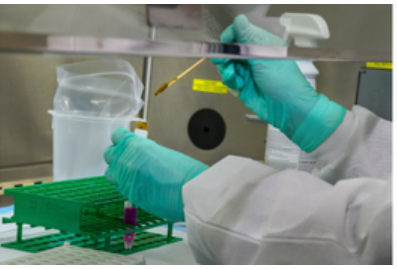
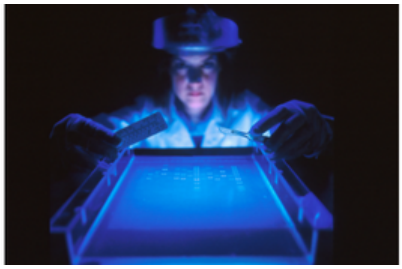
Volume 3 No 3

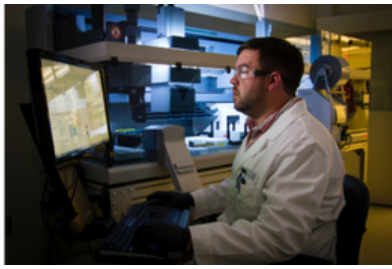
June 2025

NEWSLETTER

NEWS FROM THE RRASU

Research and Regulatory Affairs Support Unit, CIUSSS-ODIM



NEW NAGANO MODULE

LAW 5

A new module has been developed in Nagano to manage requests for access to personal information without consent (PIAs or EFVP). It will make it easier to track the status of your PIAs and submit and sign related documents. In addition, this new module will facilitate and centralize PIA requests for multi-site projects using health and social services information. The module will be activated in the coming weeks.

REPEATED PARTICIPATION

The USER (or RRASU in English) issues a warning against fraud involving repeated participation in a research project with the aim of receiving compensation multiple times. Please consult the footer of Nagano, under documentation, for the memo that explains the situation and the measures put in place to prevent it.

FRAUD

Québec

BASIC RESEARCH APPLICATION (BRAP) SUBMISSION

Please note that for BRAP studies, the submission form F11-H must be used; the submission form's title has been updated to help make this clearer. In the future, please note that you will be asked by the REB office to resubmit if the incorrect form is used. Please see the cheatsheet, 'types de projets' for more details.

RESPONSE TIME FROM THE REB AND RESEARCHERS

Did you know that the REB's response is usually given within 5 to 10 business days after the meeting for evaluations conducted in a full board committee? In addition to this period, there are 21 days between the submission of the project and the meeting. For projects undergoing delegated review, the average timeframe is 25 days. As for researchers, current standard operating procedures allow them 3 months to respond to REB comments when required. To obtain an extension, a request must be submitted to the REB.

EARLY SITE-SPECIFIC ASSESSMENT

Did you know that the start of the site-specific assessment review or the "examen de convenance" can be delayed if it is agreed that the project is at risk of being substantially modified following the REB's assessment or if the departments involved in the project have never been informed of the project and the project requires significant contributions? To this end, it is strongly recommended that you discuss the project with the departments at the time of its design to ensure their participation in advance. These discussions should not be limited to a department head, but must be validated by the person designated by the directorate to conduct the reviews.

USER WEBSITE

USER (or RRASU in English) now has a website that will host its entire regulatory framework, as well as all associated documents, including forms, guidelines, checklists, and templates. The various procedures relating to research project evaluations and approvals will be presented there, along with other relevant information. The website is scheduled to be active on July 1st. The link to the site will be communicated to you at that time.

LOOK BACK AT THE SYMPOSIUM ON ETHICS AND RESPONSIBLE CONDUCT IN RESEARCH

94 of you attended our first Symposium on Ethics and Responsible Conduct in Research; thank you! An evaluation questionnaire has been sent to you. Those eligible for accreditation will also receive an email to this effect.

RESEARCH ETHICS BOARD

Please find below the next project submission deadlines

Mental Health and Neuroscience subcommittee:
July 14 and Sept 8

Biomedical subcommittee:
June 18, July 23 and August 20

CONTACT US

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