

Documented Institutional Ethics Requirements Queen's University

Informed Consent Form Requirements:

Data Retention:

The minimum retention period is **5 years per Queen's policy**. There is no requirement to destroy data. It is suggested that the researchers consider using a data repository. **Health Canada regulated studies must retain data for a minimum of 15 years.**

SRERS Administration Queen's University

Hospital Based Researchers

For hospital-based research occurring in more than one affiliated hospital (e.g., KHSC (KGH site) and/or KHSC (HDH site) and/or PCC) with a single Principal Investigator, the Principal Investigator must specify their primary institution in the "Centre" tab of the Participating Site Initial Application (PSIA), and the additional sites in the response to question 2.5 (click "Yes" and then enter the additional Affiliated Hospital name(s) in the sub-questions).

When research is occurring in more than one Affiliated Hospital, the Kingston research team must also ensure that collaborators (as outlined in the CTO Stream section of each applicable SRERS Administration form) from each Affiliated Hospital are manually added to the PSIA.

For more information on hospital based research please refer to: <http://www.queensu.ca/traq/awards-grants-contracts/supportive-documents/>.

TRAQ

A TRAQ DSS FORM must be completed for all Research projects.

If your research is taking place within one of the Kingston area hospitals please be sure to include the Hospital Departmental Impact & Information Form with your TRAQ submission. All hospital departments impacted by your research must be checked off on the "Approvals" tab.

For more information, please consult "Hospital Based Research - Tips for Completing the TRAQ DSS FORM", "Hospital Departmental Impact & Information Form Tips", and "Hospital-Based Research Frequently Asked Questions (FAQ)" available at <http://www.queensu.ca/traq/awards-grants-contracts/supportive-documents/>.

CTO Stream

Collaborators:

The following collaborators must be given a role on the Clinical Trial Initial Application (CTIA) or Observational Study Initial Application (OSIA) and Participating Site Initial Application (PSIA) forms.

Email: kelly.blair-matuk@queensu.ca
Role: Institutional Representative

Email: meera.sidhu@queensu.ca
Role: Institutional Representative

Email: sps1@queensu.ca
Role: Institution Representative

Email: ashley.theis@queensu.ca
Role: Institution Admin

Email: efr1@queensu.ca
Role: Institutional Admin

This access is automatically granted when the PSIA is created. **When Queen's is the Lead Applicant site the research team should immediately create the PSIA for Queen' (right after creating the CTIA/OSIA).** This will ensure that access is automatically granted as required above, otherwise the research team will need to manually add these roles to the CTIA/OSIA prior to submission.

Institution Representative in application forms

The Primary Institution Representative must be indicated as follows in the applications within CTO Stream:

Title: Dr.

First Name: Stephen
Surname: Smith
Organization: Kingston General Health Research Institute
Address: Connell 4, Room 2-4-033, 76 Stuart Street
City: Kingston
Province/State: ON
Postcode/Zip: K7L 2V7
Telephone: (613) 549-6666 ext. 4287
Fax: N/A
Email: sps1@queensu.ca

The Secondary Institution Representative field must be indicated as follows:

Title: Dr.
First Name: Kelly
Second Name: Blair-Matuk
Organization: Queen's University
Address: 355 King Street West, Room 264C
City: Kingston
Province/State: ON
Postcode/Zip: K7L 3N6
Telephone: (613) 533-6000 ext. 78292
Fax: N/A
Email: kelly.blair-matuk@queensu.ca