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national health Information  
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strategy development and implementation research, as well as to health information science. An examination into the national health IM/IT plans of several countries provides knowledge into identifying the typical IM/IT priorities that selected countries are focusing upon

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for healthcare improvement. Second, a systematic literature review of the current challenges, barriers and/or issues (referred to as 'challenges' hereafter) facing IM/IT priority implementation in healthcare settings provides insight on where nations should perhaps be focusing their



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attention, in order to enable more successful healthcare IM/IT implementations. Lastly, a study on national health IM/IT priorities contributes to the body of evidence that national level IM/IT direction is necessary for better patient care and health system reform across

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theworld. In this investigation, the national health IM/IT priorities, which are reflected in the national health IM/IT strategic plans of five countries were assessed. To this end, the study:

- 1) Developed a set of measures to select four countries to study in addition to Canada?
- 2) Described the

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national health IM/IT priorities of  
Canada and four other countries? 3)  
Answers  
Performed a systematic literature  
review of the challenges to overcome  
for successful implementation of IM/IT  
into healthcare settings? 4)  
Developed and administered a  
questionnaire where participants

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were asked to give their opinions on the progress their country has achieved in dealing with such challenges? and 5) Performed an analysis of the questionnaire results with respect to the countries' national health IM/IT priorities. The systematic literature review

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uncovered a large number of challenges that the healthinformatics and healthcare community face when attempting to implement IM/IT intohealthcare settings.iiiThe priority comparison highlighted that there is no right or wrong answer for whatcountries should focus their

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energies upon.

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However, by learning about what other nations are prioritizing, a

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country can use that knowledge to help focus their own national health IM/IT priorities. The questionnaire results drew attention to the most frequently encountered challenges the five countries face in moving their national health IM/IT agendas forward. The feedback from

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the respondents provided individual reflections on how IM/IT implementations are actually progressing in their country, where problems are being encountered, including the nature of those problems, and in some cases, respondents offered insight on how to



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better deal with the challenges they face. The findings indicate that nations encounter similar problems in implementing IM/IT into healthcare settings. Currently, the world is facing many of the same healthcare system issues: shortages of healthcare professionals, long surgical and

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diagnostic imaging waitlists,  
?skyrocketing' pharmaceutical drug  
pricing, healthcare funding practices,  
and challenges with implementing  
healthcare IM/IT priorities to name a  
few. If countries are facing  
similar health system problems, then  
it would be logical to assume that

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solutions to deal with such problems would be similar across nations. Thus, it is recommended that international fora and conferences be held to further discuss the types of health system IM/IT priorities that countries are implementing at a national scale, the kinds of challenges they face and

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the solutions or conclusions that they have formulated in response to these challenges.

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outcomes. For the purposes of this guide, a patient registry is an organized system that uses observational study methods to collect uniform data (clinical and other) to evaluate specified outcomes for a population defined by a particular disease, condition, or

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exposure, and that serves one or more predetermined scientific, clinical, or policy purposes. A registry database is a file (or files) derived from the registry. Although registries can serve many purposes, this guide focuses on registries created for one or more of the following purposes: to

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describe the natural history of disease, to determine clinical effectiveness or cost-effectiveness of health care products and services, to measure or monitor safety and harm, and/or to measure quality of care. Registries are classified according to how their populations are defined.

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defined by patients having the same diagnosis, such as cystic fibrosis or heart failure. The User ' s Guide was created by researchers affiliated with AHRQ ' s Effective Health Care Program, particularly those who participated in AHRQ ' s DEcIDE (Developing Evidence to Inform

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