Development of a Mobile Application to Support Evidence-Informed, Client-Centred Community Care for Stroke Survivors with Multiple Chronic Conditions

What is the research objective?
The research objective was to collaborate with frontline homecare workers (e.g. care coordinators, nurses, physiotherapists, occupational therapists, and personal support workers) and key stakeholders (e.g. Registered Nurses' Association of Ontario, Heart and Stroke Foundation) to inform, design and build an innovative mobile health application to support communication and collaboration.

The end goals of the application are to promote evidence-informed care for older adults who have experienced a stroke and have multiple chronic conditions (MCC), and to provide a communication tool for all members of the homecare team, including healthcare providers, clients, and their family caregivers.

How was the study done?
Qualitative focus groups with 41 frontline homecare workers (end-users of the application) and consultations with 23 key stakeholders were conducted to inform the design and development of "My Stroke Team" (MYST), the web-based mobile application.

Initially, key barriers to quality homecare were identified as well as a "wish list" of application functions. Next, end-user input was used in an iterative manner to inform and provide feedback on the tool's functions and usability throughout the development of MYST.

What did the researchers find?
Stakeholders and end-users identified the following barriers to the delivery of quality homecare for stroke survivors with MCC:

1] fragmented communication between healthcare providers;
2] no consistent means to track patient status;
3] limited access to information and resources at the point-of-care,
4] safety issues resulting from communication gaps; and
5] a lack of consideration of MCC in common guidelines.
RESEARCH BRIEF

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MYST was developed to address these barriers through the following features:

1] a platform for real-time, secure communication between all homecare providers (e.g. care coordinators, nurses, physiotherapists, occupational therapists, and personal support workers) within the circle of care;

2] information for stroke survivors with MCC and their family caregivers about local community supports and online resources;

3] secure point-of-care access to patient information and other resources including practice guidelines for the circle of care; and

4] promotion of client safety through evidence-informed assessments and alerts to relevant providers.

What does this mean for practice and policy?

MYST has the potential to improve the efficiency and the quality of home-based healthcare for stroke survivors with MCC. We expect that MYST will result in:

• Improved communication between homecare providers, thereby reducing the risk of errors, facilitating seamless care, and resulting in better patient outcomes.

• Easy access to best practices for stroke management, thereby promoting evidence-informed, client-centred care.

• Enhanced client/caregiver-centred care and engagement

Where do we go from here?

The feasibility and acceptability of MYST and its effects on health-related quality of life will be evaluated in a pilot study (Study 8). Access to MYST will be provided to patients, their family caregivers, and homecare providers in their circle of care.

Who are the researchers?

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