Prospective application of implementation science theories and frameworks to inform use of PROMs in routine clinical care within an integrated pain network

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Abstract

Purpose The objective of this study is to present the implementation science approaches that were used before implementing electronic patient-reported outcome measures (ePROMs) across an integrated chronic pain network that includes primary, rehabilitation, and hospital-based care.

Methods The Theoretical Domains Framework (TDF) was used to identify potential barriers and enablers to the use of ePROMS by primary care clinicians. In rehabilitation and tertiary care, the Consolidated Framework for Implementation (CFIR) was used to guide the identification of determinants of implementations, through observation of workflow, patient and clinician surveys, and clinician interviews. A mixed-method concurrent design comprising a quantitative and qualitative analysis was used. The results were reviewed by a steering committee to iteratively inform the ePROM implementation plan. The Proctor framework of evaluation was used to guide the development of an evaluation plan for the implementation of ePROMs in the integrated chronic pain network.

Results Both frameworks provided similar results with respect to healthcare provider knowledge, behaviour, and experience interpreting PROM scores. The TDF and CFIR frameworks differed in identifying organizational-level determinants. The resultant implementation plan was structured around the adoption of PROMs to inform individual treatment planning and quality improvement. The evaluation plan focused on implementation and impact outcomes to evaluate the ePROM intervention.

Conclusions The TDF and CFIR guided the development of a multi-component knowledge translation and training intervention that will address multiple gaps and barriers to implementation of PROMs across the integrated network. The ePROM intervention will aim to increase clinicians’ knowledge and skills and foster best practices.

Keywords Implementation science · Theoretical frameworks · Barriers · Enablers · Patient-reported outcome measures · Chronic pain · Integrated care

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Background

Chronic pain (CP) is a complex condition that affects the individual’s daily activities and quality of life, and social and family environments [1]. Management of CP patients is fragmented, with patients seeking a wide variety of primary and specialty care services [2–5] resulting in poorer outcomes and higher healthcare costs as patients often receive unneeded diagnostic and medical procedures [3, 6]. Pain has the potential to impact a range of domains of a person’s health and quality of life including physical, mental, emotional, and cognitive function. Patient-Reported Outcome Measures (PROMs) enable clinicians to incorporate the patient voice in evaluating the impact of their health condition on their function and health-related quality of life (HRQL), [7–9] and in optimizing treatment planning. There is evidence to support that PROMs are necessary to capture outcomes that are meaningful to individuals with chronic pain and to support a patient-centred care approach [10, 11].

To increase their widespread adoption and to optimize their usefulness to a range of stakeholders, PROMs must be perceived as relevant, meaningful, and actionable to the patients and the clinicians using them and the organizations implementing them. Using implementation research can help identify strategies that address the contextual barriers and facilitators to using PROMs in a specific clinical context. There are over 60 frameworks to guide the implementation of healthcare innovations [12]. Using frameworks to guide the application of PROMs in clinical practice is important as they can facilitate identification of determinants of implementation, guide the selection of implementation strategies, and evaluate if implementation was effective. Frameworks can also guide the selection of domains that influence sustainability of the use of PROMs, and their impact on quality of care and health outcomes, and provide a common language to synthesize results across studies [13].

As part of a 5-year strategic plan, the McGill RUISSS Center of Expertise in Chronic Pain (CECP) developed processes to implement and evaluate the impact of PROMs across an integrated care network. Taking a grass roots approach, the CECP leadership asked chronic pain clinics about their interest in implementing PROMs. The clinics that expressed interest worked with the research team, the CECP and patients, to develop their electronic PROM (ePROM) implementation plan. Two complementary implementation science frameworks were selected to guide the process and to identify the determinants of implementation of ePROMs at the level of patients, clinicians, and the organization. The objective of this study was to present the knowledge translation (KT) [14] and implementation science approaches that we used to inform the implementation of ePROMs across an integrated chronic pain network that includes primary, rehabilitation, and hospital-based care. Specifically, we present how we applied two implementation science frameworks to evaluate determinants of implementation, to guide the selection of strategies to implement ePROMs in clinical practice, and to select implementation outcomes.

Application of theoretical frameworks for implementation in the context of the CECP network

To address the gaps in chronic pain management, the CECP developed a strategic plan to provide integrated care for individuals that included early referral to the right service based on individuals’ needs. In line with evidence-based practices, interdisciplinary teams were integrated in primary care clinics in four health regions, with referral to secondary and third lines of care for more complex chronic pain cases.

In order to successfully implement ePROMs to screen for patient service needs, develop individualized intervention plans, and monitor changes, results from two implementation research studies were integrated, one in primary care [15], and a one in tertiary and rehabilitation care [2, 11]. A steering committee reviewed the results of each study during four 1.5-h working meetings and developed and iteratively reviewed the implementation plan. Committee members consisted of a decision maker and clinician from each site, the director of the CECP, two research team members, and two patient partners met during the studies and at the end.

The choice of the theoretical implementation framework used in each study or setting was mainly driven by the specific clinical context, and the understanding of our team at the time about implementation research and its application to ePROMs (Fig. 1). In primary care, the Theoretical Domains Framework (TDF) was selected as it allowed for identifying determinants of implementation of ePROMs more at an individual clinician level and for developing theory-based behavioural change strategies to address barriers and train clinicians in the use and application of PROMs for patient care. In these settings, the main focus was to train individual team members to interpret and apply ePROM scores to treatment planning within a well-defined clinical process endorsed and supported by the healthcare organization. In tertiary and rehabilitation care, where there is more variability in care delivery processes, we needed to consider determinants related more to organizational change in addition to determinants of clinician and patient behaviour change. In these settings we used the Consolidated Framework of Implementation Research (CFIR) to evaluate clinician,
individuals with chronic pain, and organizational factors that influence implementation of ePROMs.

In this paper, we present an overview of the use of each framework (Table 1) and how the use of these frameworks resulted in a final PROM implementation plan. The steering committee integrated results across both studies to inform PROM implementation and evaluation. The Proctor framework was used to guide the development of an evaluation plan for the implementation of PROMs. The details of the methodology and results for each study are presented elsewhere [2, 15, 16]. Here, we focus on the use of the frameworks and the resulting implementation plan to feedback PROM scores to patients and clinicians. Finally, we contrast the use of the TDF and CFIR frameworks, assess whether these frameworks resulted in identifying similar barriers, and present the final PROM implementation plan, in addition to the evaluation plan that will be used across all CECP settings.

Application of the TDF in a primary care interdisciplinary program

Objectives: primary care study

The objective of this study was to identify potential barriers and enablers for the implementation of an ePROM system that will be used by clinicians working in an interdisciplinary team to deliver low back pain (LBP) care within a structured clinical program.
Study design and procedure: primary care study

To address the study objectives a four-step process proposed by French et al. (2012) [17] was used. The French approach starts with posing the question “who needs to do what differently?” and focuses on the actors implementing the intervention (ePROMs). Step 2 involves identifying the barriers and enablers to implementing the intervention. We chose the TDF framework for Step 2 as it was developed to mostly identify barriers to individual clinical behaviour change that matched the context of the chronic pain interdisciplinary teams. We also felt that the domains could be more readily mapped to behaviour change strategies (and mode(s) of delivery) to include in the ePROM implementation plan that could overcome the modifiable barriers and enhance the enablers (Step 3 of French et al.). Finally, Step 4 answers the question “How can behaviour change be measured and understood?”. We address this in the evaluation plan guided by the Proctor framework.

The TDF was developed and used to identify the barriers and enablers to clinical behavioural change among clinicians to inform the design of a theory-based ePROM implementation [18–22]. It includes 14 validated domains which are deemed to be the main contributors to behaviour change among clinicians: Knowledge, Skills, Social/Professional Role and Identity, Beliefs about Capabilities, Optimism, Beliefs about Consequences, Reinforcement, Intentions, Goals, Memory/Attention and Decision Processes, Environmental Context and Resources, Social Influences, Emotion, and Behavioural Regulation [23].

The 14 TDF domains were evaluated using an online self-administered survey that aimed to identify clinicians’ perceived barriers and enablers to using ePROMs. The survey included 30 questions adapted from validated TDF instruments [24, 25]. Items were rated on a 5-point Likert scale (strongly agree to strongly disagree). The survey also had 16 open-ended questions to obtain more information on clinicians’ opinions about the domains. A thematic analysis was conducted for open-ended responses to identify additional enablers and barriers.

Results: primary care study

Eighteen of 25 invited health professionals from primary care interdisciplinary teams responded to the survey (response rate of 72%). Respondents included two physiotherapists (PTs), three nurses, three psychologists, two occupational therapists (OTs), and two kinesiologists. The average age of the participants was 39 years (SD ± 7.7); 39% (7/18) were females, and the average
number of years practicing their respective professions was 14 years (SD ± 8.4). Determinants identified as likely to restrict the use of PROM scores included skills, social/professional role and identity, goals, decision processes, beliefs about consequences, environmental context and resources, behavioural regulation, and social influence. The barriers and enablers identified for each domain are summarized in Table 2.

Enablers reported in the open-ended question included clinicians’ motivation to use PROMs, having access to patient scores with the interpretation, and having patients fill out the questionnaires electronically. Factors that clinicians identified that may reduce the use of PROMs included lack of knowledge, limited time, and limited access to patients ePROM results if some patients choose not to complete measures or if there are a lack of resources to compile patient data. Disadvantages for using ePROMs listed by clinicians included patients’ having difficulty understanding some PROM items, clinicians having a limited understanding of what patients’ scores mean, limited usefulness of PROM scores in reflecting patients’ needs, and having to manage large volumes of ePROM data.

### Application of the CFIR in rehabilitation and tertiary care

#### Objectives: rehabilitation and tertiary-based care study

The objective was to identify the barriers and enablers for the implementation of an ePROM system that will be used in a rehabilitation and tertiary-based pain clinic to support shared decision making between patients and clinicians.

#### Study design and procedure: rehabilitation and tertiary-based care study

This project employed an intervention-mapping (IM) process [26] to inform the development of a tailored implementation plan of an ePROM system at a hospital-based pain clinic. IM consists of six steps: 1. Needs assessment; 2. Definition of program objectives; 3. Selection of theory-based intervention methods; 4. Production and pretesting; 5. Adoption, implementation, and sustainability planning; 6. Process and effect evaluation. We chose the Consolidated Framework for

### Table 2 Determinants to implementation of ePROMs in primary care using the TDF*

| Domain                          | Barrier                                                                 | Enabler                                                                 |
|---------------------------------|------------------------------------------------------------------------|------------------------------------------------------------------------|
| Knowledge                       |                                                                        |                                                                        |
| Skills                          | New skills are required to successfully use PROMs in the management of patients with LBP | Awareness of the objectives of using ePROMs in clinic                  |
| Social/Professional role and identity | Lack of clarification of clinicians’ professional role in relation to using ePROMs | Having the skills needed to interpret the results of ePROMs            |
| Optimism                        |                                                                        |                                                                        |
| Beliefs about consequences       | Using ePROMs in clinical practice is not necessary to improve patient outcomes | Believing that using ePROMs is one of the clinicians’ role in clinic for individual patient management of LBP |
| Reinforcement                   |                                                                        |                                                                        |
| Intentions                      | The plan of how to use ePROMs in clinical practice is not clear         | Expecting improved patient outcomes as a result of using ePROMs in the management of patients with LBP |
| Goals                           | The use ePROM scores is difficult in making treatment decisions        | Clinicians believe in the benefits of using ePROMs in the management of patients with LBP |
| Decision processes              |                                                                        | Having better patient health outcomes makes clinicians continue using ePROMs in the management of patients with LBP |
| Environmental Context and Resources |                                                                        | Clinicians’ commitment and intention to use ePROMs in the treatment of patients with LBP in the next three months |

*TDF Theoretical Domains Framework, LBP Low Back Pain
Implementation Research (CFIR) as a theoretical framework to guide the IM steps. CFIR serves as a practical guide for systematically assessing potential determinants in preparation for implementing an innovation [27]. The details of this study are presented elsewhere [2, 15, 16]. In this paper, we focus on the methods and results related to application of the CFIR to identify the determinants, barriers, and enablers to implementing ePROMs.

A survey with open-ended questions about enablers/barriers to the implementation of ePROMs was administered to clinicians and patients. Our clinical partner on the project, a nurse pain specialist working at the hospital-based pain clinic, recruited a convenient sample of both patients and clinicians to complete a survey adapted to both groups of participants. Patients were asked to come one hour before or immediately after their appointment with their clinician to complete the survey and answer several open-ended questions. Clinicians actively caring for patients at the pain clinic were sent an email from our clinical partner on behalf of the research team inviting them to complete an online survey. To increase participation, three email reminders were sent one week apart.

Based on the survey results, a focus group guide was developed, and focus groups were conducted with clinicians to explore their clinical practices related to the use of PROMs, and to deepen our understanding of the main perceived barriers and enablers to their use in the management of CP. Focus groups were digitally recorded and transcribed.

Quantitative survey data from clinicians and patients were analysed using descriptive statistics. The qualitative data included the open-ended questions of the patient survey and the content from the focus groups with clinicians and were analysed using deductive content analysis [28] based on core CFIR constructs. Two researchers independently conducted the analysis. All emerging themes were coded, and the two investigators continuously compared their analytical interpretations to identify the similarities and differences in the participants’ experiences to arrive at a common understanding. The results of the qualitative and quantitative analyses were reviewed with the study steering committee and three additional clinicians from the pain clinic. Together, they defined the program objectives, and synthesized the barriers and enablers identified from the survey and focus groups to be considered in the design and implementation plan of the ePROMs system.

Results: rehabilitation and tertiary-based care study

The survey was completed by 13 clinicians. Four were males and nine females. Five of the clinicians were psychologists, three nurses, two anaesthesiologists, two medical doctors, and one physiotherapist. A convenient sample of twenty-four patients, 12 males and 12 females responded to the survey with an average experience with CP of 9.8 ± 8.5 years. Overall, four focus groups were conducted, and a total of 10 clinicians were interviewed. Clinicians and patients were representative on average of those working or receiving care in the clinic.

Determinants were classified as barriers or enablers across the CFIR domains as shown in Table 3. Several enablers related to Intervention Characteristics that may facilitate implementation included designing the ePROM system in a manner that is easy for clinicians to use and meets patient needs (e.g. font, brightness), providing users with evidence about the clinical validity of PROMs, selecting PROMs that are easy to complete and interpret, and enabling access to ePROM systems using all forms of devices. Barriers related to the intervention included concerns with data privacy (e.g. who would be allowed to view PROM scores) and cost of implementing PROMs.

In the outer setting, barriers were related to patient dimensions including PROMs that matched individuals’ literacy level, the language spoken if it was not English or French, and having access to technology to be able to complete ePROMs at home or in the clinic so that scores would be readily available during the clinical encounter. There were also concerns that patients with limited cognitive abilities may not be able to understand PROM questions and responses.

Inner setting-related enablers included senior management emphasizing the importance of having onsite IT support, having human resource support to help patients complete ePROMs, providing clinicians with rapid access to results, and training on the use of ePROMs including a guide to interpret the results, and information on the validity of the measures used in the ePROM system. On the other hand, the perceived increase in workload and time associated with implementing ePROMs were considered as barriers to the use of an ePROM system.

Clinicians’ knowledge and beliefs that can facilitate implementation included perceptions that were important to use ePROMs and that sharing ePROM scores with patients will help with improving patient engagement and in turn pain management, and clinicians’ familiarity with the ePROM intervention. The counterpart barriers to these enablers included clinicians’ limited knowledge of the benefits of PROMs and how to apply them in clinical care, limited perceived usefulness of an ePROM system, and clinicians’ perception that PROMs may not be comprehensive and meet the information needs of patients. Clinicians also expressed concerns that use of PROMs may decrease patient satisfaction with health services, and that some patients may not provide accurate information on PROMs.

Finally, several determinants that were identified as likely facilitating the process of implementing PROMs included garnering support for ePROMs from all stakeholders,
### Table 3 Determinants to implementation of ePROMs in rehabilitation and tertiary care using the Consolidated Framework for Implementation Research

| CFIR Domain                             | Barrier                                                                 | Enabler                                                                 |
|-----------------------------------------|-------------------------------------------------------------------------|-------------------------------------------------------------------------|
| Intervention characteristics            |                                                                         |                                                                         |
| Evidence strength & quality             |                                                                         |                                                                         |
| Complexity                              | Filling ePROMs can be too much effort and time for some patients         | ePROMs will provide valid clinical information                           |
|                                         | Privacy/confidentiality concerns                                        | ePROM system easy to use for both patients and clinicians               |
|                                         |                                                                         | Include in ePROM, ePROMs simple to complete, short and easy to interpret |
| Design quality and packaging            |                                                                         |                                                                         |
| Cost                                    | Additional funding needed to support implementation of ePROMs (equipment, administrative) | Patients interested in completing ePROMs                                |
| Outer setting                           |                                                                         |                                                                         |
| Patient needs and resources             | Not all patients have access to IT                                      | Beliefs of senior management of the importance of ePROM Implementation  |
|                                         | Low level of literacy of patients                                      |                                                                         |
|                                         | Insufficient patient cognitive abilities to be able to complete ePROMs  |                                                                         |
|                                         | Patients not understanding English or French                            |                                                                         |
| Inner setting                           |                                                                         |                                                                         |
| Implementation climate                  | Lack of time to look at ePROMs results at each patient’s visit           | Beliefs of senior management of the importance of ePROM Implementation  |
| Readiness for implementation            |                                                                         | Onsite IT support                                                       |
|                                         |                                                                         | Having human resource support to help patients fill out ePROMs          |
|                                         |                                                                         | Training [clinicians] on the use of the ePROM system and interpretation of ePROM results |
| Characteristics of individuals          |                                                                         |                                                                         |
| Knowledge and beliefs about the intervention | Clinicians’ little familiarity with ePROMs                              | Familiarity with technology                                              |
|                                         | Usefulness and relevance of an ePROM system                            |                                                                         |
|                                         | Information provided by ePROMs will not be comprehensive                |                                                                         |
|                                         | ePROMs do not always meet the information needs of patients             |                                                                         |
|                                         | Using ePROMs may decrease patient satisfaction with healthcare services |                                                                         |
|                                         | Patients not willing to provide accurate information                    |                                                                         |
| Individual stage of change              | Reluctance of some clinicians to use the ePROM system: individual stage of change | Motivation and interest of some clinicians for an ePROM                  |
| Motivation Process Planning             |                                                                         |                                                                         |
| Planning                                | Completion of ePROMs aligned with clinical process:                     |                                                                         |
|                                         | while patient in waiting room                                          |                                                                         |
|                                         | “Top down” implementation                                               |                                                                         |
|                                         | Involvement of multidisciplinary team in development of an implementation plan (data collection and clinical process to use ePROM scores) |                                                                         |
|                                         | Project manager for data collection process and clinical use of ePROMs information |                                                                         |
| Engaging                                |                                                                         | Interest and support of stakeholders in ePROM implementation             |
deciding at the department [leadership] level to implement ePROMS (top down), involving the multidisciplinary team in the development of the ePROM system, engaging clinicians in motivating patients to complete ePROMs, aligning implementation of ePROMs to the clinical workflow, and having a project manager in the clinic to support data collection and use of ePROM information in clinical care.

**ePROM implementation plan and training from primary care, tertiary care, and rehabilitation**

The steering committee reviewed all information regarding the determinants of implementation of ePROMs identified across both studies and reviewed the objectives of the ePROM system implementation plan across the integrated pain network. The program objectives encompassed two broad categories. The first was adoption of PROMs, mainly, increasing feasibility and clinician and patient acceptability of ePROMs, and enhancing clinician and patient knowledge about ePROM score interpretation. The second category were objectives regarding quality of care and included increasing patients’ engagement and satisfaction with care, improving both patient-provider and healthcare team communication, informing shared decision making and treatment planning between patients and providers, facilitating goal setting and problem-solving, and increasing the focus of services on patient-centred outcomes.

The steering committee agreed to pursue the following steps for implementation at an organizational level. First, develop a business plan to incorporate the costs related to the ePROM system by directly integrating resources needed into the sites’ budget. This will be accomplished by examining potential cost savings associated with using ePROM scores to triage patients more efficiently and administrative time needed to complete paper-based forms for standardized and non-standardized patient information. The second organizational change recommended was to link ePROM scores to the clinic’s electronic medical record to facilitate integration of ePROM scores into the clinical workflow.

The steering committee defined the following process for facilitating implementation at the level of the clinical unit: administrative staff will send an email to patients containing the link to the ePROM questionnaire 2–4 weeks prior to the first visit or follow-up appointment. The patient will be invited to complete ePROMs at home 1 week prior to first visit or follow-up appointment. If PROMs were not completed at home patients will be encouraged by email and/or telephone to arrive before his/her appointment to complete ePROMS at the clinic.

The ePROM results will be shared with clinicians and patients through an online feedback report before the clinic visit. The feedback report includes patients’ scores over time for each measured domain, and the interpretation of each PROM score using graphs and text to indicate the current scores and the changes over time in each domain. Prior to the first visit or follow-up appointment, clinicians will be prompted to review the results in preparation to reviewing the scores with patients during the clinical encounter.

Evaluation of the determinants played a critical role in deciding on the implementation strategies that will be included in the development of a KT intervention that will increase the use of ePROMs. The process of deciding on which strategies to include in the KT intervention are presented elsewhere [2, 15, 16]. The main selected strategies were an interactive training program. The program will include educational/instructional material on the selection, application, and interpretation of PROM scores and the clinicians’ and patients’ roles in using PROMs to plan treatment and monitor changes in outcomes in collaboration with patients. The program will also include a half-day training workshop on the use of PROMs in clinical practice that will include interpretation of feedback reports of individuals’ PROM scores.; The second strategy will be designating an opinion leader to provide clinicians with one-on-one coaching when needed to support the implementation of the KT intervention components and to provide coaching to clinicians on the use of PROMs. Table 4 presents the final interactive training program that will be adopted by the McGill RUISSS CECP.

**ePROM evaluation plan across an integrated chronic pain network**

Guided by the Proctor framework of evaluation, [13] the steering committee agreed on a set of outcomes that will be used to evaluate implementation and impact of using PROMs. The PROMs that will be implemented as part of patient care are those that are part of a CORE set prioritised by patients and clinicians [29] using PROMIS measures [30]. The framework “positions implementation outcomes as preceding both service outcomes and client outcomes,
with the latter sets of outcomes being impacted by the implementa-
tion outcomes” [13]. The selection of outcomes was
guided by the CECP objectives for using ePROMs, and the
interrelationship of outcomes we expected to be impacted
by adoption of the ePROM intervention. For example, we
would expect to see the strongest impact on patient out-
comes (e.g. general health, pain interference, work ability)
as the ePROM uptake and sharing and discussion of scores
with patients increases in a clinical setting. The Proctor
framework derives service outcomes from the six quality
improvement aims set out in the reports on crossing the
quality chasm: the extent to which services are safe, effec-
tive, patient-centred, timely, efficient, and equitable [31, 32].
Table 5 presents the implementation and impact outcomes
that will be used to evaluate the ePROM intervention across
the McGill RUISSS CECP integrated network.

### Discussion

In this review of two studies, we demonstrated the use of the
TDF and CFIR as guiding frameworks to evaluate determi-
nants of implementation of ePROMs. As recommended by
Birken et al. (2017), we explicitly specified the rationale for
using the TDF or the CFIR to contribute to our understand-
ing of each framework, its scope, and its limitations of using
both frameworks to implement PROMs. Using theoretical
frameworks facilitates identifying barriers and enablers to
implement ePROMs in clinical settings and guides the de-
velopment of theory-based interventions aimed at closing the
gaps between knowledge and practice. The results of this
study provide researchers and clinicians that are planning
to monitor implementation of ePROMs an understanding
of how the TDF and CFIR can be applied and highlight
possible barriers and enablers that will likely need to be
addressed.

The contrast and integration of results from both studies
provided us with rich insight into the application of both
approaches to the implementation of PROMs in clinical
care. Similar to previous studies, our rationale for selecting
the TDF or CFIR was driven by the purpose at the time for
evaluating the determinants of implementation. The TDF
was used in the context of an interdisciplinary team perspec-
tive, where each team member had to consider use of PROM
scores for creating an intervention plan related to their disci-
pline, and to link the resulting treatment plan to the overall
program delivered by all team members in collaboration with the patient. The TDF targeted well the individual-level determinants of implementation that needed to be considered. Also, the clinical process of the LBP program was well defined and endorsed by the healthcare organizations where these programs were implemented as it was part of the provincial CECP Action Plan. Using ePROMs in these programs was endorsed and was an organizational expectation from the start. In contrast, we used the CFIR in rehabilitation and tertiary care where organizational-level factors were felt to play an important role in the adoption of ePROMs. The CFIR operationalized the individual characteristics of clinicians and patients (e.g. knowledge & beliefs), and the organizational and process factors that are critical in planning the implementation of PROMs.
Interestingly, both frameworks provided similar results with respect to healthcare providers’ knowledge and behaviour. These included the importance of considering clinician’s attitudes and perceptions of the benefits of using ePROM scores, such as believing they will have a positive impact on patients’ outcomes. The CFIR captured more barriers in relation to knowledge and beliefs including clinicians’ beliefs that PROM scores may not be comprehensive [of patients needs] and the perception that using PROMs may decrease patients’ satisfaction with care. Intention and motivation to use PROMs were identified in the TDF and CFIR study, respectively. These individual clinician characteristics identified in our studies are similar to those reported in previous work [34–38]. Both frameworks were also consistent in identifying clinicians’ knowledge and experience in using PROMs, and difficulties with interpreting PROM scores as barriers, similar to previous work [35, 39, 40].

Where the TDF and CFIR frameworks differed was in identifying organizational-level determinants, which the CFIR was able to capture because of the explicit questions asked linked to the CFIR domains. This was reported in previous studies that contrasted the use of both frameworks for implementing healthcare interventions [41–43]. Some consensus among coders in each of our studies required discussion, as some statements from the qualitative data were not easy to classify in either framework. However, this was more frequent for the TDF. For example, we felt that statements related to the relationship between clinician and patient were harder to map to the TDF framework. In the CFIR we classified “PROMs will be filled by patients if clinicians motivate patients to do so” as best fitting under ‘process’ in the CFIR, and there is no equivalent domain in the TDF.

In all settings we started with evaluating the current use of measures, what improvements in measurement were required, and asking clinicians and patients about their needs in terms of patient care to link use of PROMs to these needs [11, 44]. This was a key first step to understand the baseline practices of the sites and to tailor the knowledge translation interventions for ePROM implementation and training content to the readiness of the sites to begin using ePROMs. We also had knowledge exchange sessions with clinicians regarding the measures they were currently using and whether there were more efficient ePROMs with more comprehensive coverage of domains, stronger psychometric properties, and available evidence for interpretation of ePROM scores.

The resulting KT and training intervention elements will form a multi-component intervention, which can address multiple gaps and barriers [45], increase clinicians’ knowledge and skills, and foster best practices. A previous study [11] identified the most important PROMs as perceived by clinicians and patients that will form the core set of PROMs that will be included across all sites. As clinical programs learn from their experience with this core set, the use of ePROMs will be continually improved to include additional measures, linkages between ePROMs and electronic medical records, and optimization of the timing of administering ePROMs. The ongoing evaluation will be essential for informing ongoing improvements and sustainability of implementation and use of ePROMs.

Lastly, the importance of ePROMs feedback reports was emphasized across all our studies. All clinicians, to save time and facilitate the sharing of ePROM results among clinicians and patients, highlighted the format and integration of these reports in the electronic medical record. Feeding back the scores to clinicians and patients may increase their belief in value of using PROMs in clinical practice, improve the communication between patients and clinicians, and empower individuals with chronic pain to actively participate in managing their health condition, which may lead to better treatment adherence [46, 47]. We will qualitatively evaluate the determinants of ePROM implementation over time to iteratively improve ePROM implementation elements including feedback reports, KT and training, and integration of ePROMs across the integrated chronic pain network to facilitate uptake by each care setting that is added as we scale up the use of ePROMs to support patient-centred care.

Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

Ethical approval The project by Eilayyan et al. has been approved by the Centre for Interdisciplinary Research in Rehabilitation of Greater Montreal (#1179–1016) and the project by Simeone et al. has been approved by the McGill University Health Centre (#2017–3338).

Informed consent Informed consent was obtained from all participants.

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References

1. Dueñas, M., et al. (2016). A review of chronic pain impact on patients, their social environment and the health care system. J Pain Res, 9, 457.
2. Raffaele Simeone, M.S.-M., Adele Vizcaino, Albert Weng, Krista Brecht, Diana Zidarov, Regina Visca, Sara Ahmed, Development and Implementation of an Electronic Patient Reported Outcome Measures tool: an Intervention Mapping Protocol. Submitted PLOS One.
3. Roth, R. S., Geisser, M. E., & Williams, D. A. (2011). Interventional pain medicine: retreat from the biopsychosocial model of pain. Translational behavioral medicine, 2(1), 106–116.
4. McCarberg, B. H., et al. (2008). The impact of pain on quality of life and the unmet needs of pain management: results from pain sufferers and physicians participating in an Internet survey. Am J Therapeutics, 15(4), 312–320.
5. Dobkin, P. L., & Boothroyd, L. J. (2008). Organizing health services for patients with chronic pain: when there is a will there is a way. Pain Med, 9(7), 881–889.
6. Cheatle, M. D., Klocek, J. W., & McLellan, A. T. (2012). Managing pain in high-risk patients within a patient-centered medical home. Translational Behavioral Medicine, 2(1), 47–56.
7. Higginson, I. J., & Carr, A. J. (2001). Using quality of life measures in the clinical setting. BMJ, 322(7297), 1297–1300.
8. E., V.E.a.B., Multidimensional assessment in palliative care. 2006. London.
9. Greenhalgh, J., Long, A. F., & Flynn, R. (2005). The use of patient reported outcome measures in routine clinical practice: lack of impact or lack of theory? Social Science and Medicine, 60(4), 833–843.
10. Greenhalgh, J. (2009). The applications of PROs in clinical practice: what are they, do they work, and why? Quality of Life Research, 18(1), 115–123.
11. Zidarov, D., Visca, R., & Ahmed, S. (2019). Type of clinical outcomes used by healthcare professionals to evaluate health-related quality of life domains to inform clinical decision making for chronic pain management. Quality of Life Research, 28(10), 2761–2771.
12. Birken, S. A., et al. (2017). Combined use of the Consolidated Framework for Implementation Research (CFIR) and the Theoretical Domains Framework (TDF): a systematic review. Implementation Science, 12(1), 2.
13. Proctor, E., et al. (2011). Outcomes for Implementation Research: Conceptual Distinctions, Measurement Challenges, and Research Agenda. Administration and Policy in Mental Health and Mental Health Services Research, 38(2), 65–76.
14. Esmail, R., et al. (2020). A scoping review of full-spectrum knowledge translation theories, models, and frameworks. Implementation Science, 15(1), 1–14.
15. O Ei layyan, R.V., D Zidarov, P Ware, A Bussieres, S Ahmed. Developing Theory-Informed Knowledge Translation Strategies to Facilitate the Use of Patient Reported Outcome Measures in the Low Back Pain Clinical Practice. 2017. McGill: Montreal.
16. O Ei layyan, R.V., D Zidarov, P Ware, A Bussieres, S Ahmed. Developing Theory-Informed Knowledge Translation Strategies to Facilitate the Use of Patient-Reported Outcome Measures in the Low Back Pain Clinical Practice. Quality of Life Research. Submitted 2019 October.
17. French, S. D., et al. (2012). Developing theory-informed behaviour change interventions to implement evidence into practice: a systematic approach using the Theoretical Domains Framework. Implement Sci, 7, 38.
18. McGrady, M. E., et al. (2015). Topical Review: Theoretical Frameworks in Pediatric Adherence-Promotion Interventions: Research Findings and Methodological Implications. Journal of Pediatric Psychology, 40(8), 721–726.
19. Dobson, F., et al. (2016). Barriers and Facilitators to Exercise Participation in People with Hip and/or Knee Osteoarthritis: Synthesis of the Literature Using Behavior Change Theory. American Journal of Physical Medicine and Rehabilitation, 95(5), 372–389.
20. Mosaviyanpour, M., et al. (2016). Theoretical domains framework to assess barriers to change for planning health care quality interventions: a systematic literature review. J Multidiscip Healthc, 9, 303–310.
21. Bussières, A. E., et al. (2015). Fast tracking the design of theory-based KT interventions through a consensus process. Implementation Science, 10(1), 18.
22. French, S., et al., Developing theory-informed behaviour change interventions to implement evidence into practice: a systematic approach using the Theoretical Domains Framework. Implement Sci, 2012, 7.
23. Cane, J., O’Connor, D., & Michie, S. (2012). Validation of the theoretical domains framework for use in behaviour change and implementation research. Implement Sci, 7, 37.
24. Huijg, J. M., et al. (2014). Measuring determinants of implementation behavior: psychometric properties of a questionnaire based on the theoretical domains framework. Implement Sci, 9, 33.
25. Huijg, J. M., et al. (2014). Discriminant content validity of a theoretical domains framework questionnaire for use in implementation research. Implement Sci, 9, 11.
26. Eldredge, L.K.B., et al., Planning health promotion programs: an intervention mapping approach. 2016: John Wiley & Sons.
27. Damschroder, L. J., et al. (2009). Fostering implementation of health services research findings into practice: a consolidated framework for advancing implementation science. Implementation science, 4(1), 50.
28. Hsieh, H.-F., & Shannon, S. E. (2005). Three approaches to qualitative content analysis. Qualitative health research, 15(9), 1277–1288.
29. Zidarov, D., Zidarova-Carriété, A., Visca, R., Miller, J. M., Brecht, K., Viens, N., et al. (2020). Core patient-reported outcome domains for routine clinical care in chronic pain management: Patients’ and healthcare professionals’ perspective. Quality of Life Research, 29(7), 2007–2020.
30. Cella, D., et al. (2010). Initial Adult Health Item Banks and First Wave Testing of the Patient-Reported Outcomes Measurement Information System (PROMIS™) Network: 2005–2008. Journal of Clinical Epidemiology, 63(11), 1179.
31. Series, Q.C., Committee on crossing the quality chasm: Adapting to mental health and addictive disorders. Washington, DC: The Institute of Medicine. 2006. The National Academies Press.
32. Kohn, L.T., J. Corrigan, and M.S. Donaldson, Institute of Medicine (US). Committee on Quality of Health Care in America. (2000). To err is human: Building a safer health system.
33. Zhang, J., & Walji, M. F. (2013). The CONSORT Patient-Reported Outcomes Measurement Information System (PROMIS™) Network: 2005–2008. Journal of Clinical Epidemiology, 63(11), 1179.
34. Chang, C.-H. (2007). Patient-reported outcomes measurement and management with innovative methodologies and technologies. Quality of Life Research, 16(1), 157–166.
35. Boyce, M. B., Browne, J. P., & Greenhalgh, J. (2014). The experiences of professionals with using information from patient-reported outcome measures to improve the quality of healthcare: a systematic review of qualitative research. BMJ Qual Saf, 23(6), 508–518.
36. Jette, D. U., et al. (2003). Evidence-based practice: beliefs, attitudes, knowledge, and behaviors of physical therapists. Physical therapy, 83(9), 786–805.
37. Calvert, M., et al. (2013). The CONSORT Patient-Reported Outcome (PRO) extension: implications for clinical trials and practice. Health and quality of life outcomes, 11(1), 184.
38. Lydick, E., & Epstein, R. (1993). Interpretation of quality of life changes. *Quality of life Research, 2*(3), 221–226.
39. Calvert, M., et al. (2013). The CONSORT Patient-Reported Outcome (PRO) extension: implications for clinical trials and practice. *Health Qual Life Outcomes, 11*, 184.
40. Boyce, M. B., Browne, J. P., & Greenhalgh, J. (2014). The experiences of professionals with using information from patient-reported outcome measures to improve the quality of healthcare: a systematic review of qualitative research. *BMJ Quality & Safety, 23*(6), 508–518.
41. Gould, N. J., et al. (2014). Application of theory to enhance audit and feedback interventions to increase the uptake of evidence-based transfusion practice: an intervention development protocol. *Implementation Science, 9*(1), 92.
42. Prior, M., et al. (2014). Evaluating an audit and feedback intervention for reducing antibiotic prescribing behaviour in general dental practice (the RAPiD trial): a partial factorial cluster randomised trial protocol. *Implementation Science, 9*(1), 50.
43. Sales, A. E., et al. (2016). Implementing goals of care conversations with veterans in VA long-term care setting: a mixed methods protocol. *Implementation Science, 11*(1), 132.
44. Gogovor, A., et al. (2019). Determinants Of Patient Experience With Low Back Pain Interdisciplinary Care: A Pre-Post Interventional Study. *Journal of Pain Research, 12*, 3203.
45. Straus, S. E., J. Tetroe, and I. D. Graham, *The Knowledge-to-Action Cycle*, in Knowledge Translation in Health Care: Moving from Evidence to Practice. 2009, Blackwell Publishing Ltd. p. 59–181.
46. Antunes, B., et al. (2014). Implementing patient-reported outcome measures in palliative care clinical practice: a systematic review of facilitators and barriers. *Palliative medicine, 28*(2), 158–175.
47. Detmar, S. B., et al. (2002). Health-related quality-of-life assessments and patient-physician communication: a randomized controlled trial. *JAMA, 288*(23), 3027–3034.

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