




SOP 002\_05

Title	Confidentiality and Privacy
SOP Code	SOP 002_05
Effective Date	30-June-2023

## Site Approval/Authorization to Adopt

Name and Title of Local Personnel (Type or print)	Signature	Date dd/Mon/yyyy
<b>Neelu Sehgal</b> Director, Interprofessional Practice & Research Chief Nursing Executive, Erie Shores Health Care		
<b>Dr. Munira Sultana</b> Office of Research, Erie Shores Health Care		23/06/2023

### **1.0 PURPOSE**

This Standard Operating Procedure (SOP) describes the procedure to ensure the integrity and confidentiality of clinical data collected within the framework of a clinical study, and the privacy of clinical study participants.

### **2.0 SCOPE**

This SOP is applicable to all clinical studies undertaken at the site (Erie Shores Health Care), and to those clinical research personnel responsible for ensuring the security and confidentiality of data, and participant information.

### **3.0 RESPONSIBILITIES**

The Sponsor-Investigator or Qualified Investigator (QI)/Investigator is responsible for ensuring that the confidentiality and privacy aspects of the study meet all of the applicable regulatory, International Conference on Harmonisation (ICH) Good Clinical Practice (GCP), sponsor, and local requirements.

Any or all parts of this procedure may be delegated to appropriately trained study team members, but remain the ultimate responsibility of the Sponsor-Investigator or Qualified Investigator (QI)/Investigator.

### **4.0 PROCEDURE**

#### **4.1 General Principles**

4.1.1 Information that is disclosed in the context of a professional or research relationship must be held confidential to the extent permissible within the applicable law. Confidential information may be written, verbal, electronic, photographic or stored in any other medium (i.e. tissue, diagnostic images).

4.1.2 Protection of privacy and/or confidentiality in the context of clinical research includes prevention of disclosure, to other than authorized individuals, of any information which could identify a research participant, or any Sponsor's proprietary information.

4.1.3 Every authorized person with direct access to clinical data must comply with the Declaration of Helsinki, the directives of the ICH-GCP, regulatory requirements and applicable privacy legislation for the maintenance of confidentiality, participant identity and respect for the proprietary information of the Sponsor or the Sponsor-Investigator.

4.1.4 Authentication of the person who has access to the data constitutes the most important aspect of security. It determines the overall level of protection and is linked to key elements of data security.

## 4.2 Data Security

4.2.1 Establish a mechanism for control of access to secure premises. Document the procedure. It is recommended that the control mechanism includes use of magnetic cards or a biometric recognition system that allows tracking of movement in and out of the premises, if applicable.

4.2.2 Retain a tracking document with the signatures and initials of all persons authorized to register data, or to make corrections to the CRFs, with the essential study documentation.

4.2.3 Physical security concerns the premises where study files containing essential documents and clinical data, as well as computer equipment used for data management, such as telecommunication servers, database servers and computers are located. These rooms should be located in an area protected from possible disasters (e.g., water or fire damage, etc.), and be protected by a secure access control system.

4.2.4 Logical security concerns management of access to data, which includes identification, authentication, and authorization. In order to ensure logical security, the following measurements should be applied:

- Limit authorized access to members of the research team, and those identified by the protocol, the consent form, and the delegation of authority form;
- Grant privileges for physical or electronic access to data to personnel according to the roles and responsibilities defined by the Sponsor-Investigator or Qualified Investigator (QI)/Investigator;

4.2.5 Sponsor or Sponsor-Investigator: designate person in charge of system management (system administrator).

4.2.6 System Administrator's responsibilities include:

- Develop and enforce standardized procedures for logical security;
- Assign a different identification code to each user of the data management system;
- Ensure that users change their confidential password regularly, according to the period defined by the system administrator;
- Ensure the confidentiality of the authentication of system users, and document access tracking;
- Establish a Disaster Recovery Plan, for saving and recovering data, in the event of loss or disaster;
- Suspend the authorized access of a user after a given number of errors. Inform other users of this suspension. Update the delegation of tasks form, accordingly. Retrain user, if required. Document training; and
- Cancel access for research team members who leave the study (resignation, illness, maternity leave, etc.). Update the delegation of tasks form, accordingly.

## 4.3 Data Confidentiality

4.3.1 A participant who authorizes access to his/her data must be reasonably assured that the Sponsor or Sponsor-Investigator, QI/Investigator, their authorized representatives, Research Ethics Board (REB), and regulatory inspectors of the regulatory authorities will take precautions to ensure that verified and collected data remain confidential.

4.3.2 Include a description of the provisions and limits of confidentiality within the context of the research study, in the informed consent process and form, as follows:

- The consent form must describe to the participant who will have access to their information and for what purposes (e.g., Health Canada, US Food and Drug Administration (FDA), Research Ethics Board (REB), research sponsors and personnel monitoring/auditing the research on their behalf);
- Information collected during the research study will not be shared by the researcher without the participant's free and informed consent;
- Information pertaining to the research participant must be recorded in research case report forms (CRFs), and any other study documents that will leave the research site, including electronically captured data, in such a way as to protect participant identity. Participants will be identified with a unique identifier ;
- No study materials sent to and/or retained by the sponsor will contain any identifying information. This includes, but is not limited to, investigational drug returns, test results, medical histories, and adverse event reports,
- Participant enrolment lists, copies of prescriptions and informed consent forms, which include identifying information, will be retained at the site. All research documents with participant identifiers should be kept separate from all study documents.
- Any reimbursements or stipends paid to the participants will be paid from the research account specific to the study, and not directly from the sponsor. This will ensure sponsor does not have access to participant names; and
- The Research Ethics Board (REB) review process governs secondary use of the information gathered (for purposes other than the original intent), in accordance with the applicable regulatory guidelines.

## 5.0 REFERENCES

Health Canada, Food and Drug Regulations, Part C, Division 5, Drugs for Clinical Trials Involving Human Subjects, (Schedule 1024), June 20, 2001.

Health Canada, Guidance for Industry, Good Clinical Practice: Consolidated Guideline, ICH Topic E6, 1997.

Canadian Institutes of Health Research, Natural Sciences and Engineering Research Council of Canada, and Social Sciences and Humanities Research Council of Canada, Tri-Council Policy Statement: Ethical Conduct for Research Involving Humans, December 2014.



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Department of Justice (Canada), Personal Information Protection and Electronic Documents Act (PIPEDA), updated 2006.

Pharmaceutical Inspection Convention, Pharmaceutical Inspection Co-operation

Scheme, Annexe 11, Computerised Systems.

US Food and Drug Administration, Code of Federal Regulations, Title 21, Volume 1:

- Part 11, Electronic Records; Electronic Signatures, (21CFR11).
- Part 50, Protection of Human Subjects, (21CFR50).
- Part 54, Financial Disclosure by Clinical Investigators, (21CFR54).
- Part 56, Institutional Review Boards, (21CFR56).
- Part 312, Investigational New Drug Application (21CFR312).

US Department of Health and Human Services, Code of Federal Regulations, Title 45, Part 46, Protection of Human Subjects (45CFR46).