Science and Research Ethics Board Review Process

Important: Applications to the Holland Bloorview Research Ethics Board should identify a Bloorview Research Institute scientist, Clinical Team Investigator or Clinical Study Investigator as the Local Principal Investigator. Exceptions to this must be reviewed and approved by the Holland Bloorview VP Research/BRI Director before submitting a research ethics application.

All REB forms can be found in the "Forms" section http://research.hollandbloorview.ca/ResearchEthicsBoard/Forms

Prepare research proposal.		
		If your funded study received a
Check the <u>Science Review Process at Holland Bloorview</u> to determine the relevant requirements. Submit your research proposal to the primary science reviewer for formal review.		science review by the granting agency, or if you are a research graduate student and the protocol was approved by your supervisory committee, then an internal science review may not be required. Check with the REB Office Staff.
Receive the completed Science Review Form from the primary science reviewer.		
	+	Make further revisions as
Make the appropriate changes and submit response to the primary science reviewer.		required.
	¥	Request a Departmental
Receive approval/signoff from the primary science reviewer.		Approval for Research Involving Clients at Holland Bloorview if
		participants include Holland
Prepare TAHSN Application Form and obtain signatures from all investigators.		Bloorview clients/families. If participants include students recruited from any school including the Bloorview School
♦ Note: A research proposal that is still under review by a granting agency is usually not reviewed by the Holland Bloorview REB until after funding is awarded. Check with the REB Office Staff before you submit your proposal for ethics review.		Authority, then request a support letter from the school principal/board. Request a support letter from the Holland Bloorview Pharmacy if you require pharmaceutical services.
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Prepare Research Ethics Board Applie documents (where applicable) to the <u>BRI</u> <u>submissions</u> . BRI Operations Office will a. <u>REB Application Checklist</u>	<u>Operations Office</u> for review.	heck monthly <u>deadlines for</u> the REB Office.
b. TAHSN Application Form	i. Participant Documents	
c. Study Protocol	j. Approval letters from other REBs	
d. Informed Consent Forms	k. Investigator Brochure or Product Monograph	
e. Assent Forms	l. Health Canada NOL/ITA, FDA Approval	
f. <u>Departmental Approval Form</u>	m. Risk Assessment Documentation (Devices)	

- g. Scientific Review Form + Response
- n. Data & Biological Sample Transfer Agreements

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