Online peer support interventions for chronic conditions: a scoping review protocol

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ABSTRACT

Introduction Peer support is receiving increasing attention as both an effective and cost-effective intervention method to support the self-management of chronic health conditions. Given that an increasing proportion of Canadians have internet access and the increasing implementation of web-based interventions, online peer support interventions are a promising option to address the burden of chronic diseases. Thus, the specific research question of this scoping review is the following: What is known from the existing literature about the key characteristics of online peer support interventions for adults with chronic conditions?

Methods and analysis We will use the methodological frameworks used by Arksey and O’Malley as well as Levac and colleagues for the current scoping review. To be eligible for inclusion, studies must report on adults (≥18 years of age) with one of the Public Health Agency of Canada chronic conditions or HIV/AIDS. We will limit our review to peer support interventions delivered through online formats. All study designs will be included. Only studies published from 2012 onwards will be included to ensure relevance to the current healthcare context and feasibility. Furthermore, only English language studies will be included. Studies will be identified by searching a variety of databases. Two reviewers will independently screen the titles and abstracts identified by the literature search for inclusion (ie, level 1 screening), the full text articles (ie, level 2 screening) and then perform data abstraction. Abstracted data will include study characteristics, participant population, key characteristics of the intervention and outcomes collected.

Dissemination This review will identify the key features of online peer support interventions and could assist in the future development of other online peer support programmes so that effective and sustainable programmes can be developed.

INTRODUCTION

More than 20% Canadian adults live with one of the following chronic diseases: cardiovascular diseases, cancer, chronic respiratory diseases or diabetes. Since 2000, the prevalence rates for cardiovascular diseases and chronic respiratory diseases have increased.

Treatment of chronic diseases consumes 67% of all direct healthcare costs, and cost the Canadian economy $190 billion annually, with $68 billion related to treatment costs and the remaining costs related to lost productivity. In the USA, approximately 25% of the population has multiple chronic conditions; among Americans aged 65 years and older, approximately three in four have multiple chronic conditions. In Australia, more than 7 million people have at least one chronic condition. Thus, it is imperative to develop and implement effective interventions to manage these chronic conditions.

As a result of this increasing burden, particularly in health services and related costs, individuals with chronic conditions need assistance in learning and maintaining self-care behaviours that support healthy living, referred to as ‘chronic disease self-management’. Chronic condition self-management refers to a person’s ability to manage the symptoms, treatment, and physical, psychosocial and lifestyle changes that are associated with living with a chronic condition.

A promising intervention is peer support which is receiving increasing attention as both an effective and cost-effective intervention method to support the self-management of chronic health conditions.
There is some emerging evidence demonstrating the effectiveness of peer support for individuals with chronic conditions to self-manage their conditions. Peer support in the context of chronic disease management refers to ‘support for a person with a chronic condition from someone with the same condition or similar circumstances’. Individuals who provide peer support offer three types of support based on experiential knowledge: emotional, informational and appraisal. Emotional support involves caring, empathy and encouragement; informational support involves advice, suggestions with alternative actions and factual feedback relevant to a particular topic; and appraisal support involves affirming feelings, thoughts and behaviours and thus is motivational, encouraging the individual to continue with problem-solving attempts despite setbacks.

There are seven predominant types of peer support models in chronic condition management: professional-led groups that encourage peer interactions, structured peer-led self-management training, peer coaches, community health workers, support groups, telephone-based peer support, and web-based and email-based programmes. At the same time, it is also important to recognise that there are varied models of peer support, and that treating them as if they are fully defined and highly distinct may lead to overlooking important commonalities. The current review will focus on peer coaches and web-based and email-based programmes. Given that the majority of households have the internet (eg, 80% in Canada), and the increasing use of web-based interventions, there is an increasing need to determine the characteristics of online peer support interventions. The specific research question of this scoping review is the following: What is known about the effectiveness of peer support interventions for adults with chronic conditions?

METHODS AND ANALYSIS

We will use the methodological frameworks proposed by Arksey and O’Malley as well as Levac and colleagues for the current scoping review. These frameworks outline six different stages involved in a scoping review: (1) identifying the research question; (2) identifying relevant studies; (3) selecting studies; (4) charting the data; (5) collating, summarising and reporting the results; and, (6) consulting with relevant stakeholders. The research team has expertise in peer support across chronic conditions (SEPM, JS, SA, SNS, MLAN, SJTG, SJ), online interventions (SA, SEPM, SJ) and knowledge synthesis methods (SEPM, LP, JRT, MLAN, SJTG, SJ). Although traditionally applied to systematic review protocols, the Preferred Reporting Items for Systematic Reviews and Meta-analysis for Protocols was used to draft this protocol.

Eligibility criteria

For the purpose of this review, we will include chronic diseases identified by the Public Health Agency of Canada (PHAC), including cancer, heart disease (cardiovascular disease), hypertension, stroke, chronic respiratory diseases (asthma, chronic obstructive pulmonary disease, sleep apnoea), diabetes, inflammatory bowel diseases (Crohn’s disease, ulcerative colitis) multiple sclerosis, neurological conditions (eg, Alzheimer’s disease and other dementias, cerebral palsy, epilepsy, multiple sclerosis, Parkinson’s disease/parkinsonism, traumatic brain injury and traumatic spinal cord injury (SCI), arthritis and osteoporosis. Mental illness was excluded from the list given that peer support interventions for this group may have particularly unique features not generalisable to other patient populations with chronic disease. Similarly, in a systematic review on the effectiveness of quality improvement strategies (ie, including patient education and promotion of self-management) for coordination of care to reduce use of healthcare services, Tricco and colleagues determined that these quality improvement strategies reduced hospital admissions among patients with chronic conditions other than mental illness, indicating that different approaches are needed for mental health. To this list, we have also included HIV/AIDS (which, from our preliminary research, has a significant amount of literature on peer support interventions, and is increasingly being viewed as a chronic condition). Studies including individuals with comorbidities (including mental illness) will be accepted. Thus, to be eligible for inclusion, the studies must report on adults (≥18 years of age) with one of these PHAC chronic conditions (excluding mental illness) or HIV/AIDS. We will limit our review to peer support interventions delivered through online formats. Peer support in the context of chronic disease management is operationalised as ‘support for a person with a chronic condition from someone with the same condition or similar circumstances’. This type of support could be emotional, informational and/or appraisal. Examples of online peer interventions could include Skype-based discussions, social media peer interactions or text messages from a peer(s). Therefore, interventions that are professional-led groups that encourage face-to-face peer interactions, involve community health workers who are not peers (ie, healthcare professionals), support groups and provide telephone-based peer support will be excluded. All study designs will be included (eg, observational studies, randomised controlled trials and qualitative studies). Only studies published from January 2012 to April 2017 will be included. We have included studies from the last 5 years to ensure relevance to the current healthcare context, in technology and feasibility. Furthermore, only English language studies will be included, which may result in a predisposition in results towards English language speaking countries.
Search strategy and information sources

Literature search strategies will be developed using medical subject headings and text words related to chronic conditions and peer support interventions. Studies will be identified by searching Medline (OVID interface), CINAHL (EBSCO interface), EMBASE (OVID interface), PsycInfo (OVID interface), and Cochrane Central Register Controlled Trials (Cochrane Library) and PEDro (physiotherapy evidence database). The search strategy for Medline can be found in the online supplementary file 1. A hand search of the reference lists from reviews and selected articles will be made to ensure literature saturation. Finally, experts in the field of peer support will be contacted and consulted in order to ensure that all relevant data are obtained, including members of the research team. An information specialist (LP) who is expert in systematic and scoping reviews will conduct all of the literature searches.

Study selection

To promote the reliability of screening by the two reviewers, a pilot test of the level 1 screening form based on the criteria outlined above will be conducted on a random sample of approximately 100 articles. The κ statistic will then be calculated to determine the inter-rater reliability for study inclusion.21 If low agreement is observed, the inclusion and exclusion criteria will be simplified to increase the consistent application of the selection criteria. Two reviewers will independently screen the titles and abstracts identified by the literature search for inclusion using the screening form (ie, level 1 screening). The full text of potentially relevant articles will then be collected and screened to determine final inclusion (ie, level 2 screening). A pilot test of the level 2 screening form will be performed on approximately 1% of the articles and the inter-rater reliability for study inclusion will also be calculated.23 A third reviewer who is knowledgeable in the research area will be available to resolve conflicts, if necessary.

Data items and data collection process

Abstracted data will include study characteristics (eg, year of publication, country of study), participant population (eg, chronic disease condition, income, ethnicity, age, sex, education), key characteristics of the intervention (eg, duration, frequency, delivery setting, type of intervention, type of support provided—emotional, informational and appraisal, underlying theories for the intervention/behaviour change techniques/working mechanisms, context, etc) and type of outcome collected (if applicable) as well as results (ie, efficacy or effectiveness). Outcomes reported may include, but are not limited to, self-efficacy, depression and quality of life. In fact, we aim to identify the variety of reported impacts. Additional categories may be identified through the completion of the search and through discussions with the research team and key stakeholders. A data abstraction form will also be pilot tested and modified if poor agreement is observed. For example, any wording on the form that may be related to poor agreement will be reviewed and improved. Two reviewers will independently abstract all of the data and a discussion or the involvement of a third reviewer will resolve disagreements. Study quality will not be evaluated as the purpose of a scoping review is to identify gaps in the literature and future areas for a systematic review.15 16 DistillerSR will be used to manage the records and data throughout the review.

Synthesis

The data from this scoping review will be summarised quantitatively using numerical counts and qualitatively using thematic analysis and will be grouped by chronic condition type (eg, what kind of underlying theories are found in online peer support interventions in cancer?). These data will be coded/analysed manually. The results of this review will determine the key characteristics (eg, duration, frequency, delivery setting, type of intervention, type of support provided—emotional, informational and appraisal, underlying theories for the intervention/behaviour change techniques/working mechanisms, context, etc) of online peer support interventions for adults with chronic conditions. This scoping review will identify gaps in the literature as well as future areas for study either via implementation studies, consensus meeting or systematic review.

DISSEMINATION

Knowledge translation activities will occur at the beginning of the review and continue throughout with dissemination of the research question to key stakeholders such as the Ontario Ministry of Health and Long-term Care, the Ontario Peer Development Initiative, March of Dimes Canada, SCI Canada, Rick Hansen Institute and the Ontario Neurotrauma Foundation. End-of-grant knowledge translation could also take place through these organisations and their outlets (eg, print and online newsletters) as well as through traditional knowledge translation mechanisms (eg, peer-reviewed journals and conference). For example, the results of the scoping review will be presented at meetings locally, nationally and internationally (eg, National Conference on Peer Support, American Medical Informatics Association) and published in a peer-reviewed journal so that results are available to the appropriate academic and clinical audiences. Finally, partnerships with local clinical programmes and/or research initiatives will be made so that the results are disseminated in a timely and effective manner.

There will be expected limitations and strengths to this review. For example, this review will not include (primary) mental health conditions and a variety of other disabilities. A number of problem-solving approaches for mental health conditions are emerging as quite effective when administered by non-professionals, including in low-resource settings.22–24 The possibility of implementing these online would represent exciting advances
in the field of peer support, and thus, a future, separate systematic/scoping review on online peer support interventions for mental health conditions should be considered. Furthermore, this review will be limited to English language studies only. Lastly, our review will be limited to the published research literature. We acknowledge that we will be excluding reports on available, relevant programmes but not published in these arenas. However, the currently proposed scoping review has a number of strengths— it is guided by known methodological frameworks and all phases will be conducted in duplicate. This review will contribute to critical and emerging perspectives on peer support. For example, the results of the review will identify the key features of online peer support interventions, and in doing so, assist in the future development of other online peer support programmes (ie, so that effective and sustainable programmes can be developed). Similarly, the current review will provide considerations for programmes that are not yet in the online format so that they can be modified with features that are unique to the online environment.

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Contributors SEPM conceived of the scoping review, together with JS, LP and S.J. SEPM wrote the first draft of the protocol and is the guarantor of the review. SEPM, JS and LP were involved in the preliminary literature review. LP conducted the literature search and provided methodological expertise (knowledge synthesis). JS, SA, SNS, JRT, SJTG and MLAN provided critical content expertise on peer support that was integrated into the current protocol. SEPM, JS, LP, SA, SNS, JRT, MLAN, SJTG, SH and S.J were involved in editing and revising the protocol for important intellectual content, and approved the final version of the protocol.

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Competing interests None declared.

Patient consent Detail has been removed from these case descriptions to ensure anonymity. The editors and reviewers have seen the detailed information available and are satisfied that the information backs up the case the authors are making.

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REFERENCES
