-	X	39,77	-	0,04	0,09	45,95	0,26	0,19	12,93	0,35	99,86	1,71	0,27	86,4
KI59 3 k o2	core 1a	-	X	40,10	-	0,04	0,04	45,45	0,29	0,18	13,14	0,32	99,50	1,70	0,28	86,1
KI59 3 k o3	rim 1	-	X	39,61	-	0,04	0,08	46,12	0,29	0,14	13,01	0,33	100,19	1,71	0,27	87,5
KI59 3 k o4	~core 1b	-	X	40,20	-	0,04	0,11	45,62	0,29	0,08	12,91	0,31	99,61	1,70	0,27	87,5
KI59 3 a o4	rim 2a	-		41,27	-	0,04	0,07	45,44	0,27	0,23	13,44	0,27	99,65	1,70	0,28	87,0
KI59 3 a o5	core 2	-		39,83	-	0,06	0,04	45,69	0,29	0,18	13,09	0,29	99,61	1,71	0,27	87,4
KI59 3 b o1	core 1	-		39,96	-	0,05	0,05	45,18	0,28	0,15	13,44	0,22	99,40	1,69	0,28	87,0
KI59 3 b o2	rim 1a	-		40,05	-	0,06	0,11	46,51	0,26	0,14	11,75	0,30	98,88	1,74	0,25	88,7
KI59 3 b o3	~core 2	-		39,98	-	0,04	0,06	46,45	0,28	0,20	12,34	0,31	99,76	1,73	0,26	88,2
KI59 3 b o4	~rim 2	-		40,19	-	0,05	0,05	46,83	0,28	0,11	11,87	0,32	100,19	1,73	0,25	88,7
KI59 3 b o5	~rim 1b	-		40,14	-	0,06	0,10	47,87	0,26	0,19	10,57	0,33	100,08	1,76	0,22	90,0
KI59 3 c o1	core	-		39,77	-	0,07	0,14	47,59	0,24	0,20	10,63	0,38	99,63	1,76	0,22	89,9
KI59 3 c o2	rim 1a	-		39,85	-	0,05	0,07	47,84	0,26	0,18	10,56	0,35	99,85	1,76	0,22	90,0
KI59 3 c o3	~rim 1b	-		39,88	-	0,04	0,05	44,91	0,27	0,20	14,11	0,29	99,56	1,69	0,30	86,3
KI59 3 d o1	core 1	-		39,74	-	0,04	0,01	44,89	0,29	0,20	13,92	0,25	99,38	1,69	0,29	86,5
KI59 3 d o2	rim 1a	-		40,09	-	0,03	0,08	44,85	0,28	0,23	14,00	0,29	99,63	1,68	0,29	86,4
KI59 3 d o4	core 2	-		40,64	-	0,05	0,06	46,93	0,27	0,16	11,89	0,29	100,01	1,73	0,25	88,7
KI59 3 f o1	core	-		39,70	-	0,04	0,03	46,21	0,26	0,17	12,17	0,35	99,02	1,73	0,26	88,3

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI59 3 f o2	~rim 1a	-		39,63	-	0,05	0,10	46,44	0,26	0,15	11,88	0,37	100,29	1,71	0,25	88,6
KI59 3 f o3	rim 1b	-		39,86	-	0,03	0,00	46,75	0,28	0,14	11,80	0,36	99,87	1,73	0,25	88,7
KI59 3 g o1	core 1	-		40,36	-	0,04	0,01	46,84	0,26	0,14	11,85	0,31	99,79	1,74	0,25	88,7
KI59 3 g o2	core 2	-		39,80	-	0,06	0,12	46,32	0,28	0,19	12,44	0,33	99,90	1,72	0,26	88,1
KI59 3 g o3	rim 1	-		41,04	-	0,06	0,07	47,30	0,25	0,10	11,40	0,37	99,68	1,75	0,24	89,1
KI59 3 g o4	rim 3	-		40,51	-	0,06	0,09	47,33	0,27	0,14	11,39	0,35	100,50	1,74	0,23	89,2
KI59 3 g o5	core 3	-		40,22	-	0,04	0,07	45,46	0,28	0,17	13,25	0,33	99,52	1,70	0,28	87,2
KI59 3 h o2	rim 1a	-		40,09	-	0,04	0,06	45,29	0,29	0,13	13,33	0,27	99,39	1,69	0,28	87,1
KI59 3 h o3	core 1b	-		40,04	-	0,09	0,03	45,50	0,27	0,19	13,28	0,35	99,57	1,70	0,28	87,2
KI59 3 h o4	~rim 1b	-		40,83	-	0,06	0,03	45,60	0,27	0,22	13,17	0,33	100,50	1,68	0,27	87,3
KI59 3 n o1	core 1	-		39,80	-	0,05	0,09	47,05	0,26	0,06	11,27	0,37	99,81	1,74	0,23	89,2
KI59 3 n o2	rim 1	-		39,96	-	0,07	0,01	46,21	0,26	0,16	12,17	0,34	99,46	1,72	0,25	88,3
KI59 3 n o3	core 2a	-		39,76	-	0,05	0,08	46,38	0,25	0,10	12,51	0,30	99,72	1,72	0,26	88,0
KI59 3 n o4	~core 2b	-		40,83	-	0,04	0,10	45,87	0,27	0,14	12,61	0,34	99,22	1,72	0,26	87,8
KI59 3 i o1	core	-		40,58	-	0,05	0,03	46,17	0,27	0,14	12,89	0,30	99,85	1,72	0,27	87,6
KI59 3 i o2	rim 1a	-		40,11	-	0,05	0,13	46,40	0,26	0,20	12,51	0,32	100,37	1,72	0,26	88,0
KI59 3 i o3	rim 1b	-		40,06	-	0,07	0,14	45,73	0,27	0,14	13,00	0,35	99,49	1,71	0,27	87,4
KI59 3 j o1	core	-		39,85	-	0,05	0,08	46,25	0,28	0,21	12,52	0,35	100,01	1,72	0,26	88,0
KI59 3 j o2	rim 1a	-		40,01	-	0,05	0,09	45,61	0,28	0,18	13,02	0,33	99,16	1,71	0,27	87,4
KI59 3 j o3	~rim 1b	-		40,31	-	0,06	0,00	46,10	0,28	0,25	12,77	0,31	100,01	1,71	0,27	87,7
KI59 3 l o1	core 1	-		39,67	-	0,04	0,07	46,49	0,31	0,14	12,03	0,35	99,15	1,74	0,25	88,5
KI59 3 l o2	core 2	-		39,87	-	0,05	0,07	46,51	0,31	0,16	11,89	0,35	99,20	1,73	0,25	88,6
KI59 3 l o3	rim 2	-		40,04	-	0,04	0,03	46,37	0,27	0,16	12,17	0,36	99,43	1,73	0,25	88,3
KI59 3 l o4	rim 1	-		39,71	-	0,04	0,00	45,68	0,28	0,17	12,63	0,34	98,90	1,71	0,27	87,7
KI59 3 m o1	core	-		40,14	-	0,05	0,03	46,62	0,25	0,11	11,88	0,38	99,46	1,73	0,25	88,6
KI59 3 m o2	rim 1a	-		39,85	-	0,07	0,04	46,11	0,24	0,21	12,15	0,37	99,05	1,72	0,26	88,3
KI59 3 m o3	rim 1b	-		39,09	-	0,06	0,06	45,18	0,27	0,16	13,22	0,32	98,35	1,71	0,28	87,1

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	FeO
Margin of lake																
KI67-1-5-b1	core	1,5	X	39,22	0,01	0,10	0,14	47,65	0,26	0,22	10,98	0,46	99,07	1,78	0,23	89,6
KI67-1-5-b2	rim	1,5	X	38,43	0,01	0,09	0,08	43,27	0,25	0,25	16,62	0,39	99,45	1,65	0,36	83,7
KI67-1-5-d1	core	1,5	X	39,06	0,00	0,03	0,08	45,90	0,24	0,14	13,34	0,38	99,18	1,73	0,28	87,2
KI67-1-5-d2	rim	1,5	X	39,11	0,03	0,27	0,03	44,13	0,28	0,22	15,94	0,38	100,42	1,66	0,34	84,6
KI67-1-5-f1-1	core 1	1,5	X	39,43	0,01	0,00	0,07	47,17	0,26	0,14	12,08	0,37	99,54	1,76	0,25	88,5
KI67-1-5-f1-2	rim 1	1,5	X	39,53	0,00	0,03	0,04	47,17	0,25	0,21	11,85	0,35	99,46	1,76	0,25	88,8
KI67-1-5-a1	core	1,5		38,64	0,03	0,29	0,09	43,51	0,34	0,19	16,23	0,26	99,59	1,65	0,35	84,1
KI67-1-5-a2	rim	1,5		37,87	0,02	0,39	0,04	39,33	0,29	0,30	21,29	0,17	99,73	1,53	0,46	78,5
KI67-1-5-c1	core	1,5		39,52	0,00	0,00	0,08	45,93	0,33	0,11	13,12	0,41	99,52	1,72	0,28	87,4
KI67-1-5-c2	rim	1,5		37,67	0,00	0,09	0,00	38,10	0,29	0,27	22,91	0,24	99,59	1,49	0,50	76,7
KI67-1-5-e1	core	1,5		39,30	0,01	0,43	0,02	45,83	0,27	0,27	13,50	0,33	99,99	1,71	0,28	87,0
KI67-1-5-e2	rim	1,5		38,36	0,02	0,05	0,05	42,22	0,28	0,21	18,53	0,27	100,01	1,61	0,40	81,9
KI67-1-5-f2-1	core 2	1,5		39,84	0,00	0,08	0,09	47,38	0,26	0,19	11,86	0,41	100,14	1,75	0,25	88,8
KI67-1-5-f2-2	rim 2	1,5		38,92	0,02	0,03	0,04	45,23	0,26	0,23	14,97	0,35	100,06	1,70	0,32	85,7
KI67-1-5-g1-1	core 1	1,5		39,52	0,00	0,52	0,04	47,20	0,26	0,08	11,67	0,35	99,69	1,75	0,24	88,9
KI67-1-5-g1-2	rim 1	1,5		39,52	0,00	0,20	0,01	46,69	0,27	0,12	12,63	0,39	99,84	1,74	0,26	88,0
KI67-1-5-g2-1	core 2	1,5		39,93	0,00	0,26	0,11	47,19	0,27	0,13	11,92	0,38	100,20	1,74	0,25	88,7
KI67-1-5-g2-2	rim 2	1,5		39,85	0,00	0,00	0,07	46,83	0,26	0,11	12,35	0,41	99,90	1,74	0,26	88,3
KI67-1-5-h1	core	1,5		39,26	0,01	0,00	0,07	46,58	0,27	0,16	12,41	0,38	99,15	1,75	0,26	88,1
KI67-1-5-h2	rim	1,5		38,09	0,01	0,18	0,02	40,30	0,29	0,30	20,75	0,26	100,24	1,55	0,45	79,4
KI67-1-5-i1	core	1,5		39,36	0,03	0,40	0,09	44,70	0,28	0,21	14,77	0,30	100,16	1,67	0,31	85,7
KI67-1-5-i2	rim	1,5		38,54	0,00	0,30	0,07	41,27	0,27	0,16	18,87	0,34	99,87	1,58	0,41	81,3
KI67-1-5-j1	core	1,5		39,88	0,01	0,13	0,09	46,64	0,29	0,14	12,34	0,34	99,87	1,73	0,26	88,2
KI67-1-5-j2	rim	1,5		37,80	0,01	0,23	0,00	39,66	0,27	0,32	21,38	0,25	99,92	1,54	0,47	78,6
KI67-1-5-k1	core	1,5		39,47	0,00	0,06	0,06	46,63	0,27	0,22	12,13	0,40	99,25	1,74	0,25	88,4
KI67-1-5-k2	rim	1,5		37,61	0,01	0,08	0,03	38,48	0,27	0,40	22,57	0,30	99,74	1,51	0,50	77,2
KI67-1-20-a1	core	6,1	X	39,17	0,00	0,09	0,05	47,02	0,28	0,19	12,40	0,33	99,53	1,76	0,26	88,2
KI67-1-20-a2	rim	6,1	X	37,29	0,01	0,00	0,05	37,65	0,24	0,31	24,41	0,26	100,24	1,48	0,54	75,4
KI67-1-20-c1	core	6,1	X	40,01	0,02	0,05	0,18	47,21	0,27	0,16	12,23	0,36	100,49	1,74	0,25	88,4
KI67-1-20-c2	rim	6,1	X	38,15	0,00	0,18	0,03	39,62	0,22	0,31	21,42	0,32	100,27	1,53	0,46	78,6
KI67-1-20-e1	core	6,1	X	39,24	0,03	0,09	0,06	46,42	0,28	0,19	12,83	0,30	99,47	1,74	0,27	87,8
KI67-1-20-e2	rim	6,1	X	37,39	0,00	0,06	0,00	37,68	0,22	0,22	24,27	0,20	100,09	1,48	0,53	75,5
KI67-1-20-i1	core	6,1	X	39,71	0,00	0,04	0,04	47,38	0,23	0,17	12,20	0,41	100,18	1,75	0,25	88,5
KI67-1-20-i2	rim	6,1	X	37,73	0,02	0,09	0,02	39,81	0,22	0,34	21,15	0,31	99,73	1,55	0,46	78,8
KI67-1-20-j2-1	core	6,1	X	38,85	0,01	0,04	0,00	43,49	0,25	0,19	16,63	0,37	99,85	1,65	0,35	83,8
KI67-1-20-j2-2	rim	6,1	X	37,30	0,00	0,07	0,07	38,12	0,23	0,37	23,24	0,31	99,72	1,50	0,51	76,5
KI67-1-20-m1	core	6,1	X	39,60	0,01	0,13	0,02	45,66	0,29	0,23	13,73	0,39	100,09	1,70	0,29	86,8
KI67-1-20-m2	rim	6,1	X	37,73	0,01	0,56	0,10	38,89	0,24	0,36	21,85	0,24	99,98	1,51	0,48	77,9
KI67-1-20-n1	core	6,1	X	39,36	0,00	0,00	0,04	46,48	0,26	0,23	12,64	0,38	99,40	1,74	0,27	87,9

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI67-1-20-n2	rim	6,1	X	37,10	0,02	0,03	0,03	35,97	0,23	0,35	25,74	0,26	99,78	1,43	0,57	73,5
KI67-1-20-b1	core	6,1		39,75	0,00	0,18	0,12	46,55	0,25	0,17	12,72	0,45	100,19	1,73	0,27	87,9
KI67-1-20-b2	rim	6,1		38,26	0,02	0,17	0,01	38,58	0,22	0,26	22,57	0,27	100,37	1,49	0,49	77,2
KI67-1-20-d1	core	6,1		38,35	0,00	0,09	0,04	40,85	0,25	0,25	20,10	0,30	100,24	1,57	0,43	80,1
KI67-1-20-d2	rim	6,1		36,79	0,02	0,05	0,00	35,05	0,21	0,34	27,40	0,27	100,16	1,40	0,61	71,7
KI67-1-20-f1	core	6,1		38,96	0,01	0,16	0,11	45,90	0,27	0,12	13,55	0,42	99,54	1,72	0,29	87,0
KI67-1-20-f2	rim	6,1		37,41	0,00	0,02	0,00	37,63	0,19	0,40	24,18	0,25	100,08	1,48	0,53	75,5
KI67-1-20-g1	core	6,1		39,19	0,01	0,03	0,03	44,84	0,27	0,27	14,91	0,35	99,90	1,69	0,31	85,6
KI67-1-20-g2	rim	6,1		38,17	0,01	0,06	0,00	39,67	0,19	0,32	21,31	0,26	100,04	1,54	0,46	78,7
KI67-1-20-h1	core	6,1		38,47	0,00	0,41	0,07	41,24	0,27	0,26	19,24	0,35	100,33	1,58	0,41	80,9
KI67-1-20-h2	rim	6,1		37,94	0,00	0,00	0,00	38,27	0,23	0,30	23,54	0,27	100,59	1,49	0,51	76,3
KI67-1-20-j1-1	core	6,1		39,66	0,00	0,22	0,11	46,75	0,27	0,17	12,78	0,34	100,30	1,73	0,27	87,9
KI67-1-20-j1-2	rim	6,1		37,41	0,01	0,09	0,00	37,36	0,25	0,37	24,35	0,27	100,11	1,47	0,54	75,3
KI67-1-20-k1	core	6,1		39,45	0,01	0,08	0,03	45,41	0,26	0,22	14,11	0,33	99,93	1,70	0,30	86,4
KI67-1-20-k2	rim	6,1		37,53	0,01	0,04	0,00	37,79	0,20	0,42	24,06	0,21	100,27	1,48	0,53	75,7
KI67-1-20-l2-1	core	6,1		39,10	0,00	0,08	0,09	44,68	0,30	0,10	15,33	0,42	100,10	1,68	0,32	85,2
KI67-1-35-a1	core	10,7	X	39,59	0,01	0,14	0,08	46,94	0,24	0,23	12,10	0,41	99,78	1,75	0,25	88,5
KI67-1-35-a2	rim	10,7	X	37,28	0,01	0,18	0,04	37,33	0,21	0,27	24,83	0,29	100,49	1,46	0,55	74,9
KI67-1-35-b1-1	core 1	10,7	X	39,55	0,00	0,09	0,03	47,07	0,25	0,12	12,24	0,44	99,84	1,75	0,26	88,4
KI67-1-35-b1-2	rim 1	10,7	X	37,71	0,00	0,09	0,01	38,17	0,20	0,29	23,27	0,27	100,03	1,49	0,51	76,5
KI67-1-35-b2-1	core 2	10,7	X	39,70	0,00	0,21	0,10	47,07	0,24	0,16	12,16	0,44	100,10	1,74	0,25	88,4
KI67-1-35-b2-2	rim 2	10,7	X	37,23	0,02	0,06	0,03	37,36	0,21	0,31	24,87	0,22	100,32	1,47	0,55	74,9
KI67-1-35-b3-1	core 3	10,7	X	39,54	0,01	0,00	0,00	45,96	0,24	0,25	13,82	0,39	100,24	1,71	0,29	86,8
KI67-1-35-e1	core	10,7	X	39,35	0,00	0,08	0,04	46,44	0,28	0,20	13,23	0,39	100,03	1,73	0,28	87,4
KI67-1-35-e2	rim	10,7	X	37,38	0,02	0,00	0,00	37,25	0,21	0,40	24,81	0,26	100,34	1,46	0,55	74,9
KI67-1-35-i1	core	10,7	X	40,19	0,00	0,00	0,00	48,61	0,24	0,12	10,38	0,49	100,07	1,79	0,21	90,3
KI67-1-35-i2	rim	10,7	X	37,74	0,04	0,02	0,00	38,43	0,17	0,31	23,52	0,24	100,46	1,50	0,51	76,4
KI67-1-35-j1-1	core 1	10,7	X	39,67	0,02	0,27	0,03	47,73	0,25	0,07	11,45	0,43	99,92	1,77	0,24	89,2
KI67-1-35-j1-2	rim 1	10,7	X	37,99	0,00	0,02	0,02	39,30	0,23	0,27	22,40	0,28	100,52	1,52	0,49	77,7
KI67-1-35-j2-1	core 2	10,7	X	39,99	0,00	0,02	0,07	47,80	0,22	0,24	11,33	0,46	100,12	1,76	0,23	89,3
KI67-1-35-j2-2	rim 2	10,7	X	37,91	0,04	0,08	0,05	39,03	0,21	0,35	22,15	0,29	100,13	1,52	0,48	77,7
KI67-1-35-m1	core	10,7	X	37,72	0,00	0,00	0,00	40,73	0,26	0,29	20,42	0,27	99,74	1,58	0,44	79,8
KI67-1-35-m2	rim	10,7	X	36,85	0,02	0,14	0,02	35,39	0,22	0,41	26,98	0,19	100,24	1,41	0,60	72,2
KI67-1-35-c1	core	10,7		37,83	0,00	0,29	0,07	38,64	0,23	0,21	22,39	0,28	99,94	1,50	0,49	77,4
KI67-1-35-c2	rim	10,7		36,95	0,01	0,05	0,01	35,37	0,20	0,37	27,09	0,25	100,36	1,40	0,60	72,1
KI67-1-35-d1	core	10,7		38,85	0,00	0,04	0,06	43,88	0,26	0,21	16,20	0,32	99,86	1,66	0,34	84,3
KI67-1-35-d2	rim	10,7		36,93	0,03	0,17	0,00	36,97	0,21	0,36	24,91	0,26	99,88	1,46	0,55	74,6
KI67-1-35-g1	core	10,7		36,84	0,02	0,06	0,01	35,45	0,23	0,34	26,64	0,28	99,90	1,41	0,60	72,5
KI67-1-35-g2	rim	10,7		36,54	0,02	0,14	0,09	35,02	0,19	0,51	27,05	0,33	99,88	1,40	0,61	72,0

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI67-1-35-h1	core	10,7		37,07	0,01	0,13	0,00	35,22	0,22	0,28	26,98	0,21	100,16	1,40	0,60	72,1
KI67-1-35-h2	rim	10,7		37,00	0,02	0,46	0,04	35,15	0,16	0,39	27,31	0,23	100,78	1,39	0,61	71,8
KI67-1-35-k1	core	10,7		38,97	0,01	0,00	0,03	43,30	0,28	0,29	17,14	0,31	100,35	1,64	0,36	83,3
KI67-1-35-k2	rim	10,7		37,71	0,03	0,02	0,00	37,15	0,17	0,24	24,40	0,29	100,02	1,46	0,54	75,1
KI67-1-35-l1	core	10,7		38,83	0,02	0,14	0,09	42,35	0,26	0,24	18,16	0,33	100,42	1,61	0,39	82,2
KI67-1-35-l2	rim	10,7		37,11	0,02	0,06	0,00	35,79	0,22	0,35	26,35	0,24	100,16	1,42	0,59	72,9
KI67-1-35-n1	core	10,7		37,66	0,00	0,17	0,07	38,71	0,25	0,34	22,91	0,30	100,41	1,51	0,50	77,0
KI67-1-35-n2	rim	10,7		36,83	0,01	0,00	0,00	35,16	0,20	0,34	26,21	0,22	98,98	1,41	0,59	72,6
KI67-1-35-o1	core	10,7		37,33	0,00	0,00	0,00	36,33	0,23	0,30	25,98	0,22	100,42	1,43	0,57	73,5
KI67-1-43-b1	~core	13,1	X	38,98	0,03	0,00	0,04	43,27	0,27	0,22	17,08	0,34	100,23	1,64	0,36	83,4
KI67-1-43-b2	rim	13,1	X	37,43	0,04	0,01	0,00	38,55	0,22	0,34	23,19	0,31	100,09	1,51	0,51	76,7
KI67-1-43-a1	core	13,1		39,97	0,02	0,14	0,02	45,64	0,23	0,22	13,61	0,29	100,16	1,70	0,28	86,9
KI67-1-43-a2	rim	13,1		38,07	0,01	0,24	0,02	38,28	0,21	0,35	22,86	0,29	100,33	1,49	0,50	76,8
KI67-1-43-d1-1	core 1	13,1		39,52	0,00	1,36	0,05	46,07	0,25	0,25	13,62	0,36	101,51	1,69	0,28	87,0
KI67-1-43-d1-2	rim 1	13,1		37,59	0,02	0,30	0,01	38,04	0,21	0,39	23,53	0,24	100,38	1,48	0,52	76,2
KI67-1-43-d2-1	core 2	13,1		39,70	0,03	0,12	0,08	46,82	0,26	0,21	12,74	0,35	100,31	1,74	0,27	87,9
KI67-1-43-d2-2	rim 2	13,1		37,33	0,05	0,14	0,00	37,77	0,13	0,31	23,87	0,23	99,83	1,48	0,53	75,8
KI67-1-43-e1	core	13,1		37,57	0,01	0,10	0,01	38,14	0,22	0,33	23,75	0,25	100,41	1,49	0,52	76,1
KI67-1-43-e2	rim	13,1		37,51	0,03	0,13	0,01	37,85	0,19	0,39	23,70	0,23	100,06	1,48	0,52	76,0
KI67-1-43-f1	core	13,1		37,60	0,01	0,56	0,01	38,08	0,22	0,31	23,68	0,22	100,72	1,48	0,52	76,1
KI67-1-43-f2	rim	13,1		37,01	0,05	0,13	0,00	37,32	0,18	0,38	23,89	0,26	99,24	1,48	0,53	75,6
KI67-1-43-g1	core	13,1		38,11	0,01	0,00	0,00	39,61	0,22	0,26	22,10	0,30	100,64	1,53	0,48	78,0
KI67-1-43-g2	rim	13,1		37,60	0,05	0,12	0,02	38,77	0,18	0,37	22,95	0,23	100,30	1,51	0,50	77,0
KI67-1-43-h1	core	13,1		37,20	0,01	0,06	0,01	37,99	0,20	0,25	23,69	0,26	99,69	1,49	0,52	76,0
KI67-1-43-h2	rim	13,1		37,33	0,01	0,09	0,00	37,15	0,21	0,33	24,27	0,29	99,70	1,47	0,54	75,2
KI67-1-43-i1	core	13,1		39,26	0,00	0,12	0,05	46,05	0,28	0,17	13,34	0,36	99,66	1,72	0,28	87,2
KI67-1-43-i2	rim	13,1		37,43	0,02	0,21	0,02	38,01	0,23	0,37	23,76	0,17	100,21	1,49	0,52	76,0
KI67-1-43-j1	core	13,1		38,58	0,02	0,16	0,07	40,97	0,22	0,20	19,77	0,34	100,35	1,57	0,42	80,4
KI67-1-43-j2	rim	13,1		37,63	0,03	0,00	0,00	37,80	0,21	0,30	24,17	0,20	100,37	1,48	0,53	75,6
KI67-1-43-k1	~core	13,1		37,35	0,01	0,61	0,00	37,81	0,23	0,31	23,82	0,19	100,33	1,48	0,52	75,9
KI67-1-43-k2	rim	13,1		37,16	0,01	0,61	0,09	37,11	0,20	0,33	24,40	0,32	100,25	1,46	0,54	75,1
KI67-1-43-l1	core	13,1		37,97	0,01	0,13	0,07	39,41	0,21	0,28	21,54	0,29	99,95	1,53	0,47	78,4
KI67-1-43-l2	rim	13,1		37,70	0,03	0,29	0,00	37,44	0,18	0,31	23,34	0,22	99,58	1,47	0,51	76,1
KI67-1-43-m1	core	13,1		39,02	0,01	0,27	0,01	44,64	0,23	0,21	14,96	0,35	99,69	1,68	0,32	85,5
KI67-1-43-m2	rim	13,1		37,13	0,04	0,12	0,01	37,75	0,19	0,31	23,75	0,27	99,58	1,49	0,53	75,9
KI67-1-43-n1	core	13,1		37,73	0,01	1,38	0,04	38,53	0,24	0,30	22,29	0,30	100,85	1,48	0,48	77,4
KI67-1-43-n2	rim	13,1		37,15	0,05	0,39	0,00	37,17	0,16	0,37	24,31	0,26	99,87	1,46	0,54	75,2
KI67-1-60-a1	core	18,4	X	39,91	0,02	0,03	0,01	46,56	0,15	0,16	12,49	0,34	99,65	1,73	0,26	88,1
KI67-1-60-a2	rim	18,4	X	38,27	0,02	0,05	0,00	41,13	0,20	0,24	19,50	0,23	99,67	1,58	0,42	80,7

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI67-1-60-h1	core	18,4	X	39,93	0,01	0,07	0,05	46,67	0,23	0,15	12,73	0,33	100,19	1,73	0,27	87,9
KI67-1-60-h2	rim	18,4	X	38,56	0,01	0,00	0,02	42,81	0,17	0,32	17,63	0,34	99,91	1,63	0,38	82,8
KI67-1-60-k1	core	18,4	X	39,05	0,00	0,04	0,07	46,53	0,17	0,22	13,39	0,41	99,88	1,74	0,28	87,3
KI67-1-60-k2	rim	18,4	X	38,73	0,02	0,05	0,03	43,45	0,19	0,37	16,92	0,32	100,08	1,65	0,36	83,6
KI67-1-60-b1	core	18,4		38,73	0,02	0,07	0,02	44,53	0,18	0,20	15,31	0,28	99,37	1,69	0,33	85,2
KI67-1-60-b2	rim	18,4		38,41	0,02	0,06	0,07	41,71	0,14	0,24	18,60	0,31	99,57	1,60	0,40	81,6
KI67-1-60-c1	core	18,4		38,16	0,00	0,04	0,03	40,94	0,13	0,36	19,46	0,23	99,37	1,58	0,42	80,6
KI67-1-60-c2	rim	18,4		38,24	0,02	0,03	0,02	40,60	0,17	0,32	19,91	0,21	99,51	1,57	0,43	80,2
KI67-1-60-d1	core	18,4		38,56	0,00	0,07	0,09	42,94	0,12	0,33	17,79	0,33	100,23	1,63	0,38	82,7
KI67-1-60-d2	rim	18,4		38,73	0,00	0,06	0,06	42,65	0,16	0,34	17,55	0,35	99,91	1,62	0,38	82,8
KI67-1-60-e1	core	18,4		38,60	0,00	0,07	0,00	43,38	0,11	0,31	16,86	0,42	99,77	1,65	0,36	83,6
KI67-1-60-e2	rim	18,4		38,81	0,02	0,04	0,02	42,70	0,16	0,26	17,38	0,33	99,71	1,63	0,37	83,0
KI67-1-60-f1	core	18,4		39,21	0,01	0,06	0,11	44,05	0,12	0,18	15,96	0,37	100,06	1,66	0,34	84,6
KI67-1-60-f2	rim	18,4		38,19	0,03	0,04	0,01	41,33	0,17	0,40	19,15	0,29	99,65	1,59	0,41	81,1
KI67-1-60-g1	core	18,4		39,21	0,02	0,04	0,04	44,11	0,13	0,15	15,77	0,36	99,83	1,66	0,33	84,7
KI67-1-60-g2	rim	18,4		38,73	0,02	0,02	0,08	42,31	0,15	0,29	18,04	0,32	100,02	1,61	0,39	82,3
KI67-1-60-i1	core	18,4		38,42	0,00	0,25	0,02	42,90	0,10	0,32	17,57	0,34	99,94	1,63	0,38	82,9
KI67-1-60-i2	rim	18,4		38,34	0,05	0,02	0,00	41,84	0,15	0,40	18,68	0,25	99,75	1,61	0,40	81,6
KI67-1-60-j1	core	18,4		39,12	0,02	0,06	0,02	43,94	0,13	0,21	15,83	0,36	99,70	1,66	0,34	84,6
KI67-1-60-j2	rim	18,4		38,40	0,00	0,03	0,07	41,57	0,15	0,41	18,71	0,29	99,70	1,60	0,40	81,5
KI67-1-60-l1	core	18,4		39,59	0,00	0,02	0,04	45,67	0,15	0,18	14,02	0,32	99,99	1,71	0,29	86,6
KI67-1-60-l2	rim	18,4		38,49	0,03	0,04	0,02	41,18	0,15	0,34	19,17	0,29	99,75	1,58	0,41	81,0
KI67-1-60-m1	core	18,4		38,22	0,00	0,03	0,00	41,40	0,12	0,30	20,00	0,27	100,37	1,59	0,43	80,4
KI67-1-60-m2	rim	18,4		37,99	0,01	0,05	0,07	40,93	0,15	0,32	19,68	0,27	99,51	1,58	0,43	80,4
KI67-1-77-a1	core	23,5	X	39,41	0,00	0,05	0,00	46,56	0,22	0,17	12,63	0,36	99,42	1,74	0,27	87,9
KI67-1-77-a2	rim	23,5	X	37,60	0,01	0,02	0,06	38,72	0,19	0,22	22,39	0,28	99,51	1,52	0,49	77,4
KI67-1-77-f1	core	23,5	X	38,83	0,00	0,06	0,04	42,24	0,25	0,25	17,82	0,27	99,78	1,61	0,38	82,4
KI67-1-77-f2	rim	23,5	X	37,60	0,03	0,04	0,04	38,44	0,19	0,34	23,09	0,23	100,03	1,50	0,51	76,7
KI67-1-77-l1	core	23,5	X	37,69	0,01	0,06	0,00	38,83	0,22	0,25	22,62	0,30	100,02	1,51	0,50	77,3
KI67-1-77-l2	rim	23,5	X	37,57	0,02	0,04	0,01	37,58	0,19	0,30	24,04	0,27	100,07	1,47	0,53	75,6
KI67-1-77-b1	core	23,5		37,54	0,00	0,04	0,00	38,83	0,21	0,31	22,50	0,21	99,64	1,52	0,49	77,3
KI67-1-77-b2	rim	23,5		37,09	0,03	0,04	0,00	37,35	0,18	0,46	24,57	0,23	100,00	1,47	0,54	75,1
KI67-1-77-c1	core	23,5		37,22	0,02	0,02	0,00	37,13	0,19	0,47	24,60	0,21	99,88	1,46	0,54	74,9
KI67-1-77-c2	rim	23,5		37,13	0,03	0,04	0,00	36,86	0,20	0,43	24,87	0,18	99,78	1,46	0,55	74,6
KI67-1-77-e1	core	23,5		37,70	0,00	0,07	0,00	39,53	0,21	0,32	21,65	0,23	99,71	1,54	0,47	78,3
KI67-1-77-e2	rim	23,5		37,34	0,02	0,02	0,02	37,87	0,21	0,34	23,74	0,29	99,85	1,49	0,52	76,0
KI67-1-77-g1	core	23,5		37,59	0,00	0,03	0,00	39,74	0,21	0,29	21,61	0,25	99,73	1,55	0,47	78,4
KI67-1-77-g2	rim	23,5		37,51	0,04	0,03	0,00	37,88	0,17	0,32	23,91	0,21	100,10	1,48	0,53	75,8
KI67-1-77-h1	core	23,5		37,62	0,02	0,04	0,06	40,46	0,19	0,34	20,40	0,30	99,44	1,57	0,44	79,7

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI67-1-77-h2	rim	23,5		36,77	0,03	0,07	0,07	37,65	0,19	0,39	24,17	0,22	99,56	1,49	0,54	75,5
KI67-1-77-i1	core	23,5		37,27	0,00	0,05	0,02	37,74	0,22	0,34	23,90	0,23	99,79	1,48	0,53	75,8
KI67-1-77-i2	rim	23,5		37,22	0,03	0,05	0,00	37,14	0,17	0,37	24,60	0,23	99,85	1,47	0,54	75,0
KI67-1-77-j1	core	23,5		37,24	0,00	0,08	0,02	38,09	0,22	0,32	23,75	0,26	99,98	1,49	0,52	76,0
KI67-1-77-j2	rim	23,5		37,38	0,02	0,02	0,00	37,75	0,19	0,22	24,05	0,26	99,89	1,48	0,53	75,7
KI67-1-77-k1	core	23,5		37,47	0,00	0,04	0,02	38,93	0,20	0,33	22,87	0,27	100,12	1,52	0,50	77,1
KI67-1-77-k2	rim	23,5		37,55	0,02	0,03	0,05	38,20	0,18	0,32	23,45	0,24	100,07	1,49	0,51	76,4
KI67-1-77-m1	core	23,5		37,63	0,02	0,06	0,00	39,96	0,22	0,31	21,03	0,25	99,51	1,56	0,46	79,0
KI67-1-77-m2	rim	23,5		37,15	0,03	0,06	0,00	37,95	0,19	0,32	23,93	0,26	99,91	1,49	0,53	75,8
KI67-1-77-n1	core	23,5		37,62	0,00	0,04	0,00	38,91	0,21	0,31	22,69	0,24	100,03	1,52	0,50	77,3
KI67-1-77-n2	rim	23,5		37,04	0,03	0,06	0,03	37,89	0,20	0,39	24,26	0,25	100,16	1,49	0,54	75,6
KI75-3-5-b1	core	1,5	X	39,75	0,02	0,04	0,03	44,23	0,29	0,23	15,38	0,33	100,38	1,66	0,32	85,1
KI75-3-5-b2	rim	1,5	X	36,46	0,07	0,02	0,03	31,18	0,30	0,34	31,15	0,14	99,77	1,27	0,71	66,5
KI75-3-5-a1	core	1,5		39,67	0,00	0,03	0,02	44,70	0,28	0,20	14,77	0,33	100,00	1,68	0,31	85,7
KI75-3-5-a2	rim	1,5		38,44	0,01	0,03	0,01	38,95	0,30	0,34	21,35	0,19	99,69	1,51	0,47	78,3
KI75-3-5-c1	core	1,5		39,77	0,00	0,03	0,06	45,77	0,27	0,21	12,68	0,37	99,19	1,71	0,27	87,7
KI75-3-5-c2	rim	1,5		37,46	0,01	0,02	0,01	37,36	0,30	0,31	23,79	0,21	99,60	1,47	0,53	75,7
KI75-3-5-d1	core	1,5		39,71	0,00	0,03	0,10	44,72	0,27	0,22	13,19	0,33	98,61	1,69	0,28	87,0
KI75-3-5-d2	rim	1,5		36,60	0,01	0,02	0,00	32,51	0,27	0,32	28,92	0,24	98,96	1,32	0,66	69,0
KI75-3-5-e1	core	1,5		39,78	0,00	0,02	0,04	44,95	0,28	0,24	13,91	0,33	99,57	1,69	0,29	86,5
KI75-3-5-e2	rim	1,5		35,79	0,09	0,01	0,00	27,15	0,22	0,51	36,15	0,18	100,21	1,13	0,84	59,8
KI75-3-5-f1	core	1,5		39,58	0,02	0,05	0,08	45,71	0,25	0,03	12,79	0,40	98,93	1,72	0,27	87,6
KI75-3-5-f2	rim	1,5		37,36	0,02	0,04	0,03	36,21	0,32	0,35	24,96	0,19	99,57	1,43	0,56	74,2
KI75-3-5-g1	core	1,5		39,78	0,00	0,04	0,06	44,72	0,29	0,15	13,59	0,38	99,09	1,68	0,29	86,7
KI75-3-5-g2	rim	1,5		37,71	0,02	0,03	0,00	37,38	0,30	0,41	22,79	0,22	98,87	1,48	0,50	76,5
KI75-3-5-h1-1c	core a	1,5		39,44	0,00	0,04	0,09	43,43	0,30	0,17	15,79	0,37	99,64	1,64	0,34	84,5
KI75-3-5-h1-2c	core b	1,5		39,62	0,00	0,04	0,00	43,70	0,32	0,21	15,57	0,29	99,76	1,65	0,33	84,8
KI75-3-5-h1-3b	rim	1,5		37,44	0,09	0,03	0,00	33,89	0,30	0,35	27,88	0,24	100,25	1,35	0,62	70,7
KI75-3-5-h2-1c	core	1,5		39,19	0,01	0,02	0,07	43,69	0,30	0,21	14,95	0,29	98,80	1,66	0,32	85,3
KI75-3-5-i1	core	1,5		40,02	0,00	0,04	0,00	45,33	0,25	0,18	12,41	0,36	98,61	1,70	0,26	87,9
KI75-3-5-i2	rim	1,5		36,11	0,09	0,01	0,00	29,93	0,29	0,46	32,58	0,18	99,68	1,23	0,75	64,5
KI75-3-5-j1	core	1,5		39,85	0,00	0,02	0,10	46,28	0,25	0,11	11,38	0,46	98,50	1,73	0,24	89,0
KI75-3-5-j2	rim	1,5		36,80	0,03	0,02	0,08	33,15	0,31	0,45	28,35	0,22	99,50	1,34	0,64	69,9
KI75-3-5-k1	core	1,5		39,86	0,00	0,02	0,00	45,56	0,29	0,26	13,62	0,28	99,97	1,70	0,29	86,9
KI75-3-5-k2	rim	1,5		37,30	0,04	0,02	0,00	34,68	0,30	0,35	26,83	0,25	99,81	1,38	0,60	71,9
KI75-3-5-l1-1	core	1,5		39,73	0,00	0,02	0,05	45,60	0,29	0,24	13,50	0,36	99,82	1,70	0,28	87,0
KI75-3-5-l1-2	rim	1,5		36,17	0,04	0,01	0,00	27,63	0,27	0,46	33,95	0,11	98,69	1,15	0,80	61,7
KI75-3-5-l2-1	core	1,5		40,03	0,00	0,04	0,08	45,47	0,28	0,13	12,85	0,37	99,26	1,70	0,27	87,5
KI75-3-5-m1	core	1,5		39,88	0,00	0,02	0,07	45,95	0,29	0,16	12,47	0,38	99,29	1,72	0,26	88,0
KI75-3-5-m2	rim	1,5		38,06	0,02	0,03	0,01	37,97	0,28	0,26	22,17	0,26	99,08	1,49	0,49	77,2

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI75-3-18-a1	core	5,5	X	39,33	0,01	0,03	0,00	43,83	0,25	0,23	15,13	0,36	99,22	1,66	0,32	85,2
KI75-3-18-a2	rim	5,5	X	37,22	0,00	0,00	0,00	35,35	0,25	0,43	26,37	0,16	99,83	1,41	0,59	72,6
KI75-3-18-b1	core	5,5	X	39,74	0,01	0,03	0,13	45,54	0,24	0,14	12,88	0,37	99,11	1,71	0,27	87,5
KI75-3-18-b2	rim	5,5	X	37,28	0,00	0,04	0,06	35,60	0,25	0,24	25,07	0,22	98,79	1,42	0,56	73,8
KI75-3-18-c1	core	5,5	X	39,80	0,00	0,04	0,04	45,32	0,30	0,23	12,82	0,38	98,96	1,70	0,27	87,5
KI75-3-18-c2	rim	5,5	X	37,44	0,06	0,00	0,00	34,95	0,22	0,30	25,95	0,23	99,21	1,39	0,58	72,7
KI75-3-18-d1	core	5,5	X	39,99	0,01	0,03	0,09	45,88	0,28	0,19	12,65	0,34	99,54	1,71	0,27	87,8
KI75-3-18-d2	rim	5,5	X	37,59	0,02	0,00	0,00	36,08	0,23	0,42	25,49	0,26	100,13	1,42	0,56	73,7
KI75-3-18-g1-1	core 1	5,5	X	40,02	0,00	0,03	0,03	45,58	0,26	0,17	12,75	0,41	99,25	1,70	0,27	87,6
KI75-3-18-g1-2	rim 1	5,5	X	36,41	0,02	0,02	0,02	32,56	0,25	0,28	30,07	0,19	99,92	1,32	0,68	68,2
KI75-3-18-g2-1	core 2	5,5	X	39,85	0,00	0,05	0,10	45,96	0,27	0,12	12,20	0,40	98,94	1,72	0,26	88,2
KI75-3-18-g2-2	rim 2	5,5	X	37,95	0,02	0,03	0,04	37,07	0,25	0,30	23,50	0,26	99,47	1,46	0,52	75,8
KI75-3-18-i1	core	5,5	X	40,03	0,00	0,05	0,13	46,27	0,29	0,17	12,80	0,39	100,14	1,72	0,27	87,8
KI75-3-18-i2	rim	5,5	X	37,97	0,02	0,03	0,02	37,78	0,21	0,36	23,77	0,22	100,42	1,47	0,52	75,9
KI75-3-18-k1	core	5,5	X	39,55	0,01	0,04	0,05	43,85	0,27	0,17	14,66	0,32	98,98	1,66	0,31	85,6
KI75-3-18-k2	rim	5,5	X	37,10	0,00	0,00	0,02	33,98	0,23	0,30	27,97	0,19	99,80	1,36	0,63	70,6
KI75-3-18-e1	core	5,5		39,92	0,00	0,03	0,05	44,55	0,25	0,17	14,07	0,44	99,48	1,67	0,30	86,3
KI75-3-18-e2	rim	5,5		37,85	0,02	0,00	0,00	37,25	0,21	0,29	23,89	0,27	99,80	1,46	0,53	75,5
KI75-3-18-f1	core	5,5		39,68	0,00	0,04	0,10	45,02	0,29	0,20	13,24	0,36	98,98	1,69	0,28	87,1
KI75-3-18-f2	rim	5,5		36,91	0,02	0,01	0,03	33,31	0,22	0,32	28,02	0,22	99,09	1,35	0,64	70,2
KI75-3-18-h1	core	5,5		40,38	0,00	0,04	0,09	47,57	0,24	0,09	10,18	0,47	99,16	1,76	0,21	90,3
KI75-3-18-h2	rim	5,5		37,41	0,00	0,02	0,00	35,00	0,24	0,25	26,85	0,20	100,00	1,39	0,60	72,1
KI75-3-18-j1	core	5,5		39,17	0,00	0,03	0,01	42,66	0,29	0,24	16,96	0,30	99,77	1,62	0,36	83,3
KI75-3-18-j2	rim	5,5		36,71	0,02	0,02	0,00	32,90	0,22	0,37	28,76	0,21	99,24	1,33	0,65	69,4
KI75-3-18-l1	core	5,5		39,28	0,02	0,03	0,05	43,49	0,29	0,19	15,97	0,29	99,64	1,65	0,34	84,4
KI75-3-18-l2	rim	5,5		36,82	0,02	0,00	0,04	31,58	0,19	0,28	31,23	0,20	100,38	1,28	0,71	66,7
KI75-3-18-m1	core	5,5		39,43	0,00	0,05	0,03	43,09	0,24	0,10	16,25	0,29	99,51	1,63	0,35	84,0
KI75-3-18-m2	rim	5,5		37,58	0,01	0,02	0,03	37,42	0,17	0,32	23,83	0,28	99,68	1,47	0,53	75,7
KI75-3-18-n1	core	5,5		38,36	0,02	0,04	0,16	41,42	0,28	0,17	17,91	0,30	98,71	1,60	0,39	82,1
KI75-3-18-n2	rim	5,5		37,26	0,00	0,01	0,00	35,56	0,24	0,41	25,52	0,21	99,22	1,42	0,57	73,4
KI75-3-18-o1	core	5,5		38,52	0,02	0,04	0,01	40,57	0,26	0,27	19,46	0,23	99,43	1,56	0,42	80,5
KI75-3-18-o2	rim	5,5		36,43	0,10	0,02	0,04	31,61	0,17	0,39	30,08	0,16	99,01	1,29	0,69	67,6
KI75-3-35-e1c	core	10,7	X	39,61	0,00	0,05	0,05	45,34	0,28	0,16	13,80	0,32	99,60	1,70	0,29	86,7
KI75-3-35-e2c	core	10,7	X	39,61	0,02	0,05	0,00	45,39	0,29	0,16	14,06	0,32	99,91	1,70	0,30	86,5
KI75-3-35-e3b	rim	10,7	X	37,15	0,03	0,04	0,00	34,68	0,23	0,39	27,14	0,26	99,94	1,38	0,61	71,7
KI75-3-35-f1	core	10,7	X	39,29	0,02	0,04	0,03	43,75	0,27	0,13	15,88	0,39	99,80	1,65	0,34	84,5
KI75-3-35-f2	rim	10,7	X	37,08	0,03	0,03	0,00	33,90	0,20	0,40	28,07	0,22	99,95	1,36	0,63	70,5
KI75-3-35-k1	core	10,7	X	40,18	0,00	0,06	0,08	47,41	0,24	0,20	11,69	0,39	100,26	1,75	0,24	88,9
KI75-3-35-a1	core	10,7		37,55	0,01	0,04	0,01	35,66	0,26	0,40	26,15	0,20	100,29	1,41	0,58	73,0

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI75-3-35-a2	rim	10,7		36,98	0,03	0,05	0,03	33,39	0,22	0,26	28,89	0,22	100,06	1,34	0,65	69,6
KI75-3-35-b1	~core	10,7		37,50	0,00	0,05	0,03	35,90	0,23	0,32	25,79	0,13	99,97	1,42	0,57	73,4
KI75-3-35-b2	rim	10,7		36,97	0,01	0,03	0,04	33,50	0,22	0,33	28,32	0,24	99,71	1,35	0,64	70,1
KI75-3-35-c1	core	10,7		38,21	0,02	0,05	0,06	39,74	0,25	0,32	21,41	0,20	100,26	1,53	0,46	78,6
KI75-3-35-c2	rim	10,7		36,92	0,17	0,05	0,05	34,45	0,17	0,34	27,72	0,23	100,12	1,38	0,62	71,1
KI75-3-35-d1	core	10,7		39,96	0,00	0,04	0,09	45,92	0,25	0,19	13,09	0,34	99,88	1,71	0,27	87,4
KI75-3-35-d2	rim	10,7		37,89	0,03	0,04	0,00	36,44	0,19	0,36	25,39	0,27	100,65	1,43	0,56	74,0
KI75-3-35-g1	core	10,7		38,49	0,01	0,02	0,03	41,06	0,29	0,24	19,69	0,29	100,12	1,58	0,42	80,5
KI75-3-35-g2	rim	10,7		37,00	0,03	0,03	0,00	34,80	0,21	0,32	27,38	0,21	99,98	1,39	0,61	71,6
KI75-3-35-h1	core	10,7		37,91	0,02	0,05	0,07	38,85	0,25	0,29	22,21	0,22	99,94	1,51	0,49	77,6
KI75-3-35-h2	rim	10,7		37,08	0,02	0,03	0,05	34,76	0,21	0,29	27,61	0,20	100,25	1,38	0,62	71,4
KI75-3-35-i1	core	10,7		37,70	0,03	0,03	0,05	37,53	0,24	0,33	24,15	0,26	100,31	1,47	0,53	75,5
KI75-3-35-i2	rim	10,7		37,01	0,03	0,05	0,00	34,05	0,22	0,37	28,15	0,21	100,09	1,36	0,63	70,6
KI75-3-35-j1	core	10,7		38,07	0,01	0,02	0,02	38,64	0,25	0,20	22,89	0,31	100,43	1,50	0,50	77,0
KI75-3-35-j2	rim	10,7		36,86	0,02	0,02	0,05	33,90	0,21	0,39	28,68	0,25	100,39	1,36	0,64	70,1
KI75-3-35-l1	core	10,7		38,83	0,02	0,04	0,03	42,22	0,28	0,27	18,28	0,33	100,32	1,61	0,39	82,1
KI75-3-35-l2	rim	10,7		37,04	0,00	0,04	0,02	35,33	0,23	0,38	26,83	0,23	100,10	1,40	0,60	72,3
KI75-3-35-n1	core	10,7		37,76	0,00	0,05	0,00	36,73	0,25	0,30	24,76	0,23	100,12	1,44	0,55	74,6
KI75-3-35-n2	rim	10,7		36,98	0,03	0,02	0,00	33,75	0,21	0,32	28,45	0,23	99,98	1,35	0,64	70,2
KI75-3-46-f1	core	14,0	X	39,76	0,01	0,07	0,10	43,61	0,24	0,28	16,04	0,35	100,46	1,64	0,34	84,4
KI75-3-46-f2	rim	14,0	X	37,60	0,01	0,03	0,03	36,26	0,18	0,30	25,39	0,26	100,08	1,43	0,56	73,9
KI75-3-46-h1	core	14,0	X	40,20	0,01	0,06	0,04	47,04	0,26	0,13	11,86	0,49	100,09	1,74	0,25	88,7
KI75-3-46-h2	rim	14,0	X	37,41	0,00	0,03	0,00	36,62	0,20	0,34	25,38	0,20	100,22	1,44	0,56	74,1
KI75-3-46-m1	core	14,0	X	38,41	0,01	0,02	0,09	40,03	0,26	0,36	20,89	0,27	100,35	1,54	0,45	79,2
KI75-3-46-m2	rim	14,0	X	37,18	0,00	0,04	0,00	34,95	0,22	0,37	26,98	0,28	100,05	1,39	0,60	72,0
KI75-3-46-o1	core	14,0	X	39,06	0,00	0,08	0,04	42,58	0,24	0,23	17,52	0,30	100,04	1,62	0,37	82,8
KI75-3-46-o2	rim	14,0	X	37,28	0,02	0,04	0,06	35,09	0,23	0,35	26,56	0,21	99,85	1,40	0,59	72,4
KI75-3-46-c1	core	14,0		37,42	0,04	0,03	0,00	35,47	0,20	0,34	26,22	0,25	99,96	1,41	0,58	72,8
KI75-3-46-c2	rim	14,0		37,08	0,03	0,03	0,00	33,73	0,18	0,51	28,65	0,25	100,47	1,35	0,64	70,0
KI75-3-46-d1	core	14,0		39,14	0,01	0,03	0,03	42,86	0,27	0,16	16,73	0,32	99,61	1,63	0,36	83,6
KI75-3-46-d2	rim	14,0		37,07	0,03	0,03	0,00	34,83	0,15	0,28	27,27	0,28	99,96	1,39	0,61	71,7
KI75-3-46-e1	core	14,0		38,68	0,01	0,02	0,03	40,81	0,25	0,26	19,52	0,26	99,88	1,57	0,42	80,6
KI75-3-46-e2	rim	14,0		37,39	0,04	0,00	0,00	35,03	0,18	0,26	26,27	0,22	99,40	1,40	0,59	72,5
KI75-3-46-g1	core	14,0		38,70	0,01	0,04	0,06	41,48	0,25	0,33	18,98	0,33	100,20	1,58	0,41	81,2
KI75-3-46-g2	rim	14,0		37,56	0,02	0,01	0,07	35,98	0,21	0,36	25,54	0,24	99,99	1,42	0,57	73,6
KI75-3-46-i1	core	14,0		37,14	0,01	0,01	0,02	35,19	0,26	0,39	27,12	0,19	100,34	1,40	0,60	72,0
KI75-3-46-i2	rim	14,0		37,11	0,02	0,02	0,00	34,75	0,21	0,26	27,34	0,21	99,93	1,39	0,61	71,6
KI75-3-46-j1	core	14,0		37,39	0,01	0,03	0,00	35,63	0,23	0,30	26,37	0,25	100,22	1,41	0,59	72,8
KI75-3-46-j2	rim	14,0		37,18	0,03	0,03	0,00	34,86	0,17	0,38	27,40	0,21	100,29	1,39	0,61	71,6

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI75-3-46-k1	core	14,0		39,76	0,01	0,06	0,11	45,46	0,25	0,19	13,76	0,38	99,99	1,70	0,29	86,7
KI75-3-46-k2	rim	14,0		37,89	0,04	0,04	0,01	37,01	0,21	0,39	24,43	0,26	100,27	1,45	0,54	75,0
KI75-3-46-l1	core	14,0		38,36	0,00	0,02	0,10	40,25	0,25	0,28	20,59	0,29	100,14	1,55	0,45	79,5
KI75-3-46-l2	rim	14,0		37,49	0,03	0,04	0,00	36,10	0,22	0,35	25,76	0,24	100,24	1,42	0,57	73,5
KI75-3-46-n1	core	14,0		38,39	0,01	0,04	0,04	40,39	0,25	0,31	20,11	0,25	99,81	1,56	0,44	79,9
KI75-3-46-n2	rim	14,0		37,32	0,04	0,03	0,00	34,96	0,18	0,31	27,28	0,20	100,32	1,39	0,61	71,7
KI75-3-60-b1	core	18,3	X	37,76	0,01	0,07	0,01	37,59	0,27	0,26	23,97	0,21	100,16	1,47	0,53	75,6
KI75-3-60-b2	rim	18,3	X	37,09	0,02	0,01	0,04	34,88	0,18	0,28	27,37	0,27	100,18	1,39	0,61	71,6
KI75-3-60-f1	core	18,3	X	38,65	0,01	0,04	0,02	40,06	0,24	0,28	20,45	0,28	100,04	1,54	0,44	79,5
KI75-3-60-f2	rim	18,3	X	37,78	0,03	0,04	0,04	36,78	0,18	0,31	24,67	0,24	100,08	1,44	0,54	74,7
KI75-3-60-g1	core	18,3	X	39,69	0,02	0,05	0,06	45,65	0,27	0,18	14,01	0,35	100,31	1,70	0,29	86,6
KI75-3-60-g2	rim	18,3	X	37,89	0,03	0,02	0,00	36,81	0,19	0,39	24,83	0,26	100,43	1,44	0,55	74,6
KI75-3-60-m1	core	18,3	X	40,12	0,00	0,06	0,06	44,09	0,29	0,21	15,42	0,32	100,56	1,65	0,32	85,0
KI75-3-60-m2	rim	18,3	X	37,74	0,00	0,04	0,04	36,57	0,21	0,39	25,39	0,21	100,63	1,43	0,56	74,0
KI75-3-60-a1	core	18,3		36,96	0,01	0,02	0,02	41,05	0,16	0,36	20,23	0,37	99,18	1,60	0,44	80,1
KI75-3-60-a2	rim	18,3		37,17	0,02	0,01	0,03	35,12	0,17	0,40	27,39	0,23	100,59	1,39	0,61	71,7
KI75-3-60-c1	core	18,3		38,82	0,01	0,04	0,02	41,29	0,23	0,24	19,53	0,38	100,59	1,57	0,42	80,7
KI75-3-60-c2	rim	18,3		37,76	0,02	0,01	0,05	38,39	0,17	0,35	23,26	0,29	100,30	1,50	0,51	76,6
KI75-3-60-d1	core	18,3		38,25	0,01	0,02	0,01	39,31	0,23	0,35	22,38	0,25	100,82	1,52	0,48	77,7
KI75-3-60-d2	rim	18,3		37,70	0,03	0,03	0,03	36,78	0,21	0,37	25,05	0,27	100,47	1,44	0,55	74,4
KI75-3-60-e1	core	18,3		39,11	0,00	0,04	0,07	42,72	0,26	0,24	17,51	0,30	100,28	1,62	0,37	82,9
KI75-3-60-e2	rim	18,3		37,53	0,02	0,01	0,00	35,56	0,21	0,36	26,69	0,22	100,60	1,40	0,59	72,5
KI75-3-60-h1	core	18,3		39,99	0,01	0,05	0,09	46,34	0,27	0,17	12,70	0,37	99,98	1,72	0,26	87,9
KI75-3-60-h2	rim	18,3		38,39	0,00	0,02	0,06	38,65	0,20	0,36	22,92	0,27	100,88	1,49	0,50	76,9
KI75-3-60-i1	core	18,3		38,10	0,01	0,04	0,02	39,29	0,29	0,35	21,75	0,26	100,12	1,52	0,47	78,2
KI75-3-60-i2	rim	18,3		37,74	0,02	0,02	0,00	36,59	0,15	0,23	24,85	0,25	99,85	1,44	0,55	74,5
KI75-3-60-j1	core	18,3		38,49	0,01	0,05	0,03	40,74	0,24	0,33	20,07	0,29	100,30	1,56	0,43	80,1
KI75-3-60-j2	rim	18,3		38,08	0,04	0,03	0,00	37,48	0,15	0,27	23,73	0,28	100,05	1,47	0,52	75,8
KI75-3-60-k1	core	18,3		38,52	0,01	0,05	0,05	40,34	0,26	0,30	20,47	0,28	100,31	1,55	0,44	79,6
KI75-3-60-k2	rim	18,3		37,34	0,04	0,03	0,03	35,69	0,17	0,31	25,99	0,26	99,89	1,42	0,58	73,1
KI75-3-60-l1	core	18,3		37,60	0,01	0,02	0,00	37,35	0,23	0,28	23,69	0,28	99,46	1,47	0,52	75,7
KI75-3-60-l2	rim	18,3		37,52	0,04	0,02	0,00	36,33	0,16	0,39	24,90	0,31	99,68	1,44	0,55	74,3
KI75-3-141-a1	core	43,0	X	38,97	0,01	0,04	0,08	41,20	0,27	0,29	19,53	0,24	100,68	1,57	0,42	80,7
KI75-3-141-a2	rim	43,0	X	37,77	0,02	0,01	0,02	36,58	0,22	0,36	24,96	0,17	100,16	1,44	0,55	74,4
KI75-3-141-e1	core	43,0	X	38,53	0,00	0,05	0,02	41,13	0,27	0,29	19,33	0,29	99,92	1,58	0,42	80,8
KI75-3-141-e2	rim	43,0	X	37,97	0,02	0,03	0,00	38,31	0,22	0,31	22,43	0,26	99,54	1,50	0,49	77,2
KI75-3-141-f1	core	43,0	X	38,55	0,01	0,04	0,00	40,06	0,25	0,27	20,54	0,23	99,95	1,54	0,44	79,4
KI75-3-141-f2	rim	43,0	X	37,68	0,03	0,02	0,00	36,29	0,24	0,34	24,34	0,25	99,19	1,44	0,54	74,7

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI75-3-141-g1	core	43,0	X	38,59	0,00	0,05	0,03	40,51	0,29	0,25	19,76	0,32	99,81	1,56	0,43	80,3
KI75-3-141-g2	rim	43,0	X	37,71	0,03	0,03	0,00	36,26	0,24	0,37	25,48	0,23	100,38	1,43	0,56	73,8
KI75-3-141-j1	core	43,0	X	38,24	0,02	0,04	0,04	39,81	0,28	0,32	21,37	0,23	100,37	1,54	0,46	78,7
KI75-3-141-j2	rim	43,0	X	38,26	0,01	0,05	0,08	38,86	0,25	0,35	22,48	0,28	100,65	1,50	0,49	77,4
KI75-3-141-k1	core	43,0	X	39,66	0,01	0,06	0,04	44,80	0,24	0,07	14,72	0,37	100,03	1,68	0,31	85,8
KI75-3-141-k2	rim	43,0	X	37,86	0,03	0,03	0,07	37,94	0,22	0,38	23,85	0,29	100,67	1,48	0,52	75,9
KI75-3-141-b1	core	43,0		38,01	0,00	0,02	0,07	37,76	0,23	0,33	23,94	0,29	100,66	1,47	0,52	75,7
KI75-3-141-b2	rim	43,0		37,36	0,03	0,02	0,01	36,06	0,24	0,28	25,64	0,24	99,89	1,43	0,57	73,6
KI75-3-141-c1	core	43,0		38,55	0,01	0,04	0,02	39,90	0,29	0,22	20,90	0,29	100,25	1,54	0,45	79,1
KI75-3-141-c2	rim	43,0		37,98	0,03	0,02	0,00	37,70	0,25	0,21	23,44	0,29	99,93	1,47	0,51	76,1
KI75-3-141-d1	core	43,0		38,33	0,02	0,04	0,00	39,14	0,27	0,32	21,73	0,29	100,18	1,52	0,47	78,1
KI75-3-141-d2	rim	43,0		38,51	0,01	0,02	0,02	36,95	0,23	0,29	24,73	0,23	101,00	1,44	0,54	74,7
KI75-3-141-h1	core	43,0		38,26	0,02	0,03	0,04	38,40	0,23	0,39	22,60	0,25	100,24	1,49	0,49	77,1
KI75-3-141-h2	rim	43,0		37,85	0,03	0,02	0,05	37,00	0,23	0,31	24,37	0,22	100,09	1,45	0,54	75,0
KI75-3-141-i1	core	43,0		38,68	0,00	0,05	0,00	40,93	0,27	0,28	19,84	0,26	100,32	1,57	0,43	80,3
KI75-3-141-i2	rim	43,0		37,72	0,02	0,01	0,02	36,50	0,23	0,30	25,14	0,25	100,19	1,44	0,55	74,2
KI75-3-141-l1	core	43,0		39,09	0,00	0,04	0,02	41,55	0,25	0,26	19,04	0,29	100,55	1,58	0,41	81,2
KI75-3-141-l2	rim	43,0		38,09	0,03	0,04	0,00	38,52	0,23	0,38	22,72	0,26	100,28	1,50	0,50	77,1
KI79-5-182.4-a1	core	55,6	X	39,04	0,01	0,34	0,08	41,80	0,25	0,22	18,58	0,22	100,56	1,59	0,40	81,7
KI79-5-182.4-a2	rim	55,6	X	37,93	0,01	0,06	0,00	38,71	0,18	0,27	23,43	0,25	100,90	1,50	0,51	76,6
KI79-5-182.4-f1	core	55,6	X	37,62	0,01	0,06	0,00	38,16	0,18	0,26	23,96	0,24	100,58	1,49	0,52	75,9
KI79-5-182.4-f2	rim	55,6	X	37,91	0,03	0,11	0,00	37,51	0,19	0,35	23,10	0,28	99,50	1,47	0,51	76,3
KI79-5-182.4-g1	core	55,6	X	38,31	0,00	0,11	0,10	41,03	0,24	0,26	19,10	0,29	99,46	1,58	0,41	81,0
KI79-5-182.4-g2	rim	55,6	X	37,94	0,03	0,03	0,03	38,47	0,18	0,28	22,43	0,22	99,63	1,50	0,49	77,3
KI79-5-182.4-i1	core 1	55,6	X	38,55	0,00	0,03	0,05	41,11	0,26	0,28	19,84	0,30	100,51	1,57	0,43	80,4
KI79-5-182.4-i2	rim 1	55,6	X	37,83	0,02	0,08	0,04	37,61	0,16	0,30	22,86	0,30	99,20	1,48	0,50	76,5
KI79-5-182.4-i3	core 2	55,6	X	38,40	0,01	0,10	0,06	40,67	0,27	0,25	19,43	0,29	99,50	1,57	0,42	80,6
KI79-5-182.4-i4	rim 2	55,6	X	38,03	0,01	0,00	0,02	38,28	0,18	0,31	23,21	0,25	100,35	1,49	0,51	76,5
KI79-5-182.4-j1	core	55,6	X	38,46	0,02	0,08	0,04	40,61	0,25	0,20	19,61	0,29	99,56	1,57	0,42	80,4
KI79-5-182.4-j2	rim	55,6	X	37,94	0,02	0,02	0,04	39,56	0,18	0,27	21,59	0,28	99,94	1,54	0,47	78,4
KI79-5-182.4-b1	core	55,6		38,21	0,01	0,02	0,00	39,27	0,23	0,28	21,33	0,25	99,64	1,53	0,47	78,5
KI79-5-182.4-b2	rim	55,6		38,00	0,01	1,87	0,00	38,22	0,19	0,26	22,05	0,30	100,89	1,47	0,48	77,4
KI79-5-182.4-c1	core	55,6		38,10	0,00	0,06	0,03	39,87	0,25	0,26	20,25	0,28	99,19	1,55	0,44	79,6
KI79-5-182.4-c2	rim	55,6		37,91	0,03	0,07	0,00	38,36	0,20	0,39	22,38	0,24	99,61	1,50	0,49	77,3
KI79-5-182.4-d1	core	55,6		38,67	0,00	0,32	0,04	41,97	0,28	0,15	18,38	0,28	100,10	1,60	0,39	81,9
KI79-5-182.4-d2	rim	55,6		37,94	0,01	0,04	0,00	39,01	0,19	0,30	21,64	0,30	99,40	1,52	0,47	78,1
KI79-5-182.4-e1	core	55,6		37,76	0,03	0,00	0,00	38,54	0,19	0,38	22,76	0,22	99,89	1,50	0,50	77,0
KI79-5-182.4-e2	rim	55,6		37,63	0,03	0,05	0,02	37,96	0,16	0,21	22,66	0,28	99,05	1,49	0,50	76,9
KI79-5-182.4-h1	core	55,6		38,22	0,02	0,18	0,02	40,39	0,22	0,30	21,19	0,25	100,95	1,55	0,46	79,0
KI79-5-182.4-h2	rim	55,6		38,35	0,02	0,02	0,05	39,79	0,17	0,21	20,43	0,23	99,29	1,54	0,44	79,4

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI79-5-193.3-b1	core	58,9	X	39,00	0,00	0,05	0,05	43,15	0,25	0,12	17,31	0,31	100,30	1,63	0,37	83,2
KI79-5-193.3-b2	rim	58,9	X	37,70	0,02	0,06	0,02	38,42	0,19	0,26	22,06	0,24	98,99	1,51	0,49	77,5
KI79-5-193.3-c1	core	58,9	X	38,37	0,00	0,04	0,08	40,43	0,21	0,22	20,27	0,24	99,89	1,56	0,44	79,8
KI79-5-193.3-c2	rim	58,9	X	37,82	0,02	0,14	0,00	37,80	0,17	0,36	23,62	0,26	100,23	1,48	0,52	76,0
KI79-5-193.3-f1	core	58,9	X	38,52	0,02	0,07	0,01	41,07	0,26	0,13	19,27	0,26	99,69	1,58	0,42	80,9
KI79-5-193.3-f2	rim	58,9	X	37,67	0,01	0,01	0,00	37,67	0,18	0,32	23,10	0,24	99,30	1,48	0,51	76,4
KI79-5-193.3-g1	core	58,9	X	38,68	0,00	0,00	0,09	40,90	0,26	0,28	19,56	0,29	100,11	1,57	0,42	80,5
KI79-5-193.3-g2	rim	58,9	X	37,99	0,02	0,02	0,00	39,06	0,18	0,29	22,11	0,27	99,98	1,52	0,48	77,8
KI79-5-193.3-i1	core	58,9	X	37,88	0,00	0,02	0,01	38,11	0,23	0,31	23,11	0,26	99,92	1,49	0,51	76,6
KI79-5-193.3-i2	rim	58,9	X	37,77	0,05	0,00	0,01	37,57	0,17	0,23	23,80	0,26	99,86	1,47	0,52	75,8
KI79-5-193.3-a1	core	58,9		38,00	0,00	0,05	0,00	38,73	0,23	0,29	23,01	0,30	100,62	1,50	0,50	76,9
KI79-5-193.3-a2	rim	58,9		37,75	0,01	0,00	0,00	37,91	0,18	0,25	22,57	0,31	99,03	1,49	0,50	76,9
KI79-5-193.3-d1	core	58,9		38,00	0,01	0,26	0,04	38,94	0,22	0,31	22,51	0,26	100,58	1,51	0,49	77,4
KI79-5-193.3-d2	rim	58,9		37,76	0,03	0,02	0,04	37,57	0,16	0,30	23,06	0,27	99,25	1,48	0,51	76,3
KI79-5-193.3-e1	core	58,9		37,76	0,02	0,14	0,01	38,29	0,21	0,35	23,45	0,27	100,49	1,49	0,51	76,4
KI79-5-193.3-e2	rim	58,9		37,77	0,01	0,07	0,04	37,25	0,16	0,23	24,28	0,26	100,07	1,46	0,53	75,2
KI79-5-193.3-h1	core	58,9		38,27	0,00	0,18	0,00	39,91	0,24	0,28	20,83	0,31	100,14	1,54	0,45	79,1
KI79-5-193.3-h2	rim	58,9		37,85	0,01	0,26	0,04	38,36	0,17	0,31	23,63	0,24	100,99	1,49	0,51	76,3
KI79-5-193.3-j1	core	58,9		37,94	0,01	0,00	0,00	38,10	0,21	0,35	22,82	0,25	99,70	1,49	0,50	76,8
KI79-5-193.3-j2	rim	58,9		37,64	0,03	0,07	0,00	37,24	0,17	0,26	23,75	0,27	99,47	1,47	0,52	75,7
KI79-5-250.0-b1	core	76,2	X	38,98	0,00	0,00	0,00	42,51	0,21	0,22	18,22	0,33	100,57	1,61	0,39	82,2
KI79-5-250.0-b2	rim	76,2	X	37,98	0,02	0,01	0,00	38,90	0,19	0,36	21,94	0,23	99,63	1,52	0,48	77,8
KI79-5-250.0-d1	core	76,2	X	38,70	0,01	0,00	0,06	41,08	0,23	0,20	18,62	0,34	99,33	1,58	0,40	81,4
KI79-5-250.0-d2	rim	76,2	X	37,46	0,02	0,06	0,03	36,85	0,17	0,24	24,76	0,22	99,83	1,45	0,55	74,7
KI79-5-250.0-h1	core	76,2	X	40,02	0,02	0,01	0,03	46,96	0,29	0,22	12,55	0,37	100,50	1,74	0,26	88,1
KI79-5-250.0-h2	rim	76,2	X	37,74	0,01	0,00	0,00	38,47	0,17	0,19	22,65	0,25	99,50	1,51	0,50	77,1
KI79-5-250.0-i1	core	76,2	X	37,80	0,01	0,08	0,00	38,20	0,22	0,33	23,30	0,24	100,26	1,49	0,51	76,5
KI79-5-250.0-i2	rim	76,2	X	37,54	0,02	0,00	0,00	37,46	0,14	0,40	23,57	0,28	99,45	1,48	0,52	75,9
KI79-5-250.0-j1	core	76,2	X	37,69	0,01	0,05	0,02	38,18	0,25	0,43	22,73	0,28	99,74	1,49	0,50	76,9
KI79-5-250.0-j2	rim	76,2	X	37,22	0,03	0,03	0,00	36,28	0,18	0,39	25,10	0,21	99,53	1,44	0,56	74,1
KI79-5-250.0-a1	core	76,2		38,22	0,00	0,06	0,09	40,12	0,27	0,27	21,50	0,25	100,83	1,54	0,46	78,7
KI79-5-250.0-a2	rim	76,2		37,64	0,01	0,12	0,08	37,52	0,15	0,25	24,06	0,26	100,20	1,47	0,53	75,6
KI79-5-250.0-c1	core	76,2		38,38	0,01	0,00	0,02	40,72	0,20	0,28	19,94	0,32	99,91	1,57	0,43	80,2
KI79-5-250.0-c2	rim	76,2		37,57	0,03	0,02	0,03	36,27	0,16	0,43	24,92	0,26	99,73	1,43	0,55	74,3
KI79-5-250.0-e1	core	76,2		39,37	0,01	0,04	0,10	44,25	0,27	0,16	15,35	0,33	99,91	1,67	0,32	85,1
KI79-5-250.0-e2	rim	76,2		38,24	0,03	0,02	0,01	40,05	0,24	0,34	20,55	0,27	99,74	1,55	0,45	79,4
KI79-5-250.0-f1	core	76,2		38,01	0,01	0,41	0,05	38,70	0,19	0,34	21,74	0,27	99,71	1,50	0,47	77,9
KI79-5-250.0-f2	rim	76,2		37,95	0,01	0,01	0,04	38,58	0,17	0,22	22,19	0,26	99,44	1,51	0,49	77,5
KI79-5-250.0-g1	core	76,2		37,98	0,01	0,04	0,03	38,66	0,26	0,25	22,76	0,27	100,42	1,50	0,50	77,1
KI79-5-250.0-g2	rim	76,2		37,48	0,03	0,00	0,00	36,38	0,15	0,31	24,62	0,23	99,31	1,44	0,55	74,5

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI88-1-205.2-c1	core	62,5	X	38,10	0,00	0,05	0,06	41,58	0,26	0,25	17,99	0,32	98,67	1,61	0,39	82,1
KI88-1-205.2-c2	rim	62,5	X	38,10	0,02	0,05	0,00	39,17	0,18	0,26	20,90	0,27	98,99	1,53	0,46	78,8
KI88-1-205.2-d1	core	62,5	X	38,06	0,01	0,06	0,00	39,90	0,26	0,28	20,61	0,28	99,54	1,55	0,45	79,3
KI88-1-205.2-d2	rim	62,5	X	37,69	0,05	0,00	0,00	38,77	0,17	0,27	21,44	0,30	98,72	1,52	0,47	78,2
KI88-1-205.2-e1	core	62,5	X	37,75	0,02	0,00	0,03	39,42	0,25	0,28	20,90	0,20	98,87	1,54	0,46	78,9
KI88-1-205.2-e2	rim	62,5	X	37,83	0,03	0,12	0,01	38,02	0,18	0,28	22,76	0,22	99,48	1,49	0,50	76,8
KI88-1-205.2-g1	core	62,5	X	38,36	0,02	0,05	0,00	40,30	0,26	0,28	20,00	0,24	99,56	1,56	0,43	80,0
KI88-1-205.2-g2	rim	62,5	X	38,07	0,04	0,00	0,02	38,80	0,20	0,36	21,74	0,24	99,47	1,51	0,48	78,0
KI88-1-205.2-i1	core	62,5	X	39,20	0,00	0,15	0,07	44,22	0,24	0,14	16,07	0,33	100,42	1,66	0,34	84,5
KI88-1-205.2-i2	rim (outer)	62,5	X	37,78	0,00	0,00	0,05	38,10	0,19	0,20	23,17	0,31	99,79	1,49	0,51	76,5
KI88-1-205.2-i3	rim (inner)	62,5	X	38,89	0,01	0,20	0,01	42,16	0,25	0,25	18,30	0,26	100,35	1,60	0,39	82,0
KI88-1-205.2-a1	core	62,5		38,65	0,02	1,96	0,07	41,69	0,28	0,26	17,81	0,24	100,99	1,57	0,38	82,2
KI88-1-205.2-a2	rim	62,5		37,82	0,02	0,00	0,00	38,59	0,18	0,26	21,64	0,33	98,87	1,51	0,48	77,9
KI88-1-205.2-b1	core	62,5		38,45	0,01	0,09	0,04	41,31	0,28	0,20	18,73	0,30	99,46	1,59	0,40	81,4
KI88-1-205.2-b2	rim	62,5		37,76	0,02	0,00	0,14	39,36	0,20	0,27	21,21	0,23	99,22	1,54	0,46	78,6
KI88-1-205.2-f1	core	62,5		37,87	0,01	0,02	0,01	39,64	0,25	0,26	20,49	0,24	98,78	1,55	0,45	79,3
KI88-1-205.2-f2	rim	62,5		37,98	0,02	0,16	0,09	38,36	0,19	0,16	21,92	0,25	99,13	1,50	0,48	77,6
KI88-1-205.2-h1	core	62,5		37,97	0,01	0,05	0,00	39,05	0,25	0,21	21,44	0,23	99,25	1,52	0,47	78,3
KI88-1-205.2-h2	rim	62,5		37,83	0,04	0,13	0,00	38,28	0,20	0,30	23,01	0,27	100,11	1,49	0,50	76,7
KI88-1-205.2-j1	core	62,5		38,38	0,00	0,06	0,04	40,37	0,26	0,21	19,64	0,29	99,29	1,56	0,43	80,3
KI88-1-205.2-j2	rim 1	62,5		37,84	0,02	0,03	0,00	38,14	0,17	0,31	22,92	0,28	99,75	1,49	0,50	76,7
KI88-1-205.2-j3	rim 2	62,5		37,72	0,02	0,01	0,02	38,48	0,17	0,34	22,66	0,24	99,78	1,50	0,50	77,1
KI88-1-248.9-b1	core	75,9	X	38,58	0,00	0,04	0,02	41,67	0,26	0,18	19,48	0,32	100,55	1,59	0,42	80,9
KI88-1-248.9-b2	rim	75,9	X	38,01	0,03	0,03	0,00	39,45	0,18	0,24	21,03	0,27	99,31	1,54	0,46	78,8
KI88-1-248.9-c1	core	75,9	X	38,48	0,00	0,06	0,06	41,40	0,27	0,27	20,55	0,28	101,39	1,57	0,44	80,0
KI88-1-248.9-c2	rim	75,9	X	38,10	0,02	0,07	0,05	39,99	0,20	0,28	20,91	0,26	99,91	1,55	0,45	79,1
KI88-1-248.9-f1	core	75,9	X	38,40	0,00	0,05	0,05	41,40	0,28	0,29	18,66	0,28	99,44	1,59	0,40	81,4
KI88-1-248.9-f2	rim	75,9	X	37,98	0,01	0,05	0,06	40,18	0,21	0,31	20,72	0,29	99,91	1,55	0,45	79,3
KI88-1-248.9-g1	core	75,9	X	38,08	0,04	0,44	0,02	40,02	0,23	0,26	20,99	0,27	100,41	1,54	0,45	79,1
KI88-1-248.9-g2	rim	75,9	X	37,86	0,07	0,05	0,01	39,33	0,21	0,34	21,55	0,28	99,74	1,53	0,47	78,3
KI88-1-248.9-j1	core	75,9	X	38,11	0,00	0,05	0,04	41,93	0,27	0,26	18,36	0,26	99,35	1,61	0,40	81,9
KI88-1-248.9-j2	rim	75,9	X	37,81	0,04	0,03	0,02	40,03	0,19	0,25	20,89	0,25	99,56	1,55	0,46	79,1
KI88-1-248.9-a1	core	75,9		37,89	0,00	0,03	0,03	39,70	0,21	0,27	21,02	0,31	99,46	1,54	0,46	78,9
KI88-1-248.9-a2	rim	75,9		38,01	0,01	0,02	0,05	39,33	0,18	0,17	21,15	0,31	99,27	1,53	0,46	78,7
KI88-1-248.9-d1	core	75,9		38,55	0,01	0,02	0,00	41,70	0,26	0,16	18,67	0,29	99,67	1,60	0,40	81,6
KI88-1-248.9-d2	rim	75,9		38,02	0,01	0,09	0,00	39,92	0,18	0,24	20,58	0,26	99,40	1,55	0,45	79,4
KI88-1-248.9-e1	core 1	75,9		38,33	0,01	0,04	0,03	40,69	0,22	0,25	20,27	0,29	100,19	1,56	0,44	79,9
KI88-1-248.9-e2	core 2	75,9		38,35	0,01	0,07	0,07	40,45	0,18	0,25	20,22	0,31	99,97	1,56	0,44	79,8
KI88-1-248.9-e3	rim	75,9		38,15	0,01	0,09	0,00	40,20	0,19	0,21	20,71	0,29	99,88	1,55	0,45	79,4
KI88-1-248.9-h1	core	75,9		38,36	0,00	0,04	0,03	40,28	0,20	0,22	20,20	0,22	99,61	1,56	0,44	79,8

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI88-1-248.9-h2	rim	75,9		38,15	0,01	0,14	0,00	39,43	0,19	0,23	21,14	0,32	99,67	1,53	0,46	78,7
KI88-1-248.9-i1	core	75,9		38,58	0,00	0,06	0,00	41,85	0,29	0,25	18,71	0,29	100,03	1,60	0,40	81,6
KI88-1-248.9-i2	rim	75,9		38,20	0,01	0,01	0,00	39,74	0,20	0,30	20,55	0,29	99,35	1,54	0,45	79,3
KI88-1-298.1-b1	core	90,9	X	37,62	0,02	0,10	0,00	37,95	0,21	0,34	23,11	0,25	99,69	1,49	0,51	76,5
KI88-1-298.1-b2	rim	90,9	X	37,61	0,02	0,07	0,00	36,37	0,18	0,35	25,32	0,30	100,34	1,43	0,56	74,0
KI88-1-298.1-c1	core	90,9	X	37,38	0,02	0,11	0,04	36,60	0,22	0,33	24,69	0,23	99,65	1,45	0,55	74,6
KI88-1-298.1-c2	rim	90,9	X	37,36	0,03	0,26	0,10	35,86	0,18	0,28	24,80	0,23	99,13	1,42	0,55	74,1
KI88-1-298.1-e1	core 1	90,9	X	37,39	0,00	0,06	0,00	36,78	0,22	0,23	24,47	0,28	99,43	1,45	0,54	74,8
KI88-1-298.1-e2	rim 1	90,9	X	37,36	0,03	0,05	0,01	36,62	0,18	0,30	25,17	0,26	99,98	1,44	0,56	74,2
KI88-1-298.1-e3	core 2	90,9	X	37,64	0,01	0,06	0,02	36,83	0,20	0,38	25,07	0,29	100,49	1,44	0,55	74,4
KI88-1-298.1-e4	rim 2	90,9	X	37,61	0,04	0,03	0,02	36,59	0,19	0,36	24,26	0,25	99,45	1,45	0,54	74,9
KI88-1-298.1-f1	core	90,9	X	37,69	0,02	0,07	0,02	37,56	0,17	0,39	25,22	0,23	101,41	1,46	0,55	74,7
KI88-1-298.1-f2	rim	90,9	X	37,40	0,02	0,00	0,03	37,40	0,20	0,37	24,53	0,25	100,30	1,47	0,54	75,1
KI88-1-298.1-j1	core	90,9	X	38,96	0,00	0,08	0,03	43,19	0,26	0,26	17,08	0,36	100,26	1,63	0,36	83,3
KI88-1-298.1-j2	rim 1	90,9	X	37,43	0,02	0,70	0,02	36,93	0,17	0,34	25,07	0,25	100,92	1,44	0,55	74,5
KI88-1-298.1-j3	rim 2 (outer)	90,9	X	37,29	0,03	0,03	0,02	37,11	0,19	0,27	24,59	0,26	99,84	1,46	0,54	74,9
KI88-1-298.1-j4	rim 2 (inner)	90,9	X	38,36	0,00	0,16	0,08	41,12	0,24	0,15	19,64	0,29	100,04	1,58	0,42	80,6
KI88-1-298.1-a1	core	90,9		37,66	0,00	0,09	0,00	37,77	0,23	0,31	23,64	0,26	100,04	1,48	0,52	76,0
KI88-1-298.1-a2	rim	90,9		37,35	0,01	0,12	0,02	36,90	0,17	0,22	24,58	0,21	99,61	1,46	0,54	74,8
KI88-1-298.1-d1	core	90,9		37,20	0,01	0,09	0,03	36,90	0,19	0,31	24,91	0,27	99,97	1,46	0,55	74,6
KI88-1-298.1-d2	rim	90,9		37,45	0,02	0,11	0,00	36,25	0,19	0,33	25,48	0,32	100,25	1,43	0,56	73,8
KI88-1-298.1-g1	core	90,9		37,57	0,02	0,01	0,05	36,37	0,20	0,34	25,65	0,24	100,48	1,43	0,57	73,7
KI88-1-298.1-g2	rim	90,9		37,45	0,03	0,03	0,00	36,41	0,20	0,26	25,61	0,30	100,37	1,43	0,57	73,8
KI88-1-298.1-h1	core	90,9		37,51	0,02	0,12	0,02	37,39	0,19	0,42	23,79	0,28	99,90	1,47	0,52	75,7
KI88-1-298.1-h2	rim	90,9		37,33	0,02	0,15	0,00	36,52	0,18	0,39	25,16	0,26	100,01	1,44	0,56	74,2
KI88-1-298.1-i1	core	90,9		37,46	0,02	0,00	0,00	36,70	0,20	0,27	24,47	0,22	99,41	1,45	0,54	74,8
KI88-1-298.1-i2	rim	90,9		37,42	0,04	0,00	0,00	36,29	0,16	0,32	25,46	0,27	100,05	1,43	0,56	73,8

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fe%
Centre of lake																
K175-1-2-c1	core	0,6	X	39,62	0,01	0,05	0,09	46,25	0,28	0,14	12,75	0,30	99,47	1,73	0,27	87,8
K175-1-2-c2	rim	0,6	X	38,02	0,01	0,06	0,01	43,98	0,28	0,22	15,95	0,32	98,87	1,68	0,34	84,5
K175-1-2-d1	core	0,6	X	40,09	0,00	0,09	0,02	48,33	0,23	0,07	10,21	0,41	99,48	1,78	0,21	90,4
K175-1-2-d2	rim	0,6	X	39,25	0,02	0,07	0,02	46,72	0,24	0,20	12,59	0,40	99,52	1,75	0,26	88,0
K175-1-2-e1	core	0,6	X	38,97	0,00	0,05	0,07	45,92	0,27	0,23	13,53	0,39	99,44	1,73	0,29	87,1
K175-1-2-e2	rim	0,6	X	38,93	0,01	0,05	0,11	45,69	0,27	0,12	13,51	0,39	99,09	1,72	0,29	87,0
K175-1-2-f1	core	0,6	X	39,52	0,00	0,05	0,06	46,68	0,25	0,18	12,74	0,37	99,86	1,74	0,27	87,9
K175-1-2-f2	rim	0,6	X	39,14	0,00	0,07	0,02	44,76	0,27	0,16	15,06	0,30	99,80	1,69	0,32	85,5
K175-1-2-n1	core	0,6	X	39,28	0,01	0,05	0,03	44,66	0,28	0,18	15,22	0,32	100,04	1,68	0,32	85,3
K175-1-2-n2	rim	0,6	X	38,78	0,00	0,04	0,07	44,12	0,29	0,21	15,92	0,34	99,80	1,67	0,34	84,6
K175-1-2-o1	core	0,6	X	39,32	0,00	0,05	0,05	45,77	0,27	0,18	14,14	0,35	100,13	1,71	0,30	86,5
K175-1-2-o2	rim	0,6	X	38,66	0,00	0,04	0,06	43,77	0,29	0,20	15,83	0,37	99,23	1,67	0,34	84,6
K175-1-2-q1-1	core	0,6	X	39,50	0,00	0,07	0,09	47,49	0,24	0,14	11,80	0,41	99,77	1,76	0,25	88,8
K175-1-2-q1-2	rim	0,6	X	38,48	0,00	0,04	0,00	43,72	0,26	0,27	16,40	0,26	99,46	1,66	0,35	84,1
K175-1-2-q2-1	core	0,6	X	39,87	0,01	0,06	0,00	46,22	0,29	0,16	11,74	0,43	99,51	1,75	0,25	88,8
K175-1-2-a1	core	0,6		39,68	0,00	0,03	0,06	46,22	0,29	0,11	12,55	0,35	99,31	1,73	0,26	87,9
K175-1-2-a2	rim	0,6		38,68	0,01	0,03	0,00	42,44	0,28	0,29	17,80	0,25	99,78	1,62	0,38	82,5
K175-1-2-b1	core	0,6		39,30	0,02	0,18	0,01	46,58	0,29	0,22	12,51	0,38	99,48	1,74	0,26	88,1
K175-1-2-b2	rim	0,6		39,19	0,01	0,04	0,10	45,62	0,26	0,16	13,84	0,42	99,63	1,71	0,29	86,7
K175-1-2-g1	core	0,6		38,73	0,00	0,05	0,09	45,39	0,29	0,12	14,24	0,33	99,26	1,71	0,30	86,3
K175-1-2-g2	rim	0,6		39,33	0,01	0,06	0,03	45,76	0,26	0,17	13,41	0,35	99,41	1,72	0,28	87,1
K175-1-2-h1	core	0,6		39,30	0,02	0,06	0,05	46,86	0,27	0,17	12,72	0,36	99,82	1,75	0,27	87,9
K175-1-2-h2	rim	0,6		38,56	0,00	0,05	0,02	42,33	0,29	0,17	17,92	0,30	99,65	1,62	0,38	82,4
K175-1-2-i1	core	0,6		39,34	0,01	0,04	0,03	45,46	0,32	0,22	14,30	0,42	100,13	1,70	0,30	86,3
K175-1-2-i2	rim	0,6		38,49	0,02	0,02	0,10	45,16	0,30	0,25	14,67	0,35	99,40	1,71	0,31	85,9
K175-1-2-j1	core	0,6		39,74	0,01	0,07	0,10	46,97	0,24	0,13	11,57	0,47	99,31	1,75	0,24	88,9
K175-1-2-j2	rim	0,6		38,82	0,02	0,06	0,07	43,34	0,26	0,19	16,81	0,36	99,95	1,64	0,36	83,6
K175-1-2-k1	core	0,6		39,84	0,02	0,06	0,08	47,59	0,27	0,19	11,50	0,41	99,95	1,76	0,24	89,1
K175-1-2-k2	rim	0,6		39,56	0,01	0,05	0,07	47,16	0,26	0,13	11,84	0,41	99,50	1,76	0,25	88,8
K175-1-2-l1	core	0,6		39,29	0,01	0,05	0,02	47,49	0,26	0,03	11,57	0,45	99,19	1,77	0,24	89,0
K175-1-2-l2	rim	0,6		39,28	0,00	0,06	0,03	45,97	0,26	0,34	13,30	0,38	99,61	1,72	0,28	87,3
K175-1-2-m1	core	0,6		38,64	0,01	0,03	0,00	43,44	0,31	0,24	16,29	0,23	99,19	1,66	0,35	84,1
K175-1-2-m2	rim	0,6		38,87	0,02	0,05	0,04	43,14	0,30	0,29	17,10	0,31	100,13	1,64	0,36	83,3
K175-1-2-p1	core	0,6		38,53	0,01	0,06	0,01	42,61	0,25	0,30	17,71	0,28	99,77	1,63	0,38	82,7
K175-1-2-p2	rim	0,6		38,64	0,03	0,04	0,00	43,06	0,29	0,32	17,12	0,23	99,74	1,64	0,37	83,3
K175-1-2-r1	core	0,6		39,93	0,00	0,09	0,06	47,48	0,26	0,12	11,58	0,41	99,94	1,76	0,24	89,0
K175-1-2-r2	rim	0,6		38,98	0,02	0,06	0,06	43,58	0,27	0,24	16,26	0,34	99,82	1,65	0,35	84,2
K175-1-2-s1	core	0,6		39,24	0,01	0,06	0,05	46,97	0,26	0,12	12,40	0,37	99,49	1,75	0,26	88,2
K175-1-2-s2	rim	0,6		38,62	0,01	0,05	0,04	44,18	0,27	0,23	15,58	0,38	99,38	1,68	0,33	84,9

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI75-1-5-a1	core	1,5	X	39,26	0,00	0,04	0,05	46,03	0,26	0,12	13,58	0,40	99,76	1,72	0,29	87,0
KI75-1-5-a2	rim	1,5	X	38,65	0,01	0,02	0,00	43,10	0,28	0,22	16,61	0,33	99,26	1,64	0,36	83,7
KI75-1-5-f1	core	1,5	X	39,23	0,00	0,04	0,03	48,54	0,26	0,11	12,14	0,39	100,76	1,79	0,25	88,8
KI75-1-5-f2	rim	1,5	X	38,43	0,01	0,06	0,05	43,83	0,25	0,20	15,47	0,36	98,65	1,68	0,33	84,9
KI75-1-5-g1	core	1,5	X	38,83	0,02	0,06	0,06	44,75	0,27	0,15	14,92	0,34	99,40	1,69	0,32	85,6
KI75-1-5-g2	rim	1,5	X	37,92	0,02	0,04	0,11	43,37	0,29	0,27	16,70	0,29	99,00	1,66	0,36	83,7
KI75-1-5-h1	core	1,5	X	40,01	0,03	0,09	0,09	47,29	0,23	0,14	11,49	0,39	99,77	1,75	0,24	89,1
KI75-1-5-h2	rim	1,5	X	37,84	0,01	0,06	0,01	40,11	0,25	0,26	21,35	0,25	100,16	1,55	0,46	78,8
KI75-1-5-i1	core	1,5	X	39,42	0,00	0,07	0,02	47,38	0,26	0,14	11,51	0,45	99,26	1,77	0,24	89,1
KI75-1-5-i2	rim	1,5	X	38,29	0,00	0,06	0,03	41,67	0,26	0,27	18,34	0,28	99,20	1,60	0,40	81,8
KI75-1-5-o1	core	1,5	X	39,96	0,00	0,06	0,04	47,19	0,25	0,16	11,57	0,42	99,65	1,75	0,24	89,0
KI75-1-5-o2	rim	1,5	X	38,16	0,00	0,04	0,09	41,03	0,25	0,26	19,56	0,32	99,74	1,58	0,42	80,6
KI75-1-5-b1	core	1,5		39,11	0,02	0,05	0,12	45,67	0,27	0,15	13,50	0,32	99,24	1,72	0,29	87,0
KI75-1-5-c1	core	1,5		38,94	0,01	0,06	0,01	45,08	0,25	0,14	14,45	0,38	99,35	1,70	0,31	86,1
KI75-1-5-c2	rim	1,5		39,03	0,02	0,05	0,03	44,86	0,25	0,26	15,06	0,32	99,91	1,69	0,32	85,5
KI75-1-5-d1	core	1,5		39,59	0,01	0,05	0,12	46,94	0,26	0,16	11,63	0,37	99,15	1,75	0,24	88,9
KI75-1-5-d2	rim	1,5		37,85	0,01	0,04	0,04	40,59	0,26	0,24	19,83	0,27	99,16	1,58	0,43	80,2
KI75-1-5-j1	core	1,5		39,42	0,00	0,05	0,05	47,17	0,25	0,10	11,71	0,46	99,24	1,76	0,25	88,9
KI75-1-5-j2	rim	1,5		38,83	0,00	0,07	0,01	44,24	0,27	0,21	15,33	0,31	99,27	1,68	0,33	85,1
KI75-1-5-k1	core	1,5		39,34	0,00	0,06	0,04	46,93	0,24	0,14	11,97	0,40	99,12	1,75	0,25	88,6
KI75-1-5-k2	rim	1,5		38,40	0,01	0,07	0,03	41,59	0,26	0,29	18,78	0,29	99,73	1,60	0,40	81,4
KI75-1-5-l1	core	1,5		39,22	0,03	0,07	0,10	46,29	0,27	0,25	12,92	0,31	99,48	1,73	0,27	87,7
KI75-1-5-l2	rim	1,5		38,18	0,02	0,05	0,00	40,25	0,29	0,30	20,39	0,24	99,72	1,56	0,44	79,6
KI75-1-5-m1	core	1,5		39,81	0,00	0,05	0,07	46,86	0,26	0,11	12,48	0,26	99,94	1,74	0,26	88,1
KI75-1-5-m2	rim	1,5		38,99	0,02	0,06	0,05	43,92	0,26	0,22	15,94	0,35	99,84	1,66	0,34	84,5
KI75-1-5-n1	core	1,5		39,77	0,02	0,15	0,02	46,86	0,24	0,16	12,00	0,43	99,65	1,74	0,25	88,6
KI75-1-5-n2	rim	1,5		37,88	0,00	0,05	0,00	39,68	0,27	0,37	20,92	0,23	99,43	1,54	0,46	79,0
KI75-1-5-p1	core	1,5		39,41	0,01	0,08	0,02	47,12	0,25	0,09	11,96	0,42	99,38	1,76	0,25	88,6
KI75-1-5-p2	rim	1,5		38,32	0,00	0,08	0,05	41,87	0,25	0,24	18,54	0,32	99,68	1,61	0,40	81,7
KI75-1-11-b1	core	3,4	X	39,45	0,00	0,09	0,12	47,54	0,27	0,12	12,20	0,35	100,19	1,76	0,25	88,6
KI75-1-11-b2	rim	3,4	X	38,07	0,01	0,05	0,08	39,85	0,23	0,30	21,46	0,26	100,33	1,54	0,47	78,6
KI75-1-11-i1	core	3,4	X	39,61	0,02	0,04	0,00	45,82	0,28	0,16	13,28	0,33	99,57	1,71	0,28	87,2
KI75-1-11-i2	rim	3,4	X	36,92	0,02	0,04	0,05	35,94	0,24	0,31	25,87	0,20	99,63	1,43	0,58	73,4
KI75-1-11-k1	core	3,4	X	39,78	0,00	0,05	0,14	46,59	0,26	0,13	12,55	0,41	99,91	1,73	0,26	88,0
KI75-1-11-k2	rim	3,4	X	37,12	0,01	0,04	0,03	35,99	0,23	0,31	25,75	0,22	99,71	1,43	0,57	73,5
KI75-1-11-p1	core	3,4	X	39,17	0,02	0,04	0,00	45,03	0,26	0,19	14,11	0,40	99,25	1,70	0,30	86,4
KI75-1-11-p2	rim	3,4	X	37,20	0,00	0,06	0,00	37,18	0,23	0,25	23,92	0,25	99,18	1,47	0,53	75,5
KI75-1-11-a1	core	3,4		37,76	0,01	0,05	0,04	39,42	0,27	0,28	21,69	0,27	99,85	1,53	0,47	78,3
KI75-1-11-a2	rim	3,4		36,89	0,02	0,03	0,00	35,42	0,22	0,34	26,58	0,25	99,86	1,41	0,59	72,5

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI75-1-11-c1	core	3,4		38,47	0,01	0,03	0,05	40,48	0,29	0,25	20,80	0,28	100,68	1,55	0,45	79,4
KI75-1-11-c2	rim	3,4		37,48	0,01	0,00	0,07	36,26	0,24	0,25	25,67	0,24	100,29	1,43	0,57	73,7
KI75-1-11-d1	core	3,4		38,72	0,00	0,04	0,02	44,51	0,27	0,18	15,59	0,31	99,66	1,68	0,33	85,0
KI75-1-11-d2	rim	3,4		37,26	0,01	0,00	0,00	36,26	0,24	0,36	24,94	0,22	99,45	1,44	0,56	74,2
KI75-1-11-e1	core	3,4		39,54	0,00	0,04	0,08	46,84	0,29	0,16	13,02	0,35	100,45	1,74	0,27	87,7
KI75-1-11-e2	rim	3,4		37,08	0,01	0,03	0,01	36,77	0,22	0,46	24,85	0,21	99,64	1,46	0,55	74,6
KI75-1-11-f1	core	3,4		39,49	0,01	0,06	0,06	47,02	0,27	0,19	11,21	0,34	98,64	1,76	0,24	89,3
KI75-1-11-f2	rim	3,4		37,32	0,02	0,01	0,01	36,79	0,26	0,29	24,80	0,20	99,80	1,45	0,55	74,6
KI75-1-11-g1	core	3,4		39,43	0,01	0,00	0,13	46,85	0,28	0,23	12,59	0,38	99,90	1,74	0,26	88,0
KI75-1-11-g2	rim	3,4		37,57	0,04	0,00	0,00	37,56	0,28	0,22	23,55	0,26	99,52	1,48	0,52	75,9
KI75-1-11-h1	core	3,4		39,86	0,01	0,05	0,04	46,77	0,29	0,17	12,13	0,37	99,71	1,74	0,25	88,4
KI75-1-11-h2	rim	3,4		37,96	0,02	0,01	0,00	38,78	0,24	0,25	22,15	0,24	99,67	1,51	0,48	77,6
KI75-1-11-j1	core	3,4		39,36	0,01	0,01	0,11	45,12	0,26	0,21	14,29	0,37	99,79	1,69	0,30	86,2
KI75-1-11-j2	rim	3,4		37,37	0,01	0,04	0,06	38,04	0,25	0,24	23,10	0,28	99,43	1,50	0,51	76,5
KI75-1-11-x1	core	3,4		39,31	0,00	0,15	0,10	45,12	0,27	0,22	14,28	0,31	99,80	1,69	0,30	86,2
KI75-1-11-x2	rim	3,4		37,52	0,02	0,00	0,03	37,44	0,23	0,28	23,84	0,26	99,75	1,47	0,53	75,7
KI75-1-11-l1	core	3,4		39,54	0,01	0,09	0,11	45,07	0,26	0,20	13,73	0,33	99,42	1,69	0,29	86,7
KI75-1-11-l2	rim	3,4		37,61	0,00	0,03	0,03	36,82	0,25	0,35	23,47	0,25	98,86	1,46	0,52	75,7
KI75-1-11-n1	core	3,4		39,61	0,01	0,02	0,09	45,94	0,28	0,17	12,89	0,33	99,36	1,72	0,27	87,6
KI75-1-11-n2	rim	3,4		37,10	0,01	0,10	0,01	36,69	0,23	0,29	23,92	0,19	98,57	1,46	0,53	75,2
KI75-1-11-o1	core	3,4		39,59	0,00	0,23	0,02	45,62	0,29	0,11	12,99	0,38	99,22	1,71	0,27	87,4
KI75-1-11-o2	rim	3,4		37,12	0,01	0,00	0,03	38,48	0,23	0,28	22,31	0,18	98,66	1,52	0,49	77,4
KI75-1-25-a1	core	7,6	X	39,83	0,01	0,05	0,14	47,28	0,19	0,14	11,16	0,39	99,21	1,76	0,23	89,3
KI75-1-25-a2	rim	7,6	X	39,96	0,00	0,04	0,04	47,42	0,25	0,12	11,68	0,37	99,91	1,75	0,24	89,0
KI75-1-25-c1	core	7,6	X	39,57	0,00	0,03	0,10	46,49	0,23	0,09	12,48	0,42	99,42	1,74	0,26	88,1
KI75-1-25-c2	rim	7,6	X	38,08	0,01	0,04	0,03	39,52	0,24	0,30	20,51	0,26	99,06	1,54	0,45	79,2
KI75-1-25-j1	core	7,6	X	39,51	0,01	0,03	0,17	44,96	0,23	0,21	14,42	0,33	99,86	1,69	0,30	86,1
KI75-1-25-j2	rim	7,6	X	37,22	0,03	0,03	0,00	36,00	0,16	0,34	25,74	0,28	99,84	1,43	0,57	73,5
KI75-1-25-b1	core	7,6		38,80	0,00	0,03	0,07	44,04	0,28	0,10	15,26	0,33	98,99	1,67	0,33	85,1
KI75-1-25-b2	rim	7,6		37,89	0,00	0,02	0,01	38,16	0,24	0,41	22,62	0,27	99,62	1,49	0,50	77,0
KI75-1-25-d1	core	7,6		39,05	0,00	0,03	0,00	43,34	0,21	0,24	15,88	0,31	99,12	1,65	0,34	84,4
KI75-1-25-d2	rim	7,6		37,48	0,02	0,02	0,02	37,46	0,19	0,17	22,82	0,30	98,56	1,48	0,51	76,5
KI75-1-25-e1	core	7,6		40,05	0,00	0,09	0,12	47,03	0,35	0,13	11,13	0,43	99,33	1,75	0,23	89,3
KI75-1-25-e2	rim	7,6		38,25	0,03	0,03	0,03	39,89	0,24	0,33	19,78	0,29	98,88	1,55	0,43	80,0
KI75-1-25-f1	core	7,6		38,67	0,01	0,05	0,12	41,51	0,26	0,26	18,79	0,30	99,95	1,59	0,40	81,4
KI75-1-25-f2	rim	7,6		38,14	0,02	0,01	0,04	39,29	0,21	0,34	21,25	0,27	99,62	1,53	0,46	78,6
KI75-1-25-g1	core	7,6		37,99	0,00	0,01	0,00	40,99	0,24	0,19	19,73	0,35	99,50	1,58	0,43	80,4
KI75-1-25-g2	rim	7,6		38,62	0,01	0,03	0,04	39,40	0,21	0,19	20,35	0,35	99,27	1,53	0,44	79,3
KI75-1-25-h1	core	7,6		39,31	0,00	0,03	0,09	44,37	0,26	0,26	14,01	0,34	98,68	1,68	0,30	86,2

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI75-1-25-h2	rim	7,6		38,06	0,00	0,02	0,02	39,61	0,23	0,23	21,11	0,29	99,61	1,54	0,46	78,8
KI75-1-25-i1	core	7,6		39,57	0,00	0,01	0,00	45,85	0,28	0,09	12,63	0,34	98,81	1,72	0,27	87,8
KI75-1-25-i2	rim	7,6		38,14	0,01	0,01	0,02	39,98	0,22	0,32	20,82	0,29	99,82	1,55	0,45	79,2
KI75-1-27-c1	core	8,2	X	39,77	0,01	0,07	0,11	46,11	0,27	0,17	12,62	0,32	99,46	1,72	0,26	87,9
KI75-1-27-c2	rim	8,2	X	37,65	0,01	0,05	0,02	38,20	0,21	0,29	22,36	0,27	99,15	1,50	0,49	77,2
KI75-1-27-d1	core	8,2	X	39,56	0,00	0,15	0,11	45,85	0,27	0,15	13,36	0,34	99,86	1,71	0,28	87,2
KI75-1-27-d2	rim	8,2	X	38,04	0,00	0,00	0,07	39,71	0,20	0,27	22,20	0,26	100,81	1,53	0,48	78,0
KI75-1-27-g1	core	8,2	X	39,62	0,01	0,03	0,06	46,70	0,27	0,15	11,71	0,39	98,95	1,75	0,25	88,8
KI75-1-27-g2	rim	8,2	X	37,45	0,03	0,04	0,02	37,42	0,21	0,31	24,11	0,30	99,93	1,47	0,53	75,4
KI75-1-27-i1	core	8,2	X	39,61	0,01	0,03	0,09	46,13	0,27	0,21	13,04	0,37	99,82	1,72	0,27	87,5
KI75-1-27-i2	rim	8,2	X	38,27	0,01	0,00	0,10	38,95	0,22	0,30	21,33	0,28	99,51	1,52	0,47	78,4
KI75-1-27-k1	core	8,2	X	39,61	0,00	0,23	0,07	46,25	0,26	0,17	13,18	0,45	100,28	1,72	0,28	87,4
KI75-1-27-k2	rim	8,2	X	37,52	0,00	0,15	0,01	36,19	0,21	0,35	25,21	0,25	99,91	1,43	0,56	74,0
KI75-1-27-a1	core	8,2		39,57	0,02	0,03	0,04	45,54	0,26	0,15	13,05	0,35	99,02	1,71	0,28	87,4
KI75-1-27-a2	rim	8,2		37,63	0,01	0,09	0,00	39,51	0,22	0,31	20,64	0,26	98,77	1,55	0,45	79,1
KI75-1-27-b1	core	8,2		39,52	0,02	0,06	0,04	45,83	0,29	0,06	13,07	0,36	99,27	1,72	0,28	87,4
KI75-1-27-b2	rim	8,2		39,08	0,00	0,07	0,02	43,14	0,26	0,22	16,59	0,32	99,80	1,64	0,35	83,7
KI75-1-27-e1	core	8,2		39,70	0,01	0,02	0,03	45,84	0,29	0,16	13,03	0,35	99,48	1,71	0,27	87,5
KI75-1-27-e2	rim	8,2		37,66	0,04	0,03	0,00	36,66	0,22	0,30	23,85	0,26	99,03	1,45	0,53	75,3
KI75-1-27-f1	core	8,2		39,26	0,00	0,01	0,08	44,55	0,31	0,17	14,59	0,36	99,35	1,68	0,31	85,8
KI75-1-27-f2	rim	8,2		37,06	0,06	0,00	0,01	35,53	0,19	0,31	25,56	0,29	99,03	1,42	0,57	73,3
KI75-1-27-h1	core	8,2		39,07	0,00	0,09	0,07	43,85	0,26	0,20	15,21	0,29	99,06	1,66	0,32	85,1
KI75-1-27-h2	rim	8,2		37,90	0,00	0,04	0,00	38,23	0,25	0,30	22,78	0,22	99,76	1,49	0,50	76,9
KI75-1-27-x1	core	8,2		38,92	0,00	0,04	0,04	43,51	0,30	0,24	15,78	0,32	99,19	1,65	0,34	84,5
KI75-1-27-x2	rim	8,2		37,54	0,04	0,00	0,03	37,27	0,21	0,25	24,34	0,25	99,93	1,47	0,54	75,2
KI75-1-60-a1	core	18,3	X	37,88	0,01	0,17	0,03	37,64	0,28	0,24	23,01	0,22	99,50	1,48	0,51	76,4
KI75-1-60-a2	rim	18,3	X	36,86	0,00	0,02	0,00	34,03	0,22	0,35	27,91	0,15	99,56	1,37	0,63	70,7
KI75-1-60-c1	core	18,3	X	39,17	0,00	0,03	0,00	43,80	0,26	0,24	16,27	0,31	100,07	1,65	0,35	84,2
KI75-1-60-c2	rim	18,3	X	37,00	0,01	0,00	0,02	34,85	0,22	0,31	27,26	0,18	99,90	1,39	0,61	71,7
KI75-1-60-d1	core	18,3	X	38,09	0,00	0,04	0,00	39,52	0,30	0,37	22,03	0,24	100,62	1,53	0,48	78,0
KI75-1-60-d2	rim	18,3	X	36,67	0,00	0,01	0,00	33,51	0,24	0,22	28,75	0,21	99,63	1,35	0,65	69,8
KI75-1-60-i1	core	18,3	X	38,70	0,00	0,06	0,00	42,03	0,26	0,23	18,29	0,32	99,91	1,61	0,39	82,0
KI75-1-60-i2	rim	18,3	X	37,08	0,01	0,03	0,06	34,67	0,21	0,31	26,73	0,18	99,32	1,39	0,60	72,0
KI75-1-60-b1	core	18,3		38,50	0,03	0,05	0,04	41,25	0,27	0,23	18,97	0,28	99,63	1,59	0,41	81,2
KI75-1-60-b2	rim	18,3		37,58	0,03	0,02	0,02	36,07	0,24	0,32	24,55	0,25	99,13	1,43	0,55	74,4
KI75-1-60-e1	core 1	18,3		38,40	0,00	0,07	0,03	41,16	0,27	0,25	19,50	0,27	99,98	1,58	0,42	80,7
KI75-1-60-e2	rim 1	18,3		37,46	0,03	0,04	0,00	36,72	0,26	0,30	24,70	0,25	99,79	1,45	0,55	74,6
KI75-1-60-e3	core 2	18,3		38,52	0,00	0,03	0,02	40,63	0,29	0,25	20,25	0,28	100,30	1,56	0,44	79,9

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI75-1-60-e4	rim 2	18,3		36,55	0,01	0,03	0,03	32,16	0,23	0,34	30,59	0,21	100,19	1,30	0,69	67,5
KI75-1-60-f1	core	18,3		38,10	0,01	0,05	0,00	39,39	0,28	0,25	21,88	0,25	100,19	1,53	0,48	78,1
KI75-1-60-f2	rim	18,3		36,22	0,02	0,00	0,00	30,65	0,22	0,43	31,25	0,17	98,96	1,26	0,72	66,0
KI75-1-60-g1	core	18,3		36,93	0,00	0,01	0,02	34,50	0,25	0,32	27,72	0,21	99,99	1,38	0,62	71,2
KI75-1-60-g2	rim	18,3		35,89	0,02	0,02	0,01	29,58	0,24	0,45	33,33	0,18	99,84	1,22	0,77	63,7
KI75-1-60-h1	core	18,3		37,71	0,02	0,04	0,01	38,11	0,30	0,30	23,66	0,20	100,36	1,49	0,52	76,1
KI75-1-60-h2	rim	18,3		37,01	0,03	0,02	0,00	34,42	0,24	0,36	27,17	0,18	99,47	1,38	0,61	71,5
KI75-1-130-c1	core	39,6	X	38,30	0,01	0,00	0,00	40,16	0,28	0,22	20,98	0,27	100,29	1,55	0,45	79,1
KI75-1-130-c2	rim	39,6	X	37,12	0,03	0,00	0,01	33,33	0,21	0,33	28,31	0,22	99,60	1,34	0,64	70,0
KI75-1-130-d1	core	39,6	X	37,68	0,01	0,04	0,06	40,89	0,29	0,22	19,39	0,31	98,88	1,59	0,42	80,7
KI75-1-130-d2	rim	39,6	X	37,54	0,03	0,03	0,08	36,03	0,21	0,39	25,17	0,18	99,70	1,43	0,56	73,9
KI75-1-130-k1	core	39,6	X	38,38	0,01	0,04	0,07	41,47	0,30	0,23	19,13	0,29	99,90	1,59	0,41	81,1
KI75-1-130-k2b	rim a	39,6	X	36,25	0,04	0,00	0,02	33,45	0,23	0,41	29,19	0,24	99,93	1,35	0,66	69,4
KI75-1-130-k3b	rim b	39,6	X	36,89	0,00	0,00	0,00	35,24	0,22	0,40	26,47	0,20	99,50	1,41	0,59	72,5
KI75-1-130-a1	core	39,6		38,00	0,00	0,02	0,02	40,61	0,29	0,18	19,72	0,30	99,14	1,58	0,43	80,3
KI75-1-130-a2	rim	39,6		37,36	0,00	0,00	0,00	38,98	0,29	0,26	20,95	0,29	98,28	1,54	0,46	78,7
KI75-1-130-b1	core	39,6		37,42	0,02	0,00	0,06	38,20	0,28	0,29	22,76	0,26	99,38	1,50	0,50	76,9
KI75-1-130-b2	rim	39,6		36,11	0,02	0,00	0,00	32,63	0,20	0,36	29,53	0,18	99,09	1,33	0,68	68,6
KI75-1-130-e1	core	39,6		37,95	0,00	0,00	0,00	40,04	0,27	0,23	20,99	0,22	99,71	1,55	0,46	79,1
KI75-1-130-e2	rim	39,6		35,10	0,13	0,00	0,01	29,68	0,18	0,35	33,98	0,19	99,71	1,23	0,79	63,4
KI75-1-130-f1	core	39,6		38,39	0,01	0,03	0,09	41,47	0,23	0,28	18,57	0,30	99,39	1,60	0,40	81,5
KI75-1-130-f2	rim	39,6		36,03	0,01	0,15	0,00	30,71	0,22	0,43	31,61	0,21	99,37	1,26	0,73	65,8
KI75-1-130-g1	core	39,6		37,90	0,01	0,03	0,08	40,61	0,28	0,30	20,11	0,23	99,60	1,57	0,44	80,0
KI75-1-130-g2	rim	39,6		36,35	0,02	0,01	0,02	33,77	0,22	0,39	28,16	0,15	99,18	1,37	0,64	70,4
KI75-1-130-h1	core	39,6		38,19	0,00	0,04	0,00	40,39	0,29	0,24	20,30	0,27	99,87	1,56	0,44	79,8
KI75-1-130-h2	rim	39,6		36,81	0,02	0,00	0,03	34,48	0,22	0,36	27,95	0,20	100,11	1,38	0,63	70,9
KI75-1-130-i1	core	39,6		36,96	0,01	0,00	0,00	35,60	0,25	0,23	26,08	0,17	99,35	1,42	0,58	73,0
KI75-1-130-i2	rim	39,6		36,09	0,02	0,00	0,00	31,26	0,24	0,33	31,39	0,19	99,57	1,28	0,72	66,4
KI75-1-130-j1	core	39,6		38,06	0,00	0,17	0,00	41,68	0,25	0,21	18,49	0,31	99,21	1,61	0,40	81,7
KI75-1-130-j2	rim	39,6		36,29	0,01	0,17	0,00	32,29	0,20	0,35	30,23	0,16	99,76	1,31	0,69	67,9
KI75-1-130-l1	core	39,6		37,36	0,01	0,00	0,03	38,65	0,31	0,31	22,51	0,21	99,40	1,52	0,50	77,3
KI75-1-130-l2	rim	39,6		36,67	0,00	0,00	0,00	34,03	0,21	0,38	28,37	0,24	99,98	1,37	0,64	70,4
KI75-1-141-a1	core	43,0	X	38,28	0,00	0,00	0,03	39,31	0,29	0,34	20,82	0,27	99,44	1,53	0,45	78,9
KI75-1-141-a2	rim	43,0	X	37,83	0,02	0,02	0,04	37,95	0,28	0,17	22,81	0,22	99,36	1,49	0,50	76,7
KI75-1-141-b1	core	43,0	X	38,36	0,00	0,04	0,06	40,43	0,32	0,30	20,15	0,30	100,00	1,56	0,44	79,9
KI75-1-141-b2	rim	43,0	X	37,54	0,03	0,02	0,00	36,37	0,24	0,29	24,57	0,20	99,42	1,44	0,55	74,6
KI75-1-141-g1	core	43,0	X	38,59	0,00	0,04	0,00	41,04	0,29	0,25	19,44	0,26	100,01	1,57	0,42	80,7
KI75-1-141-g2	rim	43,0	X	38,01	0,03	0,02	0,01	37,97	0,25	0,33	23,37	0,29	100,26	1,48	0,51	76,3

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI75-1-141-i1	core	43,0	X	38,52	0,00	0,03	0,00	40,06	0,29	0,28	20,94	0,25	100,43	1,54	0,45	79,1
KI75-1-141-i2	rim	43,0	X	37,86	0,01	0,03	0,04	38,21	0,25	0,28	23,24	0,27	100,21	1,49	0,51	76,5
KI75-1-141-c1	core	43,0		38,58	0,00	0,04	0,08	40,46	0,32	0,29	19,42	0,25	99,46	1,56	0,42	80,5
KI75-1-141-c2	rim	43,0		37,77	0,02	0,02	0,00	36,99	0,26	0,30	24,42	0,20	100,08	1,45	0,54	75,0
KI75-1-141-d1	core	43,0		38,39	0,01	0,01	0,03	38,98	0,28	0,17	22,08	0,25	100,23	1,51	0,48	77,8
KI75-1-141-d2	rim	43,0		37,90	0,03	0,00	0,03	37,09	0,25	0,33	24,21	0,24	100,08	1,45	0,53	75,2
KI75-1-141-e1	core	43,0		38,36	0,00	0,02	0,06	39,58	0,30	0,23	20,54	0,18	99,29	1,54	0,45	79,2
KI75-1-141-e2	rim	43,0		37,62	0,04	0,02	0,00	36,52	0,23	0,31	25,12	0,24	100,17	1,44	0,55	74,2
KI75-1-141-f1	core	43,0		38,58	0,00	0,03	0,14	40,29	0,29	0,34	20,04	0,26	100,08	1,55	0,43	79,9
KI75-1-141-f2	rim	43,0		37,82	0,01	0,00	0,08	37,62	0,27	0,35	24,20	0,24	100,69	1,47	0,53	75,5
KI75-1-141-h1	core	43,0		38,23	0,01	0,03	0,01	38,43	0,25	0,26	22,38	0,21	99,80	1,50	0,49	77,3
KI75-1-141-h2	rim	43,0		37,75	0,05	0,02	0,02	37,35	0,24	0,35	23,67	0,27	99,75	1,47	0,52	75,8
KI75-1-141-j1	core	43,0		38,25	0,01	0,04	0,04	39,67	0,27	0,37	20,41	0,31	99,37	1,54	0,44	79,4
KI75-1-141-j2	rim	43,0		37,84	0,01	0,02	0,00	37,31	0,26	0,29	23,74	0,25	99,74	1,46	0,52	75,7
KI75-1-141-k1	core	43,0		38,23	0,01	0,05	0,01	39,40	0,28	0,26	21,02	0,24	99,50	1,53	0,46	78,8
KI75-1-141-k2	rim	43,0		37,51	0,03	0,02	0,04	35,75	0,28	0,37	25,30	0,23	99,63	1,42	0,56	73,7
KI75-1-145-d1	core	44,2	X	38,83	0,00	0,05	0,03	40,94	0,26	0,22	18,83	0,30	99,45	1,57	0,41	81,1
KI75-1-145-d2b	rim a	44,2	X	38,11	0,02	0,00	0,00	38,72	0,27	0,40	21,67	0,23	99,48	1,51	0,47	78,0
KI75-1-145-d3b	rim b	44,2	X	38,47	0,02	0,02	0,00	39,25	0,27	0,32	21,09	0,28	99,87	1,52	0,46	78,7
KI75-1-145-h1	core	44,2	X	38,30	0,00	0,00	0,07	40,16	0,28	0,22	19,75	0,25	99,06	1,56	0,43	80,1
KI75-1-145-h2	rim	44,2	X	37,98	0,02	0,01	0,06	38,19	0,27	0,31	21,47	0,22	98,59	1,50	0,47	77,9
KI75-1-145-m1	core	44,2	X	38,45	0,00	0,02	0,06	39,91	0,28	0,32	19,45	0,32	98,83	1,55	0,42	80,2
KI75-1-145-m2	rim	44,2	X	38,02	0,00	0,04	0,03	38,69	0,25	0,43	21,12	0,30	98,89	1,51	0,46	78,4
KI75-1-145-a1	core	44,2		38,48	0,00	0,03	0,05	39,74	0,28	0,22	20,72	0,29	99,88	1,54	0,45	79,2
KI75-1-145-a2	rim	44,2		38,09	0,00	0,00	0,01	38,87	0,26	0,36	21,81	0,24	99,64	1,51	0,48	77,9
KI75-1-145-b1	core	44,2		38,40	0,00	0,01	0,01	40,03	0,28	0,29	20,37	0,31	99,79	1,55	0,44	79,6
KI75-1-145-b2	rim	44,2		38,36	0,01	0,02	0,00	38,74	0,29	0,20	21,93	0,26	99,80	1,50	0,48	77,8
KI75-1-145-c1	core	44,2		38,39	0,02	0,03	0,00	39,86	0,28	0,31	20,96	0,32	100,24	1,54	0,45	79,0
KI75-1-145-c2	rim	44,2		38,30	0,04	0,04	0,10	39,12	0,28	0,31	22,31	0,23	100,77	1,51	0,48	77,6
KI75-1-145-e1	core	44,2		38,57	0,00	0,03	0,00	40,20	0,31	0,29	19,47	0,27	99,19	1,55	0,42	80,4
KI75-1-145-e2	rim	44,2		38,11	0,02	0,04	0,00	38,93	0,27	0,35	20,86	0,28	98,90	1,52	0,46	78,7
KI75-1-145-f1	core	44,2		38,24	0,01	0,01	0,04	39,56	0,29	0,29	19,83	0,28	98,55	1,54	0,43	79,8
KI75-1-145-f2	rim	44,2		38,18	0,02	0,04	0,03	39,25	0,27	0,28	21,70	0,25	100,10	1,52	0,47	78,2
KI75-1-145-g1	core	44,2		38,17	0,01	0,02	0,00	39,39	0,26	0,26	21,09	0,27	99,47	1,53	0,46	78,7
KI75-1-145-g2	rim	44,2		38,21	0,01	0,01	0,10	38,47	0,25	0,23	22,37	0,27	99,92	1,50	0,49	77,3
KI75-1-145-i1	core	44,2		38,58	0,00	0,05	0,15	41,33	0,27	0,23	18,22	0,28	99,18	1,59	0,39	81,8
KI75-1-145-i2	rim	44,2		38,25	0,02	0,03	0,09	38,98	0,27	0,26	20,95	0,30	99,25	1,52	0,46	78,6
KI75-1-145-j1	core	44,2		38,55	0,00	0,04	0,06	40,66	0,25	0,28	19,05	0,21	99,12	1,57	0,41	80,9
KI75-1-145-j2	rim	44,2		38,12	0,04	0,01	0,01	38,11	0,25	0,28	21,90	0,29	99,02	1,49	0,48	77,5

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI75-1-145-k1	core	44,2		38,22	0,01	0,02	0,04	39,69	0,29	0,26	19,93	0,30	98,83	1,55	0,44	79,8
KI75-1-145-k2	rim	44,2		38,05	0,03	0,00	0,02	38,55	0,27	0,26	22,32	0,27	99,82	1,50	0,49	77,4
KI75-1-145-l1	core	44,2		38,50	0,01	0,02	0,12	39,98	0,27	0,30	20,89	0,30	100,38	1,54	0,45	79,1
KI75-1-145-l2	rim	44,2		38,07	0,03	0,01	0,05	38,18	0,26	0,25	21,56	0,25	98,73	1,50	0,48	77,8
KI75-1-145-n1	core	44,2		38,61	0,02	0,00	0,03	40,17	0,29	0,27	19,14	0,21	98,75	1,56	0,42	80,6
KI75-1-145-n2	rim	44,2		38,06	0,05	0,03	0,00	39,04	0,26	0,32	22,05	0,30	100,14	1,52	0,48	77,8
KI75-1-145-o1	core	44,2		38,29	0,00	0,04	0,04	39,37	0,28	0,28	20,50	0,30	99,20	1,53	0,45	79,2
KI75-1-145-o2	rim	44,2		38,12	0,01	0,03	0,05	38,90	0,28	0,25	21,20	0,25	99,14	1,52	0,46	78,4
KI81-1-200.2-c1	core	61,0	X	38,69	0,01	0,00	0,03	41,40	0,28	0,22	18,76	0,30	99,75	1,59	0,40	81,4
KI81-1-200.2-c2	rim	61,0	X	38,58	0,03	0,07	0,00	40,87	0,27	0,29	19,12	0,29	99,60	1,57	0,41	80,9
KI81-1-200.2-d1	core	61,0	X	38,55	0,00	0,03	0,00	41,89	0,26	0,19	18,66	0,28	99,89	1,60	0,40	81,7
KI81-1-200.2-d2	rim	61,0	X	38,49	0,00	0,13	0,00	40,85	0,27	0,25	19,14	0,30	99,51	1,57	0,41	80,9
KI81-1-200.2-h1	core	61,0	X	38,52	0,01	0,07	0,00	41,18	0,27	0,32	19,18	0,29	99,82	1,58	0,41	81,0
KI81-1-200.2-h2	rim	61,0	X	38,45	0,01	0,19	0,00	40,98	0,27	0,25	19,45	0,27	99,91	1,57	0,42	80,7
KI81-1-200.2-j1	core	61,0	X	38,49	0,01	0,10	0,10	41,73	0,29	0,31	18,06	0,28	99,36	1,60	0,39	82,1
KI81-1-200.2-j2	rim	61,0	X	38,14	0,01	0,01	0,00	40,57	0,27	0,32	18,97	0,23	98,52	1,58	0,41	80,9
KI81-1-200.2-k1	core	61,0	X	38,57	0,00	0,17	0,07	42,31	0,27	0,28	17,89	0,29	99,88	1,61	0,38	82,4
KI81-1-200.2-k2	rim	61,0	X	38,21	0,01	0,00	0,00	40,71	0,26	0,37	19,60	0,29	99,53	1,57	0,43	80,4
KI81-1-200.2-l1	core	61,0	X	38,83	0,00	0,00	0,05	43,22	0,26	0,13	16,32	0,33	99,19	1,65	0,35	84,0
KI81-1-200.2-l2	rim 1	61,0	X	38,27	0,01	0,01	0,01	40,62	0,25	0,24	19,47	0,27	99,20	1,57	0,42	80,5
KI81-1-200.2-l3	rim 2	61,0	X	38,23	0,02	0,17	0,00	41,13	0,27	0,16	19,06	0,32	99,44	1,59	0,41	81,0
KI81-1-200.2-a1	core	61,0		38,50	0,01	0,50	0,00	41,55	0,24	0,31	18,89	0,26	100,29	1,58	0,40	81,3
KI81-1-200.2-a2	rim	61,0		38,17	0,03	0,04	0,00	40,46	0,28	0,27	19,12	0,27	98,68	1,57	0,42	80,7
KI81-1-200.2-b1	core	61,0		38,55	0,01	0,08	0,09	41,72	0,27	0,22	18,59	0,28	99,82	1,60	0,40	81,6
KI81-1-200.2-b2	rim	61,0		38,56	0,03	0,00	0,00	41,21	0,27	0,27	19,13	0,19	99,67	1,58	0,41	81,0
KI81-1-200.2-e1	core	61,0		38,10	0,01	0,34	0,02	40,48	0,26	0,34	19,31	0,25	99,13	1,57	0,42	80,6
KI81-1-200.2-e2	rim	61,0		38,28	0,03	0,24	0,00	40,69	0,28	0,19	19,53	0,29	99,55	1,57	0,42	80,5
KI81-1-200.2-f1	core	61,0		38,46	0,00	0,68	0,08	40,85	0,24	0,24	18,95	0,28	99,86	1,56	0,41	81,0
KI81-1-200.2-f2	rim	61,0		38,10	0,02	0,12	0,00	40,61	0,27	0,15	19,74	0,26	99,28	1,57	0,43	80,3
KI81-1-200.2-g1	core	61,0		38,36	0,04	0,07	0,00	40,78	0,26	0,24	19,37	0,30	99,45	1,57	0,42	80,7
KI81-1-200.2-g2	rim	61,0		38,20	0,05	0,07	0,03	40,69	0,27	0,30	19,68	0,27	99,57	1,57	0,43	80,4
KI81-1-200.2-i1	core	61,0		38,42	0,01	0,05	0,00	41,46	0,28	0,17	18,23	0,32	99,01	1,60	0,39	81,8
KI81-1-200.2-i2	rim	61,0		38,40	0,00	0,02	0,00	40,64	0,27	0,31	19,51	0,32	99,50	1,57	0,42	80,5
KI81-1-249.7-j1	core	76,1	X	38,89	0,01	0,04	0,02	43,24	0,24	0,23	16,19	0,34	99,20	1,65	0,35	84,1
KI81-1-249.7-j2	rim	76,1	X	38,54	0,01	0,20	0,04	41,94	0,27	0,13	17,80	0,28	99,22	1,61	0,38	82,3
KI81-1-249.7-i1	core	76,1	X	38,61	0,00	0,07	0,03	42,20	0,26	0,27	18,77	0,32	100,60	1,61	0,40	81,7
KI81-1-249.7-i2	rim	76,1	X	38,72	0,00	0,08	0,00	41,78	0,28	0,20	17,40	0,32	98,84	1,61	0,38	82,6
KI81-1-249.7-h1	core	76,1	X	38,26	0,00	0,17	0,00	41,95	0,25	0,28	17,88	0,32	99,12	1,61	0,39	82,3

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI81-1-249.7-h2	rim	76,1	X	38,30	0,00	0,04	0,02	41,85	0,27	0,18	18,15	0,27	99,08	1,61	0,39	82,0
KI81-1-249.7-g1	core	76,1	X	38,78	0,00	0,09	0,11	44,21	0,24	0,15	14,92	0,36	98,95	1,68	0,32	85,4
KI81-1-249.7-g2	rim	76,1	X	38,60	0,03	0,00	0,02	41,95	0,27	0,15	17,93	0,23	99,18	1,61	0,39	82,3
KI81-1-249.7-c1	core	76,1	X	38,41	0,02	0,02	0,00	41,92	0,25	0,23	18,64	0,24	99,75	1,61	0,40	81,7
KI81-1-249.7-c2	rim	76,1	X	38,69	0,01	0,02	0,04	42,19	0,26	0,25	18,66	0,26	100,43	1,61	0,40	81,8
KI81-1-249.7-k1	core	76,1		38,48	0,01	0,06	0,05	42,08	0,28	0,23	18,12	0,30	99,64	1,61	0,39	82,1
KI81-1-249.7-k2	rim	76,1		38,51	0,00	0,54	0,07	41,76	0,25	0,21	17,66	0,30	99,32	1,60	0,38	82,4
KI81-1-249.7-f1	core	76,1		38,17	0,01	0,00	0,08	41,91	0,26	0,29	18,31	0,28	99,37	1,61	0,40	81,9
KI81-1-249.7-f2	rim	76,1		38,15	0,03	0,00	0,00	41,73	0,27	0,30	18,19	0,26	99,00	1,61	0,39	82,0
KI81-1-249.7-e1	core	76,1		38,47	0,01	0,00	0,01	42,07	0,26	0,28	18,55	0,22	99,89	1,61	0,40	81,8
KI81-1-249.7-e2	rim	76,1		38,54	0,01	0,00	0,08	41,97	0,27	0,29	18,68	0,30	100,19	1,60	0,40	81,7
KI81-1-249.7-d1	core	76,1		38,61	0,01	0,04	0,06	42,18	0,26	0,28	18,38	0,31	100,16	1,61	0,39	82,0
KI81-1-249.7-d2	rim	76,1		38,57	0,02	0,00	0,05	41,91	0,27	0,29	18,29	0,30	99,68	1,60	0,39	81,9
KI81-1-249.7-b1	core	76,1		38,71	0,03	0,00	0,07	41,93	0,26	0,21	18,71	0,32	100,26	1,60	0,40	81,6
KI81-1-249.7-b2	rim	76,1		38,56	0,02	0,05	0,12	42,22	0,27	0,15	18,49	0,30	100,20	1,61	0,40	81,9
KI81-1-299.9-c1	core	91,4	X	37,70	0,00	0,10	0,02	38,23	0,23	0,39	22,90	0,30	99,90	1,49	0,50	76,8
KI81-1-299.9-c2	rim	91,4	X	37,73	0,03	0,80	0,00	37,91	0,23	0,32	22,95	0,25	100,23	1,47	0,50	76,6
KI81-1-299.9-d1	core	91,4	X	38,01	0,02	0,06	0,01	38,20	0,21	0,30	22,98	0,23	100,10	1,49	0,50	76,7
KI81-1-299.9-d2	rim	91,4	X	37,80	0,01	0,02	0,00	38,09	0,25	0,37	23,34	0,22	100,23	1,49	0,51	76,4
KI81-1-299.9-e1	core	91,4	X	37,72	0,00	0,01	0,03	37,90	0,22	0,30	22,68	0,26	99,14	1,49	0,50	76,8
KI81-1-299.9-e2	rim	91,4	X	37,65	0,00	0,00	0,00	37,84	0,25	0,36	23,04	0,26	99,44	1,49	0,51	76,5
KI81-1-299.9-f1	core	91,4	X	37,95	0,00	0,07	0,04	39,37	0,23	0,34	20,96	0,31	99,27	1,53	0,46	78,8
KI81-1-299.9-f2	rim	91,4	X	37,67	0,01	0,02	0,05	37,93	0,24	0,22	23,33	0,30	99,81	1,49	0,51	76,3
KI81-1-299.9-h1	core	91,4	X	39,03	0,01	0,03	0,09	44,14	0,25	0,18	15,80	0,31	99,88	1,67	0,34	84,7
KI81-1-299.9-h2	rim (outer)	91,4	X	37,83	0,01	0,00	0,00	38,43	0,23	0,26	22,14	0,28	99,24	1,51	0,49	77,5
KI81-1-299.9-h3	rim (inner)	91,4	X	38,52	0,00	0,04	0,02	41,32	0,23	0,21	18,86	0,26	99,54	1,59	0,41	81,3
KI81-1-299.9-j1	core	91,4	X	37,91	0,01	0,56	0,08	39,18	0,23	0,26	20,80	0,24	99,32	1,52	0,45	78,9
KI81-1-299.9-j2	rim	91,4	X	37,87	0,02	0,03	0,01	38,06	0,22	0,25	23,17	0,24	99,88	1,49	0,51	76,5
KI81-1-299.9-a1	core	91,4		37,81	0,00	0,24	0,00	38,70	0,24	0,29	22,07	0,26	99,61	1,51	0,48	77,6
KI81-1-299.9-a2	rim	91,4		37,57	0,04	0,03	0,05	37,99	0,26	0,34	22,99	0,27	99,67	1,49	0,51	76,6
KI81-1-299.9-b1	core	91,4		37,69	0,02	0,05	0,00	38,24	0,23	0,29	23,14	0,23	99,88	1,50	0,51	76,6
KI81-1-299.9-b2	rim	91,4		37,93	0,03	0,00	0,03	38,25	0,25	0,38	23,16	0,21	100,26	1,49	0,51	76,6
KI81-1-299.9-g1	core	91,4		37,77	0,01	0,07	0,06	38,95	0,23	0,34	22,03	0,26	99,84	1,52	0,48	77,8
KI81-1-299.9-g2	rim	91,4		37,72	0,00	0,16	0,01	38,00	0,25	0,30	22,94	0,29	99,73	1,49	0,50	76,6
KI81-1-299.9-i1	core	91,4		37,81	0,03	0,19	0,03	37,92	0,21	0,26	23,48	0,28	100,31	1,48	0,51	76,2
KI81-1-299.9-i2	rim	91,4		37,85	0,03	0,00	0,05	37,72	0,27	0,32	23,84	0,19	100,27	1,47	0,52	75,8
KI81-1-299.9-k1	core	91,4		38,53	0,01	0,08	0,00	41,40	0,26	0,20	18,96	0,32	99,87	1,59	0,41	81,2

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI81-2-185.7-b1	core	56,6	X	38,58	0,01	0,05	0,00	40,77	0,29	0,16	20,01	0,30	100,22	1,56	0,43	80,2
KI81-2-185.7-b2	rim	56,6	X	37,58	0,03	0,05	0,05	37,77	0,22	0,32	23,96	0,27	100,28	1,48	0,53	75,7
KI81-2-185.7-c1	core	56,6	X	38,51	0,00	0,12	0,04	41,23	0,27	0,27	19,55	0,28	100,30	1,58	0,42	80,7
KI81-2-185.7-c2	rim	56,6	X	37,71	0,02	0,04	0,03	37,89	0,23	0,33	23,36	0,25	99,87	1,48	0,51	76,3
KI81-2-185.7-e1	core	56,6	X	38,20	0,00	0,10	0,07	38,82	0,25	0,45	22,26	0,19	100,40	1,50	0,48	77,5
KI81-2-185.7-e2	rim	56,6	X	37,80	0,02	0,04	0,00	37,45	0,25	0,31	24,09	0,21	100,23	1,47	0,53	75,5
KI81-2-185.7-f1	core	56,6	X	38,92	0,01	0,07	0,00	42,63	0,27	0,19	16,83	0,31	99,27	1,63	0,36	83,4
KI81-2-185.7-f2	rim 1	56,6	X	37,89	0,00	0,15	0,10	38,89	0,24	0,21	22,59	0,26	100,36	1,51	0,49	77,3
KI81-2-185.7-f3	rim 2 (outer)	56,6	X	37,61	0,02	0,03	0,03	37,72	0,23	0,29	24,03	0,26	100,21	1,48	0,53	75,6
KI81-2-185.7-f4	rim 2 (inner)	56,6	X	38,57	0,00	0,09	0,09	41,67	0,29	0,32	18,60	0,30	99,97	1,59	0,40	81,6
KI81-2-185.7-i1	core	56,6	X	39,09	0,00	0,11	0,00	43,17	0,23	0,16	16,32	0,34	99,46	1,64	0,35	84,0
KI81-2-185.7-i2	rim	56,6	X	38,09	0,02	0,39	0,00	39,57	0,21	0,27	21,00	0,31	99,86	1,53	0,46	78,9
KI81-2-185.7-j1	core	56,6	X	38,51	0,00	0,05	0,01	41,30	0,27	0,27	18,96	0,31	99,76	1,59	0,41	81,2
KI81-2-185.7-j2	rim	56,6	X	37,85	0,02	0,12	0,06	38,65	0,24	0,33	22,32	0,22	99,82	1,51	0,49	77,4
KI81-2-185.7-a1	core	56,6		38,29	0,00	0,13	0,12	41,08	0,29	0,27	19,87	0,30	100,37	1,57	0,43	80,4
KI81-2-185.7-a2	rim	56,6		38,22	0,00	0,03	0,00	39,47	0,24	0,23	21,27	0,28	99,77	1,53	0,46	78,6
KI81-2-185.7-d1	core	56,6		38,29	0,00	0,02	0,00	41,58	0,26	0,23	19,18	0,31	99,89	1,60	0,41	81,1
KI81-2-185.7-d2	rim	56,6		38,00	0,01	0,05	0,00	39,57	0,24	0,31	21,41	0,23	99,84	1,54	0,47	78,5
KI81-2-185.7-g1	core	56,6		37,88	0,02	0,62	0,00	40,07	0,24	0,38	20,24	0,30	99,79	1,55	0,44	79,7
KI81-2-185.7-g2	rim	56,6		37,97	0,02	0,01	0,00	39,35	0,22	0,26	21,62	0,29	99,74	1,53	0,47	78,3
KI81-2-185.7-h1	core	56,6		38,17	0,00	0,46	0,03	41,14	0,28	0,21	19,31	0,25	99,93	1,58	0,42	80,8
KI81-2-185.7-h2	rim	56,6		37,66	0,00	0,00	0,00	38,15	0,23	0,34	22,80	0,19	99,40	1,50	0,50	76,8
KI81-2-185.7-k1	core	56,6		37,94	0,01	0,02	0,00	40,19	0,27	0,24	20,97	0,30	99,99	1,56	0,46	79,2
KI81-2-185.7-k2	rim	56,6		37,72	0,02	0,01	0,10	38,44	0,23	0,37	23,19	0,23	100,38	1,50	0,51	76,6
KI88-2-197.7-c1	core	60,3	X	38,63	0,00	0,02	0,03	41,76	0,27	0,20	18,53	0,28	99,83	1,60	0,40	81,7
KI88-2-197.7-c2	rim	60,3	X	37,73	0,02	0,13	0,00	38,53	0,22	0,37	21,99	0,25	99,26	1,51	0,48	77,6
KI88-2-197.7-d1	core	60,3	X	38,79	0,00	0,02	0,09	42,10	0,26	0,26	17,83	0,30	99,72	1,61	0,38	82,4
KI88-2-197.7-d2	rim	60,3	X	37,74	0,02	0,01	0,00	37,77	0,18	0,28	23,29	0,25	99,56	1,48	0,51	76,2
KI88-2-197.7-e1	core	60,3	X	38,06	0,00	0,07	0,04	40,05	0,28	0,27	20,42	0,20	99,40	1,55	0,44	79,5
KI88-2-197.7-e2	rim	60,3	X	37,43	0,02	0,00	0,00	37,27	0,18	0,16	23,86	0,28	99,25	1,47	0,53	75,6
KI88-2-197.7-i1	core	60,3	X	38,69	0,00	0,20	0,04	41,18	0,26	0,25	18,78	0,31	99,74	1,58	0,40	81,3
KI88-2-197.7-i2	rim	60,3	X	37,63	0,00	1,47	0,01	37,51	0,20	0,28	22,80	0,29	100,23	1,46	0,50	76,5
KI88-2-197.7-j1	core	60,3	X	39,37	0,02	0,08	0,07	43,70	0,23	0,15	15,96	0,31	99,91	1,65	0,34	84,4
KI88-2-197.7-j2	rim 1	60,3	X	38,16	0,00	0,02	0,00	40,10	0,24	0,18	19,99	0,27	98,96	1,56	0,44	79,9
KI88-2-197.7-j3	rim 2	60,3	X	37,80	0,02	0,05	0,03	38,81	0,21	0,34	21,46	0,23	99,03	1,52	0,47	78,2
KI88-2-197.7-a1	core	60,3		38,16	0,01	0,07	0,02	39,55	0,26	0,26	20,58	0,27	99,18	1,54	0,45	79,2
KI88-2-197.7-a2	rim	60,3		37,62	0,00	0,00	0,02	38,04	0,20	0,38	23,32	0,23	99,83	1,49	0,51	76,3
KI88-2-197.7-b1	core	60,3		38,28	0,00	0,07	0,00	40,91	0,27	0,21	19,04	0,32	99,14	1,58	0,41	81,0
KI88-2-197.7-b2	rim	60,3		38,05	0,02	0,05	0,01	38,88	0,21	0,26	21,89	0,27	99,70	1,51	0,48	77,9

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI88-2-197.7-f1	core	60,3		38,09	0,01	0,47	0,01	38,32	0,22	0,36	23,43	0,31	101,29	1,48	0,51	76,4
KI88-2-197.7-f2	rim	60,3		37,58	0,03	0,00	0,03	37,15	0,15	0,33	23,89	0,22	99,39	1,47	0,53	75,5
KI88-2-197.7-g1	core	60,3		38,61	0,00	0,03	0,11	41,85	0,26	0,23	18,14	0,25	99,52	1,60	0,39	82,0
KI88-2-197.7-g2	rim	60,3		37,90	0,02	0,14	0,00	38,62	0,22	0,40	22,31	0,28	99,92	1,50	0,49	77,4
KI88-2-197.7-h1	core	60,3		37,83	0,00	0,02	0,00	39,06	0,24	0,30	21,48	0,27	99,26	1,53	0,47	78,3
KI88-2-197.7-h2	rim	60,3		37,49	0,02	0,05	0,00	36,50	0,16	0,33	25,27	0,32	100,22	1,44	0,56	74,1
KI88-2-249.5-a1	core	76,0	X	38,57	0,00	0,02	0,00	40,81	0,26	0,32	19,66	0,31	99,98	1,57	0,42	80,5
KI88-2-249.5-a2	rim	76,0	X	38,16	0,02	0,09	0,00	40,33	0,20	0,28	20,56	0,28	99,99	1,56	0,45	79,5
KI88-2-249.5-c1	core	76,0	X	39,14	0,01	0,08	0,05	43,85	0,23	0,22	15,91	0,35	99,85	1,66	0,34	84,5
KI88-2-249.5-c2	rim	76,0	X	38,30	0,02	0,00	0,03	39,95	0,19	0,24	20,33	0,33	99,40	1,55	0,44	79,6
KI88-2-249.5-e1	core	76,0	X	38,30	0,02	0,10	0,00	40,56	0,24	0,21	19,87	0,23	99,54	1,57	0,43	80,2
KI88-2-249.5-e2	rim	76,0	X	38,21	0,01	0,16	0,00	39,65	0,20	0,28	20,80	0,25	99,58	1,54	0,45	79,1
KI88-2-249.5-h1	core	76,0	X	38,71	0,01	0,08	0,00	41,78	0,26	0,22	18,46	0,27	99,79	1,60	0,40	81,8
KI88-2-249.5-h2	rim	76,0	X	38,50	0,02	0,04	0,05	40,71	0,21	0,21	20,65	0,24	100,67	1,56	0,44	79,6
KI88-2-249.5-k1	core	76,0	X	38,47	0,01	0,07	0,02	41,28	0,27	0,34	18,87	0,30	99,65	1,59	0,41	81,2
KI88-2-249.5-k2	rim	76,0	X	38,19	0,03	0,04	0,04	40,22	0,20	0,18	20,85	0,32	100,12	1,55	0,45	79,3
KI88-2-249.5-l1	core	76,0	X	38,62	0,00	0,05	0,05	41,37	0,27	0,21	18,48	0,30	99,40	1,59	0,40	81,6
KI88-2-249.5-l2	rim	76,0	X	38,12	0,02	0,02	0,04	40,64	0,20	0,32	20,57	0,32	100,28	1,57	0,44	79,7
KI88-2-249.5-b1	core	76,0		38,54	0,01	0,15	0,00	40,96	0,24	0,31	19,81	0,29	100,33	1,57	0,43	80,4
KI88-2-249.5-b2	rim	76,0		38,17	0,02	0,10	0,00	40,31	0,21	0,28	20,64	0,25	100,04	1,56	0,45	79,4
KI88-2-249.5-d1	core	76,0		38,57	0,01	0,03	0,04	41,37	0,27	0,19	19,01	0,30	99,84	1,59	0,41	81,2
KI88-2-249.5-d2	rim	76,0		38,25	0,02	0,14	0,00	39,78	0,22	0,33	20,69	0,22	99,67	1,54	0,45	79,2
KI88-2-249.5-f1	core	76,0		38,65	0,00	0,08	0,08	41,56	0,27	0,28	18,31	0,29	99,53	1,59	0,39	81,8
KI88-2-249.5-f2	rim	76,0		37,97	0,00	0,05	0,05	40,24	0,22	0,30	20,05	0,28	99,21	1,56	0,44	79,9
KI88-2-249.5-g1	core	76,0		38,43	0,00	0,05	0,02	40,79	0,24	0,20	19,56	0,27	99,61	1,57	0,42	80,5
KI88-2-249.5-g2	rim	76,0		38,31	0,03	0,00	0,03	40,29	0,21	0,26	20,43	0,31	99,91	1,55	0,44	79,6
KI88-2-249.5-i1	core	76,0		37,88	0,00	0,05	0,00	39,47	0,19	0,21	20,75	0,29	98,88	1,54	0,46	79,0
KI88-2-249.5-i2	rim	76,0		38,07	0,01	0,03	0,07	39,66	0,19	0,31	20,47	0,25	99,13	1,54	0,45	79,3
KI88-2-249.5-j1	core	76,0		38,25	0,00	0,00	0,00	39,93	0,20	0,29	20,57	0,23	99,49	1,55	0,45	79,4
KI88-2-249.5-j2	rim	76,0		38,30	0,02	0,02	0,00	40,03	0,21	0,23	21,38	0,25	100,44	1,54	0,46	78,8
KI88-2-296.1-a1	core	90,3	X	37,47	0,03	0,15	0,00	37,02	0,22	0,35	24,96	0,25	100,44	1,45	0,55	74,6
KI88-2-296.1-a2	rim	90,3	X	37,62	0,01	0,01	0,00	36,63	0,23	0,31	24,15	0,28	99,25	1,45	0,54	75,0
KI88-2-296.1-b1	core	90,3	X	38,12	0,01	0,01	0,06	38,94	0,21	0,25	20,77	0,25	98,63	1,52	0,46	78,8
KI88-2-296.1-b2	rim	90,3	X	37,50	0,01	0,27	0,02	37,05	0,21	0,31	25,38	0,28	101,10	1,45	0,56	74,3
KI88-2-296.1-f1	core	90,3	X	37,38	0,00	0,03	0,05	36,81	0,22	0,27	24,30	0,23	99,36	1,46	0,54	75,0
KI88-2-296.1-f2	rim	90,3	X	37,32	0,04	0,09	0,00	36,32	0,20	0,36	24,42	0,27	99,09	1,44	0,54	74,7
KI88-2-296.1-h1	core	90,3	X	37,42	0,01	0,06	0,02	36,73	0,21	0,28	24,69	0,30	99,72	1,45	0,55	74,7
KI88-2-296.1-h2	rim	90,3	X	37,23	0,01	0,04	0,01	36,59	0,23	0,28	24,94	0,23	99,62	1,45	0,55	74,4

Annexe 2 (suite)

Sample	Spot	Depth	Deformed	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	NiO	Total	MgO	FeO	Fo%
KI88-2-296.1-i1	core	90,3	X	38,89	0,01	0,08	0,00	43,17	0,25	0,22	16,64	0,32	99,64	1,64	0,36	83,7
KI88-2-296.1-i2	rim	90,3	X	37,54	0,02	0,07	0,00	36,98	0,20	0,38	23,71	0,26	99,16	1,46	0,53	75,6
KI88-2-296.1-c1	core	90,3		37,44	0,00	0,00	0,04	36,98	0,22	0,31	24,64	0,26	100,03	1,46	0,54	74,8
KI88-2-296.1-c2	rim	90,3		37,63	0,00	0,18	0,01	36,96	0,21	0,41	25,05	0,24	100,73	1,45	0,55	74,5
KI88-2-296.1-d1	core	90,3		37,78	0,01	0,00	0,01	37,39	0,21	0,32	24,48	0,26	100,47	1,46	0,54	75,2
KI88-2-296.1-d2	rim	90,3		37,48	0,01	0,69	0,05	36,68	0,21	0,30	24,53	0,33	100,30	1,44	0,54	74,8
KI88-2-296.1-e1	core	90,3		37,51	0,00	0,00	0,01	37,65	0,20	0,40	23,83	0,23	99,87	1,48	0,53	75,8
KI88-2-296.1-e2	rim	90,3		37,40	0,02	0,07	0,00	36,70	0,22	0,30	24,61	0,26	99,63	1,45	0,55	74,7
KI88-2-296.1-g1	core	90,3		37,72	0,01	0,06	0,03	37,79	0,21	0,24	23,17	0,25	99,49	1,48	0,51	76,4
KI88-2-296.1-g2	rim	90,3		37,48	0,01	0,00	0,00	36,96	0,22	0,26	24,68	0,19	99,89	1,46	0,55	74,8
KI88-2-296.1-j1	core	90,3		38,03	0,00	0,05	0,00	40,00	0,19	0,29	20,60	0,32	99,54	1,55	0,45	79,4
KI88-2-296.1-j2	rim	90,3		38,06	0,00	0,09	0,00	40,02	0,19	0,32	20,46	0,28	99,45	1,55	0,45	79,5

FeO*: total Fe; Fo% = $100 \times [\text{MgO} / (\text{MgO} + \text{FeO})]_{\text{mol}}$

ANNEXE 3

Compositions en éléments majeurs et mineurs de la matrice vitreuse de certains échantillons de Mauna Ulu et Kilauea Iki.

	(wt%)												
Sample	P ₂ O ₅	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	Na ₂ O	K ₂ O	F	Total
Mauna Ulu													
MU70-1 2v1	0,235	50,04	2,550	13,26	0,017	6,11	11,77	0,160	10,25	2,15	0,39	0,16	97,08
MU70-1 3v1	0,259	50,19	2,499	12,74	0,051	6,87	11,51	0,164	10,51	2,15	0,40	0,00	97,33
MU70-1 3v2	0,223	50,27	2,561	12,58	0,051	6,37	11,96	0,139	10,66	2,21	0,41	0,14	97,58
MU70-1 3v3	0,238	50,13	2,480	12,53	0,037	7,45	11,30	0,184	10,86	2,22	0,42	0,05	97,90
MU70-1 4v1	0,200	50,80	2,526	12,88	0,071	7,37	11,16	0,147	10,60	2,27	0,43	0,14	98,58
MU70-1 5v1	0,230	50,01	2,498	12,71	0,084	7,02	11,47	0,150	10,69	2,29	0,40	0,00	97,56
MU70-1 6v1	0,190	50,13	2,514	12,37	0,000	6,80	11,51	0,149	10,64	2,25	0,43	0,05	97,04
MU70-1 7v1	0,241	50,64	2,414	12,39	0,054	7,93	11,60	0,189	10,38	1,70	0,40	0,06	98,00
MU70-1 8v1	0,230	49,29	2,712	13,21	0,074	5,58	11,74	0,198	10,62	1,93	0,39	0,14	96,11
MU70-1 12v1	0,220	50,35	2,478	12,58	0,050	7,20	11,40	0,154	10,97	2,13	0,42	0,01	97,96
MU70-1 13v1	0,208	50,33	2,501	12,37	0,054	7,28	11,40	0,146	11,05	2,23	0,42	0,00	97,99
MU70-1 14v1	0,246	49,97	2,520	12,86	0,104	7,11	11,53	0,135	10,81	2,30	0,41	0,14	98,12
MU70-1 15v1	0,216	50,17	2,510	12,56	0,047	7,34	11,40	0,127	10,96	2,26	0,42	0,06	98,08
MU70-1 15v2	0,216	50,33	2,424	12,47	0,074	7,15	11,45	0,153	10,90	2,24	0,43	0,04	97,87
MU70-1 17v1	0,224	49,98	2,611	12,58	0,044	7,03	11,63	0,212	10,61	2,33	0,41	0,01	97,68
MU70-1 18v1	0,276	50,41	2,479	12,49	0,007	7,20	11,40	0,199	10,50	2,17	0,42	0,23	97,79
MU70-1 19v1	0,205	50,33	2,589	12,81	0,061	6,74	11,61	0,188	10,70	2,02	0,39	0,00	97,64
MU72-2 1v1	0,191	50,19	2,464	12,94	0,024	7,39	11,31	0,179	10,25	2,30	0,42	0,04	97,68
MU72-2 2v1	0,219	50,18	2,456	12,38	0,013	7,00	11,33	0,179	10,54	2,27	0,43	0,00	96,99
MU72-2 3v1	0,240	50,22	2,464	12,21	0,030	7,04	11,51	0,173	10,38	2,26	0,42	0,02	96,95
MU72-2 4v1	0,204	50,52	2,542	12,65	0,000	7,22	11,32	0,144	10,42	2,24	0,42	0,00	97,67
MU72-2 5v1	0,207	50,46	2,502	12,52	0,054	7,22	11,35	0,168	10,61	2,36	0,42	0,16	98,04
MU72-2 6v1	0,256	50,27	2,460	11,61	0,037	7,08	11,62	0,183	10,66	2,19	0,42	0,00	96,77
MU72-2 6v2	0,226	50,01	2,571	12,27	0,013	7,19	11,39	0,189	10,19	2,22	0,42	0,00	96,69
MU72-2 7v1	0,197	50,57	2,395	12,63	0,024	7,16	11,45	0,131	10,71	2,27	0,45	0,16	98,15
MU72-2 8v1	0,246	49,95	2,558	12,53	0,071	7,17	11,45	0,176	10,20	2,33	0,45	0,15	97,28
MU72-2 9v1	0,194	50,13	2,573	12,48	0,020	6,95	11,44	0,187	10,41	2,30	0,45	0,04	97,16
MU72-2 10v1	0,222	49,94	2,552	12,27	0,118	7,04	11,56	0,149	10,58	2,25	0,43	0,11	97,22
MU72-2 11v1	0,231	50,25	2,474	11,88	0,007	7,28	11,44	0,150	10,36	2,24	0,43	0,21	96,93
MU72-2 12v1	0,197	50,05	2,515	12,52	0,037	7,03	11,44	0,164	10,57	2,23	0,42	0,01	97,17
MU72-2 13v1	0,241	49,40	2,452	13,10	0,064	7,19	11,22	0,158	10,41	2,32	0,43	0,13	97,09
MU72-2 14v1	0,226	50,21	2,512	12,37	0,081	7,19	11,35	0,143	10,37	2,23	0,43	0,05	97,16
MU72-2 15v1	0,204	50,30	2,527	12,70	0,054	6,96	11,34	0,185	10,39	2,21	0,42	0,00	97,29
MU72-2 16v1	0,160	50,20	2,511	12,68	0,020	7,16	11,33	0,161	10,49	2,21	0,40	0,13	97,44
MU72-2 17v1	0,221	50,35	2,448	12,44	0,017	7,14	11,33	0,191	10,68	2,20	0,43	0,09	97,52
MU72-2 17v2	0,216	50,35	2,492	12,42	0,081	7,26	11,49	0,159	10,53	2,28	0,44	0,02	97,72
MU72-2 17v3	0,229	50,37	2,531	12,20	0,013	7,20	11,43	0,174	10,50	2,23	0,43	0,15	97,46
MU73-1a 1v1	0,232	50,48	2,546	12,57	0,020	6,78	11,48	0,132	10,43	2,29	0,45	0,07	97,48
MU73-1a 2v1	0,223	50,15	2,381	12,88	0,040	6,53	11,76	0,156	10,51	2,22	0,44	0,00	97,29
MU73-1a 3v1	0,210	47,49	2,689	12,58	0,054	6,82	11,52	0,162	10,55	2,17	0,52	0,05	94,81
MU73-1a 4v1	0,213	49,87	2,562	12,92	0,040	6,68	11,51	0,162	10,49	2,26	0,45	0,10	97,26
MU73-1a 5v1	0,189	50,50	2,667	12,91	0,017	6,73	11,44	0,165	10,58	2,18	0,44	0,00	97,82
MU73-1a 6v1	0,220	50,87	2,647	12,73	0,030	6,52	11,32	0,154	11,31	2,27	0,46	0,00	98,52

Annexe 3 (suite)

Sample	P ₂ O ₅	SiO ₂	TiO ₂	Al ₂ O ₃	Cr ₂ O ₃	MgO	CaO	MnO	FeO*	Na ₂ O	K ₂ O	F	Total
MU73-1a 7v1	0,213	50,97	2,590	12,61	0,094	6,58	11,99	0,175	10,70	2,24	0,39	0,03	98,59
MU73-1a 8v1	0,204	50,33	2,565	12,94	0,000	6,73	11,43	0,200	10,56	2,25	0,47	0,08	97,75
MU73-1a 9v1	0,183	51,09	2,693	12,36	0,000	6,79	11,72	0,181	11,04	2,34	0,41	0,02	98,83
MU73-1a 10v1	0,216	50,51	2,595	12,63	0,061	6,05	12,06	0,177	10,76	2,21	0,45	0,07	97,78
MU73-1a 11v1	0,234	50,22	2,567	12,51	0,000	6,75	11,55	0,146	10,84	2,29	0,45	0,00	97,55
MU73-1a 12v1	0,196	50,53	2,599	12,73	0,067	6,65	11,53	0,142	10,46	2,27	0,45	0,10	97,72
MU73-1a 13v1	0,191	50,46	2,569	12,19	0,061	6,24	11,87	0,166	10,74	2,23	0,41	0,02	97,14
MU73-1a 14v1	0,232	50,23	2,683	12,81	0,030	6,51	11,61	0,168	10,41	2,26	0,46	0,00	97,39
MU73-1a 15v1	0,198	49,88	2,630	12,81	0,088	6,52	11,61	0,170	10,55	2,25	0,44	0,14	97,28
MU73-1a 16v1	0,207	50,46	2,586	13,01	0,000	6,88	11,53	0,155	10,65	2,33	0,47	0,11	98,39
MU73-1a 17v1	0,215	50,31	2,624	12,67	0,010	6,70	11,46	0,203	10,41	2,29	0,47	0,00	97,37
MU74-2a n verre	n.a.	50,47	n.a.	13,24	0,072	6,91	11,61	0,153	10,19	n.a.	n.a.	n.a.	92,63
Kilauea Iki													
KI75-1-145-a-v1	0,04	52,29	4,66	12,94	0,03	4,91	0,45	0,15	8,84	2,65	0,02	0,21	87,20
KI75-1-145-c-v2	0,01	52,24	4,69	13,06	0,00	5,04	0,44	0,18	8,77	2,68	0,05	0,03	87,19
KI75-1-145-g-v3	0,03	52,20	4,76	12,92	0,00	4,99	0,43	0,16	8,81	2,58	0,04	0,03	86,96
KI75-1-145-j-v4	0,03	52,22	4,67	12,98	0,00	4,98	0,47	0,16	8,67	2,56	0,03	0,11	86,88
KI75-1-145-m-v5	0,06	51,90	4,67	12,95	0,00	4,90	0,47	0,15	9,14	2,61	0,04	0,17	87,06
KI75-3-141-e-v1	0,52	56,53	3,43	13,53	0,00	3,52	6,54	0,13	10,45	2,68	1,62	0,05	99,02
KI75-3-141-h-v2	0,51	55,98	3,43	13,42	0,00	3,57	6,56	0,16	11,23	2,58	1,63	0,17	99,24
KI75-3-141-j-v3	0,54	56,08	3,59	13,25	0,00	3,54	6,50	0,16	10,91	2,66	1,65	0,22	99,10
KI75-3-141-l-v4	0,50	56,29	3,45	13,49	0,00	3,45	6,47	0,14	10,77	2,64	1,70	0,15	99,04
KI81-1-200.2-a-glass	0,45	50,47	4,13	13,38	0,00	5,49	9,15	0,15	10,71	2,56	0,86	0,16	97,49
KI81-1-200.2-d-glass	0,48	50,28	4,19	12,95	0,01	5,52	9,13	0,16	10,74	2,65	0,87	0,00	96,97
KI81-1-200.2-g-glass	0,48	50,47	4,16	13,40	0,00	5,48	9,23	0,13	10,46	2,60	0,89	0,17	97,49
KI81-1-200.2-l-glass	0,42	50,10	4,23	13,21	0,00	5,46	9,16	0,15	10,73	2,66	0,87	0,24	97,23
KI81-1-249.7-i-glass	0,36	49,63	3,93	12,00	0,00	5,85	9,51	0,16	9,65	2,69	0,84	0,13	94,75
KI81-1-249.7-f-glass	0,36	50,56	3,95	12,82	0,07	5,89	9,73	0,17	9,87	2,53	0,83	0,13	96,90
KI81-1-249.7-c-glass	0,40	50,29	3,89	11,05	0,01	6,03	9,75	0,14	10,49	2,80	0,84	0,19	95,87
KI81-1-299.9-e-glass	0,71	53,45	4,01	13,09	0,00	4,37	7,47	0,19	10,35	2,53	1,44	0,16	97,78
KI81-1-299.9-h-glass	0,74	53,27	3,86	13,33	0,00	4,26	7,43	0,14	10,40	2,55	1,45	0,22	97,65
KI81-1-299.9-i-glass	0,73	53,10	4,10	13,11	0,00	4,24	7,56	0,15	10,20	2,60	1,45	0,11	97,34
KI81-2-185.7-b-glass	1,23	59,49	2,25	13,61	0,03	2,35	4,88	0,12	7,98	2,09	2,37	0,00	96,39
KI81-2-185.7-d-glass	1,26	59,41	2,35	14,77	0,01	2,35	4,94	0,11	7,97	1,76	2,30	0,30	97,53
KI81-2-185.7-i-glass	1,26	59,02	2,39	14,25	0,00	2,46	4,98	0,12	8,03	2,27	2,24	0,30	97,30
KI81-2-185.7-k-glass	1,27	58,77	2,46	14,04	0,04	2,49	4,99	0,12	8,31	2,28	2,23	0,00	96,98

FeO*: total Fe; n.a. means 'Not Analysed'

ANNEXE 4

Compositions en éléments majeurs et mineurs de chromites de quelques échantillons de Mauna Ulu et Kilauea Iki.

	(wt%)														
Sample	SiO ₂	TiO ₂	Al ₂ O ₃	V ₂ O ₃	Cr ₂ O ₃	Fe ₂ O ₃	MgO	CaO	MnO	FeO	CoO	NiO	Na ₂ O	Total	
Kilauea Iki															
KI59-1 b c1	0,10	2,37	14,40	0,23	45,34	6,94	12,28	0,01	0,00	17,07	0,00	0,27	0,01	99,03	
KI59-1 b c2	0,10	2,36	16,90	0,26	41,87	7,84	12,58	0,01	0,00	17,06	0,07	0,19	0,01	99,24	
KI59-1 b c3	0,09	2,55	14,56	0,33	43,82	8,05	11,37	0,02	0,00	18,91	0,31	0,19	0,00	100,19	
KI59-1 b c4	0,08	1,95	13,78	0,18	46,93	7,21	11,77	0,02	0,00	17,66	0,20	0,12	0,03	99,93	
KI59-1 c c1	0,08	1,89	13,28	0,26	46,68	7,71	10,93	0,00	0,00	18,80	0,00	0,10	0,00	99,72	
KI59-1 j c1	0,10	1,76	14,47	0,19	47,19	6,72	10,73	0,00	0,00	19,56	0,00	0,26	0,01	100,99	
KI59-1 j c2	0,09	1,86	14,12	0,20	47,78	6,44	12,39	0,00	0,00	16,68	0,05	0,21	0,00	99,82	
KI59-1 n c1	0,07	2,04	13,33	0,28	45,45	8,17	9,65	0,06	0,00	21,02	0,29	0,27	0,02	100,64	
KI59-1 o c1	0,09	1,69	13,96	0,20	48,19	6,82	11,79	0,02	0,00	17,72	0,05	0,19	0,00	100,74	
KI59-1 o c2	0,07	1,69	13,89	0,20	48,82	6,49	12,77	0,01	0,00	16,05	0,00	0,37	0,00	100,36	
KI59-2a e c1	0,09	1,59	13,98	0,20	48,75	5,79	11,77	0,00	0,00	17,37	0,49	0,29	0,01	100,31	
KI59-2a e c2	0,10	1,63	14,03	0,20	47,72	5,52	11,60	0,05	0,00	17,17	0,00	0,23	0,01	98,25	
KI59-2a h c1	0,09	2,34	13,44	0,28	45,10	7,98	11,52	0,01	0,00	18,06	0,68	0,09	0,02	99,61	
KI59-2a h c2	0,10	2,29	13,84	0,28	43,10	7,62	8,85	0,08	0,00	21,90	0,00	0,17	0,01	98,25	
KI59-2a l c1	0,07	1,72	13,96	0,20	48,16	6,16	12,39	0,01	0,00	16,37	0,14	0,26	0,02	99,45	
KI59-2a l c2	0,05	1,74	13,95	0,15	48,68	5,94	12,46	0,00	0,00	16,43	0,09	0,20	0,01	99,67	
KI59-2a m c1	0,08	1,88	13,28	0,22	45,76	6,79	7,85	0,05	0,00	23,42	0,00	0,14	0,01	99,47	
KI59-2a m c2	0,12	2,08	14,44	0,19	45,25	7,39	12,13	0,02	0,00	17,01	0,00	0,25	0,01	98,90	
KI59-3 b c1	0,08	2,59	13,65	0,26	44,30	9,33	11,78	0,00	0,00	18,42	0,00	0,38	0,02	100,79	
KI59-3 b c2	0,04	2,20	13,65	0,28	44,87	8,68	11,30	0,02	0,00	18,55	0,31	0,14	0,01	100,06	
KI59-3 c c1	0,06	2,22	13,55	0,27	46,10	7,90	11,21	0,00	0,00	18,91	0,22	0,20	0,04	100,67	
KI59-3 c c2	0,05	2,15	13,22	0,29	46,40	8,06	11,36	0,00	0,00	18,49	0,00	0,10	0,00	100,11	
KI59-3 e c1	0,08	1,62	13,98	0,16	49,42	5,74	12,35	0,00	0,00	16,68	0,11	0,29	0,02	100,46	
KI59-3 e c2	0,10	1,56	13,98	0,19	49,80	6,11	13,22	0,01	0,00	15,37	0,00	0,11	0,00	100,44	
KI59-3 h c1	0,10	1,79	14,09	0,19	47,71	7,42	12,67	0,00	0,00	16,45	0,00	0,21	0,01	100,63	
KI59-3 j c1	0,09	1,80	13,62	0,27	46,71	8,26	12,35	0,00	0,00	16,62	0,00	0,20	0,04	99,95	
KI59-3 j c2	0,10	2,16	13,72	0,22	45,42	9,05	12,45	0,01	0,00	16,89	0,00	0,22	0,01	100,24	
KI59-3 k c1	0,10	2,47	14,34	0,28	42,84	9,41	11,53	0,10	0,00	18,48	0,70	0,14	0,02	100,41	
KI59-3 k c2	0,07	1,86	14,01	0,20	46,19	8,33	11,85	0,02	0,00	17,69	0,00	0,13	0,02	100,37	
Mauna Ulu															
MU74- 2a a c1	0,10	2,48	14,72	0,35	41,66	8,66	10,16	0,18	0,00	20,30	0,34	0,21	0,01	99,15	
MU74- 2a a c2	0,12	2,83	15,14	0,29	40,52	9,39	11,29	0,01	0,00	19,01	0,00	0,20	0,01	98,80	
MU74- 2a a c3	0,10	2,35	15,47	0,29	41,63	9,02	11,34	0,01	0,00	18,66	0,72	0,22	0,02	99,82	
MU74- 2a e c1	0,06	2,47	14,85	0,35	42,17	9,16	10,61	0,03	0,00	20,05	0,00	0,20	0,01	99,96	
MU74- 2a f c1	0,04	2,49	11,97	0,29	46,41	8,18	10,57	0,01	0,00	19,69	0,23	0,15	0,02	100,05	
MU74- 2a o c1	0,06	1,90	12,71	0,32	42,55	11,74	8,49	0,01	0,00	22,62	0,36	0,03	0,00	100,78	
MU74- 2a v c1	0,10	3,24	16,49	0,60	36,81	10,18	9,83	0,01	0,00	21,90	0,00	0,11	0,01	99,27	

ANNEXE 5

Compositions roche totale en éléments majeurs, traces et Terres Rares des échantillons de surface du Kilauea Iki.

Sample	KI-59-1	KI-59-2a	KI-59-3
<i>(XRF)</i>			
SiO ₂	46,92	46,36	46,35
TiO ₂	2,176	2,043	2,049
Al ₂ O ₃	10,58	9,82	9,89
Fe ₂ O ₃ *	13,30	13,48	13,49
MnO	0,179	0,175	0,175
MgO	15,81	17,90	17,83
CaO	9,39	8,68	8,75
Na ₂ O	1,81	1,68	1,69
K ₂ O	0,45	0,43	0,43
P ₂ O ₅	0,221	0,207	0,208
Total	100,84	100,78	100,86
Mg#	72,3	74,5	74,4
BaO	77	69	80
Co	112	122	118
Cr	1080	1193	1202
Cu	147	147	153
Ni	698	848	837
Sc	20	12	30
V	247	230	229
Zn	67	63	63
<i>(ICP-MS)</i>			
Ga	15,2	13,6	13,9
Nb	14,9	13,8	14,0
Pb	nd.	nd.	nd.
Rb	8,9	8,4	8,3
Sr	296,8	280,4	278,1
Th	1,6	1,2	nd.
U	1,0	nd.	nd.
Y	19,9	19,4	18,4
Zr	137,4	128,5	128,6
Y	17,7	16,6	16,6
La	11,7	10,9	10,9
Ce	27,9	25,9	25,9
Pr	3,99	3,70	3,69
Nd	18,3	17,0	17,0
Sm	4,42	4,19	4,30
Eu	1,54	1,47	1,41
Gd	4,49	4,30	4,29
Tb	0,685	0,640	0,646
Dy	3,98	3,66	3,68
Ho	0,721	0,676	0,677
Er	1,87	1,73	1,79
Tm	0,242	0,224	0,229
Yb	1,48	1,40	1,37
Lu	0,195	0,189	0,198
Olivine abundance	12,7	16,6	14,6

* Fe₂O₃ is total iron. n.d. means 'Not Detected'. Oxides are in wt %; trace elements are in µg/g.; Mg# is Mg/(Mg+Fe²⁺); Olivine abundance is in vol. %.

