

RESEARCH ARTICLE

The French adaptation and validation of the Partners in Health (PIH) scale among patients with chronic conditions seen in primary care

Émilie Hudon^{1,2}, Maud-Christine Chouinard^{2,3}, Cynthia Krieg¹, Mireille Lambert², Heithem Joober¹, Sharon Lawn⁴, David Smith⁴, Sylvie Lambert⁵, Catherine Hudon^{1,6,7*}

1 Faculté de Médecine et des Sciences de la Santé, Université de Sherbrooke, Québec, Canada, **2** Centre intégré universitaire de Santé et de Services sociaux du Saguenay-Lac-Saint-Jean, Chicoutimi, Québec, Canada, **3** Département des sciences de la santé, Université du Québec à Chicoutimi, Chicoutimi, Québec, Canada, **4** Flinders Human Behaviour & Health Research Unit (FHBHRU), Flinders University College of Medicine and Public Health, Bedford Park, South Australia, Adelaide, Australia, **5** École des sciences infirmières, Université McGill, Montréal, Québec, Canada, **6** Département de médecine de famille et de médecine d'urgence, Université de Sherbrooke, Sherbrooke, Québec, Canada, **7** Centre de recherche du Centre hospitalier de l'Université de Sherbrooke, Québec, Canada

* Catherine.Hudon@Usherbrooke.ca



OPEN ACCESS

Citation: Hudon É, Chouinard M-C, Krieg C, Lambert M, Joober H, Lawn S, et al. (2019) The French adaptation and validation of the Partners in Health (PIH) scale among patients with chronic conditions seen in primary care. PLoS ONE 14(10): e0224191. <https://doi.org/10.1371/journal.pone.0224191>

Editor: Gregor Stiglic, University of Maribor, SLOVENIA

Received: April 26, 2019

Accepted: October 8, 2019

Published: October 23, 2019

Copyright: © 2019 Hudon et al. This is an open access article distributed under the terms of the [Creative Commons Attribution License](https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Data Availability Statement: All relevant data are within the paper and its Supporting Information files.

Funding: This work was supported by Faculté de médecine et des sciences de la santé de l'Université de Sherbrooke [2017-2018] and Fondation de ma vie, Centre intégré universitaire de santé et de services sociaux du Saguenay-Lac-Saint-Jean [2016-2017] to ÉH. MCC was supported by Réseau de recherche en

Abstract

Objective

Measuring self-management helps identify the degree of participation of people in the management of their chronic conditions and guides clinicians in determining person-centred priorities for providing support. The *Partners in Health* scale, a self-report generic questionnaire, was developed to capture the self-management of patients with chronic conditions. This study aimed to translate the *Partners in Health* scale into French and to examine its psychometric properties in French-speaking people with chronic conditions followed in primary care.

Methods

The *Partners in Health* scale was translated into French using Hawkins and Osborne's method (2012). Content validity was evaluated through cognitive interviews (Think Aloud Method). Internal consistency was measured at baseline with Cronbach's alpha. Test-retest reliability was evaluated at baseline and two weeks later using the intraclass correlation coefficient. Concurrent validity was measured at baseline with the Self-efficacy for Managing Chronic Disease (SEM-CD) and the Patient Activation Measure (PAM), using Spearman correlations.

Results

Cognitive interviews were conducted with 10 participants. During these interviews, most items were clearly understood and accepted as formulated; only a few terms were modified. To evaluate the psychometric properties of the French-language version of the Partners in Health scale, 168 participants (male = 34.5%; mean age = 58 years; mean number of

interventions en sciences infirmières du Québec [2016–2018]. The funding agencies were not involved in the design of the study nor the collection, analysis, and interpretation of data and writing the manuscript.

Competing interests: The authors have declared that no competing interests exist.

chronic conditions = 4.1) completed the questionnaire at baseline and 47 of them completed the questionnaire two weeks later by telephone. Cronbach's alpha for internal consistency was 0.85 (95% confidence interval: 0.81–0.88). The intraclass correlation coefficient for test-retest reliability was 0.77 (95% confidence interval: 0.58–0.87). Concurrent validity with spearman's coefficient correlation of Self-efficacy for Managing Chronic Disease and Patient Activation Measure was 0.68 and 0.61 respectively.

Conclusion

The French-language version of the *Partners in Health* scale is a reliable and valid questionnaire for the measure of self-management in persons with chronic conditions seen in primary care.

Introduction

Chronic conditions are the leading cause of mortality and are responsible for 70% of mortality rates worldwide; representing 40 million deaths per year [1]. Self-management is a complex process in which the person actively participates in the management of his or her chronic conditions. It leads to important benefits both at the individual level, such as improvement in self-control, well-being and quality of life, and at the societal level, such as a decrease in use and costs of health services [2]. Self-management involves medical/behavioral, decision-making, emotional and cognitive strategies [3, 4].

Clinicians need tools to measure self-management in order to assess the participation of the person with chronic conditions in the management of his or her health, to adjust their self-management support interventions to the needs, preferences and priorities of the person in order to maximise engagement and motivation for change, and to evaluate the impact of these interventions [5]. The Partner in Health (PIH) scale [5], developed by researchers in Australia, approximately 20 years ago, offers primary healthcare providers collaborating with people with chronic conditions a generic tool to measure the self-management of this client population. The fourth and last version of the questionnaire was published in 2016 by Smith, Harvey, Lawn, Harris and Battersby [5–8]. This version of the PIH comprises 12 items, with responses rated using a 9-point Likert-type scale. It presents a variety of interesting characteristics: 1) it targets adults with at least one chronic condition; 2) it is completed by self-report; 3) it is short (<25 items); 4) items are short (<20 words/item); 5) the psychometric properties are good; and 6) it measures all strategies of the self-management concept (behavioral/medical, cognitive/decision-making, emotional and social strategies) [3, 4, 9, 10]. Previous factor analysis identified four factors: 1) knowledge; 2) partnership in treatment; 3) recognition and management of symptoms; and 4) coping [7]. The PIH has been translated into Spanish [11], Dutch [12], and Chinese [13] and adapted for populations with diabetes [14], chronic renal disease [15, 16], mental illness [17], and for older adults with hearing loss [18]. To date, no French-language version of the PIH is available.

Aim

The main purpose of this study was to translate the PIH into French version (PIH-Fv) and to evaluate its psychometric properties (internal consistency, test-retest reliability and concurrent

validity) among a French-speaking population with at least one chronic condition and followed in primary care.

Materials and methods

French-language cross-cultural adaptation

The cross-cultural adaptation was conducted following the Hawkins and Osborne method (2012) [19, 20], involved the authors of the PIH, and included the following steps:

1. *Translation*: A professional native French-speaking translator translated the original version into a French version.
2. *Back-translation*: A professional native English-speaking translator back-translated the French version into English without seeing the original English version of the PIH.
3. *Committee evaluation*: An expert panel composed of researchers specialized in primary care, one developer of the PIH (SL), two translators, healthcare providers (nurses and family physicians) and one bilingual patient met to compare the original version with the back-translated version. The purpose was to clarify any inconsistencies between the two English versions and to come to an agreement in the labeling of items in French, while preserving the same meaning as the original version.
4. *Pretest*: Cognitive interviews were conducted by the first author (ÉH) with persons with various chronic conditions until data saturation was reached ($n = 10$). The recruitment process was the same as the one used for the validation part of this study (see next section). Interviews based on the Think Aloud Method lasted from 30 to 45 minutes [21, 22]. This consisted of having the participant read the questionnaire out loud and answering without the help of the research assistant. Any issues raised by the participants were then reviewed by the team to modify any term that lacked clarity or was confusing.
5. *Validation of the French-language version of the questionnaire*: details are provided in the methods section below.

Settings, participants and recruitment procedures

For the evaluation of the psychometric properties of the PIH-fv, recruitment was carried out in two Family Medicine Groups from two cities located in two different regions of Québec, Canada (Saguenay-Lac-Saint-Jean and Estrie), using a convenient sampling procedure. These regions were selected to represent rural and urban areas. The inclusion criteria were: 1) being a patient in the participating clinic; 2) 18 years and older; 3) native French-language speaker; and 4) suffering from at least one chronic condition (regardless of type, time since diagnosis, or treatment). Pregnant women and patients with an acute exacerbation of their chronic condition were excluded, because these states require a more frequent follow-up that may interfere with self-management and study outcomes [23].

Recruitment was conducted May 9–26, 2016, while the patients were waiting for their appointment with a primary healthcare provider. The clinic's receptionist provided a description of the project with a list of inclusion criteria to each patient. Two authors (EH and CK) approached patients to explain the project and assess their eligibility for participation against the inclusion criteria. Eligible persons provided consent and completed the questionnaire (T1). Two weeks later, some participants completed the questionnaire (T2) once again by telephone, without the sociodemographic and Disease Burden Morbidity Assessment (DBMA) sections. This timeframe is considered adequate to assess test-retest reliability [24]. The

research protocol was approved by the ethics review board of the Centre intégré universitaire de la santé et des services sociaux. Informed consent was obtained from all individual participants in the study.

The required sample size was 120 participants, based on Polit recommendations [25]. To evaluate test-retest reliability by telephone at T2, 50 participants was deemed sufficient [26].

Measures

The questionnaire was divided into five sections. It included a sociodemographic section (gender, date of birth, place of birth, first language, education, occupation, income and marital status), the French-language versions of the DBMA [27], the PIH, the Patient Activation Measure (PAM) [28] and the Self-efficacy for Managing Chronic Disease (SEM-CD) [29]. Because self-management is a concept associated to activation and self-efficacy as Richard and Shea [30] stated in their concept analysis, the PAM and the SEM-CD were judged to be appropriate to measure their concurrent validity with the PIH.

Partners in Health (PIH) scale. The PIH scale [7] is a self-report questionnaire that includes 12 items, which are answered using 9-point Likert-type scales (Items 1 and 2: 0 “very poor” to 8 “a lot”; Items 3 to 8: 0 “never” to 8 “always”; Items 9 to 12: 0 “not very well” to 8 “very well”). The total score ranges from 0 to 96, 0 representing poor self-management and 96 representing a greater self-management. Factor analysis identified four related self-management constructs, but in this study the scale was treated as unidimensional because dimensions contained less than three items and the questionnaire is considered as a whole [31] (S1 Appendix).

French-language version of the Patient Activation Measure (PAM). We explored the concurrent validity of the PIH with the PAM because activation levels are significantly associated with self-management behaviors [32]. The PAM [32] is a self-report questionnaire that measures the degree of activation based on a global score transformed into a one-dimensional scale ranging from 0 to 100 (a higher score indicating higher activation). The questionnaire contains 13 items with a 4-point Likert scale (disagree strongly, disagree, agree or agree strongly). The PAM was adapted for use in French and showed good psychometric properties [28].

French-language version of the Self-Efficacy for Managing Chronic Disease (SEM-CD). We also explored the concurrent validity of the PIH with the SEM-CD which measures self-efficacy, a concept also related to self-management as cited by Wilde and Garvin [33]. The SEM-CD is a self-report questionnaire measuring a person’s perception of their self-efficacy [34]. The questionnaire contains 6 items and uses an 11-point Likert scale (0–10), ranging from “not at all confident” to “totally confident”. The final score is a mean ranging from 0 to 10 of all items (a higher score indicating higher self-efficacy). The French version of the SEM-CD demonstrated good psychometric properties [29].

French-language version of the Disease Burden Morbidity Assessment (DBMA). The Disease Burden Morbidity Assessment (DBMA) [35] was used to identify the number of chronic conditions, out of a total of 21. The DBMA was translated into French and demonstrated good psychometric properties [27].

Analysis

Data were analyzed using the Statistical Package for the Social Sciences (SPSS) version 23.0 [36]. Participant characteristics, such as age and number of chronic conditions, were described using means and standard deviations (SD) for continuous variables and frequencies (%) for categorical data as presented in Table 1. Internal consistency was measured using Cronbach’s

alpha (α). A Cronbach alpha between 0.70 and 0.90 indicates good internal consistency; a value below 0.70 indicates poor internal consistency, whereas above 0.90 indicates redundancy among item [37]. Test-retest reliability was measured with an interclass correlation coefficient (ICC). An ICC above 0.75 indicates good reliability, whereas an ICC below 0.75 indicates a low to moderate reliability [38]. Concurrent validity was measured with Spearman's correlations, as data were not normally distributed.

Results

Translation, back-translation and committee evaluation

A French version of the PIH was obtained after comparing the original English version, the translated French version and the back-translated version. Modifications were minor, for example, the term "worse" became "worsen".

Pretest

Cognitive interviews were conducted with 10 persons with chronic conditions. Some terms were adapted to the French language. For example, the item "I am able to deal with health professionals to get the services I need that fit with my culture, values and beliefs" was modified to "I am able to compose with health professionals to get services I need that fit with my culture, values and beliefs". Participants were not sure they understood the meaning of the term "deal", so they suggested "compose". Following the first four steps of the French translation of the PIH a French version was approved by the authors of the PIH (S2 Appendix).

Validation of the French-language version of the questionnaire

Sample characteristics. For the study, 719 patients with at least one chronic condition were approached at T1 (Fig 1). Most refusals to participate ($n = 164$) were due to lack of time before the medical appointment. Among the 232 patients who completed the questionnaire, 46 were incomplete (less than half of the questionnaire was completed), 8 were not valid (two answers to the same question or completed by another person) and 10 questionnaires presented 16 missing data (PIH section), particularly for the item one. These questionnaires ($n = 64$) were excluded from the analysis. Finally, 168 participants (23.4%) completed the questionnaire at T1. At T2, 47 participants completed the questionnaire after 82 of the 168 participants of T1 were contacted (Table 1). Those samples were used for the analysis.

Participants were diagnosed with a mean average of four chronic conditions. Almost half of participants were retired (48.2%), had an annual income of \$50,000 CAD (47.2%) and were married or lived with a partner (54.5%).

PIH analysis. Mean score at T1 was 81.1 (40.0–96.0). Cronbach's alpha was 0.85 (95% CI: 0.81–0.88). The intraclass correlation coefficient between T1 and T2 was 0.77 (CI at 95%: 0.58–0.87). Spearman's correlation was 0.61 with the PAM, and 0.68 with the SEM-CD.

Discussion

The results of this study demonstrated good psychometric properties of the PIH-Fv, which are comparable to the original version and to other cross-cultural adaptations (Dutch and Chinese). To our knowledge, this is the first study of the translation and adaptation of the PIH into French. We used Hawkins and Osborne's method. The authors of the cross-cultural adaptation of the Chinese and Dutch versions used the Brislin (1970) [39] and the Guillemin (1993) [40] methods [12, 13] respectively. These two translation methods require an evaluation

Table 1. Sociodemographic characteristics of participants (n = 168).

Age: \bar{x} (SD)	58 (15.8)
Number of chronic conditions: \bar{x} (SD)	4.1 (2.3)
Region: n (%)	
Rural	49 (29.2)
Urban	110 (70.8)
Male: n (%)	58 (34.5)
Chronic condition: n (%)	
High blood pressure (HBP)	69 (42.6)
Overweight	68 (41.5)
Osteoarthritis	59 (36.2)
Depression or anxiety	59 (36.2)
Education completed: n (%)	
Less than high school	39 (24.1)
Completed high school	47 (29.0)
College or post-secondary school	41 (25.3)
University	35 (21.6)
Occupation: n (%)	
Employed	57 (34.8)
Unemployed	25 (15.2)
Retired	79 (48.2)
Annual family income: n (%)	
Less than \$ 20 000 CAD	22 (13.5)
\$ 20 000 to \$ 49 999 CAD	64 (39.3)
\$ 50 000 CAD or more	77 (47.2)
Marital status: n (%)	
Married, living with a partner	91 (54.5)
Separated, divorced	30 (18.0)
Widowed	22 (13.2)

n: sample size, SD: standard deviation, \bar{x} : mean, %: percentage of the population

<https://doi.org/10.1371/journal.pone.0224191.t001>

committee for the translation. However, they did not mention if patients or authors of the original questionnaires participated [39, 40].

This study is also the first evaluation of the psychometric properties of the PIH-Fv. The average score for T1 (82.17) tended towards good self-management, similarly to the Dutch adaptation (\bar{x} = 78.1) [12]. Internal consistency of previous initial English versions [5, 6] and the Dutch adaptation [12] of the PIH yielded Cronbach's alphas varying from 0.82 to 0.86. The internal consistency in this study remained similar after the translation process (α = 0.85). Test-retest reliability was also documented in the Chinese adaptation [13] with an ICC of 0.82, but the confidence interval was not mentioned. The coefficient (ICC = 0.77) in our study is slightly lower than the Chinese adaptation. In our study the ICC could have been affected by a medical visit between T1 and T2 [38] which might have provided some self-management support explaining an increase in total score (81.1 at T1 versus 84.7 at T2). This result may also be linked to the change in the administration procedure of the PIH-Fv at T1 (in person) and T2 (by telephone) [41].

This is the first study to measure the concurrent validity of the PIH with the SEM-CD and the PAM. The concurrent validity of the PIH-Fv measuring self-management showed a

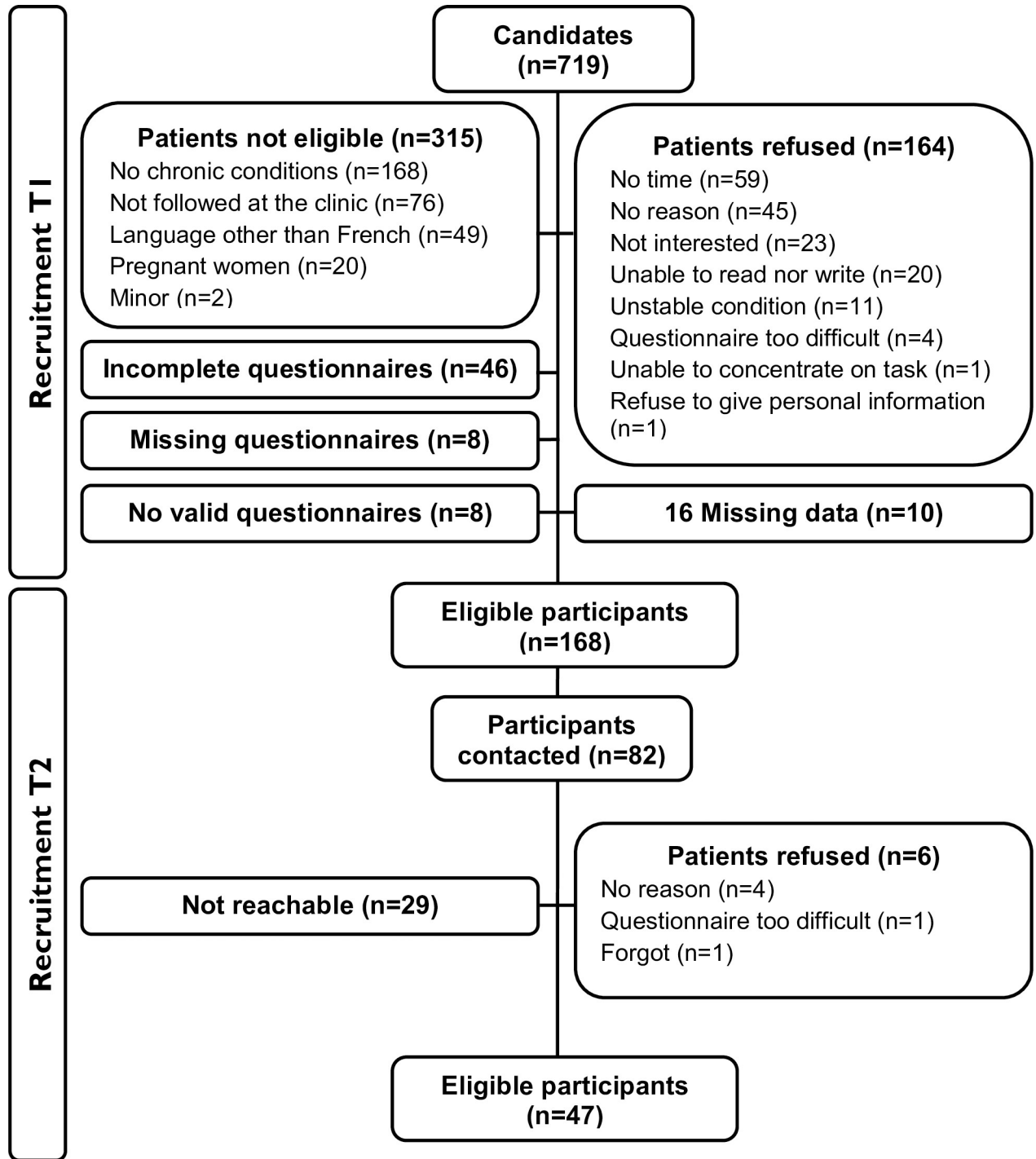


Fig 1. Participation in the study at T1 and T2. Definition of terms: "not eligible" represents people who did not meet selection criteria; "incomplete questionnaires" corresponds to questionnaires with an incomplete section or a section with more than half of the section missing responses; "missing questionnaires" represents the questionnaires lost during recruitment; "invalid questionnaire" represents a questionnaire completed by another person or a questionnaire with two answers to the same question; "missing data" corresponds to a questionnaire without answers at some questions; and "unreachable" represents people whose telephone was out of service when called or the people who did not answer.

<https://doi.org/10.1371/journal.pone.0224191.g001>

moderate correlation with the concept of activation as measured by the PAM and with the concept of self-efficacy as measured by SEM-CD. This is explained by the fact that these concepts are related [9].

Strengths and limitations

Strengths. The study was conducted in two different regions of Quebec, which covers a rural population ($n = 49$) as well as an urban one ($n = 119$), making the sample more representative of the population.

The study was performed rigorously, using Hawkins and Osborne's method, which allowed us to conduct the cross-cultural French adaptation of the questionnaire while preserving the original meaning of the questions by consulting the authors of the PIH. Finally, all information collected and entered into the database was validated by a second person, independently, ensuring accuracy.

Limitations. Participants had to complete the questionnaire while sitting in the clinics' waiting rooms. If the patient's appointment with the clinician was ahead of time and the participants did not have enough time to complete the questionnaire, it was rarely completed after that appointment. This may explain the high number of incomplete questionnaires ($n = 45$).

To further appraise PIH-Fv psychometric properties, a next study should recruit a larger sample to evaluate PIH-Fv structural validity. In addition, it may be appropriate to conduct a sensitivity to change analysis in a clinical intervention study to assess the capacity of the PIH-Fv to capture clinical improvement of self-management level [7]. Finally, missing data were particularly associated with item one of the PIH-Fv. This could be explained by the configuration of the questionnaire; the first question was presented directly below examples of answer options. To reduce the number of missing data, it would be preferable to separate examples of answers options more clearly at the beginning to the PIH-FV. Finally, we suggest conducting a pre-test to evaluate cultural differences before using the French version with other French populations.

Conclusions

The PIH-Fv demonstrated good internal consistency, good test-retest reliability and moderate concurrent validity with the PAM and the SEM-CD. The PIH-Fv can be used to measure level of self-management among French-speaking adults with chronic conditions.

Supporting information

S1 Appendix. Factors and items of the PIH questionnaire. PIH with a 9-point Likert scale; 12 questions assessing four factors: Knowledge (items 1–2), partnership in treatment (items 3–6), recognition and management of symptoms (items 7–8) and coping (items 9–12). Licence tool: +61 8 8404 2607. For more information's contact original authors at: ccm@flinders.edu.au. (DOCX)

S2 Appendix. Factors and items of the PIH-Fv questionnaire. French-language version of the PIH with a 9-point Likert scale; 12 questions assessing four factors: Knowledge (items 1–2), partnership in treatment (items 3–6), recognition and management of symptoms (items 7–8) and coping (items 9–12). (DOCX)

Acknowledgments

The authors would like to thank the Australian developers for permission to use the PIH for this research. We would also like to thank Ms. Catherine Brown for the translation, the bilingual participant Mrs. Maureen Morris, the two participating GMF-U and the participants to the project. Finally, we would like to thank Susie Bernier for her help in editing this paper.

Author Contributions

Conceptualization: Émilie Hudon, Maud-Christine Chouinard, Catherine Hudon.

Data curation: Catherine Hudon.

Formal analysis: Émilie Hudon, Maud-Christine Chouinard, Mireille Lambert, Catherine Hudon.

Funding acquisition: Émilie Hudon, Maud-Christine Chouinard.

Investigation: Émilie Hudon, Maud-Christine Chouinard, Sharon Lawn, Catherine Hudon.

Methodology: Émilie Hudon, Catherine Hudon.

Project administration: Émilie Hudon, Maud-Christine Chouinard, Catherine Hudon.

Resources: Cynthia Krieg, Mireille Lambert, Heithem Joobar, Sharon Lawn, David Smith, Catherine Hudon.

Supervision: Émilie Hudon, Maud-Christine Chouinard, Catherine Hudon.

Validation: Émilie Hudon, Cynthia Krieg, Mireille Lambert, Heithem Joobar, Sharon Lawn.

Visualization: Maud-Christine Chouinard.

Writing – original draft: Émilie Hudon, Maud-Christine Chouinard, Catherine Hudon.

Writing – review & editing: Émilie Hudon, Maud-Christine Chouinard, Cynthia Krieg, Mireille Lambert, Heithem Joobar, Sharon Lawn, David Smith, Sylvie Lambert, Catherine Hudon.

References

1. World Health Organization. Noncommunicable diseases Switzerland: World Health Organization; 2017 [Available from: <http://www.who.int/mediacentre/factsheets/fs355/en/>].
2. Blok AC. A middle-range explanatory theory of self-management behavior for collaborative research and practice. *Nursing Forum*. 2017; 52(2):138–46. <https://doi.org/10.1111/nuf.12169> PMID: 27438773
3. Barlow J, Wright C, Sheasby J, Turner A, Hainsworth J. Self-management approaches for people with chronic conditions: A review. *Patient Education and Counseling*. 2002; 48(2):177–87. [https://doi.org/10.1016/s0738-3991\(02\)00032-0](https://doi.org/10.1016/s0738-3991(02)00032-0) PMID: 12401421
4. Schulman-Green D, Jaser S, Martin F, Alonzo A, Grey M, McCorkle R, et al. Processes of self-management in chronic illness. *Journal of Nursing Scholarship*. 2012; 44(2):136–44. <https://doi.org/10.1111/j.1547-5069.2012.01444.x> PMID: 22551013
5. Battersby MW, Ask A, Reece MM, Markwick MJ, Collins JP. The Partners in Health scale: The development and psychometric properties of a generic assessment scale for chronic condition self-management. *Australian Journal of Primary Health*. 2003; 9(3):41–52.
6. Petkov J, Harvey P, Battersby M. The internal consistency and construct validity of the partners in health scale: Validation of a patient rated chronic condition self-management measure. *Quality of Life Research*. 2010; 19(7):1079–85. <https://doi.org/10.1007/s11136-010-9661-1> PMID: 20437206
7. Smith D, Harvey P, Lawn S, Harris M, Battersby M. Measuring chronic condition self-management in an Australian community: Factor structure of the revised Partners in Health (PIH) scale. *Quality of Life Research*. 2016; 26(1):149–59. <https://doi.org/10.1007/s11136-016-1368-5> PMID: 27432251

8. Battersby M, Harris M, Smith D, Reed R, Woodman R. A pragmatic randomized controlled trial of the Flinders Program of chronic condition management in community health care services. *Patient Education and Counseling*. 2015; 98(11):1367–75. <https://doi.org/10.1016/j.pec.2015.06.003> PMID: 26146240
9. Miller WR, Lasiter S, Ellis RB, Buelow JM. Chronic disease self-management: A hybrid concept analysis. *Nursing Outlook*. 2015; 63(2):154–61. <https://doi.org/10.1016/j.outlook.2014.07.005> PMID: 25241136
10. Unger WR, Buelow JM. Hybrid concept analysis of self-management in adults newly diagnosed with epilepsy. *Epilepsy & Behavior*. 2009; 14(1):89–95.
11. Penarrieta-de Cordova I, Barrios FF, Gutierrez-Gomes T, Pinonez-Martinez Mdel S, Quintero-Valle LM, Castaneda-Hidalgo H. Self-management in chronic conditions: Partners in health scale instrument validation. *Nurs Manage (London)*. 2014; 20(10):32–7.
12. Lenferink A, Effing T, Harvey P, Battersby M, Frith P, Van Beurden W, et al. Construct validity of the Dutch version of the 12-Item Partners in health scale: Measuring patient self-management behaviour and knowledge in patients with chronic obstructive pulmonary disease. *PloS One*. 2016; 11(8): e0161595. <https://doi.org/10.1371/journal.pone.0161595> PMID: 27564410
13. Chiu TML, Tam KTW, Siu CF, Chau PWP, Battersby M. Validation study of a Chinese version of Partners in health in Hong Kong (C-PIH HK). *Quality of Life Research*. 2017; 26(1):199–203. <https://doi.org/10.1007/s11136-016-1315-5> PMID: 27216940
14. Battersby MW, Kit JA, Prideaux C, Harvey PW, Collins JP, Mills PD. Implementing the flinders model of self-management support with aboriginal people who have diabetes: Findings from a pilot study. *Australian Journal of Primary Health*. 2008; 14(1):66–74.
15. Baxter C, Morello A, Smith D, Norton L, Bentley D. A preliminary investigation of the Partners in Health scale measurement properties in patients with end stage renal disease. *Australian Journal of Primary Health*. 2017; 23(3):288–93. <https://doi.org/10.1071/PY16095> PMID: 28076747
16. Walker RC, Marshall MR, Polaschek NR. A prospective clinical trial of specialist renal nursing in the primary care setting to prevent progression of chronic kidney: A quality improvement report. *BMC Family Practice*. 2014; 15(1):155.
17. Fotu M, Tafa T. The Popao model: A Pacific recovery and strength concept in mental health. *Pacific Health Dialog*. 2009; 15(1):164–70. PMID: 19585747
18. Convery E, Meyer C, Keidser G, Hickson L. Assessing hearing loss self-management in older adults. *International journal of audiology*. 2018; 57(4):313–20. <https://doi.org/10.1080/14992027.2017.1390268> PMID: 29081257
19. Hawkins M, Osborne RH. Health literacy questionnaire: Translation and cultural adaptation procedure. Australia: Deakin University; 2012.
20. Maindal HT, Kayser L, Norgaard O, Bo A, Elsworth GR, Osborne RH. Cultural adaptation and validation of the Health literacy questionnaire (HLQ): Robust nine-dimension Danish language confirmatory factor model. *SpringerPlus*. 2016; 5(1):1232. <https://doi.org/10.1186/s40064-016-2887-9> PMID: 27536516
21. Ericsson KA, Simon HA. Verbal reports as data. *Psychological review*. 1980; 87(3):215.
22. Hofmeyer A, Sheingold BH, Taylor R. Do you understand what I mean? How cognitive interviewing can strengthen valid, reliable study instruments and dissemination products. *International Journal of Educational Research*. 2015; 11(4):261.
23. Kilpatrick SJ, Papile L-A, Macones GA. Guidelines for Perinatal Care. 8 ed. Washington, SC: American Academy of Pediatrics; 2017.
24. Terwee CB, Bot SD, Boer MRd, Windt DAvd, Knol DL, Dekker J, et al. Quality criteria were proposed for measurement properties of health status questionnaires. *Journal of Clinical Epidemiology*. 2007; 60(1):34–42. <https://doi.org/10.1016/j.jclinepi.2006.03.012> PMID: 17161752
25. Polit DF. Statistics and data analysis for nursing research. 2nd ed. Boston, Canada: Pearson; 2010. xii, 440 p. p.
26. Streiner DL, Norman GR, Cairney J. Health measurement scales: A practical guide to their development and use. 5 ed. Oxford, England: Oxford University Press; 2015.
27. Poitras M-E, Fortin M, Hudon C, Haggerty J, Almirall J. Validation of the disease burden morbidity assessment by self-report in a French-speaking population. *BMC Health Services Research*. 2012; 12:35. <https://doi.org/10.1186/1472-6963-12-35> PMID: 22333434
28. Hudon C, Lambert M, Almirall J, Cook M, editors. The reliability of the french version of the Patient Activation Measure. 40th Annual meeting of North American Primary Care Research Group 2012; New Orleans.

29. Hudon C, Chouinard M-C, Bélanger A, Fortin M, Freund T, Almira J, et al. The Self-efficacy for managing chronic disease scale—French version: A validation study in primary care. *European Journal for Person Centered Healthcare*. 2014; 2(4):533–8.
30. Richard AA, Shea K. Delineation of self-care and associated concepts. *Journal of Nursing Scholarship*. 2011; 43(3):255–64. <https://doi.org/10.1111/j.1547-5069.2011.01404.x> PMID: 21884371
31. Hinkin TR. Scale development principles and practices. *Research in organizations: Foundations and methods of inquiry*. 2005:161–79.
32. Hibbard JH, Stockard J, Mahoney ER, Tusler M. Development of the Patient Activation Measure (PAM): Conceptualizing and measuring activation in patients and consumers. *Health services research*. 2004; 39(4p1):1005–26.
33. Wilde MH, Garvin S. A concept analysis of self-monitoring. *Journal of Advanced Nursing*. 2007; 57(3):339–50. <https://doi.org/10.1111/j.1365-2648.2006.04089.x> PMID: 17233653
34. Lorig KR, Sobel DS, Ritter PL, Laurent D, Hobbs M. Effect of a self-management program on patients with chronic disease. *Effective Clinical Practice*. 2001; 4(6):256–62. PMID: 11769298
35. Bayliss EA, Ellis JL, Steiner JF. Subjective assessments of comorbidity correlate with quality of life health outcomes: Initial validation of a comorbidity assessment instrument. *Health and Quality of Life Outcomes*. 2005; 3(1):51.
36. Statistic IS. IBM SPSS Statistics for Windows, Version 23.0. IBM Corp2015.
37. Terwee CB, Bot SD, de Boer MR, van der Windt DA, Knol DL, Dekker J, et al. Quality criteria were proposed for measurement properties of health status questionnaires. *Journal of Clinical Epidemiology*. 2007; 60(1):34–42. <https://doi.org/10.1016/j.jclinepi.2006.03.012> PMID: 17161752
38. Portney LG, Watkins MP. *Foundations of clinical research: Applications to practice*. 3rd revised ed. Upper Saddle River, NJ: Pearson; 2015. xix, 892 pages p.
39. Brislin RW. Back-translation for cross-cultural research. *Journal of Cross-Cultural Psychology*. 1970; 1(3):185.
40. Guillemin F, Bombardier C, Beaton D. Cross-cultural adaptation of health-related quality of life measures: Literature review and proposed guidelines. *Journal of clinical epidemiology*. 1993; 46(12):1417–32. [https://doi.org/10.1016/0895-4356\(93\)90142-n](https://doi.org/10.1016/0895-4356(93)90142-n) PMID: 8263569
41. Leppink J, Pérez-Fuster P. We need more replication research: A case for test-retest reliability. *Perspectives on Medical Education*. 2017; 6(3):158–64. <https://doi.org/10.1007/s40037-017-0347-z> PMID: 28390030