

A Mixed-Methods Evaluation of Diabetic Retinopathy Screening Supported by Provincial Healthcare Administrative Data

JAMES M. BOWEN^{1,2,3}, ALEKSANDRA STANIMIROVIC^{1,2,3}, OLIVERA SUTAKOVIC^{3,4}, CONRAD POW^{3,6}, DEBBIE SISSMORE^{3,7}, MALCOLM SISSMORE^{3,7}, JENNIFER RAYNER⁸, SARA BHATTI⁸, REBECCA MERRITT^{3,9}, BAIJU R. SHAH^{2,3,10}, MICHAEL H. BRENT^{3,4,5}, VALERIA E. RAC^{1,2,3}

¹Program for Health System and Technology Evaluation, Toronto General Hospital Research Institute, UHN, ²IHPME, University of Toronto, ³Diabetes Action Canada SPOR Network ⁴Donald K Johnson Eye Institute, Toronto Western Hospital, University Health Network (UHN), ⁵Dept. of Ophthalmology & Vision Sciences, Faculty of Medicine, University of Toronto. ⁶North York General Hospital, ⁷Patient Partner, ⁸Alliance for Healthier Communities, ⁹South Riverdale Community Health Centre, ¹⁰ICES & Sunnybrook Health Sciences Centre, Toronto, Ontario, Canada.

Background

- Diabetic retinopathy is a public health issue impacting the lives of 3 million or more Canadians.
- Systematic identification of those requiring screening for retinopathy is not possible within electronic medical records.

Methodology

- Provincial-level identification through a list of unscreened individuals to community health centres (CHCs) (Figure 1)
- Convergent mixed-methods implementation-effectiveness hybrid evaluation approach. (Figures 2 & 3)
- Evaluating effectiveness, cost-effectiveness, implementation and policy barriers and key success factors (Figure 4).

Methodology

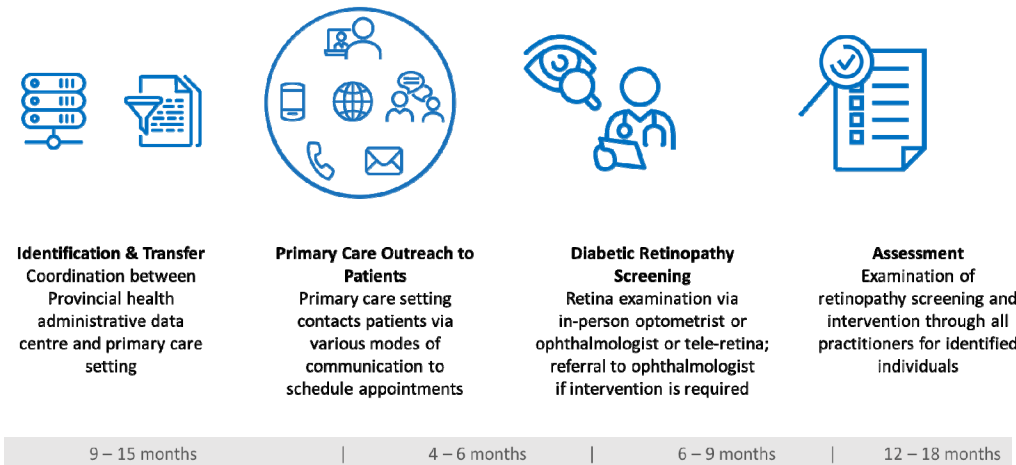


Figure 1. Stages of diabetic retinopathy screening intervention and evaluation

Results

- Initial evaluation of contact lists across three CHCs found between 575 and 900 individuals unscreened within 425 days.

Conclusions

- A “top-down” approach using healthcare administrative data provided to primary care settings can raise awareness of need for diabetic retinopathy screening.
- A step towards increased screening and reducing vision loss.

Acknowledgement

This study is supported through a generous private donation, Fighting Blindness Canada and by Diabetes Action Canada through funds provided by the Canadian Institutes of Health Research (CIHR) Strategy for Patient-Oriented Research (SPOR) Networks. ClinicalTrials.gov: NCT05074342.

References 1. Creswell JW, Plano Clark VL. Designing and Conducting Mixed Methods Research. Third ed. Los Angeles, CA: Sage Publications; 2017; 2. Curran GM, Bauer M, Mittman B, Pyne JM, Stetler C. Effectiveness-implementation hybrid designs: combining elements of clinical effectiveness and implementation research to enhance public health impact. Med Care. 2012;50(3):217-26.

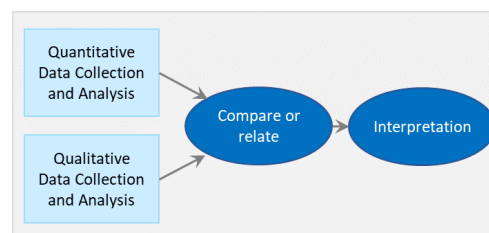


Figure 2. The Convergent Parallel Mixed-methods Design (Figure adopted from Creswell and Clark, 2017)

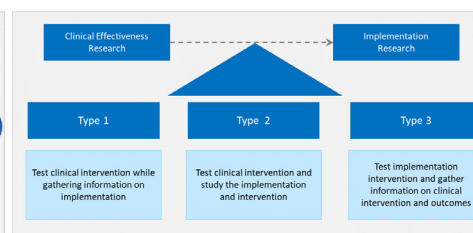


Figure 3. Effectiveness-Implementation Hybrid Study Designs (Figure adopted from Curran et al., 2012)

Implementation Study Quantitative & Qualitative	Clinical Evaluation Study Quantitative
<ul style="list-style-type: none"> • Data transfer feasibility & information completeness • Qualitative study using ethnographic fieldwork & semi-structured interviews • Descriptive analysis of intervention uptake 	<ul style="list-style-type: none"> • Comparative study of clinical outcomes and healthcare resource utilization for screened patients compared with concurrent controls
Policy Study Quantitative & Qualitative	Economic Analysis Study Quantitative
<ul style="list-style-type: none"> • Qualitative study using semi-structured interviews of relevant stakeholders and analysis of relevant documentary sources • Descriptive analysis of the overall DRS patterns 	<ul style="list-style-type: none"> • Descriptive implementation cost analysis • Budgetary impact analyses • Cost-effectiveness analysis

Figure 4. Study Components and Designs