

Queen's REB  
Guidelines on Case  
Report  
Studies/Series

# Queen's REB Guidelines on Case Report Study/Series

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**Research Compliance, Training and Ethics**

Queen's University is situated on [traditional Anishinaabe and Haudenosaunee Territory](#).

## Purpose

The purpose of this guideline is to:

- Provide guidance on the requirements for submitting a medical case report study/series and chart reviews for research purposes.

## Background

For HSREB, a case report study/series is a detailed report of the symptoms, signs, diagnosis, treatment, and follow-up of an individual patient/person.

For GREB, a case report is a detailed report of the situation, events, and intervention or support provided in a person's file.

Certain types of case report studies require REB approval.

### Please note:

**Queen's REBs do not issue retroactive approval for applications (i.e., if a submission requires REB approval, the application is required to be submitted before the participant consents and any data are collected. If a submission requires REB exemption, it is preferred that the application is to be submitted before the participant consents but may be submitted after).**

## Types of case report studies that require REB approval

### Research case report studies:

Research case report studies will be reviewed under the TCPS 2, Article 2.1. A research case report satisfies the following:

- Includes a research objective and/or question (i.e., What, how, or why).
- Makes a conceptual and theoretical contribution to the discipline.
- This could include the development of a research instrument.
- It could have theoretical propositions.
- It will be submitted to an external publication or conference.

### Chart review studies:

Chart review studies are a type of research in which pre-recorded, patient-centered data are used to answer one or more research questions. This type of study can be used to answer specific clinical questions in a less resource-intensive manner. A chart review has a clear objective and looks for trends or commonalities among participants' charts.

A research case report and chart review studies are required to be submitted for approval from the REB before the case report/chart review begins (i.e. before consenting the participant). Please refer to the applications listed in TRAQ and submit the most appropriate application type for a case study/series vs. a chart review.

## Types of case report studies that do not require REB approval

### Teaching case report studies:

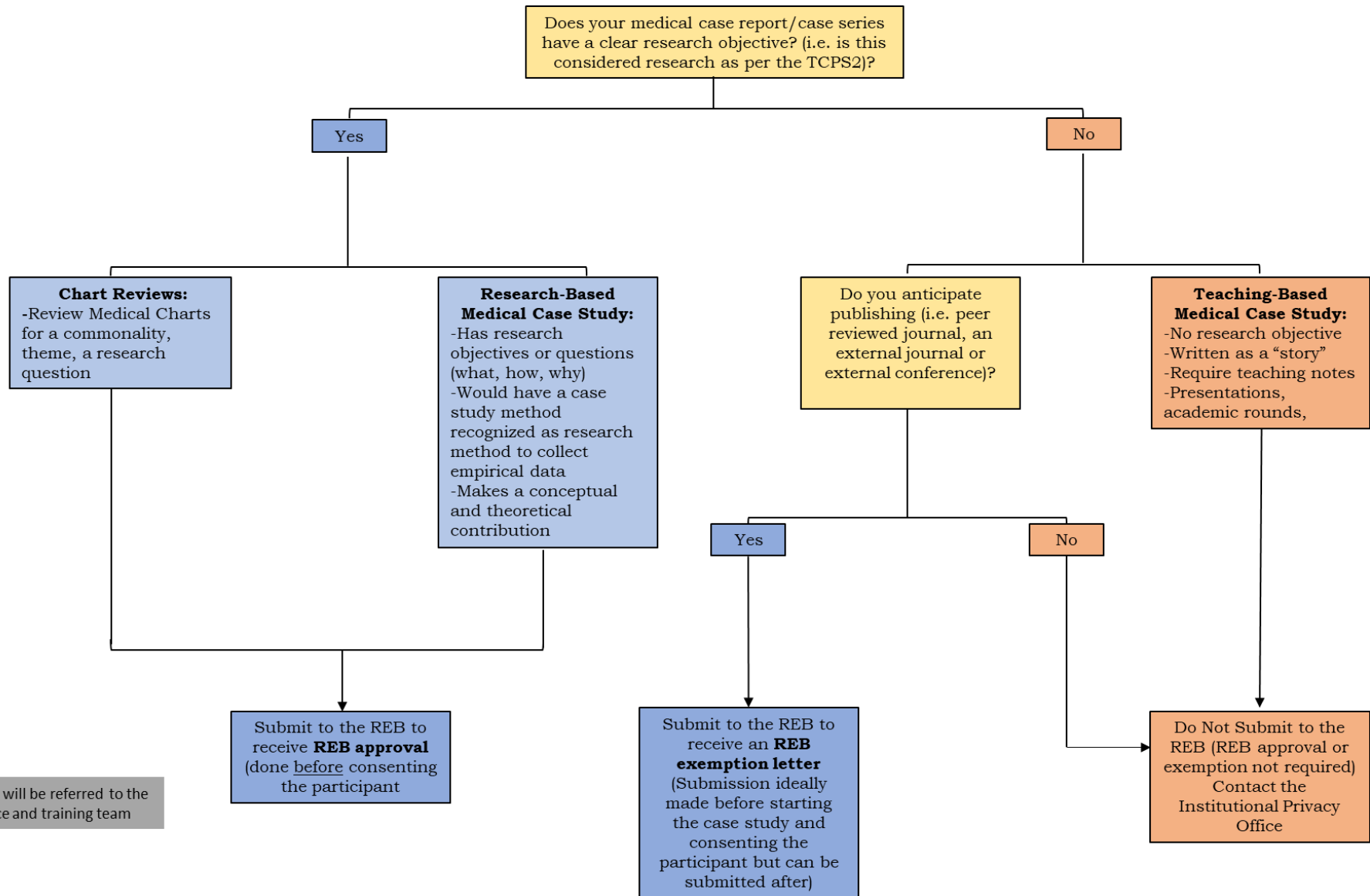
Teaching cases are exempt from ethics review based on the TCPS 2, Article 2.5. The 'intent or purpose' of a teaching case report study is for educational or learning purposes rather than research and, therefore, does not fall under the scope of the TCPS 2. A teaching case report will satisfy the following:

- It would be written as a "story."
- It would be written to support problem-based learning.
- It would require teaching notes.
- It would value practical implications more than theoretical knowledge.

## Decision Tree for determining what requires REB approval

Please see the decision tree below to determine if submission to the REB is required. **Note: Queen's REBs will not issue a retroactive approval.** If REB review is required, submission and approval by the REB are required before the participant consents and data is collected. If REB exemption is required, ideally this application is submitted prior to consenting and collecting data, but may be submitted after for an exemption letter.

Any non-compliance will be referred to the research compliance and training team.



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## Critical considerations for research case report studies

- Consent **will be obtained** from the participant, parent/legal guardian/substitute decision maker. Use of the Queen's REB consent form template for case studies is strongly recommended. The case report consent form template will be used to obtain consent (a qualified physician must also sign the consent form if the report is being authored by students/residents/fellows).
- There is no intention to test various therapies/treatments/interventions prospectively or retrospectively.
- Assent will be obtained for those who can assent, where consent has already been obtained by the parent/legal guardian/substitute decision maker.
- The consent/assent process will be documented and kept on file for 5 years per Queen's University guidelines.
- Justification of personal information/Personal Health Information (PHI) is required. It is best practice to limit the information collected (e.g., age, gender) (i.e., not using the full date of birth (DOB)/date of death (DOD) or any other information used in combination could lead to the identification of individuals). All measures will be taken to minimize the risk of re-identification through publication from the participant/friends/family members.
- All images/photographs will be de-identified (i.e., do not include name, medical record number, DOB) and do not include pictures with faces/facial features.
- The investigator, sub-investigators, or anyone connected to them through their interpersonal relationships (including their partners, family members, or former or current professional associates) will not receive any personal financial benefit from the report.
- Students/Residents/Fellows ensure that case reports are co-authored by a qualified physician who has appropriate credentials, is aware of, and shall make all reasonable efforts to comply with the applicable laws, guidelines, policies, and professional obligations.
- A TRAQ DSS form has been submitted to obtain hospital approval.
- Approval has been obtained from your departmental research committee.
- Refer to the [CARE Guidelines](#) and the [CARE Checklist](#) when writing your case report.

If you have any questions about your case report study/series, please get in touch with HSREB or GREB at [HSREB@queensu.ca](mailto:HSREB@queensu.ca) or [chair.greb@queensu.ca](mailto:chair.greb@queensu.ca)