A Research Protocol for Implementation and Evaluation of a Patient-Focused eHealth Intervention for Chronic Kidney Disease

Maoliosa Donald1 · Heather Beanlands2 · Sharon Straus3 · Lori Harwood4 · Gwen Herrington5 · Blair Waldvogel5 · Maria Delgado5 · Dwight Sparkes5 · Paul Watson5 · Meghan Elliott1 · Kerry McBrien6 · Aminu Bello7 · Brenda Hemmelgarn7

Received: 19 November 2021 / Accepted: 13 January 2022 / Published online: 30 January 2022
© The Author(s) 2022

Abstract
Self-management in chronic kidney disease (CKD) can slow disease progression; however, there are few tools available to support patients with early CKD. My Kidneys My Health is a patient-focused electronic health (eHealth) self-management tool developed by patients and caregivers. This study will investigate the implementation of My Kidneys My Health across primary care and general nephrology clinics. The study aims to: (1) identify and address barriers and facilitators that may impact implementation and sustainability of the website into routine clinical care; (2) evaluate implementation quality to inform spread and scale-up. We will conduct a multi-stage approach using qualitative methods, guided by the Quality Implementation Framework and using a qualitative content analysis approach. First, we will identify perceived barriers and facilitators to implementation and considerations for sustainability through interviews with clinicians, based on the Readiness Thinking Tool and the Long Term Success Tool. Analysis will be guided by the Consolidated Framework for Implementation Research and the Theoretical Domains Framework. Appropriate implementation strategies will be identified using the Expert Recommendations for Implementing Change compilation, and implementation plans will be developed based on Proctor’s recommendations and the Action, Actor, Context, Target, Time framework. Finally, we will explore implementation quality guided by the RE-AIM framework. There is limited literature describing systematic approaches to implementing and sustaining patient-focused self-management tools into clinical care, in addition to employing tailored implementation strategies to promote adoption and sustainability. We aim to generate insights on how My Kidneys My Health can be integrated into clinical care and how to sustain use of patient-centric eHealth tools in clinical settings on a larger scale.

Keywords Implementation science · eHealth · Chronic kidney disease · Quality implementation framework · Person-centered care

Introduction
Chronic kidney disease is a progressive condition, often characterized by multi-morbidity that impacts treatment and management. To slow disease progression, persons with chronic kidney disease are required to balance medical management of their disease and other chronic conditions (e.g., diabetes, cardiovascular disease, depression) with the demands of daily life (Tonelli et al., 2015). Self-management is a promising approach to slow disease progression and improve health outcomes important to patients. It aims to facilitate an individual’s ability to make positive changes by focusing on illness needs, activating resources, and developing coping skills to live with...
the biopsychosocial demands of the disease and related comorbidities (Richard & Shea, 2011).

International chronic kidney disease research priority setting activities have identified a need to develop optimal strategies to support patients to self-manage their chronic kidney disease and related comorbidities (Hemmelgarn et al., 2016; National Kidney Foundation, 2017; Tong et al., 2015). Our research team (i.e., patient partners [patients and caregivers], clinicians, researchers, and policy makers), supported by a pan-Canadian, patient-oriented research (POR) initiative (Can-SOLVE CKD Network), is investigating opportunities to enhance chronic kidney disease self-management support for patients (Levin et al., 2018). We are conducting a multi-phased project examining strategies and tools for chronic kidney disease self-management most valued by patients and those who care for them. Important to these individuals are pertinent, tailored, timely information and resources to live well with chronic kidney disease, especially for those in early stages of the disease process (i.e., not on dialysis) or recently diagnosed (Donald et al., 2019a, 2019b). An electronic health (eHealth) platform has potential to address these priorities.

While there are numerous eHealth self-management interventions available for patients with chronic kidney disease, they typically focus on later stages of chronic kidney disease and lack key characteristics important in providing person-centered care (Donald et al., 2018; Smekal et al., 2019). First, they have been predominantly designed by health care professionals without patient input (Bonner et al., 2018; Donald et al., 2018; Shen et al., 2019). Second, only a few self-management interventions have been theory-informed for the planning, design, and evaluation of the intervention (Donald et al., 2018; Shen et al., 2019). Finally, there is sparse literature in regard to planning, measuring, and reporting implementation and sustainability efforts for chronic kidney disease self-management eHealth interventions (Ghimire et al., 2015; Shen et al., 2019).

To address these knowledge gaps, we undertook a theory-informed, person-centered approach to develop an eHealth tool, the My Kidneys My Health website, that is concordant with patients’ values, needs, and preferences for chronic kidney disease self-management support (Donald et al., 2021a). The focus of this paper is to describe the process of implementation and evaluation of the My Kidneys My Health website.

The key objectives are to identify and address provider and organizational barriers and facilitators that may impact implementation and sustainability of the My Kidneys My Health website into routine clinical care for individuals with chronic kidney disease; and explore implementation quality within primary care and general nephrology clinics to understand and inform spread and scale-up in these settings.

Design and Methods

We will conduct a multi-stage approach using qualitative methods to undertake the implementation (i.e., integration into practice) and evaluation of the My Kidneys My Health website guided by the quality implementation framework (QIF) (Meyers et al., 2012). Reporting of data will be guided by the Consolidated Criteria for Reporting Qualitative Research guidelines (Tong et al., 2007).

Website Overview

The evidence-informed My Kidneys My Health website (www.mykidneysmyhealth.com) was co-developed using a systematic, user-centered design process previously described (Donald et al., 2021a). The results of our exploratory sequential mixed-methods feasibility study demonstrated that the My Kidneys My Health website had high user acceptance, with a statistically significant increase in perceived self-efficacy sub-scale, gaining information (Donald et al., 2021b). It is an open-access, patient-centered web application, accessible by computer or mobile device. My Kidneys My Health was designed based on a conceptual framework identifying optimal design features for eHealth interventions (i.e., interactivity between social context and support, direct contact with the intervention, tailored information, and self-management) (Morrison et al., 2012), patient preferences (Donald et al., 2019b), and behavior change theory (Baey et al., 2019). The website supports self-management by “informing” (chronic kidney disease related information); “activating” (prompts/tools to encourage action to manage chronic kidney disease and enhance quality of life); and “collaborating” (links/tools that lead to interaction and engagement with health care professionals and peers) (Donald et al., 2021a; Gagliardi et al., 2015).

Specific elements of the website include interactive tools; personalized food list and health care professionals question list; and multimedia education and resources. While the website is targeted for patients, clinicians play a key role in offering and engaging patients through this resource. For example, the website can be used opportunistically, or at the point of care, when individuals are diagnosed with chronic kidney disease or share a concern related to supporting chronic kidney disease self-management.

Quality Implementation Framework

Research supports the use of implementation science in improving the quality of implementation and ultimately achieving desired patient outcomes (Durlak & DuPre, 2008). The Quality Implementation Framework (QIF) is an implementation process model providing mechanisms and...
strategies for successful implementation of an intervention (Blanchard et al., 2017). It is based on 25 implementation frameworks and consists of six themes with 14 steps, grouped into four phases that are practical and intuitive to our clinical context (see Fig. 1) (Meyers et al., 2012). The four phases entail site assessments, implementation, evaluation, and sharing lessons learned.

**System Engagement and Implementation Team**

Guided by the Knowledge-to-Action Framework, key stakeholders (i.e., patient partners, clinicians, policy makers, and organizations involved in the care of individuals with chronic kidney disease) were engaged in the co-design of the evidence-informed *My Kidneys My Health* website (Donald et al., 2021a). We will continue to work with these stakeholders to achieve quality implementation, applying the Interactive Systems Framework (ISF) to identify stakeholder roles and to situate the QIF within the clinical and research context. The ISF considers who is in the delivery system (clinicians in the primary care setting and general nephrology clinics), support system (implementation team), and synthesis and translation system (research team) (see Fig. 2) (Wandersman et al., 2012). The delivery system is responsible for carrying out the activities needed to integrate the website into clinical practice. The implementation team will oversee and support implementation efforts of the *My Kidneys My Health* website at the sites throughout all stages of implementation. It is recommended that members of the team have knowledge about the intervention, as well as implementation experience (Fixsen et al., 2007). Specifically, team members will help inform, prepare, and support the sites to effectively integrate the website into practice. The team members will include: an implementation lead with experience in the co-design of the website and knowledge in implementation science; a support coach (project coordinator) and patient partner with knowledge of the intervention components; two clinicians (a nephrologist and a primary care physician with experience in managing patients with chronic disease, in addition to understanding primary care organizational functioning); and a knowledge broker with skills in knowledge translation. Our research team is considered part of the synthesis and translation system, co-creating the *My Kidneys My Health* website.

**Setting**

For the purposes of this study, we plan to implement the *My Kidneys My Health* website in two distinct settings within the Canadian context: primary care and general nephrology. Care for patients with chronic kidney disease varies depending on etiology of chronic kidney disease, disease severity, and presence of comorbidities. For example, patients with early stage, stable, or slowly progressive chronic kidney disease with other chronic conditions are usually cared for by primary care physicians. Furthermore, primary care physicians play a key role in identification and management of patients with chronic kidney disease, in addition to coordinating care with specialists such as nephrologists (Levin et al., 2013).

Working with our current stakeholders and networks in Alberta (i.e., Alberta Kidney Care, Primary Care Networks, 3 months (July–September 2021) 9 months (October 2021–June 2022) 3 months (July–September 2022) 3 months (October–December 2022)

![Fig. 1 Quality Improvement Framework guiding implementation and sustainability of My Kidneys My Health website. Note AACTT action, actor, context, target, time, CFIR consolidated framework for implementation research, ERIC expert recommendations for implementing change, LTST long term success tool, RE-AIM reach, effectiveness, adoption, implementation, and maintenance, RTT readiness thinking tool, TDF theoretical domains framework](image)
Medicine Strategic Clinical Network, Primary Health Care Integration Network) we will identify six primary care clinics and two general nephrology clinics. The networks comprise patients, clinicians, researchers, and decision-makers who are knowledgeable about specific areas of health, with goals to achieve better outcomes and better value for Albertans (Noseworthy et al., 2015). We will approach sites that see patients with chronic kidney disease and associated comorbidities. We will strive for diversity with respect to geographic clinic location (i.e., rural/urban), services provided (i.e., team-based/individual PCP) and clinics that serve diverse populations (e.g., to gain a better understanding of how context affects implementation and sustainability).

**Proposed Phases and Processes**

The following activities will be completed during the four phases: site assessments, implementation, evaluation, and learning from the implementation experience (see Fig. 1). An overview of the theory-based methods supporting this work is provided (see Table 1).

**Phase 1: Site Assessments**

The aim of this phase is to identify perceived barriers and facilitators (determinants) to implementation (i.e., integration into practice by offering and/or engaging patients with the website), as well as considerations for sustainability of the My Kidneys My Health website for use within the primary care and general nephrology settings by assessing readiness and sustainability. Identified determinants will inform our selection of appropriate implementation strategies. The findings from this phase will be used to inform the implementation plan for the clinical sites.

Using a purposive sampling strategy at the eight clinics, we will invite staff (e.g., primary care physicians, nephrologists, allied health) who provide direct clinical care and education to patients with chronic kidney disease, as well as clinic administrators who oversee site operations with key decision-making roles as the potential participant group. There are no exclusion criteria. To obtain maximum variation in professional roles, genders, and practice settings we will set quotas for the minimum number of participants that should be represented in each of the specified groups. We anticipate approximately 25 to 30 interviews based on previous similar studies (Eldh et al., 2020; Greer et al., 2019).
Telephone interviews will be 30-min, and the semi-structured interview guide will include a series of questions that are based on the Readiness Thinking Tool (RTT) (Wandersman et al., 2012) and the Long Term Success Tool (LTST) (see Supplementary File 1—pre-implementation interview guide). The RTT consists of multiple constructs (e.g., motivation and capacity) that can be applied across various levels, including the individual, team, and organization (Wandersman et al., 2012). The LTST is relevant for the healthcare setting and consists of 12 key factors to identify risks and prompt actions to increase chances of sustainability over time (Lennox et al., 2017).

The interviews will be conducted in the summer and fall of 2021 by the implementation lead (MD) and/or implementation support team member, both with qualitative research experience. Participants will also complete a short, structured survey describing their role and practice arrangements (e.g., professional background, years of experience, clinical site structure, familiarity with self-management resources for chronic kidney disease, etc.) (see Supplementary File 2—demographic questionnaire). Interviews will be digitally recorded and uploaded to a secure server. A trained transcriptionist will provide a verbatim, deidentified transcript (i.e., removal of participant and clinic names) of the interview. A research staff member will review transcripts for accuracy and confidentiality before uploading to NVivo Version 12 (QRS International, Inc.) for storing and organizing the data.

The analysis will focus on understanding individual and system barriers and facilitators that are associated with implementing and sustaining the My Kidneys My Health website as a self-management support tool for patients, in primary care and general nephrology clinic settings. We will conduct a qualitative directed content analysis using a deductive coding approach to coding guided by two frameworks (Hsieh & Shannon, 2005). One, applying the Consolidated Framework for Implementation Research (CFIR) (Damschroder et al., 2009), comprised five major domains: the intervention, inner and outer setting in which it is implemented, the individuals involved in the implementation, and the process by which implementation is accomplished (Damschroder et al., 2009). Two, using the Theoretical Domains Framework (TDF), a comprehensive, theory-informed framework to identify individual determinants of behavior (Michie et al., 2014). The TDF can be used to assess implementation problems and support implementation planning (Atkins et al., 2017). Two research team members will independently familiarize themselves with the data, then review all transcripts line-by-line to identify sections of text that reflect the domains of the CFIR and TDF. After independently coding the first 3 interviews, the two members will meet and compare the labels they have applied and agree on the set of codes to apply to all subsequent transcripts. They will then chart the data by creating a matrix that presents codes categorized to individual, system, and intervention barriers and facilitators, along with illustrative quotations (Gale et al., 2013). The analysis will be iterative and as a final step the first author (MD) will review the coding schema with the original transcripts. For the interpretation of the data the following questions will be considered: (1) How do inner factors (e.g., organizational culture, leadership) influence implementation and sustainment?; (2) How do outer context factors (e.g., financial) influence implementation and sustainment?; (3) What are the perceptions
of clinicians and decision-makers to implementation and sustainment?; (4) To what extent do factors and perceptions vary by site (e.g., organization and geographic)?

To address barriers and enhance facilitators to implementation we will identify relevant implementation strategies (tools, activities, and/or actions) that will support adoption of the website. Acknowledging that different strategies can be used at multiple levels within the delivery system (i.e., clinic, team, individual health care professionals) we will refer to the Expert Recommendations for Implementing Change (ERIC) compilation (Powell et al., 2015). The ERIC taxonomy has the following categories: (1) engage consumers, (2) use evaluative and iterative strategies, (3) change infrastructure, (4) adapt and tailor to the context, (5) develop stakeholder interrelationships, (6) use financial strategies, (7) support clinicians, (8) provide interactive assistance, and (9) train and educate stakeholders with 73 discrete implementation strategies, which can be combined to form a multifaceted strategy (Waltz et al., 2015).

Phase 2: Implementation

The aim of this phase is to design a detailed implementation plan that includes describing the implementation strategies, operationalizing these strategies, and designating implementation supports needed to deliver the website.

We will describe and operationalize the implementation strategies based on Proctor’s recommendations for specifying and reporting implementation strategies (Proctor et al., 2013) and the Action, Actor, Context, Target, Time framework (Presseau et al., 2019). Figure 3 describes the implementation strategy, specifically who needs to do what differently, when, where, and for whom. We will also consider what types of implementation supports will be needed for the clinician/administrator to offer and engage patients with chronic kidney disease with the website. Based on the Evidence-Based System for Innovation Support framework for implementing innovations with quality, these supports can include tools, training, technical assistance, and quality assurance to address initial needs and problems that arise during the implementation period (Wandersman et al., 2012). For example, these supports may include check-in calls and site visits, or identifying a “site implementer” to provide ongoing support. In addition, we will develop a process to monitor and provide supportive feedback to sites based on their needs.

Phase 3: Evaluation

The aim of this phase is to explore implementation quality guided by the RE-AIM framework (reach, effectiveness, adoption, implementation, and maintenance) to understand the adoption, implementation, and sustained use of the My Kidneys My Health website. The RE-AIM framework is an evaluation framework that measures success of an intervention implemented within a given context (Glasgow
et al., 2019). For this component, we will focus on the following elements: reach, adoption, implementation, and maintenance.

We will conduct 30-min telephone interviews 9 months post-implementation using a purposive sampling strategy at the eight clinics, similar to pre-implementation. We will invite staff that both had an interest or lack thereof in the use of the *My Kidneys My Health* website. A semi-structured interview guide will include a series of questions that are based on the RE-AIM Qualitative Guide (see Supplementary File 3—post-implementation interview guide) (Hol-trop et al., 2018; RE-AIM, 2021). Reach will provide us with information about health care professionals’ level of interest in the intervention and in engaging patients to use the website (e.g., attitudes toward addressing chronic kidney disease self-management with patients). Adoption will be assessed by understanding how health care professionals engaged patients (e.g., what informal criteria they used for selecting which patients) and how they integrated the *My Kidneys My Health* website into their management for patients with chronic kidney disease. We will explore the implementation dimension by investigating to what extent the website is implemented as intended (e.g., consistency of promoting website; challenges to embedding into clinical care). We will investigate the dimension of maintenance by exploring the extent to which the website will be sustained in routine care going forward (e.g., intended sustained use). In addition, participants will complete a short, structured survey describing their role and practice arrangements (see Supplementary File 2).

Interview data will be analyzed using a deductive qualitative directed content analysis approach similar to the activities outlined in Phase 1. Data will be examined based on the CFIR and TDF domains to provide structure for the factors identified and to fully understand the underpinnings behind the results (Damschroder et al., 2009). Data that does not fall within the framework will be captured to ensure ideas are not being missed. Interpretation of the findings will focus on implementation successes, failures, and when, where, and what types of adaptations are needed to overcome future implementation issues and ensure sustainability.

**Phase 4: Learning from the Implementation Experience**

The aim of this phase is to share learnings from our research findings to broader communities (e.g., patients and families, clinicians, researchers, decision-makers) through public engagement forums, webinars, publications, and conference presentations with the intention to improve future implementation efforts of patient-focused eHealth tools in similar settings. In addition, we anticipate that we will build stronger inter-organizational relationships based on the co-planning, implementation, and evaluation of the *My Kidneys My Health* website. As part of this project, we will work appropriate partners to ensure that evaluation is a continuous process within their settings.

**Discussion**

Implementation science uses established theories and frameworks to understand contextual and other factors related to the use of an innovation (Bauer & Kirchner, 2020). To successfully implement a patient-focused eHealth intervention tailored for patients in primary care and general nephrology we need to understand the potential implementation problems, desired implementation behaviors, and contextual factors, and identify the appropriate evidence-based implementation strategies. To the best of our knowledge, this is the first implementation science study of the integration of a patient-focused eHealth tool tailored for patients in primary care and general nephrology context. We are proposing to test a “formula for success” where we are implementing an evidence-informed intervention (*My Kidneys My Health* self-management website), with quality implementation (QIF), in an enabling context (primary care and general nephrology clinics). The goal of this study is to generate knowledge about implementing and sustaining an eHealth tool in the primary care and general nephrology setting. Applying the QIF to guide this work will support the achievement of implementation outcomes based on a strategic approach to identifying implementation strategies for the sites.

The knowledge gained from this study will strengthen the implementation and sustainment of eHealth self-management interventions in chronic kidney disease care and clinical care settings by identifying barriers and facilitators, as well as implementation strategies appropriate to real-world clinical contexts. Anticipated impacts of this study include moving toward enhanced chronic kidney disease self-management and the application of a standard approach to the development and customization of transferrable implementation strategies and tools for use in different care settings. It is anticipated that the use of QIF activities will support successful implementation with the intention to promote self-management strategies and enhance desired health outcomes, in addition to informing how to implement patient-centric eHealth tools in clinical settings on a larger scale.

There are limitations to this study. This study is limited to eight participating sites and the health care professionals within these clinical settings. The participants may not be representative of health care professionals in other contexts. A potential limitation of the evaluation process may be recall bias when obtaining interview data after the implementation period (post 9 months) as adherence may fluctuate.
However, we believe the use and integration of several different theoretical perspectives is an approach that will extend the literature in implementation science and help identify how such theoretical perspectives can be combined to offer a comprehensive approach to implementing evidence-informed interventions into other settings.

Supplementary Information The online version contains supplementary material available at https://doi.org/10.1007/s43477-022-00038-3.

Acknowledgements We would like to acknowledge our patient partners and stakeholders who are integral in the development of this protocol and the success of this study.

Author Contributions MD, HB, SS, LH, GH, BW, MD, DS, PW, ME, KM AB, and BH contributed to the research idea and study design. MD and BH wrote the initial protocol with each author contributing important intellectual content during manuscript drafting and revisions. MD, HB, SS, LH, GH, BW, MD, DS, PW, ME, KM AB, and BH had the opportunity to read and approve the final manuscript.

Data Availability The datasets used and/or analyzed during the current study will be available from the corresponding author on reasonable request.

Declarations

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

Ethical Approval Ethical approval has been obtained from the University of Calgary Conjoint Health Research Ethics Board (REB21-0930).

Consent to Participate All participants will be consented into the study as per CHREB approved protocol: oral consent will be obtained from participants for interviews and written consent will be obtained for surveys, as approved by the ethics committee.

Consent for Publication Not applicable.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

References

Atkins, L., Francis, J., Islam, R., O’Connor, D., Patey, A., Ivers, N., Foy, R., Duncan, E. M., Colquhoun, H., Grimshaw, J. M., Lawton, R., & Michie, S. (2017). A guide to using the Theoretical Domains Framework of behaviour change to investigate implementation problems. Implementation Science, 12, 77. https://doi.org/10.1186/s13012-017-0605-9

Baay, S., Hemmelgarn, B., Tam-Tham, H., Finlay, J., Elliott, M. J.,Straus, S., Beanlands, H., Herrington, G., & Donald, M. (2019). Understanding adults with chronic kidney disease and their caregivers’ self-management experiences: A qualitative study using the Theoretical Domains Framework. Canadian Journal of Kidney Health and Disease. https://doi.org/10.1177/2054358119848126

Bauer, M. S., & Kirchner, J. (2020). Implementation science: What is it and why should I care? Psychiatry Research, 283, 112376. https://doi.org/10.1016/j.psychres.2019.04.023

Blanchard, C., Livet, M., Ward, C., Sorge, L., Sorensen, T. D., & McClurg, M. R. (2017). The active implementation frameworks: A roadmap for advancing implementation of comprehensive medication management in primary care. Research in Social & Administrative Pharmacy, 13(5), 922–929. https://doi.org/10.1016/j.sapharm.2017.05.006

Bonner, A., Gillespie, K., Campbell, K. L., Corones-Watkins, K., Hayes, B., Harvie, B., Kelly, J. T., & Havas, K. (2018). Evaluating the prevalence and opportunity for technology use in chronic kidney disease patients: A cross-sectional study. BMC Nephrology, 19(1), 28. https://doi.org/10.1186/s12882-018-0830-8

Damschroder, L. J., Aron, D. C., Keith, R. E., Kirsh, S. R., Alexander, J. A., & Lowery, J. C. (2009). Fostering implementation of health services research findings into practice: A consolidated framework for advancing implementation science. Implementation Science, 4, 50. https://doi.org/10.1186/1748-5908-4-50

Donald, M., Beanlands, H., Straus, S., Ronksley, P., Tam-Tham, H., Finlay, J., MacKay, J., Elliott, M., Herrington, G., Harwood, L.,Large, C. A., Large, C. L., Waldvogel, B., Sparkes, D., Delgado, M.,Tong, A., Grill, A., Novak, M., James, M. T., et al. (2019a). Identifying needs for self-management interventions for adults with CKD and their caregivers: A qualitative study. AJKD, 74(4), 474–482. https://doi.org/10.1053/j.ajkd.2019.02.006

Donald, M., Beanlands, H., Straus, S., Ronksley, P., Tam-Tham, H., Finlay, J., Smeakal, M., Elliott, M. J., Farragher, J., Herrington, G., Harwood, L.,Large, C. A., Large, C. L., Waldvogel, B., Delgado, M. L., Sparkes, D., Tong, A., Grill, A., Novak, M., et al. (2019b). Preferences for a self-management e-health tool for patients with chronic kidney disease: Results of a patient-oriented consensus workshop. CMAJ Open, 7(4), E713–E720. https://doi.org/10.9778/cmajopen.20190081

Donald, M., Beanlands, H., Straus, S. E., Smeakal, M., Gil, S., Elliott, M. J., Herrington, G., Harwood, L., Waldvogel, B., Delgado, M. S., Sparkes, D., Tong, A., Grill, A., Novak, M., James, M. T., Brimble, K. S., Samuel, S., Tu, K., Farragher, J., & Hemmelgarn, B. R. (2021a). A web-based self-management support prototype for adults with chronic kidney disease (My Kidneys My Health): Co-design and usability testing. JMIR Form Res. https://doi.org/10.2196/22220

Donald, M., Kahlon, B. K., Beanlands, H., Straus, S., Ronksley, P.,Herrington, G., Tong, A., Grill, A., Waldvogel, B., Large, C. A., Large, C. L., Harwood, L., Novak, M., James, M. T., Elliott, M., Fernandez, N., Brimble, S., Samuel, S., & Hemmelgarn, B. R. (2018). Self-management interventions for adults with chronic kidney disease: A scoping review. British Medical Journal Open, 8(3), e019814. https://doi.org/10.1136/bmjopen-2017-019814

Donald, M., Straus, S., Smeakal, M., Herrington, G., Waldvogel, B.,Sparkes, D., Delgado, M., & Hemmelgarn, B. R. (2021b).
POSt-479 Acceptability and feasibility of a web-based self-management support intervention for adults with chronic kidney disease: A mixed methods study. *Kidney International Reports*, 6(4), S207.

Durlak, J. A., & DuPre, E. P. (2008). Implementation matters: A review of research on the influence of implementation on program outcomes and the factors affecting implementation. *American Journal of Community Psychology*, 41(3-4), 327–350. https://doi.org/10.1007/s10464-008-9165-0

Eldh, A. C., Sverker, A., Bendtsen, P., & Nilsson, E. (2020). Health care professionals’ experience of a digital tool for patient exchange, anamnesis, and triage in primary care: Qualitative study. *JMIR Human Factors*. https://doi.org/10.2196/21698

Fixsen, D. L., Blase, K. A., Timbers, G. D., & Wolf, M. M. (2007). In search of program implementation: 792 replications of the teaching-family model. *The Behavior Analyst Today*, 8(1), 96–110. https://doi.org/10.1037/h0101004

Gagliardi, A. R., Legare, F., Brouwers, M. C., Webster, F., Bradley, E., & Straus, S. (2015). Patient-mediated knowledge translation (PKT) interventions for clinical encounters: A systematic review. *Implementation Science*. https://doi.org/10.1186/s13012-016-0389-3

Gale, N. K., Heath, G., Cameron, E., Rashid, S., & Redwood, S. (2013). Using the framework method for the analysis of qualitative data in multi-disciplinary health research. *Bmc Medical Research Methodology*. https://doi.org/10.1186/1471-2288-13-117

Ghimire, S., Castelino, R. L., Lioufas, N. M., Peterson, G. M., & Zaidi, S. T. (2015). Nonadherence to medication therapy in haemodialysis patients: A systematic review. *PLoS ONE*. https://doi.org/10.1371/journal.pone.0144119

Glasgow, R. E., Harden, S. M., Gaglio, B., Rabin, B., Smith, M. L., Porter, G. C., Ory, M. G., & Estabrooks, P. A. (2019). RE-AIM planning and evaluation framework: Adapting to new science and practice with a 20-year review. *Frontiers in Public Health*, 7, 64. https://doi.org/10.3389/fpubh.2019.00064

Greer, R. C., Liu, Y., Cavanaugh, K., Diamantidis, C. J., Estrella, M. M., Sperati, C. J., Soman, S., Abdel-Kader, K., Agrawal, V., Plantinga, L. C., Schell, J. O., Simon, J. F., Vassalotti, J. A., Jaar, B. G., Choi, M. J., & Committee, N. K. F. E. (2019). Primary care physicians’ perceived barriers to nephrology referral and co-management of patients with CKD: A qualitative study. *Journal of General Internal Medicine*, 34(7), 1228–1235. https://doi.org/10.1007/s11606-019-04975-y

Hemmelgarn, B. R., Pannu, N., Ahmed, S. B., Elliott, M. J., Tam-Tham, H., Lillie, E., Straus, S. E., Donald, M., Barnieh, L., Chong, G. C., Hillier, D. R., Huffman, K. T., Lei, A. C., Villanueva, B. V., Young, D. M., Fowler, E. A., Manns, B. J., & Laupacis, A. (2016). Determining the research priorities for patients with chronic kidney disease not on dialysis. *NDT*. https://doi.org/10.1093/ndt/gfw065

Holtrop, J. S., Rabin, B. A., & Glasgow, R. E. (2018). Qualitative approaches to use of the RE-AIM framework: Rationale and methods. *BMC Health Services Research*, 18(1), 177. https://doi.org/10.1186/s12913-018-2938-8

Hsieh, H. F., & Shannon, S. E. (2005). Three approaches to qualitative content analysis. *Qualitative Health Research*, 15(9), 1277–1288. https://doi.org/10.1177/104973305276687

Lennon, L., Doyle, C., Reed, J. E., & Bell, D. (2017). What makes a sustainability tool valuable, practical and useful in real-world healthcare practice? A mixed-methods study on the development of the long term success tool in northwest London. *BMJ Open*. https://doi.org/10.1136/bmjopen-2016-014417

Levin, A., Adams, E., Barrett, B. J., Beanlands, H., Burns, K. D., Chiu, H. H., Chong, K., Dart, A., Ferera, J., Fernandez, N., Fowler, E., Garg, A. X., Gilbert, R., Harris, H., Harvey, R., Hemmelgarn, B., James, M., Johnson, J., Kappel, J., et al. (2018). Canadians seeking solutions and innovations to overcome chronic kidney disease (Can-SOLVE CKD): Form and function. *Canadian Journal of Kidney Health and Disease*. https://doi.org/10.1177/2054358119863091

Levin, A., Stevens, P. E., Bilous, R. W., Coresh, J., De Francisco, A. L., De Jong, P. E., Griffith, K. E., Hemmelgarn, B. R., Ikizler, T. A., & Lamb, E. J. (2013). Kidney disease: Improving global outcomes (KDIGO) CKD work group. KDIGO 2012 clinical practice guideline for the evaluation and management of chronic kidney disease. *Kidney International Supplements*, 3(1), 1–150. https://doi.org/10.1038/kisup.2012.73

Meyers, D. C., Durlak, J. A., & Wandersman, A. (2012). The quality implementation framework: A synthesis of critical steps in the implementation process. *American Journal of Community Psychology*, 50(3–4), 462–480. https://doi.org/10.1007/s10464-012-9522-x

Michie, S., Atkins, L., & West, R. (2014). *The behaviour change wheel: A guide to designing interventions* (1st ed.). Siverback Publishing. www.behaviourchangewheel.com

Morrisson, L. G., Yardley, L., Powell, J., & Michie, S. (2012). What design features are used in effective e-health interventions? A review using techniques from critical interpretive synthesis. *Telemedicine and e-Health*, 18(2), 137–144. https://doi.org/10.1089/tmj.2011.0062

National Kidney Foundation. (2017). *National research priorities for kidney disease identified by patients*. https://www.kidney.org/news/national-research-priorities-kidney-disease-identified-patients

Noseworthy, T., Wasylik, T., & O’Neill, B. (2015). Strategic clinical networks in Alberta: Structures, processes, and early outcomes. *Healthcare Management Forum*, 28(6), 262–264. https://doi.org/10.1177/0840470415600130

Powell, B. J., Waltz, T. J., Chinman, M. J., Damschroder, L. J., Smith, J. L., Matthieu, M. M., Proctor, E. K., & Kirchner, J. E. (2015). A refined compilation of implementation strategies: Results from the expert recommendations for implementing change (ERIC) project. *Implementation Science*, 10, 21. https://doi.org/10.1186/s13012-015-0209-1

Presseau, J., McCleary, N., Lorenzatto, F., Patey, A. M., Grimshaw, J. M., & Francis, J. J. (2019). Action, actor, context, target, time (AACCTT): A framework for specifying behaviour. *Implementation Science*, 14(1), 102. https://doi.org/10.1186/s13012-019-0951-x

Proctor, E. K., Powell, B. J., & McMillen, J. C. (2013). Implementation strategies: Recommendations for specifying and reporting. *Implementation Science*, 8, 139. https://doi.org/10.1186/1748-9098-8-139

RE-AIM. (2021). *RE-AIM: Reach, effectiveness, adoption, implementation, maintenance*. www.RE-AIM.org

Richard, A. A., & Shea, K. (2011). Delineation of self-care and associated concepts. *Journal of Nursing Scholarship*, 43(3), 255–264. https://doi.org/10.1111/j.1547-5069.2011.01404.x

Shen, H., van der Kleij, R., van der Boog, P. J. M., Chang, X., & Chavannes, N. H. (2019). Electronic health self-management interventions for patients with chronic kidney disease: Systematic review of quantitative and qualitative evidence. *Journal of Medical Internet Research*, 21(11), e12384. https://doi.org/10.2196/12384

Smeak, M., Gil, S., Donald, M., Beanlands, H., Strauss, S., Herrington, G., Spakes, D., Harwood, L., Tong, A., Grill, A., Tu, K., Waldvogel, B., Large, C., Large, C., Novak, M., James, M., Elliott, M., Delgado, M., Bramble, S., et al. (2019). Content and quality of websites for patients with chronic kidney disease: An environmental scan. *Canadian Journal of Kidney Health and Disease*, 6, 2054358119863091. https://doi.org/10.1177/2054358119863091

Tonelli, M., Wiebe, N., Guthrie, B., James, M. T., Quan, H., Fortin, M., Klarenbach, S. W., Sargiouis, P., Straus, S., Lewanczuk, R.,
Ronksley, P. E., Manns, B. J., & Hemmelgarn, B. R. (2015). Comorbidity as a driver of adverse outcomes in people with chronic kidney disease. *Kidney International, 88*(4), 859–866. https://doi.org/10.1038/ki.2015.228

Tong, A., Crowe, S., Chando, S., Cass, A., Chadban, S. J., Chapman, J. R., Gallagher, M., Hawley, C. M., Hill, S., Howard, K., Johnson, D. W., Kerr, P. G., McKenzie, A., Parker, D., Perkovic, V., Polkinghorne, K. R., Pollock, C., Strippoli, G. F., Tugwell, P., et al. (2015). Research priorities in CKD: Report of a national workshop conducted in Australia [Research Support, Non-U.S. Gov’t]. *American Journal of Kidney Diseases, 66*(2), 212–222.

Tong, A., Sainsbury, P., & Craig, J. (2007). Consolidated criteria for reporting qualitative research (COREQ): A 32-item checklist for interviews and focus groups. *International Journal for Quality in Health Care, 19*(6), 349–357. https://doi.org/10.1093/intqhc/mzm042

Waltz, T. J., Powell, B. J., Matthieu, M. M., Damschroder, L. J., Chinman, M. J., Smith, J. L., Proctor, E. K., & Kirchner, J. E. (2015). Use of concept mapping to characterize relationships among implementation strategies and assess their feasibility and importance: Results from the expert recommendations for implementing change (ERIC) study. *Implementation Science. https://doi.org/10.1186/s13012-015-0295-0*

Wandersman, A., Chien, V. H., & Katz, J. (2012). Toward an evidence-based system for innovation support for implementing innovations with quality: Tools, training, technical assistance, and quality assurance/quality improvement. *American Journal of Community Psychology, 50*(3–4), 445–459. https://doi.org/10.1007/s10464-012-9509-7