## Memorandum of Understanding for a Systematic Review of the Literature Project number:

A systematic review must demonstrate that sufficient effort has been made to identify as many methodologically sound studies as possible to answer a clinical question. As such, the search is a critical step in the systematic review process.

RESPONSIBILITIES OF EACH PARTY		
The lead investigator will:		
	Provide direction and feedback for the search	
	Keep the librarian updated on progress in the research team's work	
	Complete screening and analysis using established standards (e.g. multiple screeners)	
	Send the final manuscript to the librarian for comments and feedback prior to submission	
	Give notice before final documentation and manuscript are submitted to a publisher	
The MUHC librarian will:		
	* Design the search strategy and have it peer reviewed	
	* Run searches in the bibliographic databases outlined on page 2	
	* Create an EndNote library with all retrieved publications and remove duplicates	
	Search the grey literature (including conference abstracts)	
	Search trials registries	
	Handsearch selected journals (number: )	
	* Compile the PRISMA flow diagram of the selection process	
	Search reference lists of eligible studies	
	Conduct citation search of eligible studies	
	* Write the "sources and search methodology" section	
	Rerun the Medline search strategy prior to submitting for publication	
Other:		

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<sup>\*</sup> If responsible for these steps, the MUHC librarian will be granted co-authorship of the final publication.

## **Selected Databases and Other Resources**

Bibliographic databases				
Medicine African Index Medicus Biosis Previews Cochrane Library / CENTRAL Embase Global Health LILACS (Latin America) Medline (OvidSP) PsycINFO (Ovid) PubMed (ePub records only)	Nursing/Allied Health  AMED CINAHL OT Seeker (Occupational therapy) PEDro (Physiotherapy) RehabDATA (Rehabilitation)  Multidisciplinary Scopus (Elsevier) Web of Science (ISI)			
Other types				
Research in progress  ClinicalTrials.gov International Clinical Trials Registry Platform International Standard Randomised Controlled Trial Number Registry UK Clinical Trials Gateway  Conference proceedings: conferences to be search	Dissertations/Theses ProQuest Dissertations and Theses Guidelines CMA Infobase National Guideline Clearinghouse National Institute for Clinical Excellence dindividually			
Grey literature: literature published outside the traditional publishing ecosystem				
Health Technology Assessments (HTA)  Agency for Healthcare Research and Quality (AHRQ)  Canadian Agency for Drugs and Technologies in Health  Centre for Reviews and Dissemination (CRD)  Institut National d'excellence en santé et en services sociaux (INESSS)  MUHC TAU	Drug and device regulatory approvals:  Food and Drug Administration (FDA)  Health Canada  Other:  Google Scholar  Open Grey			

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## Date of the initial meeting with a librarian: Timeframe for delivery of search results:

Research teams should allow for approximately of 2 months from the initial meeting with a librarian to the delivery of search results, with the understanding that delays outside of the control of the librarian, such as delayed feedback from the research team and/or changes to the research question, will affect the originally agreed upon timeframe.

Lead investigator:			
Name (capital letters):			
Signature:			
Date:			
Librarian			
Name (capital letters):			
Signature:			
Date:			

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