Monitoring physical functioning as the sixth vital sign: evaluating patient and practice engagement in chronic illness care in a primary care setting—a quasi-experimental design

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Abstract

Background: In Canada, one in three adults or almost 9 million people report having a chronic condition. Over two thirds of total deaths result from cardiovascular disease, diabetes, cancer and respiratory illness and 77% of persons ≥65 years have at least one chronic condition. Persons with chronic disease are at risk for functional decline; as a result, there is an increased awareness of the significance of functional status as an important health outcome. The purpose of this study was to determine whether patients who receive a multi-component rehabilitation intervention, including online monitoring of function with feedback and self-management workshops, showed less functional decline than case matched controls who did not receive this intervention. In addition, we wanted to determine whether capacity building initiatives within the Family Health Team promote a collaborative approach to Chronic Disease Management.

Methods: A population-based multi-component rehabilitation intervention delivered to persons with chronic illnesses (≥ 44 yrs) (n = 60) was compared to a group of age and sex matched controls (n = 60) with chronic illnesses receiving usual care within a primary healthcare setting. The population-based intervention consisted of four main components: (1) function-based individual assessment and action planning, (2) rehabilitation self-management workshops, (3) on-line self-assessment of function and (4) organizational capacity building. T-tests and chi-square tests were used for continuous and categorical variables respectively in baseline comparison between groups.

Results: Two MANOVA showed significant between group differences in patient reported physical functioning (Λ = 0.88, F = (2.86) = 5.97, p = 0.004) and for the physical performance measures collectively as the dependent variable (Λ = 0.80, F = (6.93) = 3.68, p = 0.0025). There were no within group differences for the capacity measures.

Conclusion: It is feasible to monitor physical functioning as a health outcome for persons with chronic illness in primary care. The timeline for this study was not sufficient to show an increase in the capacity within the team; however there were some differences in patient outcomes. The short timeline was likely not sufficient to build the capacity required to support this approach.

Trial registration: NCT00859638

Keywords: Rehabilitation, Primary care, Chronic disease, Physical functioning, Self-management, Self-monitoring
Background
Chronic disease is the leading cause of death (60%) and disability (43%) worldwide [1,2]. In Canada, one in three adults or almost 9 million people report having at least one of seven high impact, high prevalence chronic conditions and 77% of persons ≥65 years have at least one chronic condition [3]. Over two thirds of total deaths result from cardiovascular disease, diabetes, cancer and respiratory illness. Canadians with chronic conditions account for over 70% of all nights spent in hospital and half of Canadians with multiple chronic conditions report moderate to severe disability in daily living [3,4].

Optimizing and preserving physical functioning is a central goal for all persons with chronic illness. The results of the Medical Outcomes study of ambulatory patients (mean age 46 years) showed that persons with eight out of the nine chronic conditions studied had worse function across physical, social and mental domains compared to persons without the conditions [5]. Comorbidity was associated with greater decrements in functioning [5].

There is increasing awareness of the importance of functional status as a major health outcome as well as an emphasis on cost effective interventions for its enhancement [6,7]. There is growing evidence that functional status data are vital to clinical practice and substantiate health system performance [8]. Functional status as an outcome of care is a major concern for persons of all ages with chronic illness who are trying to self-manage their conditions. Loss of self-management abilities has been associated with loss in different functioning domains, including physical functioning [9]. Changes in physical functioning are a better predictor of loss of self-management skills than chronological age. Self-management skills associated with function are needed by persons with chronic disease to prevent and manage functional decline and loss of resources [10]. There is a lack of documentation in health records in hospital and primary care settings about the functional status of patients [7]. It has been suggested that functional status should be the sixth vital sign monitored, in addition to the conventional vital signs of temperature, pulse, respiration rate, blood pressure, and blood oxygen saturation [7]. Physicians do not routinely monitor or assess functional status as part of a patient's management even when prompted [11,12]. Thus, providing awareness about the relationship between physical function and the patient's overall health status in addition to assisting physicians and other members of the health care team to build their capacity in assessing functional status is likely to improve the care for persons with chronic illness. Emerging technology such as electronic health records linked to in situ technology will assist in understanding the dynamic nature of chronic diseases, the management to maximize health and the efficient use of healthcare resources in this population [13].

The Chronic Care Model (CCM) http://www.improvingchroniccare.org is an approach to planned, proactive, population-based, evidenced based and patient-centered care to manage chronic illness rather than reactive acute oriented care [14-16]. The six dimensions of the CCM are organization of health care (health and design systems), clinical information systems, self-management support, delivery system support, and community resources. It represents enhancements in organizations and practices that contribute to productive interactions between providers and patients [16,17]. Effective care using this model requires regular interaction between caregivers and patients and the Institute of Medicine (IOM) recommends alternate methods of interaction rather than solely face-to-face visits [15]. The Expanded Chronic Care Model (ECCM) enlarges the community portion of the CCM and demonstrates how the patient fits within the concepts of population health [17]. This model forms the basis for the framework for chronic disease management in Ontario [17]. Canada has adopted the population health approach as a health care strategy for health policy and program development [18-20].

Allied health professionals provide services to approximately 18% of patients within primary care settings [3]. However physiotherapists and occupational therapists have not been integrated into most primary health care settings in Canada and there is a gap in rehabilitation service provision for persons with chronic illness. This study examined a population-based approach that involves an intervention targeted at patients’ self monitoring of physical functioning and capacity building within a primary care setting to support the integration of physical functioning within the Chronic Care Model.

The overall goals for the project were:
1. To assess whether adopting a population-based, rehabilitation self-management approach that focused on physical functioning as a major health outcome in a primary care setting improves the process and outcome of care for patients with chronic conditions.
2. To evaluate the extent to which members of a Family Health Team (FHT) integrate the assessment and monitoring of physical function, and implement of interventions to maintain physical function of their patients within the process of delivering chronic illness care.

Research objectives:
1. To determine whether patients who received a multi-component intervention, namely rehabilitation assessment, action planning, access to online functional
assessment with feedback and rehabilitation self-management workshops, showed less functional decline as measured by the Physical Functioning Inventory [21], the Two Minute Walk Test, [22] Grip Strength [23] and lower extremity function tests [24] than case matched controls who did not receive the intervention.

2. To determine whether a multi-component intervention increased patient self-management within a Family Health Team as measured by the Self-Efficacy for Chronic Disease Scale [25] and the Rapid Assessment of Physical Activity [26].

3. To determine whether capacity building initiatives within the FHT increased the collaborative approach to Chronic Disease Management by the Patient Care Provider (PCP) as measured by The Assessment of Primary Care Resources and Supports for Chronic Disease Self-Management (PCRS) [27] and the Patient Assessment of Chronic Illness Care (PACIC) [28].

Methods
Study design
A before-after design was used to compare a population-based rehabilitation intervention delivered to persons with chronic disease compared to a group of age and sex matched controls receiving usual care. The population-based intervention consisted of four main components: (1) function-based individual assessment and action planning, (2) rehabilitation self-management workshops, (3) on-line self-assessment of function (4) organizational capacity building. These are described below and summarized in Figure 1. Outcome measures for each of the four components of the intervention were administered at baseline and 6 months follow-up. The study was conducted with approval from the Hamilton Health Sciences/McMaster University Research Ethics Board (REB #08-177).

Sample
Two academic family practice units, the Stonechurch Family Health Centre (SFHC) and the McMaster Family Practice (MFP) that together comprise the McMaster Family Health Team (FHT), participated in the project. Family Health Teams are health care organizations that include a team of family physicians, nurse practitioners, registered nurses, social workers, dietitians, and other professionals who work together to provide health care for their community [29]. The intervention group was recruited from the Stonechurch Family Health Centre (SFHC), with 15 physicians and 30 residents on three teams that serve approximately 13,500 patients. The control group was selected from the McMaster Family Practice (MFP), which provides care to 12,000 patients, attended by 13 physicians and 36 residents. The sites are very similar in composition and organization and operate as one Family Health Team although they are geographically at different sites and have different Medical Directors. Ten family physicians and ten allied health team members from SFHC agreed to participate in the capacity building component of the study. The study used a convenience sample of providers. Patients were randomly selected from the practice database who met the following inclusion criteria: age ≥44 years; having one or more chronic conditions; at least 3 physician visits in the past year; able or willing to access an email address. The ICD-9 billing codes for these chronic diseases were used to select the sample: rheumatoid arthritis, back pain, cardiac arrest, heart failure, chronic obstructive pulmonary disease, stroke, diabetes, emphysema, hypertension, multiple sclerosis, osteoarthritis, osteoporosis, Parkinson’s disease, and cerebral vascular accident. Case matched controls, matched on age, gender, type and number of chronic conditions selected from the McMaster Family Practice formed the control group. Control group participants received baseline and final physical functioning assessments from the PT and OT and usual care from their health care team. Sixty patients were recruited to both the intervention and control groups. We recruited 20 patients from three defined stages of functional ability. The levels of functioning were: (1) no difficulty in physical functioning but the patient has made modifications to the tasks they complete or have changed the frequency with which they complete these tasks; (2) early changes or difficulty in physical functioning; (3) established difficulty in physical functioning, experiencing significant or longstanding difficulties with physical functioning, mobility or activities of daily living [30]. The study coordinator, a physiotherapist, reviewed each patient’s Electronic Medical Record (EMR), and assigned a physical functioning rating based on the patient’s age, number and type(s) of chronic condition(s), length of time chronic conditions were present, work status, disability insurance, mobility status, activities of daily living, use of gait aids, recent surgeries and hospitalizations, use of a disability parking pass, falls, Worker Safety and Insurance Board claims, level of physical activity and smoking. The patient’s physician agreed with the investigators’ rating in 88% of the cases, the final level was determined by the physician’s rating.

Assessments
Evaluation of physical functioning
The Physical Functioning Inventory (PFI) was designed to assess pre clinical functional impairment and physical functioning in older adults [21]. It contains 21 items from 4 different subscales: activities of daily living (ADLs), instrumental activities of daily living (IADL), mobility activities, and moderate/strenuous activities.
Probe questions are used to determine whether the person experiences difficulty in completing a task, and changes the method and/or frequency of task performance. Agreement between raters in at least 19 of 22 tasks on the probes has been reported with the PFI to be greater than 80% [21]. The 2-Minute Walk test is a reproducible measure of functional exercise capacity that is measured as the distance walked in 2 min [22]. Correlations between 2- and 6- min walk tests indicate that they are similar measures of exercise tolerance (r = 0.89) [22]. Test-retest reliability ranges from 0.82 to 0.89 in frail elderly individuals. Grip strength was measured using a JAMAR hand-held dynamometer averaged over 3 trials with each hand. It is an indicator of overall strength and a strong predictor of mortality in adults 65-90 years with moderate to severe disabilities [23]. Lower extremity function was measured by the 8-foot timed walk test and standing balance test. The performance of each of these tests and a summed score over several tests, have been associated with self-reported disability [24]. The Timed Up and Go (TUG) was completed only during the patients’ final assessment. The TUG is designed to measure balance and basic functional mobility [31]. It is the time in seconds to rise from a regular chair with arms, walk 3 m at a comfortable safe pace, turn, walk back to the chair and sit down. High sensitivity (87%) and specificity (87%) for the TUG’s ability to identify community dwelling adults at risk for falls has been reported [32,33]. The inter-rater reliability is 0.98 [31].

**Evaluation of Rehabilitation Self-Management (RSMP)**

Three measures, developed at Stanford Patient Education Research Centre for Chronic Disease, [33] were used to evaluate the workshops. The Self Efficacy for Chronic Disease Scale (SECDS) has 6 items rated on a 10-point scale [24,34,35]. It covers the domains of symptom control, role functioning, emotional functioning and communicating with physicians. The internal consistency ranged from 0.8-0.9 and test re-test reliability r = 0.7-0.9 [24]. The Health Utilization Scale, a 4 item questionnaire that asks about the emergency room use, hospital use and physician visits in the past 6 months, [34-36] and a single question on self-rated health were also administered the first and last day of the program [34]. The Rapid Assessment of Physical Activity (RAPA) was considered an indication of self-management for this study since it was postulated that patients in the intervention group might increase their level of physical activity.

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**Figure 1** Population-Based Model for Rehabilitation in Primary Care for Persons with Chronic Illness.
activity as a result of increased awareness about physical functioning. The RAPA is a nine-item self-report measure that assesses frequency and intensity of aerobic physical activity as well as frequency of strengthening and flexibility exercises. It has been validated in adults older than 50 years [26]. The RAPA is more highly correlated \( (r = 0.54) \) with moderate caloric expenditure than two other commonly used measures of physical activity, the Patient-Centred Assessment and Counseling for Exercise (PACE) \( (r = 0.44) \) and the Behavioural Risk Factor Surveillance System (BRFSS) \( (r = 0.40) \) [26,37]. It has a reported sensitivity (81%) and positive predictive value (77%) [26].

**Evaluation of capacity building (CB) intervention**

**Process measures** The Assessment of Primary Care Resources and Supports for Chronic Disease Self-Management (PCRS) was used to evaluate the CB intervention [27]. It assesses patient self-management based on the process and structural level of care. It has acceptable psychometric properties; Cronbach’s alpha for individual and organizational support subscales were 0.94 and 0.90 indicating a high level of internal consistency [27]. The Patient Assessment of Chronic Illness Care (PACIC) assesses the patient’s perspective on chronic illness care received based on the Chronic Care Model [27,35]. It has 5 scales and an overall summary score. It has demonstrated moderate test-retest reliability \( (r = 0.58 \text{ over 3 months}) \) and correlated moderately with measures of primary care and patient activation \( (r = 0.32-0.60, \text{median 0.50 p < 0.001}) \) [35].

**Focus group** A focus group evaluation was held at the end of the project to investigate further the Patient Care Providers’ (PCPs’) experiences of the CB process, and how this influenced their practice. It was also used to discuss whether the results of the functional assessments were incorporated into their discussions with the patient and the care plan.

**Patient intervention**

**Function-based individual assessment and action planning**

The intervention is depicted in Figure 1. At entry to the study, the patient engaged in a joint physiotherapy (PT) and occupational therapy (OT) assessment to identify their concerns and goals related to physical function. The PT and OT conducted a performance-based assessment of physical functioning [22,23,38] using standardized tests and the results were discussed with the patient in the context of age- and sex-based norms. During the initial assessment, patients participated in collaborative goal setting with the therapist, identifying a personal functional goal for the next 3-6 months. Patients then created weekly action plans related to the goal using a self-management behaviour that addressed the functional goal. A copy of the action plan was given to the patient. The therapists documented the results of the performance tests, the problem list and action plan in the EMR for the PCPs to review. The patients received the intervention for a period of 6 months, and then were reassessed by the PT and OT.

**Rehabilitation self-management workshops**

An evidence-based 5-week Rehabilitation Self-Management Program (RSM) was delivered to increase patients’ self-management behaviours and self-efficacy in the monitoring of physical functioning. The goals of the program were to build self-management skills, identify goals, set action plans and engage in problem solving approaches in a group setting. It was a derivative of the Stanford model of Chronic Disease Self-Management Program [39] but was based on rehabilitation principles aimed at maintaining physical function. The topics of the workshops included: changes in physical functioning associated with various stages of aging; the impact of various impairments; managing fatigue and relaxation techniques; the effect of different environments on functioning; how to assess changes in one’s function and mobility; how to promote and maintain function including the use of specific exercise and activities; approaches to safety, falls and injury prevention; the use of assistive devices; coping with the impact of pain on mobility and function; the effect of impaired balance on mobility and function; and the effect of medications on mobility and function. All participants who attended the RSM also received a Personal Health Record (PHR) booklet [40], which they learned to use as part of their self-management to record details of physical functioning and other aspects of their health.

**On-line self-monitoring of function**

The purpose of on-line self-monitoring of function was to increase individual patients’ knowledge of their functional status. Patients were able to complete self-assessments of physical functioning, the Physical Functioning Inventory (PFI) [21] and the Rapid Assessment of Physical Activity (RAPA), independently [26]. The assessments were available on MyOSCAR, a web-based system http://myoscar.org. It is a secure personal health record with a secure messaging system for patients to view the results of their online assessments and to communicate with the therapists. The patient was sent a message to their email address indicating when there were updates in the personal health record. The study coordinator summarized the results of the assessment in terms of age- and sex-based norms within the record and responded to the participants’ questions using the messaging system. The results of these assessments were also documented in the patient’s EMR. Patients were asked to complete the assessments online three times over the course of the study.
Organizational capacity building
The goal of the capacity building (CB) was to develop PCP skills and organizational supports to address physical functioning as a major health outcome. Monitoring of physical function was prioritized for persons with chronic illness by promoting increased collaboration between 20 PCPs and patients. The capacity building targeted the improvement of chronic illness care through increased understanding of self-management principles. Capacity building was undertaken through a series of activities including: (i) two workshops, (ii) a problem-based learning module, (iii) case reviews of selected patients participating in the study, and (iv) development of a flow sheet for monitoring changes in physical functioning (PF) to be used within the EMR.

Workshops
The first workshop introduced SM strategies for chronic disease and the second workshop introduced strategies to monitor PF and provide SM support. The PCPs were educated on how to use the flow sheet to monitor changes in physical functioning and how to integrate this within the patient’s assessment.

The problem based learning module
“Physical Functioning in Patients with Chronic Disease: the Sixth Vital Sign?” was developed for use in a problem based tutorial. The module was based on a patient case at three different stages of functional decline. The objectives of the module were to examine the impact of chronic disease on physical functioning, rapidly identify patients at high risk of developing a disability, monitor the patient’s functional status over time, provide self-management support in the area of physical functioning, explore the role of various team members in this area, and participate in proactive, preventative strategies with an interdisciplinary team.

Case reviews
The study coordinator met with PCPs to review the patients’ progress in the study activities and to involve them in the process of supporting the patient in the SM of physical functioning. Information about the patients’ level of physical functioning, self-identified goals and problem lists, self-management skills and performance test results were discussed with the PCP. The study coordinator discussed what actions the PCP would take as a result of this information and what the coordinator could do to support the patient.

Development of a self-management flow sheet
To encourage the monitoring of physical functioning, a decision support and tracking flow sheet that focused on physical functioning was developed for use within the EMR by the authors. Level of physical activity, falls history and Timed Up & Go (TUG) results were included [31]. Information boxes turn red or yellow if the data entered does not meet recommended parameters. To provide decision support, a colour coding system cued the PCP to possible actions if the patient’s results were outside the parameters. A list of actions was provided for PCPs to consider including various referral sources. The PCP is notified of areas that require reassessment within 6 months.

Analysis
T-tests and chi square tests were used for continuous and categorical variables respectively in the baseline comparison between groups. Paired t-tests were used to assess whether there was a significant difference between baseline and follow-up scores for the intervention group on the Self Efficacy for Chronic Disease Scale, Health Care Utilization and Self-Rated Health measures. Paired t-tests were used to assess whether there was a significant difference between baseline and follow-up scores on the PACIC and the PCRS. One-way analysis of variance (α = .05) was used to test for differences between groups in each of the following variables: RAPA, PACIC, Self Efficacy for Chronic Disease Scales, Self-Rated Health. We used a one-way multivariate analysis of covariance (MANOVA) to compare the group effect for the performance measures which included: the two-minute walk test, grip strength, eight foot walk and balance using age as a covariate. We also used a MANOVA to examine group effect for the measure of physical activity, the RAPA and the Physical function Inventory including age as a covariate. STATA/IC 10.1 for Macintosh was used for the analysis [41].

Results
A total of 200 participants from SFHC were eligible; 65 people (32.5%) agreed to participate in the study and 55 patients completed the final assessment in both the intervention and control group (Figure 2). There were no differences between the two groups at baseline on the variables assessed (Table 1).

At follow-up, there were no significant difference between groups on the PFI overall score or subscale scores (Table 2). The ANOVA showed a significant difference between the groups in the level of physical activity as measured by the RAPA. The MANOVA showed a significant difference between the groups for physical function (PFI) and physical activity (RAPA) (Λ = 0.88, F = (2.86) = 5.97, p = 0.004). There was no significant difference between groups on Self-Rated Health or Self-Efficacy for Chronic Disease Scale scores (Table 2).

Forty six participants completed the online self-monitoring of physical function independently at baseline and 33 at follow-up using MyOSCAR. Seven more patients completed the follow-up with support of the study coordinator. The average amount of time spent assisting patients who did not use online methods...
independently was approximately 1.5 h per patient over the course of the study. Although 48 out of 60 (80%) of the intervention group participants had access to an email address and were accustomed to using online services, several of these participants required support. There was a significant difference between the groups in favour of the intervention group in grip strength (both dominant and non-dominant hands) (Table 3). The MANOVA also showed a significant between group effect when the physical performance measures were examined collectively as the dependent variable ($\Lambda = 0.80$, $F = (6,93) = 3.68$, $p = 0.0025$). In the one way ANOVA grip strength was the only outcome that showed a between group difference.

There was no significant difference between the groups for Health Care Utilization (visits to the physician, emergency room visits and hospitalizations) (Table 4).

There were no significant differences or within group change in the capacity building outcomes or the Primary Care Resources and Supports scores (Table 5). The Patient Assessment of Chronic Illness (PACIC) also showed no significant differences for between group change as a result of CB initiatives (Table 6). The PACIC was used to measure changes in practice from the patient’s perspective as a result of capacity building initiatives.

Self-Rated Health and Self-Efficacy for Chronic Disease were administered to the intervention group before and after the Rehabilitation Self-Management Program. There was a significant within group difference for the intervention group in Self-Efficacy for Chronic Disease Scale scores. These scores increased by 0.54 points after RSMP, $p = 0.01$ (Table 7). In the RSMP, the mean number of sessions attended was 3.45; 75% of participants attended $\geq 3/5$ sessions.

**Results of focus group**

Nine PCPs from a variety of disciplines attended the focus group. Several PCPs stated they became more intentional in addressing the patients’ physical functioning as a result of the CB workshops. They reported an increase in the level of the patients’ physical activity, improved goal setting and problem solving, and greater focus in their interactions with PCPs. PCPs felt their own efforts at integrating the self-management aspects of care could be improved. The main barrier
identified by the PCPs was the lack of time to address the multiple concerns that patients with chronic conditions face and SM goals were sometimes given a lower priority.

Results of capacity building process measures
Forty-eight PCPs from SFHC participated in the capacity building initiatives. Thirty three PCPs from 6 different disciplines attended the workshops (Table 8) and 16

Table 1 Baseline Characteristics for Intervention and Control Group

|                          | Intervention group (n = 60) | Control group (n = 59) | t/X² | P       |
|--------------------------|----------------------------|------------------------|------|---------|
| Age, years mean (SD)     | 62.78 (11.04)              | 62.62 (10.29)          | t = 0.08 | P = 0.93 |
| Gender, mean (SD)        |                            |                        |      |         |
| Female                   | 42 (70%)                   | 43 (72.9%)             | X² = 0.16 | P = 0.69 |
| Male                     | 18                         | 17                     |      |         |
| Level of function, frequency (%) |                     |                        |      |         |
| Level 1                  | 22 (36.67%)                | 22 (33.90%)            | X² = 0.0   | P = 0.10 |
| Level 2                  | 20 (33.33%)                | 20 (36.21%)            |      |         |
| Level 3                  | 18 (30%)                   | 16 (27.12%)            |      |         |
| Use of gait aid, frequency (%) |                     |                        |      |         |
| No Gait aid              | 55 (91.67%)                | 51 (87.93%)            |      | P = 0.71** |
| Cane                     | 4 (6.67%)                  | 5 (8.62%)              |      |         |
| Walker                   | 1 (1.67%)                  | 2 (3.45%)              |      |         |
| Number of chronic conditions, frequency (%) |                     |                        |      |         |
| 1                        | 21 (35%)                   | 14 (23.72%)            | X² = 6.22 | P = 0.18 |
| 2                        | 264 (43%)                  | 22 (37.29%)            |      |         |
| 3                        | 9 (15%)                    | 18 (30.51%)            |      |         |
| 4                        | 3 (5%)                     | 5 (8.47%)              |      |         |
| 5                        | 1 (1.66%)                  | 0                      |      |         |
| Self-Rated Health, mean (SD) (1: Excellent - 5: Poor) |                     |                        |      |         |
| 2.94 (0.74)              | 3.05 (1.13)                | t = -0.60              | P = 0.55 |
| Self-Efficacy for Chronic Disease Scale, mean (SD) (max score = 60) |                     |                        |      |         |
| 6.92 (1.84)              | 6.83 (2.24)                | t = 0.22               | P = 0.83 |
| Physical Activity Level (1: Sedentary - 7: Active), mean (SD) |                     |                        |      |         |
| 4.58 (1.61)              | 4.32 (1.79)                | t = 0.84               | P = 0.40 |

*Chi-square tests and t-test were used for categorical and continuous variables respectively
**Fishers exact test

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Table 2 Results of Self-Report Measures by Group: Physical Functioning Inventory, Rapid Assessment of Physical Activity, Self-Rated Health and Self-Efficacy for Chronic Disease

|                          | Intervention | Control | F    | P    |
|--------------------------|--------------|---------|------|------|
| Physical Functioning Inventory Overall Score | 81.85 (19.74) | 85.89 (17.13) | 79.39 (20.43) | 83.07 (18.36) | 0.64 | 0.42 |
| ADL Subscale             | 21.76 (4.88) | 22.68 (4.07) | 20.97 (5.01) | 21.96 (4.30) | 0.77 | 0.38 |
| IADL Subscale            | 22.53 (3.60) | 22.74 (3.43) | 21.90 (4.44) | 23.19 (2.57) | 0.42 | 0.52 |
| Mobility Subscale        | 20.28 (5.44) | 21.08 (5.21) | 19.57 (5.5)  | 19.87 (5.50) | 1.28 | 0.26 |
| Moderate Activity Subscale| 17.45 (9.46) | 20.2 (7.52)  | 15.61 (10.43) | 17.5 (10.06) | 2.33 | 0.13 |
| Level of Physical Activity - RAPA (1: Sedentary - 7: Active) | 4.58 (1.61) | 5.09 (1.47)  | 4.32 (1.79)  | 4.05 (1.58)  | 12.95 | 0.0005 |
| Self-Rated Health (1: Excellent - 5: Poor) | 2.94 (0.74) | 2.87 (0.85)  | 3.05 (1.13)  | 3.03 (0.94)  | 0.83  | 0.36 |
| Self-Efficacy for Chronic Disease | 6.92 (1.84) | 7.44 (1.66)  | 6.83 (2.24)  | 7.04 (2.33)  | 1.60  | 0.31 |

The Self-Efficacy for Chronic Disease Scale score ranges from 0-10, with lower scores indicating lower confidence at completing tasks or managing symptoms related the chronic illness. Self-Rated Health is rated on a scale from 1-excellent, to 5-poor
individuals, which included a group of residents and a group of allied health professionals attended the PBLM. The residents noted that specific tools and handouts within the module would assist them in applying the information in clinical practice.

Results of case review process
Case reviews about patients’ physical functioning during the study were undertaken with 8 PCPs. The PCPs discussed whether they would take a specific action as a result of the case review (e.g. referral to a community program or a CCAC day program, a Chronic Pain Program or a dietitian) as well as plans for ongoing support for self-management activities with the participant. The PT met with PCPs to review a total of 21 patient cases. Each case took approximately 10 min to complete.

Discussion
The purpose of this study was to determine whether a population-based rehabilitation intervention for persons with chronic conditions in a primary care setting resulted in decreased functional decline and improved self-management compared to case-matched controls. The goal of the capacity building initiatives was to increase PCP collaboration in chronic disease management. Self-efficacy for managing chronic conditions improved as seen by a significant increase in self-efficacy scores immediately after the RSMP. There was greater improvement in the intervention group than in the control group in level of physical activity as measured by the RAPA (p = 0.0005) and bilateral grip strength (p = 0.0008). The current intervention is a feasible method to increase self-efficacy for physical activity through RSMP workshops together with reinforcement from members of the primary care team. This approach has the potential to impact on physical functioning. Counseling by primary care physicians over 6 weeks has been reported to increase levels of physical activity; however this effect was not sustained at 8 months [42]. Sustained improvement in physical activity levels requires more intensive patient contact [43]. Physically active individuals with and without chronic conditions have been reported to be less likely to develop a disability when compared to less active individuals [44]. This population-based approach with a restricted time frame resulted in elevated levels of physical activity.

The improvements in grip strength may be the result of increased levels of physical activity. Grip strength is a measure that reflects generalized muscle strength and acts as a predictor of disability [38]. Older men and women with chronic illness have shown 30% improvement in upper and lower extremity strength as a result of stretching, strengthening and aerobic exercise [45].

The capacity building initiatives were well received with participation from 48 PCPs from 5 disciplines. There was no significant change in the amount or type of information relating to physical functioning that was documented in patients’ EMRs by PCPs during the study period. The impact of PCP self-management support on patient care as measured by the PCRS and PACIC did not change in the 6 month time period but focus group results indicated that PCPs recognized the

Table 4 Results of Health Care Utilization by Group: Physician Visits, Emergency Room Visits, Hospitalisations

| Outcome Measure | Intervention | Control | F | P |
|-----------------|--------------|---------|---|---|
| Number of physician visits | 4.88 (7.29) | 4.24 (6.073) | 0.05 | 0.82 |
| Number of emergency room visits | 0.19 (0.99) | 0.22 (0.57) | 0.02 | 0.89 |
| Number of hospitalizations | 0.19 (0.86) | 0.05 (0.23) | 0.11 | 0.73 |
| Number of total nights spent in hospital | 0.28 (0.17) | 0.2 (1.11) | 0.06 | 0.81 |

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importance of these concepts. Time availability was identified as a significant barrier to collaborating around the maintenance of physical functioning; however PCPs supported the integration of rehabilitation professionals into the FHT to address physical functioning.

The feasibility of the intervention was demonstrated throughout the study. In the RSMP, the mean number of sessions attended was 3.45, while 75% of participants attended ≥3/5 sessions. The attrition rate of 25% in this study was comparable to 28%-50% attrition rates reported in other studies of CDSMP [14,46,47]. These data suggest that retention of participants in this type of program is an ongoing challenge. We tracked the feasibility of using MyOSCAR to administer outcome measures for the self-monitoring of physical function. The average amount of time required over the 6 months to support patients with online questionnaires was 1.5 h/patient. Additional support maybe required for persons who do not access the internet or who do not operate a computer regularly. Healthcare reform includes the use of technology by both health care professionals and patients [13], however it is likely there will always be a gap between the ability of the technology and the ability of some patients to engage in the monitoring or usage. Further understanding of the patients’ difficulties in interacting with the technology to maximize this self-monitoring approach is required. Recent research has shown that increasing adherence to monitoring health indices such as blood pressure and blood glucose does not necessarily improve health outcomes [48]. It is possible that the technology associated with the intervention might have deterred participants from completing assessments, however similar cohorts in the future will be more experienced with incorporating computer assisted technology into health related management. Future work needs to evaluate whether an increase in adherence to monitoring physical function is accompanied by a concomitant increase in performance, health status or quality of life outcome.

The capacity building strategies to improve self-management support of physical functioning were feasible in the FHT. The PCPs indicated that the workshops increased their awareness of functional status as an important health outcome. Incentives such as Continuing Medical Education (CME) credits for the physicians linked to workshop attendance may have increased participation and a higher rate of participation would have been desirable. Involving PCPs in any educational activity during clinical hours is always going to be a challenge. Multiple strategies and opportunities for participation, as implemented in this study and recommended in knowledge translation approaches, will likely maximize involvement [49]. PCPs reported a lack of time for management and follow-up with patients with complex chronic issues as a barrier to implementing principles discussed in CB initiatives. They did report patients’ behavioral changes such as increased levels of physical activity observed in outcome measures such as the RAPA. Improved self-efficacy and increased initiative taken by patients was also noted in those self-managing their conditions. There was no change in health care utilization.

A population-based approach to rehabilitation for persons with chronic illness who had varying levels of physical functioning was successfully implemented in a compressed time period. The current intervention was designed to take a comprehensive approach to improve chronic disease management by addressing the characteristics of the patients as well as the health care environment in which patients receive services. The use of multiple strategies to address barriers to change in PCP behaviour is a strength of this study and over an extended period is likely to effect organizational change [50].

| Table 6 Results of Patient Assessment of Chronic Illness Care (PACIC) with Subscale Scores by Group |
|-------------------------------------------------------------|-----------------|------------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| Outcome Measure                                             | Intervention    | Follow-up        | Control         | Follow-up       | F    | p    |
| PACIC overall score                                         | 2.57 (0.17)     | 2.9 (1.11)       | 2.52 (0.11)     | 2.7 (0.91)      | 0.91 | 0.34 |
| Patient activation subscale                                 | 2.87 (1.42)     | 3.36 (1.33)      | 3.11 (1.34)     | 3.29 (1.22)     | 0.07 | 0.79 |
| Delivery system design/decision support subscale            | 3.24 (0.15)     | 3.24 (1.10)      | 2.98 (0.14)     | 2.98 (1.03)     | 1.57 | 0.21 |
| Goal setting subscale                                       | 2.36 (0.19)     | 2.67 (1.31)      | 2.31 (0.13)     | 2.56 (1.06)     | 0.22 | 0.64 |
| Problem solving/contextual counseling subscale              | 3.02 (0.20)     | 3.14 (1.24)      | 3.07 (0.15)     | 3.13 (1.22)     | 0.00 | 0.98 |
| Follow-up/coordination subscale                             | 2.44 (0.16)     | 2.44 (1.19)      | 2.13 (0.31)     | 2.13 (0.96)     | 2.06 | 0.15 |

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This approach uses limited resources to benefit a larger number of patients than would normally be possible with care by a single provider. A group-based intervention is a feasible way to address the demand for rehabilitation professionals and to provide patients with chronic conditions access to services that may not be available to them [51]. This study generated modest results which would be consistent with other rehabilitation interventions that typically produce small to moderate effect sizes [52]. However, this trend is offset by small associated risks, an important consideration in a population with multiple morbidity where many of the interventions have their own associated risks and patients and physicians are required to make judgments about which risks they would prefer to undertake [53,54]. To date, PTs have not been integrated into FHT practices in Ontario [55] while funding for OTs is being implemented by the Ministry of Health and Long Term Care on a case by case basis [55]. With a shift in chronic disease management from disease recovery to functional health, the expertise of rehabilitation professionals in considering the functional consequences of disease can make a valuable contribution to primary care [56].

The principal limitations of this study were the short duration of the study period and the non-randomized design. Six months was likely insufficient time to result in a change in self-rated health, health care utilization and physical functioning. Changes in organizational structure and practices can also take much longer than 6 months to occur [57]. Quality of life maybe a more appropriate outcome of interest than health status since it may take years to change health status through lifestyle interventions. However, it is possible that patients could experience an important improvement in their quality of life as a result of the intervention. Physical functioning is a core component of a person’s health status and quality of life. Persons with chronic disease will experience episodic illness which along with the symptoms of a chronic disease is likely to negatively influence their physical functioning. It is imperative that physical functioning is optimized and that the physiological reserve required for daily activities is maintained.

**Conclusion**

It is feasible to monitor physical functioning as a health outcome for persons with chronic illness in primary care. The timeline for this study was not sufficient to show an increase in the capacity within the team; however there were some differences in patient outcomes. The short timeline was likely not sufficient to build the capacity required to support this approach. This intervention needs to be tested by a large scale randomized controlled trial.

**Abbreviations**

CCM: Chronic Care Model; ECCM: Expanded Chronic Care Model; FHT: Family Health Team; PCP: Patient Care Provider; PCRS: The Assessment of Primary Care Patient Care Resources and Supports; PACIC: Patient Assessment of Chronic Illness Care; SFHC: Stonechurch Family Health Centre; MFP: McMaster Family Practice; PFI: The Physical Functioning Inventory; ADL: Activities of Daily Living; IADL: Instrumental Activities of Daily Living; RAPA: Rapid Assessment of Physical Activity; TUG: Timed Up and Go; SECDS: The Self Efficacy for Chronic Disease Scale; EMR: Electronic Medical Record; PT: Physiotherapy; OT: Occupational Therapy; RSMR: Rehabilitation Self-Management Program; PCP: Patient Care Provider.

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Authors’ contributions
JR, LL, DC and LMC contributed to the design of the study. JR and LL were responsible for obtaining funding for the study. AO and SW participated in the acquisition of data. JR, LL, AO contributed to the analysis and interpretation of the data. DO, AM, LMC, DP, SK provided administrative, technical or material support. JR drafted the manuscript and all authors critically reviewed the manuscript for important intellectual content. All authors approved the final version of the manuscript submitted for publication.

Competing interests
The authors declare that they have no competing interests.

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