



GHC/SAH REB GUIDANCE ON CASE REPORTS

Background

The purpose of this document is to provide guidance on when publication/presentation of case report(s) constitutes human subjects research and requires prospective REB review. The Tri-Council Policy Statement defines *research* as "an undertaking intended to extend knowledge through a disciplined inquiry or systematic investigation." A case report is a description of (a) the course of medical treatment with one or more patients that has a unique outcome or (b) the handling of a unique clinical case; which in either case did not involve the investigator having any research intent at the time of the intervention [i.e., no prospective plan to systematically evaluate the outcome for purposes other than treating the particular patient(s)]. A case report simply describes the course of medical treatment, a unique outcome, or a unique case. In an academic teaching hospital it is not unusual for unique and interesting clinical cases to be written up as case reports for publication in medical journals or presentation at medical or scientific meetings.

GHC/SAH REBs' Position on Case Reports

A case report with three or fewer patients does not meet the definition of research on the basis that the information in the case report has not been obtained through a systematic investigation and was not collected with a prior research intent. Furthermore, the information presented in a case report is not considered to be generalizable. Therefore, case studies that involve three or fewer patients do not require GHC/SAH REB review.

Conversely, if anything was done in the course of care with a research intent, the case report becomes research. If more than three patients are included in the project, it becomes a case series, is considered research, and requires GHC/SAH REB review and approval. The GHC/SAH REB expects that patients will be made aware that the author / investigator plans to create a report which may be published, about their case. Patients or their family members/substitute decision makers should provide written consent to have the patients' personal health information included in a case report. Clinicians should be sensitive to the "small cell problem": the existence of individuals with such unique or unusual diagnoses or illnesses, that it might be possible for others (or patients and families themselves) to identify the individuals in case reports or medical text books based upon limited information, such as city of residence, age and diagnosis.

The GHC/SAH REB does not grant "retroactive" approval after research is done. Researchers are advised to consult with the REB when uncertainty exists and when formal and systematic collection of human subjects' research will be occurring.





	CASE REPORT	RESEARCH
	(REB approval NOT needed)	(REB approval needed)
Medical	Three or fewer patients	Reviewing four or more patient records
Record Review		would meet the definition of research
		(disciplined inquiry or systematic
		investigation to extend knowledge).
Clinical	Each patient is treated as clinically	Clinical Trials: Research protocol
Treatment	indicated - treatment is not directed	dictates how all patient/subjects are
	by a research protocol nor are	treated. Specific research
	specific research	interventions/interactions scheduled.
	interventions/interactions required.	Other research (e.g. retrospective
		review of clinical records): treatment
		occurs clinically or as part as part of
		REB approved protocol.
Generalizable?	NOT GENERALIZABLE or	GENERALIZABLE. Hypotheses can be
	scientifically rigorous. Information is	proved/disproved. Data is collected
	considered to be educational by	and analyzed in a rigorous manner
	sharing a very limited number of	(may include statistics).
	interesting cases (3 or less) and how	
	they were treated. No statistical	
	analysis.	

NOTE: Investigators asked by a journal to provide documentation of REB approval or REB waiver for approval prior to publication for a case report of three (3) or less patients should provide the journal with a copy of this GHC/SAH REB guidance document.