1. PURPOSE
The purpose of this standard operating procedure (SOP) is to describe the Health Sciences Research Ethics Board (HSREB) submission requirements, and the administrative review procedures conducted by the Office of Human Research Ethics (OHRE). This SOP applies to all submissions including but not limited to: new research projects for initial review, amendments or modifications to approved research, and any new information.

2. GENERAL POLICY STATEMENT
HSREB members must rely on the documentation provided by the principal investigator for initial and continuing review. Therefore, the materials submitted must provide sufficient information to conduct the review and make the required determinations.

The REB is supported by administrative procedures that assure that HSREB members not only have adequate time for assessment of proposed research, but that the materials they receive allow them to adequately assess whether the research meets the criteria for HSREB approval.

3. RESPONSIBILITY
This SOP applies to the all HSREB members including the Chair and Vice-Chair(s) and to all OHRE staff.

4. DEFINITIONS
See Glossary of Terms

5. SPECIFIC POLICIES AND PROCEDURES.
The OHRE is responsible for maintaining the submission requirements and for making such information available to Investigators. The instructions to Investigators regarding submission requirements, including deadlines and meeting dates, are available on the OHRE webpages or by contacting the OHRE.

5.1. Submission Requirements
5.1.1. The required documents, checklists, number of copies, format, and submission procedures are outlined on the OHRE webpages including but not limited to:
- Letter of Information/Consent (LOI/C) Guidance Document
- Amendment Form
- Continuing Ethics Review (CER) Form
• Updated Approvals
• Request for Acknowledgement (FYI Form)
• End of Study Form

5.2. **Administrative Review Procedures**

5.2.1. All applications can be viewed in the online system by the OHRE staff as soon as they are submitted by the Investigator or designee;

5.2.2. The unique system-generated number is assigned automatically to each submission at the time the application is created;

5.2.3. The review type (Delegated, Full Board review) is initially done by the Investigator by filling out the appropriate online application form;

5.2.4. Once the application has been created and submitted through the online system the OHRE Administrative Staff will inform the triage Ethics Officer (EO) that a new submission has been submitted;

5.2.5. Upon receipt of the submission, the triage EO conducts a preliminary review to ensure completeness of the application, including validation of the appropriate attachments and to ensure that the review type is correct;

5.2.6. If the submission is incomplete (e.g., documents are missing, incorrect documents were uploaded) or if there are clear problems with the submission (i.e., coherence of the submission, major methodology concerns, etc) that will preclude timely review by either the Full Board or delegated review), the triage EO may route it back to the Investigator/study staff within the online system to make the necessary changes and to resubmit the revisions, time permitting;

5.2.7. Upon receipt of a complete submission, the submission is then assigned to an EO and, appropriate reviewer(s) for either Full Board or delegated review;

5.2.8. For submissions required Full Board review, the submissions are posted to the agenda of the next Full Board meeting and an EO and primary reviewer is assigned once the agenda is completed;

5.2.9. For submissions requiring Delegated review, the triage EO routes the submission in the online system to the assigned reviewers (i.e., to the HSREB Chair or Vice-Chair(s), EO, and to other reviewers as directed by the HSREB Chair, Vice Chair(s), or designee).

6. **REFERENCES**


6.2. The International Conference on Harmonization Guidelines for Good Clinical Practice, Section 3;


6.5. OHRP Guidance on Written IRB Procedures;
6.6. FDA Information Sheets.

7. SOP HISTORY

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