OHSN-REB

Breaking News

for studies
under
the jurisdiction of
both
CHEO and OHSN
Research Ethics
Boards

Effective Date: December 18th, 2019

To whom does this apply?

To all investigators and researchers whose studies fall under the jurisdiction of both CHEO and OHSN REB; an institutional agreement for streamlining the REB review process has now been finalized.

What does this mean?

- Studies falling under the jurisdiction of both CHEO and OHSN REB (i.e., due to Investigator affiliations and/or study activity at both sites) will no longer require two separate ethics applications.
- Instead, CHEO REB and OHSN REB will work collaboratively to review and approve the research studies via <u>one</u> ethics application. There will be a single Research Ethics Board of record assigned.
- Note, the studies must be registered with the opposite REB (not acting as Board of record) and Institutional approval at each institution (OHRI or OHIRC and CHEO/CHEO RI) is still required prior to study start.
- If in addition to CHEO and or TOH/UOHI, the research will be conducted at multiple sites in Ontario who are participating sites in Clinical Trials Ontario (CTO), the research study should be submitted through CTO Stream.

What is the process?

- 1) Study teams will complete a REDCap Survey: TOH & CHEO REB Harmonization
- 2) The Survey will be reviewed to determine the following, which will need to be submitted simultaneously:
 - the review system (ROMEO at CHEO/CHEO RI or IRIS at TOH/OHRI) to be used for ethics review which will be used for the life of the study

and

- the system for study registration (TOH/OHRI Clinical Research Registration Form (CRRF) via IRIS or CHEO Administration application via ROMEO)
- 3) The study team will receive research <u>ethics</u> review and decision making from a single REB (OHSN or CHEO) for the life of the study.
- 4) If CHEO REB is the Board of record, the study team will be required to continue with the registration process for the life of the study at TOH/OHRI.