

STANDARDS OF PRACTICE

Human Health Research

Under Review: No

Issued By: Council: September 1, 2012 (*Health Human Research Ethics Review*)

Reissued by Council: October 1, 2015 (*Human Health Research*)

The **Standards of Practice** of the College of Physicians & Surgeons of Alberta (“CPSA”) are the **minimum** standards of professional behavior and ethical conduct expected of all regulated members registered in Alberta. Standards of Practice are enforceable under the *Health Professions Act* and will be referenced in the management of complaints and in discipline hearings. CPSA also provides **Advice to the Profession** to support the implementation of the Standards of Practice.

1. This standard applies to any regulated member involved in human health research as identified through a current and recognized screening tool¹.
2. A regulated member who intends to conduct human health research **must** comply with the [Health Information Act](#), including to submit a proposal for review by a research ethics board in the Province of Alberta. Such boards include:
 - a. Health Research Ethics Board of Alberta ([HREBA](#))
 - b. Conjoint Health Research Ethics Board ([CHREB](#)), University of Calgary
 - c. Health Research Ethics Board ([HREB](#)), University of Alberta
3. A regulated member **must** have approval from a research ethics board before commencing human health research.
4. A regulated member participating in human health research **must**:
 - a. ensure the welfare of any patient involved in the research study is the primary concern throughout the duration of the study;
 - b. [disclose to patients](#) that the study has been reviewed by an ethics board and relevant conditions imposed;

¹ As of this date, the recommended tool is the [ARECCI \(A pRoject Ethics Community Consensus Initiative\)](#).

Terms used in the Standards of Practice:

- “Regulated member” means any person who is registered or who is required to be registered as a member of this College. The College regulates physicians, surgeons and osteopaths.
- “Must” refers to a mandatory requirement.
- “May” means that the physician may exercise reasonable discretion.
- “Patient” includes, where applicable, the patient’s legal guardian or substitute decision maker.

- c. comply with the requirements of the research ethics board as it relates to initial and ongoing review of the research study; and
- d. disclose any potential or actual [conflicts of interest](#) to the research ethics board.

RELATED STANDARDS OF PRACTICE

- [Code of Ethics & Professionalism](#)
- [Conflict of Interest](#)
- [Informed Consent](#)

COMPANION RESOURCES

- [Alberta Clinical Research Consortium](#)
- [ARECCI Ethics Screening Tool](#)
- [Tri-Council Policy Statement: Ethical Conduct for Research Involving Humans \(CIHR, NSERC, SSHRCC\)](#)

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