Enhancing community health through patient navigation, advocacy, and social support: A community health navigator pilot study

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Key highlights:

- 1) Interventions using Community Health Navigators (CHNs), community members that are hired and trained to navigate the healthcare system, are not well studied in the Canadian primary care context.
- 2) In the preliminary results from this pilot study, the CHN intervention appeared to improve patients' social and financial support and satisfaction with care.
- 3) The CHN intervention is feasible to implement in primary care for patients with multiple chronic conditions.

Abstract

The healthcare system is complex and difficult to navigate, particularly for patients with multiple chronic conditions and complex care plans. Patient adherence to care plans and patient health outcomes can be negatively impacted by language, financial, and other social barriers. Community Health Navigators (CHNs) are community members that are hired and trained to navigate the healthcare system, who work with patients to overcome barriers to care and support patient self-management by providing services tailored to needs. While these types of interventions can improve access to care in other settings, they are not well studied in Canada nor in Canadian primary care settings. For this pilot study, we aimed to determine the feasibility of a CHN intervention for patients with multiple chronic conditions. Our secondary objective was to assess the potential impact of a CHN intervention on patient-reported outcome measures. We used an observational single arm pre-post study design. Using interviewer-administered patient surveys, we assessed patient-reported outcomes at baseline (pre-enrolment), and 6-months and 12-months post-enrolment. The survey included instruments to assess quality of life (EQ-5D-5L), patient chronic disease care experience (PACIC), social support (mMOS-SS), and cost-related adherence to care (i.e. financial security to pay for care-related costs). Descriptive analysis was performed on survey data, and the sample was restricted to participants who completed both follow-up survey data, and the sample was restricted to participants who completed both follow-up survey data. And the sample was restricted to participants of conditions a patient reported was 5.4 (SD 2.3, range 3-11). Of the sample enrolled, 14 (67%) patients completed both follow-up surveys. Mean social support (scale: 0-100) did not change over time. Mean patient experience with chronic disease care (scale: 0-100) did not change over time. Mean patient experience with chronic disease care (scale: 1-3) was 2.01 at baseline; 2.24 at 6

Key words

Patient navigator, community health navigators, primary care, chronic disease, social determinants of health.



Cite as: Pritchard C., MacDonald C., Ludlow N., Fabreau G., McBrien KA. 2019. Enhancing community health through patient navigation, advocacy, and social support: A community health navigator pilot study. Alberta Academic Review, Vol 2 (3) 22, CASCH Special Issue (not peer-reviewed), DOI: 10.29173/aar113.

