

## CLINICAL TRIAL RESEARCH ETHICS REVIEW NAVIGATOR

### Background

Multi-site studies and clinical trials play a vital role in child health research in Canada. One of the challenges for study teams in conducting multicentre research is efficiently obtaining research ethics review across all participating sites. When a single research ethics board (REB) review is not possible for all participating sites, navigating through the various ethics review platforms and systems across the country can be a challenge. Each platform has a unique application process and the local REB requirements, including protocol and consent templates that are often different.

### Solution Offering

Working alongside the Canadian Collaboration for Child Health: Efficiency and Excellence in the Ethics Review of Research ([CHEER](#)) initiative and institutional REBs, MICYRN is offering a pediatric clinical research ethics submission consultation service. The aim of the consultation service is to facilitate, enhance, and expedite multi-site, multi-jurisdictional pediatric research, and position Canada as a global leader in children's health.

### Role of the Clinical Trial Research Ethics Review Navigator

The MICYRN Clinical Trial Research Ethics Review Navigator will serve as the repository of knowledge with a rich understanding and functionality of the various existing streamlined ethics review processes across Canada. The Navigator will:

- Assist study teams in determining if a multi-site submission, a single-site submission, or a combination of the two is the most efficient path for their research study
- Provide guidance on who the lead study team should be for a multi-site submission and what it means to act as a lead site versus a participating site
- Identify the appropriate and applicable consent forms required for a research project
- Collaborate with the CHEER project manager to help lead and participating institutions benefit from the CHEER initiative
- Assist study teams in obtaining local REB acknowledgement and approval

## Scope of Service

Clinical research projects should meet the inclusion criteria below to be supported by the Clinical Trial Research Ethics Review Navigator:

- Multicentre study sites must be located in at least two provinces and use more than one streamlined system (e.g., a provincial system and CHEER, or two or more provincial systems)
- Must be prospective observational clinical research studies or interventional clinical trials
- Must be scientific or peer reviewed by a designated steering committee or granting body (i.e., CIHR)

*If you or an investigator at your institution is seeking support for a streamlined REB submission consultation, please email Wenli Xie at [wenli.xie@micyrn.ca](mailto:wenli.xie@micyrn.ca).*