Évaluation de quelques paramètres de la qualité du bois affecté par l’épidémie de la tordeuse des bourgeons de l’épinette

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<td>Tordeuse des bourgeons de l’épinette</td>
</tr>
<tr>
<td>SB</td>
<td>Spruce budworm</td>
</tr>
<tr>
<td>DBH</td>
<td>Diameter at breast height</td>
</tr>
<tr>
<td>EWD</td>
<td>Earlywood density</td>
</tr>
<tr>
<td>LWD</td>
<td>Latewood density</td>
</tr>
<tr>
<td>MC</td>
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<td>RWC</td>
<td>Relative water content</td>
</tr>
<tr>
<td>Ψwp</td>
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</tr>
<tr>
<td>VPD</td>
<td>Vapour pressure deficit</td>
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</table>
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INTRODUCTION


Les peuplements de sapins et d’épinettes deviennent progressivement défoliés au cours des années en période épidémique de la TBE (Blais 1958; Dorion 2016). Les
dommages apparaissent d'abord sur la cime supérieure de l'arbre, limités à une perte partielle de nouveau feuillage, puis se propagent vers les branches inférieures (Carisey et Bauce 1997; Garcia 2015). Ces dommages sont intrinsèques au cycle de vie de l'insecte qui se déroule sur un an. Il compte un stade œuf, six stades larvaires, un stade chrysalide et un stade adulte (Neau 2014). La défoliation se produit durant les stades larvaires deux à six (Simard 2011). Carisey et Bauce (1997) ont démontré que lorsque l'épidémie devient sévère, les larves se nourrissent des nouvelles aiguilles et même de celles âgées d'un an. À partir de la quatrième à la cinquième année de perturbation consécutive et sévère, le développement radial de certains arbres peut être totalement interrompu, et les arbres les plus faibles peuvent mourir (Archambault et al. 1989; Garcia 2015).


Le temps nécessaire pour que la défoliation cause la mort des arbres diffère entre le sapin baumier et l’épinette noire. Normalement, la mortalité chez le sapin baumier est attendue après 4-5 ans de défoliation sévère, alors que c’est plutôt après 6-7 ans chez l’épinette noire (Blais 1965; MacLean 1980; Pothier 2012). La mortalité des peuplements

important pour les propriétés du bois de structure (Bowyer et al. 2003; Forest Products Laboratory 2010; Passarini 2011).


Holusa (2006) ont observé que la défoliation consécutive des mélèzes par *Cephalcia lariciphila* avait causé une réduction de la croissance et également une réduction de la densité du bois final à cause des changements anatomiques du bois. Axelson et al. (2014) ont détecté le même phénomène chez les Douglas défoliés par la *Choristoneura occidentalis*.

Dans ce contexte, il devient essentiel d’évaluer les changements physiologiques de l’arbre selon un gradient d’intensité et de durée de défoliation. Établir une relation entre ces modifications et la qualité du bois est de première importance pour l’industrie. Pour ce faire, deux facteurs relatifs à la physiologie des arbres défoliés (potentiel hydrique et contenu relatif en eau) et cinq paramètres de qualité du bois (taux d’humidité, proportion d’aubier, volume annuel, densité et anatomie du bois) ont fait l’objet de cette étude. Ceci permettra d’établir le lien entre la défoliation par la TBE, les impacts physiologiques et les paramètres de qualité du bois.

Plusieurs hypothèses ont été émises : suite à la diminution de la biomasse des aiguilles par la défoliation, il y aurait une réduction dans la translocation de l’eau et des nutriments des branches et des aiguilles restantes (1). La réduction du besoin en eau diminuerait alors la translocation en eau dans la tige, se traduisant ainsi par des réductions au niveau de l’aubier le long de la tige et du taux d’humidité du bois. La baisse de la quantité d’eau sera plus importante avec une augmentation de la durée de la défoliation (2). De plus, la diminution de la biomasse foliaire mènera à une baisse de la croissance radiale, particulièrement dans le bois initial, lieu de transport de l’eau (3). Les modifications de la croissance radiale changeront la masse volumique moyenne du bois (4). Le manque de sucres changera aussi l’épaisseur des parois cellulaires qui influence la masse volumique du bois (5).
Ce mémoire est divisé en deux chapitres qui exposent les résultats obtenus sous forme d’articles scientifiques.

Chapitre I: Potentiel et état hydrique des arbres défoliés par la tordeuse des bourgeons de l’épinette et leur influence sur le taux d’humidité et la proportion d’aubier.

Ce chapitre s’intéresse à l’effet que peut avoir la défoliation sur des paramètres physiologiques des arbres défoliés (potentiel et état hydrique) et de la qualité du bois de la tige (taux d’humidité et proportion d’aubier) d’épinette noire et de sapin baumier. Quarante sites ont été échantillonnés dans la région du Saguenay Lac-Saint-Jean, avec des arbres défoliés d’une à quatre années consécutives et d’intensité de défoliation faible à élevée.

Chapitre II: Évaluation de quelques paramètres de la qualité du bois affecté par l’épidémie de la tordeuse des bourgeons de l’épinette.

Ce second chapitre porte sur les effets de la défoliation sur la croissance et les propriétés du bois des tiges d’épinette noire et du sapin baumier après la défoliation consécutive. L’accroissement annuel en volume a été calculé pour les mêmes arbres que chapitre I. La densité du bois et l’anatomie du bois ont été mesurées à partir d’une rondelle prise à 1.3 m.
Spruce budworm (Choristoneura fumiferana Clem.) is one of the most destructive defoliator agent of coniferous forests in eastern North America. In Canada, balsam fir (Abies balsamea L. (Mill)) and black spruce (Picea mariana B.S.P. (Mill)) are the most important and widely distributed hosts. Little is known of the impact of defoliation on tree physiology and moisture content for the two species. Thirty-six infested stands, varying from one to four years of defoliation, were sampled in the Quebec boreal forestry in 2016 and 2017 to determine whether modifications had occurred on physiological parameters of defoliated trees (potential and relative water status) and on the quality of wood (moisture content and sapwood proportion) of spruce black and balsam fir. The results show that defoliation has no effect on water status and water potential for both species. No changes were observed between defoliated and non-defoliated trees in terms of moisture content and sapwood proportion neither. Wood moisture content has been pointed out as an important parameter during the wood transformation and based on these results no changes are expected for defoliated balsam fir and black spruce.

Keywords: Spruce budworm outbreak, wood quality, Picea mariana, Abies balsamea, moisture content, sap- and heartwood proportion, relative water content.
1.1 INTRODUCTION

Boreal forests are widely distributed across the northern hemisphere (Bergeron et Fenton 2012; Strid et al. 2014). In eastern North America, two major types of natural disturbances periodically influence the natural boreal forest: insect outbreaks and fires (Cogbill 1985; MacLean 1985; Stocks 1987; MacLean 2016). The spruce budworm (Choristoneura fumiferana Clem.) is one of the most widely distributed defoliator agents in these ecosystems (Morris 1963; Blais 1983; Lapointe 2013; MacLean 2016). In the last century, there have been three major budworm outbreaks, causing dramatic growth reductions and stand mortality (Morin et al. 2007). Recurrent outbreaks mostly affect balsam fir (Abies balsamea (L.) Mill)) the preferred host, and with the population explosion, other coniferous trees such as black spruce (Picea mariana B.S.P. (Mill)) are damaged (MacLean 1984; Hennigar et al. 2008). However, it has been suggested that insect populations have moved further north (Morin et al. 2007; MacLean 2016), where black spruce is dominating. Thus, black spruce (Picea mariana (Mill.) BSP), which has a crucial role in the forest industry in Eastern Canada (MFFP 2015), is more and more attacked. During a typical outbreak, the insect larvae consume the new foliage of the trees, starting on the upper crown and then spreading towards the lower branches (Carisey et Bauce 1997; Garcia 2015). The outbreaks tend to occur every 30-40 years (Morin et al. 2007; Simard 2011; Boulanger et al. 2012) and last for 10-15 years (Barrette et al. 2015). An outbreak is currently affecting the Eastern boreal forest of Canada.

Several studies have shown a decrease in radial growth after defoliation periods, as a result of the repeated loss of new foliage over successive years of infestation (Blais 1962; MacLean 1984; Krause et al. 2003; Krause et al. 2012). The complete tree mortality can be observed following 4-5 years of severe infestation in balsam fir and 6-7 years in black spruce.
(Blais 1965; MacLean 1980; Pothier 2012). However, after tree death, besides the direct loss in volume, the quality of wood harvested and of the end-use products can be altered by the infestation (Binotto et Locke 1981; Basham 1984; Koran et Nlombi 1994).

Forest product properties depend strongly on different wood characteristics that are affected by genetics, environmental and silvicultural aspects of tree growth (Zhang et al. 1996; Barnett et Jeronimidis 2003; Downes et Drew 2008). Moisture content is one of the most important properties to monitor for wood quality. It directly affects harvesting and logging operations in the field, the efficiency of numerous industrial processes, as well as the characteristics of the end-products (Ip et al. 1996; Zhang 2004; Barrette et al. 2015).

As sapwood transports water and nutrients, the moisture content of this part is higher than that of heartwood (Woo et al. 2005; Lowell et al. 2010). Thus the proportion of sapwood and heartwood has a significant impact on the moisture content of the stem (Barnett et Jeronimidis 2003). Additionally, moisture content increases with height because of the decreasing proportion of heartwood to sapwood throughout the stem (Barnes et Sinclair 1983; Ip et al. 1996).

Overall, hydraulic efficiency in a tree can be related to two factors: the sapwood efficacy and the water acquisition by the roots (Smith et Hinckley 1995). The first is directly influenced by the largest cell diameters produced in the earlywood (Deslauriers 1999; Domec et Gartner 2002; Lupi et al. 2010); and the second by the water potential gradient between the soil and the atmosphere (Tyree et Zimmermann 2002). Foliage removal disturbs various physiological processes in plants (Reich et al. 1993; Clancy et al. 1995). Studies on several tree species have shown that defoliation or browsing can have contradictory effects on water availability in plants (Kolb et al. 1999; Hart et al. 2000; Lavigne et al. 2001; Quentin et al. 2012; Jacquet et al. 2014). Water potential and relative water
content have been used in these studies to express water relations because it reflects changes in the balance between transpiration and water uptake (Stephens et al. 1972; Pallardy et al. 1995; Gieger et Thomas 2002).

For wood moisture content, in the last three decades, numerous studies have examined the changes arising from defoliation only after the tree death (Barnes et Sinclair 1983; Van Raalte 1983; Ip et al. 1996; Barrette et al. 2015). The relationship between duration of defoliation and wood moisture content is still poorly understood. Thus, the main objective of this study was to evaluate the effects of one-to-four years of defoliation on water relations and moisture content for balsam fir and black spruce trees during a spruce budworm outbreak. Water relations will be expressed by examining the water potential and the relative water content in the branches of defoliated trees, and wood moisture content along the stem in sapwood and heartwood. The hypotheses were that: a significant reduction in the water potential and relative water content would be observed with the increase duration of the defoliation event (1); the plant with lower water demand would present less sapwood, first in the upper crown and then in the other parts of the stem, and this would be paralleled by a decreased moisture content in the sapwood (2). The findings of this study will contribute to our understanding of the effects of spruce budworm on wood quality properties and the resilience and plasticity of the two main harvested species in Eastern North America.

1.2 METHODOLOGICAL APPROACH

1.2.1 STUDY AREA

The study area covers approximately 10 000 km² at the transition between northern temperate and boreal forest in Quebec, Canada (48°25’N 71°04’W). The region is
characterized by hills that rarely exceed 500 m, on thick and undifferentiated glacial till deposits. This area is part of balsam fir-white birch, balsam fir-yellow birch, and spruce-moss bioclimatic domain (Figure 1). The climate is continental and has a short warm summer, without a dry season. The 30-year mean annual temperature is 2.8° C and total precipitation of 930 mm, with 5-month covert by snow (Government of Canada 2017).

![Figure 1 - Study site location in Saguenay, Lake Saint-Jean area, Quebec, Canada](image)

1.2.2 TREE SELECTION

The infested areas was assessed using the spruce budworm annual aerial detection survey data from the government of Quebec (MFFP 2016; 2017). In order to delimit the sampling area, some factors were considered, such as: public lands that were not harvested in the last years, stands defoliated between one to four years of defoliation, stands within
150 kilometers from the Lake Saint-Jean. Finally, twenty natural stands dominated by balsam fir (Abies balsamea (L.) Mill.) and twenty-one dominated by black spruce (Picea mariana (Mill.)) were selected. At each site, representative defoliated individuals were chosen randomly among healthy trees. A visually assessment of the defoliation was made with binoculars before harvesting. The trees were sampled in the summers of 2016 and 2017 (June and early July).

Stands were divided according to the duration of defoliation in five classes: from no defoliation to up to four continuous years of defoliation. D0 represents control trees without defoliation, D4 four years of consecutive defoliation. D1, D2, and D3 represent one to three years of continuous defoliation. Additionally, a visual assessment of defoliation intensity was carried out for each sampled tree in the field. Three intensities of defoliation were used: light (25% needle loss), moderate (from 26% to 70% needle loss), and severe (more than 71% needle loss). Thus, the goal of this study is to understand the combined effects of timing and intensity of defoliation on each parameter, where each defoliation class represents the duration of consecutive defoliation, and intensity of defoliation, the degree of needle loss. Preliminary analysis showed more variation in the data between the trees with a longer defoliation lasting. For this reason, the number of harvested trees varied between the defoliation classes, with more trees to a longer defoliation period (Table 1).

Stand characterization was done including information and measurements of stand composition, slope, drainage, and soil type. Height, diameter at breast height (DBH) and crown length were also measured on 15 live trees (si) (Table 1). Soil profiles were classified into three broad categories: well drained (64 % of the study sites), moderately (20 %) and poorly drained (16 %). Two types of soils were found: podzolic (73 %) and organic soils (23
Tree age ranged from 24- to 132-year-old for the balsam fir and 29- to 172-year-old for black spruce stands (Table 1).

Table 1 - Structure and characteristics of the five classes of defoliation by species (D0 to D4). The number of harvested trees within classes is reported in parentheses. The data from the stand inventory (si) including 15 trees per stand is also shown.

<table>
<thead>
<tr>
<th>Specie</th>
<th>Defoliation class (Number of Trees)</th>
<th>Number of Stands per defoliation class</th>
<th>Mean Age (years)*</th>
<th>Mean DBH (cm)*</th>
<th>Mean Height (m)*</th>
<th>Mean DBHfi (cm)</th>
<th>Mean Hfi (m)</th>
</tr>
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<tbody>
<tr>
<td>Spruce</td>
<td>D0 (14)</td>
<td>3</td>
<td>74 ± 31</td>
<td>15.9 ± 4.2</td>
<td>12.6 ± 2.8</td>
<td>13.7 ± 3.4</td>
<td>10.9 ± 2.6</td>
</tr>
<tr>
<td></td>
<td>D1 (16)</td>
<td>3</td>
<td>108 ± 43</td>
<td>13.3 ± 2.2</td>
<td>13.0 ± 1.6</td>
<td>13.4 ± 3.1</td>
<td>12.8 ± 2.5</td>
</tr>
<tr>
<td></td>
<td>D2 (21)</td>
<td>5</td>
<td>74 ± 24</td>
<td>15.7 ± 4.1</td>
<td>14.4 ± 3.1</td>
<td>15.6 ± 6.6</td>
<td>13.8 ± 4.8</td>
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<td>D3 (33)</td>
<td>5</td>
<td>71 ± 26</td>
<td>13.1 ± 2.4</td>
<td>11.5 ± 1.4</td>
<td>12.9 ± 3.0</td>
<td>11.8 ± 1.8</td>
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<tr>
<td></td>
<td>D4 (35)</td>
<td>5</td>
<td>78 ± 23</td>
<td>16.3 ± 5.6</td>
<td>13.5 ± 3.2</td>
<td>14.5 ± 5.5</td>
<td>12.8 ± 3.6</td>
</tr>
<tr>
<td>Fir</td>
<td>D0 (10)</td>
<td>2</td>
<td>85 ± 21</td>
<td>16.5 ± 2.1</td>
<td>16.0 ± 2.6</td>
<td>18.9 ± 8.6</td>
<td>14.2 ± 5.6</td>
</tr>
<tr>
<td></td>
<td>D1 (20)</td>
<td>4</td>
<td>78 ± 29</td>
<td>15.9 ± 5.2</td>
<td>14.1 ± 3.1</td>
<td>15.7 ± 5.3</td>
<td>12.8 ± 2.6</td>
</tr>
<tr>
<td></td>
<td>D2 (20)</td>
<td>4</td>
<td>58 ± 7</td>
<td>14.6 ± 2.3</td>
<td>12.3 ± 1.3</td>
<td>13.3 ± 3.1</td>
<td>11.0 ± 2.2</td>
</tr>
<tr>
<td></td>
<td>D3 (34)</td>
<td>5</td>
<td>55 ± 14</td>
<td>13.8 ± 2.4</td>
<td>12.5 ± 2.1</td>
<td>11.9 ± 4.5</td>
<td>10.9 ± 2.8</td>
</tr>
<tr>
<td></td>
<td>D4 (35)</td>
<td>5</td>
<td>54 ± 14</td>
<td>16.2 ± 3.6</td>
<td>14.3 ± 2.0</td>
<td>15.0 ± 5.6</td>
<td>14.1 ± 3.1</td>
</tr>
</tbody>
</table>

* Means and standard deviations per defoliation class

A total of 119 black spruce and 119 balsam fir trees were sampled. Five stem discs were collected from each tree, located at 0%, 25%, 50%, 75%, and 100% tree height (15-30 cm from terminal branch end). Two branches were also collected from each tree (Figure 2). Both discs and branches were stored in polyethylene bags and kept refrigerated until moisture content, relative water content, and water potential measurements were made.

1.2.3 MEASUREMENTS

Water potential and Relative water content

The leaf water potential (Ψwp) was obtained by a Scholander pressure chamber (Scholander et al. 1965). To determine the relative water content (RWC), measurements of fresh (immediately weighed after collecting), saturated (24 hours immersed in water) and
dry mass (48 hours in an oven at 80°C) were taken. The RWC could be expressed as a percentage of the dry mass (Morabito et al. 2006):

\[
RWC = \frac{(\text{fresh mass} - \text{dry mass})}{(\text{saturated mass} - \text{dry mass})} \times 100
\]

**Moisture Content (MC)**

Moisture content was measured by separating sap- and heartwood from ¼ sections (Figure 2). The wet and dry weight of each sample was taken with a precision of ±0.1 g. Then MC was calculated as follows (Parham 1983):

\[
MC = \frac{(\text{wet weight} - \text{dry weight})}{\text{dry weight}} \times 100
\]

For balsam fir, heartwood was divided into wet heartwood (wood > 100 % of MC) and dry heartwood (wood < 100 % of MC).

**Sapwood and heartwood proportions**

On each concentric disc with regular sapwood, the diameter under the bark and the heartwood diameter were measured in the one-half disc, and the sectional cross-area was calculated (Figure 2). For the lower stem part, where the limit between sapwood and heartwood was irregular, and in the cases where decay was present, the discs were scanned to measure precisely the surface using the ImageJ software (ImageJ 2012). The sapwood-heartwood boundary was established visually on fresh discs, based on the darker color of the heartwood.
Age

Standard method was used to archive dendrochronological data (Stokes 1968; Holmes 1983). The number of tree rings was counted for the discs at 0 % tree height.

**1.2.4 STATISTICAL ANALYSES**

The moisture content of defoliated (treated) and non-defoliated trees (control) were compared using a repeated measures analysis of variance (ANOVA). The MIXED procedure in SAS was used, with the estimation of the restricted maximum likelihood, and the “UN” covariance structure was applied to the model. The SLICE option of the LSMEANS statement was used when the interaction term defoliation × height was found to be significant to identify which heights differed for the control and for treated trees. Mixed-
models were performed to test for differences in water measurements and sapwood/heartwood ratio between defoliated and control trees. For these mixed-models, sites, defoliation lasting, and intensity of defoliation were used as fixed factors. For the random effects sites were nested to defoliation lasting. Data were log-transformed when necessary to meet the normality and homoscedasticity assumptions. All the analysis was also performed by specie. Differences between mean values were considered significant when p-value was <0.05. Statistical analyses were performed using SAS 9.4 software (SAS Institute Inc. 2013).

1.3 RESULTS

1.3.1 RELATIVE WATER CONTENT AND WATER POTENTIAL

A non-significant trend was observed in relative water content (RWC) for both species. For water potential (Ψwp), only balsam fir showed a non-significant trade (Table 2, Figure 3 AB). Balsam fir showed the mean RWC varied between 72% and 66% in defoliated trees, and water potential from -1.2 and -1.6 MPa (Table 2, Figure 3A). For black spruce, RWC increased from D0 (62%) to D2 (76%), and after decreased to 64% (D4). Water potential did not present a clear trend for black spruce (Table 2, Figure 3B).

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Effect</th>
<th>DF</th>
<th>F</th>
<th>Pr &gt; F</th>
<th>DF</th>
<th>F</th>
<th>Pr &gt; F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Relative Water Content</td>
<td>Defoliation (D)</td>
<td>3</td>
<td>3.14</td>
<td>0.0926</td>
<td>3</td>
<td>0.23</td>
<td>0.8764</td>
</tr>
<tr>
<td></td>
<td>Intensity (IN)</td>
<td>2</td>
<td>0.65</td>
<td>0.5278</td>
<td>2</td>
<td>0.76</td>
<td>0.4733</td>
</tr>
<tr>
<td></td>
<td>Age</td>
<td>1</td>
<td>0.03</td>
<td>0.8626</td>
<td>1</td>
<td>0.53</td>
<td>0.4672</td>
</tr>
<tr>
<td></td>
<td>D x IN</td>
<td>5</td>
<td>0.57</td>
<td>0.7224</td>
<td>5</td>
<td>1.37</td>
<td>0.2444</td>
</tr>
<tr>
<td>Water Potential</td>
<td>Defoliation (D)</td>
<td>3</td>
<td>1.47</td>
<td>0.2757</td>
<td>3</td>
<td>1.68</td>
<td>0.2158</td>
</tr>
<tr>
<td></td>
<td>Intensity (IN)</td>
<td>2</td>
<td>0.62</td>
<td>0.5423</td>
<td>2</td>
<td>0.06</td>
<td>0.9390</td>
</tr>
<tr>
<td></td>
<td>Age</td>
<td>1</td>
<td>0.71</td>
<td>0.4016</td>
<td>1</td>
<td>0.12</td>
<td>0.7284</td>
</tr>
<tr>
<td></td>
<td>D x IN</td>
<td>5</td>
<td>0.94</td>
<td>0.4619</td>
<td>5</td>
<td>0.10</td>
<td>0.9923</td>
</tr>
</tbody>
</table>
1.3.2 MOISTURE CONTENT

The average moisture content (MC) in control and defoliated trees varied between 117% and 123% in the sapwood of black spruce and between 144 and 159% for balsam fir (Table 3). Mean MC of the heartwood of black spruce was stable with a percentage varied...
from 44% to 46%. More variability was obtained for balsam fir with MC heartwood ranging from 67% to 136%. In some trees, MC of heartwood was similar to sapwood in balsam fir and named wetwood.

Table 3 - Mean moisture content (MC) and standard deviation (SD) in sapwood, heartwood, and wetwood of control (D0) and defoliated trees (D1 to D4)

<table>
<thead>
<tr>
<th>Defoliation class</th>
<th>Number of trees</th>
<th>% MC</th>
<th>SD</th>
<th>% MC</th>
<th>SD</th>
<th>Number of trees</th>
<th>% MC</th>
<th>SD</th>
<th>% MC</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Black Spruce</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sapwood</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Heartwood</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Wetwood</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Balsam Fir</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sapwood</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Heartwood</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Wetwood</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Significant changes in MC were observed depending on the tree height for both species and in all the defoliation classes (Table 4, 5). The pattern of black spruce sapwood was more variable between defoliation classes (Table 4), but the general pattern showed the lowest MC at 0% stem height, followed by a gradual increased up to 75% and lower values at 100% stem height. MC values for black spruce heartwood were higher at 0%, decrease at 25% and increase gradually up to 75%. No trend was observed by increasing the duration and intensity of defoliation.

Table 4 - Mean moisture content for control (D0) and defoliated black spruce (D1 to D4) at 0%, 25%, 50%, 75% and 100% stem height for sapwood and heartwood.

<table>
<thead>
<tr>
<th>Defoliation class</th>
<th>Sapwood stem height (% tree height)</th>
<th>Heartwood stem height (% tree height)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0 25 50 75 100</td>
<td>0 25 50 75</td>
</tr>
<tr>
<td>D0</td>
<td>110 a 125 ab 133 bd 128 cd 121 ad</td>
<td>45 a 40 ab 48 ac 41 ab 51 ad</td>
</tr>
<tr>
<td>D1</td>
<td>93 a 128 ab 139 c 126 c 141 bc</td>
<td>43 a 36 ab 39 ab 30 a</td>
</tr>
<tr>
<td>D2</td>
<td>112 a 118 ab 121 bd 136 c 122 ad</td>
<td>53 a 39 c 42 b 50 a</td>
</tr>
<tr>
<td>D3</td>
<td>109 a 122 ab 128 bc 127 c 114 a</td>
<td>45 a 39 b 43 a 48 a</td>
</tr>
<tr>
<td>D4</td>
<td>118 a 120 ab 125 a 132 c 119 ab</td>
<td>53 a 40 c 43 b 49 a</td>
</tr>
</tbody>
</table>

Means in the same horizontal row followed by different letters are significantly different (p≤0.05).
For balsam fir, significant changes in MC were observed depending on the tree height for both heartwood and sapwood of control and defoliated trees (Figure 4A, B). All the classes presented the same pattern, the lowest MC at 0% stem height, followed by a gradual increase up to the middle of the stem and lower values at the top tree (Table 5). The trees attacked by the spruce budworm had a slightly higher non-significant trade of sapwood moisture content than the control trees at all measured heights (Figure 4A). The highest difference was found at 0% stem height, varying between 123% (D0) and 149% (D2) of moisture content.

Table 5 - Mean moisture content for control (D0) and defoliated balsam fir (D1 to D4) at 0%, 25%, 50%, 75% and 100% stem height for sapwood and heartwood.

<table>
<thead>
<tr>
<th>Defoliation class</th>
<th>Sapwood stem height (% tree height)</th>
<th>Heartwood stem height (% tree height)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0</td>
<td>25</td>
</tr>
<tr>
<td>D0</td>
<td>123&lt;sup&gt;a&lt;/sup&gt;</td>
<td>149&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>D1</td>
<td>133&lt;sup&gt;a&lt;/sup&gt;</td>
<td>163&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>D2</td>
<td>149&lt;sup&gt;a&lt;/sup&gt;</td>
<td>162&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>D3</td>
<td>136&lt;sup&gt;a&lt;/sup&gt;</td>
<td>159&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>D4</td>
<td>145&lt;sup&gt;a&lt;/sup&gt;</td>
<td>154&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

Means in the same horizontal row followed by different letters are significantly different (p≤0.05). Heartwood over 100% moisture content is not presented.

Despite the changes observed for both species in the moisture content along the stem in the sapwood and heartwood, the statistical analyses do not show an interaction of the term defoliation × height (Table 6). Statistical analyses pointed out only significant differences in heartwood of balsam fir. However, the changes were not observed between control and defoliated trees, but between the different defoliation classes. Tree age was significant (P=0.0493) only in black spruce in the sapwood.
Table 6 - Repeated measure of moisture content (MC) in a mixed model procedure (Using SAS software, PROC MIXED)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Effect</th>
<th>Black Spruce</th>
<th>Balsam Fir</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>DF</td>
<td>F Value</td>
<td>Pr &gt;F</td>
</tr>
<tr>
<td>Sapwood</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Defoliation (D)</td>
<td>3</td>
<td>0.95</td>
<td>0.4452</td>
</tr>
<tr>
<td>Intensity (IN)</td>
<td>2</td>
<td>0.22</td>
<td>0.7992</td>
</tr>
<tr>
<td>Height (H)</td>
<td>4</td>
<td>33.07</td>
<td>&lt;.0001*</td>
</tr>
<tr>
<td>Age</td>
<td>1</td>
<td>3.98</td>
<td>0.0493*</td>
</tr>
<tr>
<td>D*H</td>
<td>12</td>
<td>1.36</td>
<td>0.1930</td>
</tr>
<tr>
<td>D<em>IN</em>H</td>
<td>20</td>
<td>1.38</td>
<td>0.1394</td>
</tr>
<tr>
<td>Heartwood</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Defoliation (D)</td>
<td>3</td>
<td>1.00</td>
<td>0.4094</td>
</tr>
<tr>
<td>Intensity (IN)</td>
<td>2</td>
<td>0.79</td>
<td>0.4569</td>
</tr>
<tr>
<td>Height (H)</td>
<td>3</td>
<td>31.05</td>
<td>&lt;.0001*</td>
</tr>
<tr>
<td>Age</td>
<td>1</td>
<td>0.01</td>
<td>0.9191</td>
</tr>
<tr>
<td>D*H</td>
<td>9</td>
<td>0.84</td>
<td>0.5849</td>
</tr>
<tr>
<td>D<em>IN</em>H</td>
<td>15</td>
<td>0.62</td>
<td>0.8524</td>
</tr>
</tbody>
</table>

Figure 4 - Comparison of MC for balsam fir trees in (A) sapwood, (B) heartwood (>100%), and wetwood (<100%) between control trees (D0) and four years of defoliation (D4) by height (% stem height).
1.3.3 SAPWOOD AND HEARTWOOD RATIO

The same pattern observed in MC was also noted in the sap-heartwood ratio, the values were significantly different depending on the stem height for both species, but no effect of the interaction of the stem height was found associated with defoliation (Table 7).

Table 7 - Comparisons among defoliation class for sapwood/heartwood area ratio (Using SAS software, PROC MIXED)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Effect</th>
<th>Black Spruce</th>
<th>Balsam Fir</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>DF</td>
<td>F Value</td>
<td>Pr &gt;F</td>
</tr>
<tr>
<td>Defoliation (D)</td>
<td>3</td>
<td>1.67</td>
<td>0.2147</td>
</tr>
<tr>
<td>Intensity (IN)</td>
<td>2</td>
<td>0.03</td>
<td>0.9707</td>
</tr>
<tr>
<td>Height (H)</td>
<td>3</td>
<td>37.56</td>
<td>&lt;.0001*</td>
</tr>
<tr>
<td>Age</td>
<td>1</td>
<td>26.95</td>
<td>&lt;.0001*</td>
</tr>
<tr>
<td>D*H</td>
<td>9</td>
<td>0.38</td>
<td>0.9436</td>
</tr>
<tr>
<td>D<em>IN</em>H</td>
<td>15</td>
<td>0.41</td>
<td>0.9760</td>
</tr>
</tbody>
</table>

Overall for black spruce, sapwood area was low at 0%, 25% and 50% of the stem height and much higher at 75%, except for D1 class (Figure 5). Whereas for balsam fir the pattern was more variable at 75%, with lowest values for D0 and D4 classes (Figure 5).

![Figure 5 - Ratio Sapwood/Heartwood area for black spruce and balsam fir by height (% stem height).](image-url)
1.4 DISCUSSION

A consecutive defoliation, lasting from one to four years, in different intensities had no effect in balsam fir and black spruce on either moisture content or sapwood/heartwood ratio. Moreover, repeated defoliation of the current-year needles had no effect on relative water content and water potential.

1.4.1 RELATIVE WATER CONTENT AND WATER POTENTIAL

Herbivore causes a variety of negative physiological responses in most plants (Reich et al. 1993; Clancy et al. 1995). The first immediate effect of defoliation is the reduction of plant’s capability to obtain carbon (Smith et Hinckley 1995; Deslauriers et al. 2015), which reduces the amount of available carbohydrates (Li et al. 2002). Severe defoliation can also deplete the plant’s carbohydrates reserves, so tree growth may decline, and in some cases, the tree death can occur (Reich et al. 1993; Krause et Morin 1995a; Rossi et al. 2009).

Depending on the publications, negative or positive effects were reported for water availability after defoliation (Kolb et al. 1999; Hart et al. 2000; Lavigne et al. 2001; Jacquet et al. 2014). An increase of water availability was associated with a compensatory photosynthesis mechanism following defoliation by several authors (Stephens et al. 1972; Wright et Berryman 1978; Gieger et Thomas 2002; Quentin et al. 2012). This compensatory photosynthesis mechanism in plants has been greatly discussed (Heichel et Turner 1983; Nowak et Caldwell 1984; Hoogesteger et Karlsson 1992; Ovaska et al. 1992). While it is clear that some plants can compensate in part the damages for defoliation by improved water relations, the extent and timing of this response are poorly understood (Meinzer et Grantz 1990; Ovaska et al. 1992; Hart et al. 2000). The photosynthesis stimulation after
defoliation has been attributed to either improved water relations, reduced foliar carbohydrate concentrations, or increased allocation of mineral nutrients (Ovaska et al. 1992; Clancy et al. 1995; Lavigne et al. 2001).

In our study the defoliation of black spruce and balsam fir did not affect water relations. However, it is possible that the water potential in this study did not represent adequately all variations in the defoliated trees since the measurements were taken at different moments during the day, over a six-week period. Inherent differences entailed by temperature and relative humidity changes along the day (expressed by the VPD) may have confused the interpretation of the data.

The typical pattern expected in plants is decreasing stomatal conductance with increasing VPD (Zhao et Running 2010). By dropping stomatal conductance to water vapor, plants can diminish water deficit and maintain the hydration of plant cells as VPD increases (Ocheltree et al. 2014). The response of stomatal aperture to VPD varies with environmental factors as light, air, and soil temperature, and soil water potential (Graham et Running 1984; Beadle et al. 1985; Goldstein et al. 1985; Pallardy et al. 1995). All those factors together become challenging to control under natural forestry conditions in northern areas defoliated by spruce budworm.

Furthermore, the influence of VPD on water potential can differ among different conifers species (Grossnickle 2000; Zhao et Running 2010; Ocheltree et al. 2014). Black spruce is described as having a direct influence on the water potential and stomatal conductance with the daily change in VPD (Grossnickle et Blake 1985; Fan et Grossnickle 1998; Grossnickle 2000). However, the balsam fir behavior in field conditions to different VPD has not been yet described. Lavigne et al. (2001) observed for the balsam fir seedling suffering artificial defoliation in controlled greenhouses conditions, no significant changes
in water potential to defoliation. The author suggests that these results might be due to a response of increased photosynthetic rate. In this study, for the water potential and relative water content, balsam fir presented a non-significant reduced trend when defoliation duration increased (Figure 3A).

1.4.2 SAPWOOD AND HEARTWOOD: MC AND AREA RATIO

On living balsam fir and black spruce trees, no effects of defoliation were observed on neither MC nor in the sapwood/heartwood ratio. The average moisture content (MC) in black spruce was in agreement with the literature (Krause et Gagnon 2006; Savard 2010) oscillating around 120% in the sapwood and 45% in the heartwood. Balsam fir presented slightly higher MC values, around 151% in sapwood and from 67% to 136% in the heartwood in accordance with Etheridge et Morin (1962) and Jeremic et al. (2004). The sapwood proportion increased exponentially at 75% tree height for both species. These findings are coherent with previously published results for sound trees in both species (Shmulsky et Jones 2011). Thus, the hypothesis that defoliated trees would form less sapwood area, and therefore moisture content would decrease in sapwood by the fact of lower water transported was rejected.

As expected, the MC of the sapwood was higher than that of heartwood (except for some balsam fir trees that presented wetwood). The sapwood has several primordial functions, as water storage and water transport from the roots to the foliage (Gartner 1995; Pallardy et al. 1995) whereas heartwood has only mechanical function (Woo et al. 2005; Lowell et al. 2010).

From a hydraulic point of view, trees that suffer defoliation are at a disadvantage considering they have smaller carbon reserves available for allocation to fine root
production. Fine roots develop an important role for water and nutrients acquisition in the tree (Pallardy et al. 1995; Schäfer et al. 2013; Deslauriers et al. 2015). Further, the amount of foliage on a tree is often strongly correlated to the amount of sapwood (Long et Smith 1988; Ryan 1989; Berthier et al. 2001). However, the period from one to four years of consecutive defoliation presented in this study apparently does not have effect in the water transport, and consequently, in the moisture content and sapwood/heartwood ratio.

After a consecutive defoliation period, the loss of foliage normally translates into changes in the formation of latewood cells, whereas the formation of earlywood cells remains unaltered (Krause et Morin 1995a; Schmitt et al. 2003). Vejpustkova et Holusa (2006) and Axelson et al. (2014) studied different insect-trees interactions in conifers species and noted important reductions in latwood anatomical features, while the cellular characteristics of the earlywood remained fairly constant. The cells of earlywood are directly related to the hydraulic role of sapwood (Deslauriers 1999; Domec et Gartner 2002; Lupi et al. 2010). This may suggest that the changes observed in the anatomical features after defoliation are not significant to reduced hydraulic efficiency in the stem, because they mainly occur in latewood cells (Vejpustkova et Holusa 2006; Axelson et al. 2014).

Furthermore, conifers species in the boreal forests have been evolving to manage challenging strategies to survive in harsh and extreme conditions by millennia (Gartner 1995; Domec et Gartner 2002; Eder et al. 2009; West 2014). It can be argued that mature balsam fir and black spruce have developed strategies to reduce the effects of the foliage loss in some physiological aspects such water relations, water transport and storage into the tree during the first four years of defoliation.
1.5 CONCLUSION

In light of the results obtained, it can be suggested that the period from one to four years of consecutive defoliation by the spruce budworm does not have any effect on the tree water relations (water potential and relative water content), sapwood proportion and stem moisture content in balsam fir and black spruce. Thus the hypothesis of the reduction in foliage would be compensated by decreases in the water potential or relative water content into the tree, and consequently in sapwood formation and stem moisture content was rejected.

Despite the limited number of sites sampled, the analyses provided detailed information on the temporal changes in sapwood proportion and wood moisture content at different tree heights. However, we are aware that our results require additional investigations to be validated with extensive samplings involving more trees and sites. Further, future studies on the effects of defoliation in the water potential in mature trees, especially in black spruce, should be taken in pre-dawn to minimize the effects of VPD in the results.
ÉVALUATION DE QUELQUES PARAMÈTRES DE LA QUALITÉ DU BOIS AFFECTÉ PAR L’ÉPIDÉMIE DE LA TBE

ABSTRACT

Spruce budworm (Choristoneura fumiferana Clem.) is one of the most widely distributed and impressive defoliator agent of coniferous forests in eastern North America. In Canada, balsam fir (Abies balsamea L. (Mill)) and black spruce (Picea mariana B.S.P. (Mill)) are the most important and widely distributed hosts. Thirty-six infested stands, varying from one to four years of defoliation, were sampled in the Quebec boreal forestry in 2016 and 2017 to determine whether modifications had occurred in the wood quality of the infected trees. Ring growth, wood density, and anatomical characteristics of stem wood formed during the outbreak years was analyzed for both species. We determined that rings formed during the spruce budworm outbreak had a significantly and progressively loss of volume with the lasting of defoliation, reduced latewood density in the second and third year of defoliation in black spruce, and third and fourth year in balsam fir. A reduced average wood density only for black spruce after four years of defoliation was also measured. These changes were related with changes in the anatomical features. While the cellular characteristics of the earlywood remained fairly constant, significant reductions in cell wall occurred only after three years of defoliation. Our study shows that spruce budworm outbreak not only reduce annual radial growth, but also temporarily modify cellular characteristics in latewood cells, which has implications for wood density and quality in black spruce and balsam fir.

Keywords: Spruce budworm outbreak, wood quality, Picea mariana, Abies balsamea, growth reduction, wood density, lumen area, cell wall thickness, early- and latewood.
2.1 INTRODUCTION

The boreal forest biome covers much of the landmass of the northern hemisphere and constitutes a very large pool of the global carbon stock (Melillo et al. 1993; Dixon et al. 1994). The dynamics of the boreal forest are periodically influenced by two principal natural disturbances: fires (Cogbill 1985; MacLean 2016) and insect epidemics (MacLean 1985; Stocks 1987). In particular, the spruce budworm (Choristoneura fumiferana Clem.) is one of the most widely distributed and impressive defoliator agents in the Canadian boreal forest (Morris 1963; Blais 1983; Lapointe 2013; MacLean 2016). Since 1909 there have been three major spruce budworm outbreaks destroying hundreds of thousands of hectares of valuable forest stands (Morin et al. 2007). Damages occur at the larval stage. Spruce budworm feeds on needles and buds of balsam-fir (Abies balsamea L. (Mill)), the preferred host, and with the population explosion, other coniferous trees are damaged, such as black spruce (Picea mariana B.S.P. (Mill)) (Blais 1983; Hennigar et al. 2008). A new outbreak is currently developing in the eastern boreal forest following its usual cycle, but now it occurs in more northern areas than previous outbreaks, where black spruce, a species that is not recognized as a privileged host, is dominant (Gouvernement du Québec 2014).

Once the forest is attacked by the spruce budworm, the infested trees become progressively defoliated (Blais 1958; Dorion 2016). The damages first appears on the upper crown of the tree, restricted to a partial loss of new foliage, and then spreads towards the lower branches (Carisey et Bauce 1997; Garcia 2015). Several studies have shown a decrease in radial growth after defoliation periods as a result of repeated losses of new foliage over a number of years of infestation (Blais 1962; MacLean 1984; Krause et al. 2003; Krause et al. 2012), and consequently tree mortality following four to five years of severe infestation in the case of balsam fir, and a little longer for black spruce (Blais 1965; MacLean
Outbreaks play a significant role in forest yield (MacLean 2016), spruce budworm typically results in an average of 42-50% stand mortality (MacLean 1980; Bergeron et al. 1995) and volume losses varying between 32 and 48% (Archambault 1983; Ostaff et MacLean 1995). From 1975 to 2000, spruce budworm outbreaks defoliated a total area exceeding 450 M ha (MacLean 2016). In Quebec, from 139 to 238 million m$^3$ of spruce and fir timber was lost due to budworm during the last epidemic period between 1967 and 1992 (Boulet et al. 1996; Marmen 2014). From an economic viewpoint, volume losses are crucial. However, after tree death, besides direct losses, insect attacks can also result in reductions of quality of wood harvested from budworm-infested-stands (Binotto et Locke 1981; Basham 1984; Koran et Nlombi 1994).

Forest product properties depend strongly on wood characteristics such as ring density, modulus of elasticity (MOE) and microfibril angle (Zhang et al. 1996; Downes et Drew 2008). More specifically, wood density has a major impact on the yield, quality and value of wood-based composites and solid wood products (Shi et al. 2007). From an anatomical point of view, the major factor explaining the variation of wood density is the proportion of latewood and, specifically, cell wall thickness (Zobel et Buijtenen 1989; Ivkovich et al. 2002). This suggests that differences in cell wall thickness, as well as the percentage of latewood, should translate into changes in mechanical wood properties (Krause et al. 2010). Since the formation of the latewood tracheids is mainly dependent on the growing conditions of the current year, it may be directly affected by defoliation (Schweingruber 1979; Krause et Morin 1995a). Vejpustkova et Holusa (2006) observed in larch trees defoliated by Cephalcia lariciphila in the Czech Republic a decrease in radial growth, followed by a decrease in latewood density, as a result of latewood cell-wall thickness reduction. Axelsson et al. (2014) also detected that Douglas-fir rings formed during
a western spruce budworm outbreak in Canada had significantly reduced ring width followed by decreases in latewood cell wall thickness and cell width.

Over the last two decades, several studies have focused on quantifying and understanding the growth losses caused by spruce budworm in balsam fir (Piene et al. 1989; Krause et al. 2003), and in black spruce (MacLean et MacKinnon 1997; Krause et Morin 1999; Krause et al. 2012), and how defoliation might affect cambium activity (Krause et Morin 1995a; Rossi et al. 2009; Deslauriers et al. 2015). However, the degree to which tree defoliation alter wood formation and in turn the wood density of balsam fir and black spruce remains unclear.

This paper aims to assess the volume loss, wood density and the variation of anatomical characteristics in black spruce and balsam-fir trees affected by the current spruce budworm outbreak. This was done by examining inter-annual changes in xylem characteristics formed during the defoliation period. The hypotheses were that: a significant volume loss would occur after the beginning of defoliation, and would be more pronounced in balsam fir than in black spruce (1); the reduced radial growth of the stem following defoliation would occur as a result of changes in anatomical structure (lumen area and cell wall thickness in earlywood and latewood) (2); thus wood density properties would also decrease following defoliation since they are highly correlated with the latewood proportion and ratio lumen area/cell wall thickness (3). The findings of the research will contribute to our understanding the effects of spruce budworm on wood quality, and the resilience and plasticity of the two main harvested species in Eastern North America.
2.2 METHODOLOGICAL APPROACH

2.2.1 STUDY AREA

The study area covers approximately 10 000 km\(^2\) at the transition between northern temperate and boreal forest in Quebec, Canada (48°25'N 71°04'W). The region is characterized by hills that rarely exceed 500 m, on thick and undifferentiated glacial till deposits. This area is part of balsam fir-white birch, balsam fir-yellow birch, and spruce-moss bioclimatic domain (Figure 1). The climate is continental and has a short warm summer, without a dry season. The 30-year mean annual temperature is 2.8° C and total precipitation of 930 mm, with 5-month covert by snow (Government of Canada 2017).

2.2.2 TREE SELECTION

The infested areas was assessed using the spruce budworm annual aerial detection survey data from the government of Quebec (MFFP 2016; 2017). In order to delimit the sampling area, some factors were considered, such as: public lands that were not harvested in the last years, stands defoliated between one to four years of defoliation, stands within 150 kilometers from the Lake Saint-Jean. Finally, twenty natural stands dominated by balsam fir \textit{(Abies balsamea (L.) Mill.)} and twenty-one dominated by black spruce \textit{(Picea mariana (Mill.))} were selected. At each site, representative defoliated individuals were chosen randomly among healthy trees. A visually assessment of the defoliation was made with binoculars before harvesting. The trees were sampled in the summers of 2016 and 2017 (June and early July).

Stands were divided according to the duration of defoliation in five classes: from no defoliation to up to four continuous years of defoliation. D0 represents control trees without defoliation, D4 four years of consecutive defoliation. D1, D2, and D3 represent one to three
years of continuous defoliation. Additionally, a visual assessment of defoliation intensity was carried out for each sampled tree in the field. Three intensities of defoliation were used: light (25% needle loss), moderate (from 26% to 70% needle loss), and severe (more than 71% needle loss). Thus, the goal of this study is understand the combined effects of timing and intensity of defoliation in each parameter, where each defoliation class represents the duration of consecutive defoliation, and intensity of defoliation, the degree of needle loss. Preliminary analysis showed more variation in the data between the trees with a longer defoliation lasting. For this reason, the number of harvested trees varied between the defoliation classes, with more trees to a longer defoliation period (Table 1).

Stand characterization was done including information and measurements of stand composition, slope, drainage, and soil type. Height, diameter at breast height (DBH) and crown length were also measured on 15 live trees (si) (Table 1). Soil profiles were classified into three broad categories: well drained (64 % of the study sites), moderately (20 %) and poorly drained (16 %). Two types of soils were found: podzolic (73 %) and organic soils (23 %). Tree age ranged from 24- to 132-year-old for the balsam fir and 29- to 172-year-old for black spruce stands (Table 1).

A total of 119 black spruces and 119 balsam firs were harvested (Table 1). Five stem discs were collected from each tree, located at tree base (0m), DBH (1.30m), 25%, 50%, 75% tree height.

2.2.3 MEASUREMENTS

Age and Growth estimation

A standard method was used to archive dendrochronological data (Stokes 1968); discs were cut in half, air-dried and transversal surface cut or sanded (Figure 6). Tree-ring
widths were measured to an accuracy of 0.01 mm using a WinDendro measuring system (Régent Instruments Inc 2011) along two paths per disc. Crossdating was performed visually and by using the COFECHA computer program for the discs at 0 m (Holmes 1983).

To standardize the data, radial growth in the defoliated and control trees was evaluated based on the average of the growth in the last 20 years, for the respective height and defoliation class. The ring width index was calculated as follows:

\[
\text{Ring width index} = \frac{\text{Annual ring width}}{\text{20-years mean annual ring width}}
\]

To assess the growth response to defoliation, the current (CAI) and periodic (PAI) annual volume increments were assessed for each tree 15 years prior to the outbreak. The volume of a tree for one particular year was estimated as the sum of the volumes of cone sections delimited by two consecutive discs whose diameters were calculated from a quadratic mean of two radii. A linear equation of the form \(y_1=ax_1+b\) was used to express the relationship between the CAI of the stem before the infestation and the age of each defoliated tree. The mean correlation coefficient of these equations was 0.92 for black spruce and 0.95 for balsam fir. Theoretical growth (without infestation) was assessed by extrapolating this equation and the growth loss was obtained by subtracting observed growth from theoretical growth (Archambault 1983). In order to minimize the probability of overestimating the theoretical increment, only trees that had not reached their maximum current annual increment in volume were used in the analysis. Furthermore, trees with equation correlation lower than 0.85 were disregarded.
**Wood density and Anatomical features**

Wood density and anatomical measurements are time-consuming, we limited these analyses to trees with three and four years of consecutive defoliation. Furthermore, eighty breast height discs (40 fir and 40 spruce trees) were randomly chosen from 137 discs sampled. Strips of 1.63 mm thick (longitudinal) and 25 mm wide (tangential) were obtained from radial segments (Figure 6). The strips were dried to 12% moisture content and measurements were made from bark to pith at intervals of two mm, using an X-ray densitometer available at Université Laval, Quebec City, Canada. The transition from early-to late-wood within a ring was defined as the point where the maximum change in density was reached (Mothe et al. 1998; Lemay et al. 2016). Earlywood, latewood, and average ring density were measured for every annual ring of each sample. Only the data from the 11 years prior to the defoliation until the last year of defoliation were kept for the analyses.

The same eighty discs were used for the wood anatomical analyses. Wood samples containing the last five annual rings were collected. The samples were embedded in paraffin (Leica TP1020), cut into sections of 7μm with a rotary microtome (Leica RM2145), fixed on slides, and stained with an aqueous solution of 1% safranin. A camera fixed on an optical microscope was used to record numerical images at a magnification of 20×. Lumen area, cell wall thickness, cell width, and cell numbers were measured in each ring on three radial files per ring using Wincell v.2010 (Deslauriers et al. 2003).
2.2.4 STATISTICAL ANALYSES

All statistical analyses were performed using the software SAS, version 9.4 (SAS Institute Inc. 2013). The MIXED procedure in SAS was used, with the estimation of the restricted maximum likelihood, and the “UN” covariance structure was applied to the model. For the growth analysis, mixed-models were performed to test for differences in growth between individual years of defoliation (2013-2016), the eleven years prior to defoliation (2002-2012) and the last fifteen years in non-defoliated control trees (2002-2016). Individual years had to be significantly different from both, control and non-defoliated periods, to reject the null hypothesis. The following fixed effect factors were used in the mixed-models: defoliation class (D), intensity of defoliation (IN), year of the growth ring, the respective year of defoliation during the formation of growth ring (Y), and the stem height (H) for ring width.
When the interaction term defoliation class \( (D) \times \text{Year of Defoliation} \) was found significant, the SLICE option of the LSMEANS statement was used. This statement allowed to identify in the cumulative period of defoliation which years differed between the control and defoliated trees.

The same mixed-models were performed for wood density and anatomical properties. However, the control period used was only the eleven years prior to defoliation (2002-2012) for wood density and the last year before defoliation (2012) for anatomical analysis. Because only trees with more than 3 years of cumulative defoliation were used, the defoliation class effect was no more used. The SLICE option of the LSMEANS statement was also performed to identify the years \((Y)\) and the intensities \((IN)\) that differed between the control and treated trees.

Data were log-transformed when necessary to meet the normality and homoscedasticity assumptions. Differences between mean values were considered significant when the \( p\)-value was <0.05.

2.3 RESULTS

2.3.1 RING WIDTH

There was a clear negative effect of the defoliation on growth (Figure 7). Comparing the pre-outbreak period with the post-outbreak, 95 % of the defoliated black spruce trees showed a radial growth decrease after defoliation on the average of the four measured heights, whereas this percentage was nearly 90 % for balsam fir trees. For black spruce, there was a mean radial growth loss of 34 % (maximum 78 % loss) while balsam firs had an average of 31 % (maximum 83 % loss).
In the pre-outbreak period, the ring width index of the four heights was similar to that of the control trees for both species (statistical analysis not shown) but a decreasing nonsignificant trend was observed after each consecutive year of defoliation for balsam fir, and significant for Black spruce in all D3 class (Table 8A, Figure 7). The decrease in ring width on the average of the four heights was, respectively, 22, 39, 36, and 33 % from the period of one to four years of consecutive defoliation for black spruce and 31, 18, 40, 28 % for balsam fir.

By looking at the defoliation intensity of black spruce, the second year of defoliation was significantly lower in the moderate and severe intensities in the defoliation class D2. In D3, the severe defoliation intensity was different from control trees from the first until the third year of defoliation; for the last year, the three defoliation intensities presented differences among them. In D4, only the severely defoliated trees were significantly lower in the third and fourth year compared to control trees (Figure 8).
Figure 7 - Average radial growth index for the four stem heights for balsam fir and black spruce for control trees and the four defoliation classes; each different symbol represents a year of defoliation. Control class are shown in the purple row. The colored area represents the length of defoliation period. Asterisks inside shaded areas indicate a significant difference between defoliated and control trees for a given year, as determined by a slice test.

![Diagram of balsam fir and black spruce growth index]

Figure 8 - Ring width index by defoliation class for black spruce. Only significant results are presented for black spruce at the 0.05 levels. L (Light), M (Moderate), and S (Severe). "N" in parentheses showed the number of trees for each defoliation intensity.

<table>
<thead>
<tr>
<th>Defoliation class</th>
<th>D2</th>
<th>D3</th>
<th>D4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Year of defoliation</td>
<td>I → II → M → S</td>
<td>I → II → III → M → S</td>
<td>I → II → III → IV → S</td>
</tr>
<tr>
<td>Intensity</td>
<td>L (9) M (9) S (3)</td>
<td>L (20) M (8) S (5)</td>
<td>L (18) M (7) S (10)</td>
</tr>
<tr>
<td>N</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Figure 8 - Ring width index by defoliation class for black spruce. Only significant results are presented for black spruce at the 0.05 levels. L (Light), M (Moderate), and S (Severe). "N" in parentheses showed the number of trees for each defoliation intensity.
For both species, there was a decrease of radial growth along the stem (Figure 9). The radial growth loss for black spruce varied on average from 26 to 37 % and for balsam fir from 13 to 33 %. However, statistical analysis did not show a significant difference associated with duration and intensity of defoliation.

Figure 9 - Radial growth by height for balsam fir and black spruce. (A) Control trees and (B) D4 defoliation class (four years of consecutive defoliation). The shaded area represents the defoliation period, with I being the first year of defoliation.

2.3.2 MEAN ANNUAL INCREMENT VOLUME (MAI)

There was a visible negative effect of defoliation duration in the mean annual volume increment between defoliated and control trees for both species (Figure 10). Defoliated black spruce trees had a significant volume loss of 13, 27, 39, and 46 % respectively, from the first to the fourth year of defoliation compared to control trees. Annual increment volume decreased significantly for balsam fir trees with losses of 10, 20, 34, and 38 % for the same period. All the years showed a significant difference between control and defoliated trees, except the first year in D4 defoliation class for balsam fir (Figure 10). The evaluated intensity
of defoliation for all defoliation classes was not significant for either species compared to control trees (Table 8B).

![Diagram](image)

**Figure 10** - Annual volume loss by defoliation class for balsam fir and black spruce before and after a consecutive period of defoliation. Each different symbol represents a year of defoliation. The colored area represents the length of defoliation period. Asterisks inside shaded areas indicate a significant difference between defoliated and control trees for a given year, as determined by a slice test.

### 2.3.3 WOOD DENSITY

Only trees defoliated for three and four years were analyzed for wood density and cellular features. Comparing the pre-outbreak period (2002-2012) with the last 4-year post-outbreak period (2013-2016) showed that black spruce presented clearly higher wood densities and slightly higher cumulative changes than balsam fir (Figure 11). Despite this
pattern, the outbreak period did not affect latewood proportion and earlywood density in either species (Table 8C,E, Figure 11A,C). For the ring density, a decreasing trend was observed for black spruce in the outbreak period, with significant changes only after four years of defoliation, and no changes were calculated for balsam fir (Table 8D, Figure 11B).

In black spruce, latewood density continuously decreased over the defoliation period from years one to three, with significant values for the second and third years of defoliation (Table 8F, Figure 11D). Balsam fir showed a decrease over the entire period, with significant reductions in the third and fourth years of defoliation (Table 8F, Figure 11D). The defoliation intensities affected only balsam firs in latewood density for severely defoliated trees (Table 8F).

### Table 8 - Mixed model ANOVA results of the measured growth and wood quality parameters for black spruce and balsam fir (Using SAS software, PROC MIXED)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Effect</th>
<th>Black spruce</th>
<th>Balsam fir</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>DF</td>
<td>F</td>
<td>Pr &gt; F</td>
</tr>
<tr>
<td>(A) Ring width</td>
<td>Defoliation (D)</td>
<td>3</td>
<td>0.74</td>
</tr>
<tr>
<td></td>
<td>Year (Y)</td>
<td>4</td>
<td>25.45</td>
</tr>
<tr>
<td></td>
<td>Intensity (IN)</td>
<td>2</td>
<td>1.29</td>
</tr>
<tr>
<td></td>
<td>Height (H)</td>
<td>3</td>
<td>53.41</td>
</tr>
<tr>
<td></td>
<td>Age</td>
<td>1</td>
<td>302.13</td>
</tr>
<tr>
<td></td>
<td>D x Y</td>
<td>6</td>
<td>2.38</td>
</tr>
<tr>
<td></td>
<td>D x Y x IN</td>
<td>11</td>
<td>2.51</td>
</tr>
<tr>
<td></td>
<td>D x Y x IN x H</td>
<td>33</td>
<td>0.49</td>
</tr>
<tr>
<td>(B) Volume loss (%)</td>
<td>Defoliation (D)</td>
<td>3</td>
<td>3.86</td>
</tr>
<tr>
<td></td>
<td>Year (Y)</td>
<td>4</td>
<td>72.54</td>
</tr>
<tr>
<td></td>
<td>Intensity (IN)</td>
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<td>2.5</td>
</tr>
<tr>
<td></td>
<td>Age</td>
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<td>7.01</td>
</tr>
<tr>
<td></td>
<td>D x Y</td>
<td>6</td>
<td>7.48</td>
</tr>
<tr>
<td></td>
<td>D x Y x IN</td>
<td>11</td>
<td>1.75</td>
</tr>
<tr>
<td>(C) Latewood %</td>
<td>Year (Y)</td>
<td>3</td>
<td>0.20</td>
</tr>
<tr>
<td></td>
<td>Intensity (IN)</td>
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<td>1.03</td>
</tr>
<tr>
<td></td>
<td>Age</td>
<td>1</td>
<td>19.95</td>
</tr>
<tr>
<td></td>
<td>Y x IN</td>
<td>6</td>
<td>1.39</td>
</tr>
<tr>
<td>(D) Ring density</td>
<td>Year (Y)</td>
<td>3</td>
<td>3.22</td>
</tr>
<tr>
<td></td>
<td>Intensity (IN)</td>
<td>2</td>
<td>0.15</td>
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<tr>
<td></td>
<td>Age</td>
<td>1</td>
<td>9.06</td>
</tr>
<tr>
<td></td>
<td>Y x IN</td>
<td>6</td>
<td>1.68</td>
</tr>
<tr>
<td>(E) Earlywood density</td>
<td>Year (Y)</td>
<td>3</td>
<td>2.22</td>
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<td>Intensity (IN)</td>
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<td></td>
<td>Age</td>
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<td>5.96</td>
</tr>
<tr>
<td></td>
<td>Y x IN</td>
<td>6</td>
<td>1.03</td>
</tr>
<tr>
<td>(F) Latewood density</td>
<td>Year (Y)</td>
<td>3</td>
<td>3.01</td>
</tr>
<tr>
<td></td>
<td>Intensity (IN)</td>
<td>2</td>
<td>0.86</td>
</tr>
<tr>
<td></td>
<td>Age</td>
<td>1</td>
<td>6.01</td>
</tr>
<tr>
<td></td>
<td>Y x IN</td>
<td>6</td>
<td>1.91</td>
</tr>
</tbody>
</table>

**Significant results are presented in bold.**
Figure 11 – (A) Latewood proportion, (B) average ring density, (C) earlywood density, (D) latewood density for black spruce and balsam fir stems before and after defoliation. The shaded area represents the defoliation period; with I being the first year of defoliation. Significant differences between the year before and a given year after defoliation are represented with asterisks for balsam fir and with “x” for black spruce.
2.3.4 ANATOMICAL FEATURES

Figure 12 shows the lumen area, mean cell wall thickness, radial cell width, and number of xylem cells in early and latewood for 2012, the year before the defoliation, and the consecutive defoliation period (2013-2016). For all the anatomical features balsam fir showed higher values than black spruce, however, the changes observed were quite similar during the outbreak period for both. In lumen area, both species demonstrated the same pattern, an increasing trend in earlywood and decreasing in latewood. Balsam fir showed significant differences (Table 9A,E, Figure 12A) in earlywood, however statistical analyses could not pinpoint a single year and the defoliation intensity where it was different, and in latewood the differences period was found in the first, third and four years of defoliation.

A significant decreasing trend in latewood for all the defoliation period for black spruce and only after three years of consecutive defoliation for balsam fir was detected in cell wall thickness and cell width (Table 9F,G, Figure 12B,C). In earlywood, the two species presented significant differences only after three years of consecutive defoliation (Table 9B,C), but with different patterns. The defoliation period tends to decrease the cell wall and cell width in black spruce, while balsam fir does not have a stable pattern in cell wall thickness and a slight increasing trend for cell width (Figure 12B,C). The cell number also decrease significantly with the increase of duration of defoliation for both species in early and latewood (Table 9D,H, Figure 12D). The defoliation intensities affect balsam fir in early and latewood and black spruce only in latewood for cell wall thickness (Table 9C,E,F, Figure 13).
Table 9 - Mixed model ANOVA results of the lumen area, cell wall thickness, cell thickness, and cell number in early and latewood for black spruce and balsam fir.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Effect</th>
<th>Black Spruce</th>
<th>Balsam Fir</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Earlywood</strong></td>
<td></td>
<td>DF</td>
<td>F</td>
</tr>
<tr>
<td>(A) Lumen Area</td>
<td>Year (Y)</td>
<td>3</td>
<td>0.77</td>
</tr>
<tr>
<td></td>
<td>Intensity (IN)</td>
<td>2</td>
<td>6.86</td>
</tr>
<tr>
<td></td>
<td>Age</td>
<td>1</td>
<td>4.27</td>
</tr>
<tr>
<td></td>
<td>Y x IN</td>
<td>6</td>
<td>1.50</td>
</tr>
<tr>
<td>(B) Average Wall Thickness</td>
<td>Year (Y)</td>
<td>3</td>
<td>3.29</td>
</tr>
<tr>
<td></td>
<td>Intensity (IN)</td>
<td>2</td>
<td>0.39</td>
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<tr>
<td></td>
<td>Age</td>
<td>1</td>
<td>0.66</td>
</tr>
<tr>
<td></td>
<td>Y x IN</td>
<td>6</td>
<td>0.38</td>
</tr>
<tr>
<td>(C) Cell Thickness</td>
<td>Year (Y)</td>
<td>3</td>
<td>4.63</td>
</tr>
<tr>
<td></td>
<td>Intensity (IN)</td>
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<td>3.02</td>
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<tr>
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<td>Age</td>
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<tr>
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</tr>
<tr>
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<tr>
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<tr>
<td><strong>Latewood</strong></td>
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<td>F</td>
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<tr>
<td>(E) Lumen Area</td>
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</tr>
<tr>
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</tr>
<tr>
<td>(F) Average Wall Thickness</td>
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<tr>
<td>(G) Cell Thickness</td>
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<tr>
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<tr>
<td></td>
<td>Y x IN</td>
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<td>0.69</td>
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Figure 12 – (A) Lumen area, (B) cell wall thickness, (C) cell thickness, and (D) cell number in early and latewood for black spruce and balsam fir. Before represents the last year before the beginning of defoliation and the shaded background, the first four years of defoliation (2013-2016). Significant differences between the year before and a given year after defoliation are represented with asterisks for balsam fir and with “x” for black spruce.
**DISCUSSION**

### 2.4.1 GROWTH RESPONSE

Results confirmed that growth losses are a function of the duration of the defoliation for black spruce and balsam fir. Spruce budworm defoliation affects annual ring width, to the point that volume increment of affected trees is also impacted. In our study, balsam fir presented on average 31% of the loss of growth in volume after a four-year defoliation period and black spruce 34%. Several studies have already shown a reduced growth response following sustained defoliation over a few years by the spruce budworm (Archambault 1983; Krause et Morin 1995a; Krause et al. 2012) or with other types of insect pests outbreaks in North America (Alfaro et al. 1985; MacLean 1985; Hennigar et al. 2008).

Generally, significant volume losses were observed for balsam fir and black spruce following the first year of defoliation (Figure 10). This immediate growth response to defoliation of current year’s foliage was also observed for Piene (1980) and Pothier et al. (2005) in balsam fir stands defoliated by spruce budworm. However, a lag of 1 to 3 years between the beginning of defoliation and the appearance of the loss of volume growth has been also described for balsam fir and black spruce stands (Archambault 1983; Krause et
al. 2003; Krause et al. 2012) and other species defoliated by insects outbreaks (Stark et Cook 1957; Axelson et al. 2014). These differences in the synchronous of defoliation and growth loss probably reflect an inherent imprecision of the defoliation map surveys used to detect the onset of defoliation (Krause et al. 2003; Pothier 2012). Differences of tree size and defoliation intensity have been also described as major factors, which could influence the growth and lag response of plants to defoliation (MacLean 1980; Alfaro et al. 1985; Rossi et al. 2009).

Two opposite observations of growth reduction caused by spruce budworm have been also described: one stating that growth loss occurs throughout the entire stem in the same year (Blais 1958; Piene 1980) and the other reported a growth loss starts in the upper crown and a delay at the base of the stem (Piene et al. 1989; Krause et Morin 1995b). In New Brunswick and more southern areas with longer growing seasons, where the defoliation occurs over a longer period, Krause et al. (2003) observed in balsam fir stands that the delay between defoliation and growth loss was variable depending upon stand age. In immature trees, ring reduction occurred simultaneously along the entire stem, whereas in mature trees, the growth reduction occurred one to three years earlier in the crown than at the stem base. The tree-ring widths in our study exhibited a synchronous decrease along the whole stem for both species, independently of stand age.

Further, it is interesting to note that the volume loss was more important for black spruce than for balsam fir in this study. In the past spruce budworm outbreaks, many studies showed a decreased growth after defoliation for both species, however balsam fir was the most vulnerable specie (Blais 1958; MacLean 1984; MacLean et MacKinnon 1997). These results may suggest that, in the current epidemic period, black spruce could be as impacted as balsam fir was in the last outbreak.
2.4.2 WOOD PROPERTIES OF BALSAM FIR X BLACK SPRUCE BEFORE
DEFOLIATION

Balsam fir is known for the low quality of its wood, especially in terms of mechanical
properties (Koga et Zhang 2002). This was confirmed in our study, with an average ring
density 26 % lower than black spruce for the period before defoliation. However, latewood
proportion is similar in both species. Given this result, the difference in the wood density is
likely due to anatomical differences. The tracheids of xylem of both species are the same
width on average, the cell wall thickness is almost similar in earlywood and latewood.
However, lumen area is 32 % higher in the balsam fir than black spruce in earlywood and
19 % in latewood. Consequently, balsam fir has a smaller cell wall-lumen ratio, which could
explain the lower density of balsam fir in relation to black spruce. These anatomical
differences between the two species may be explained for the better capacity for
physiological adjustments of balsam fir (such as higher light-use efficiency, longer seasonal
period of active photosynthesis, lower respiration rate) (Messier et al. 1999; Pothier et
Prévost 2002; Pothier 2012). Further, black spruce is known to be a slow-growing species,
which requires a smaller lumen area for supporting water transport needs than balsam fir
(Krause et al. 2010).

2.4.3 WOOD PROPERTIES OF BALSAM FIR X BLACK SPRUCE AFTER
DEFOLIATION

For balsam fir, very little changes were observed in wood density data during years
with growth loss. However, latewood density decreased after three and four years of
consecutive defoliation. These results might be explained by the anatomical characteristics.
A significant inter-annual decreased trend after three years of defoliation in latewood
tracheids was detected for cell wall thickness when compared to a non-outbreak year, which
seems to result in a decreasing latewood density. Lumen area has been also impacted by defoliation in balsam fir trees in the third and four years of defoliation, however, these changes are less important than those observed in the cell wall.

For black spruce, a similar decreasing trend to balsam fir was observed in latewood density. However, latewood anatomical features in black spruce have shown to be more sensitive to foliage loss than balsam fir during the wood formation. Significant decreases in latewood cell wall thickness were observed from the first year of defoliation. As balsam fir is known to adapt more rapidly to new environmental conditions than black spruce (Krause et al. 2010), which may suggest that the two species can differ in physiological and anatomical characteristics.

In this study, both species presented changes in earlywood cell wall thickness only after three years of consecutive defoliation. Since the formation of the earlywood tracheids is mainly dependent on the reserves (Schweingruber 1979; Krause et Morin 1995a), we can hypothesize that the cumulative defoliation stress culminates in the depletion of reserves, and now also influencing negatively the earlywood cell formation. In conifer species, although a negative relationship generally exists between wood density and growth rate, there were many exceptions, from non-significant or even a weak positive relationship (Zhang et al. 1996; Koga et Zhang 2002; Vincent et al. 2011). Furthermore, a severe defoliation tends do decrease firstly the density of latwood, and only later on earlywood density (Schweingruber 1979). Vejpustkova et Holusa (2006) and Axelson et al. (2014) also studied different insect-tree interactions noted similar results, a reduction in tree growth, followed by a reduction in latewood anatomical features, while the cellular characteristics of the earlywood remained fairly constant.
2.4.4 REDUCTIONS IN WOOD GROWTH X WOOD QUALITY

Duration and intensity of wood formation is influenced by various exogenous and endogenous factors (Savidge 1996). Several studies clearly showed that less vigorous trees had a reduced cambium activity, resulting in reduced lasting of wood formation (Bauch et al. 1986; Schmitt et al. 2003; Deslauriers et al. 2015). We suggest that the changes identified in the wood quality in this study are related to a reduced cambium activity, and mainly expressed to the formation of latewood cells.

In balsam fir and black spruce, cambium reactivation and, consequently, the onset of cell differentiation start normally from early April to early June in the study zone (Deslauriers et al. 2003; Lupi et al. 2010). The larvae of spruce budworm emerge from over-wintering at the end of May and feed on the new shoots throughout June (Carisey et Bauce 1997; Rossi et al. 2009), firstly on balsam fir, its main host, and posteriorly black spruce (Morin et al. 2007; Hennigar et al. 2008). By this time, for both species, earlywood formation is ongoing and latewood differentiation is starting, thus latewood formation, including cell wall thickening, follows when the trees have undergone the maximum foliage loss. Evergreen trees store a higher proportion of C in the leaves rather in the wood, as occur in broadleaf trees (Fajardo et al. 2013). In consequence of repeated and heavy defoliation, the remaining leaves provide energy only to maintain metabolism and growth of the subsequent leaves (Li et al. 2002). Despite the continuous requirement of energy to accomplish radial growth, the trees have lower carbon sources, so reducing amounts of photosynthates available for tracheids formation and the production of reserves (Reich et al. 1993; Deslauriers et al. 2015).

Trees have to manage challenging optimization strategies during growth to keep the ideal proportion of latewood to ensure mechanical stability, as well as the production of
earlywood to maintain the water transport efficiency (Gartner 1995; Domec et Gartner 2002; Eder et al. 2009; West 2014). As the tree is in a saving carbon mode by producing fewer tracheids (Deslauriers et al. 2015), the fact that probably the defoliated trees have already had a sufficient number of tree-rings to support the mechanical stability makes latewood less essential. In contrast, the cells produced during the beginning of growing period belong to earlywood, which performed an essential and efficient hydraulic role in the tree (Domec et Gartner 2002).

In the wood formation of conifers, latewood wall thickening, and lignification are an expensive and long process for completing maturation, being the duration of the deposition of material within the cell wall (cellulose and lignin) the factor that causes the ticker cell walls (Deslauriers et al. 2003; Lupi et al. 2010). However, in defoliated trees, the cambium activity is shortened (Bauch et al. 1986; Schmitt et al. 2003; Deslauriers et al. 2015), reducing the cell wall thickness (Vejpustkova et Holusa 2006; Axelson et al. 2014).

2.5 CONCLUSIONS

In this study we focused attention on the wood quality impact of a single spruce budworm outbreak in the balsam fir and black spruce. We demonstrated that this outbreak resulted in statistically significant annual reductions in volume for both species, more important in black spruce than in balsam fir. Reductions in wood density was observed only for black spruce after four years of defoliation, whereas latewood density changed in the second and third year for black spruce, and third and fourth years for balsam fir. These changes were correlated with changes in the anatomical features. Significant reductions in cell wall thickness were observed from the first year in black spruce and third year for the balsam fir. Future research should examine the wood quality of multiple outbreaks in samples collected at different heights on the tree bole. This approach would provide
additional insight into the cumulative effects of outbreak disturbances on wood structure during the epidemic period and after in the recuperation of the tree stress. Further, mechanical tests could provide insights into the response of these changes in structural properties of the wood.
CONCLUSIONS ET IMPLICATIONS

Actuellement, la perte de bois de l’épinette noire et du sapin baumier due à l'infestation de la tordeuse des bourgeons de l’épinette est considérable. Dans le but de réduire les pertes, le Ministère des Forêts, de la Faune et des Parcs établit une récolte de récupération des peuplements fortement défoliés et morts. Ce bois est transformé dans des usines de sciage et de pâte et papier. Toutefois, les industries ont observé des pertes plus importantes de la biomasse récoltée et une diminution de la qualité du bois qui nuit à la rentabilité de leurs opérations. Cependant, très peu d'information est disponible sur l'altération de la qualité du bois des arbres défoliés par la TBE avec la progression de la durée et de l'intensité de défoliation.

Dans le chapitre I, sur la physiologie des arbres défoliés, les résultats ont montré que la défoliation n’a pas eu d’effet sur le potentiel hydrique et l’état hydrique. De plus, aucun changement n’a été observé au niveau du taux d’humidité et de la proportion d’aubier entre les arbres défoliés et non défoliés pour les deux espèces. Cependant la défoliation a réduit la croissance de façon significative et la perte de volume s’intensifie avec la durée de défoliation (chapitre 2). Les analyses de densitométrie ont montré que les arbres défoliés ont une densité du bois final inférieure aux arbres sains. La densité moyenne du bois n’a présenté des réductions que pour l’épinette noire à la quatrième année de défoliation. Les analyses anatomiques ont démontré des réductions importantes au niveau de l’épaisseur de la paroi cellulaire dans le bois initial et final chez les deux espèces. Néanmoins, la surface du lumen n’a pas démontré des changements si importants au long de la période de défoliation. Ainsi, il est possible de suggérer que les changements observés au niveau de la densité du bois final sont dus aux altérations au niveau de l’épaisseur de la paroi cellulaire chez les deux espèces.
De cette façon, nos hypothèses de départ ont été en partie confirmées, les tiges étaient affectées au niveau de leur accroissement radial alors que la densité du bois a été impactée plus au niveau de la densité du bois final pour les deux espèces, résultat des changements au niveau de l’anatomie du bois. Toutefois, les hypothèses que la défoliation allait réduire la demande en eau, et par conséquent, entraînerait une réduction de la proportion d’aubier et du taux d’humidité dans la tige, ont été rejetées.

L’étude a permis de décrire l’évolution temporelle de la présente épidémie de la TBE sur certains paramètres de qualité du bois chez l’épinette noire et le sapin baumier dans la région du Saguenay Lac-Saint-Jean. Lorsqu’on intègre les informations relatives à la perte de croissance à celles sur la perte de densité du bois, on peut obtenir une représentation du temps dont on dispose pour procéder à la récolte des peuplements touchés par TBE afin de garder la qualité du bois récolté proche de celle des arbres sains. Chez l’épinette noire, les pertes plus importantes en volume ont été observées à 4 ans de défoliation et pour la densité du bois à 2-3 ans de défoliation. Chez le sapin baumier, les pertes en volume plus importantes ont été observées à 4 ans de défoliation, suivi par des réductions de densité du bois final à 3-4 ans de défoliation. Intégrant la perte en volume et la diminution de la paroi cellulaire dans le bois final, on suggère l’amorce de la récolte des peuplements défoliés de l’épinette noire à partir de deux à trois ans du début de la défoliation. Chez le sapin baumier cette récolte est préférable entre la troisième et quatrième année de défoliation.

Le taux d’humidité est une variable importante pour l’industrie de pâtes à papier. Comme nous n’avons mesuré aucune modification de ce paramètre avec la durée ou l’intensité de la défoliation, nous conseillons une récolte avant la mort de l’arbre. Cependant, le temps dont on dispose pour récolter les arbres affectés proposé par cette
étude peut varier selon les procédés de transformation et les exigences du marché. Les résultats obtenus lors de cette étude sont une référence utile pour planifier une récolte en cours d’épidémie. Toutefois, pour que son application soit adéquate, des validations doivent être faites sur le terrain et des études similaires en incluant d’autres sites dans la province.
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