Additional Resources

Primary sources of information and regulation

- Tri-Council Policy Statement: Ethical Conduct for Research Involving Humans, 2nd edition (TCPS2)
- Declaration of Helsinki - World Medical Association (WMA), 1964
- Drugs for Clinical Trials involving Human Subjects - Health Canada Food and Drug Regulations, Part C, Division 5
- Medical Devices For Investigational Testing Involving Human Subjects - Health Canada: Food And Drug Act: Medical Device Regulations Part 3
- E6 - Good Clinical Practice: Consolidated Guideline
- US Code of Federal Regulation Title 56 on Protection of Human Subjects
- US Code of Federal Regulations Title 21 parts 50 and 56

Other useful links

- The Interagency Advisory Panel on Research Ethics (PRE) offers more information on the ethics requirements of the Tri-Council including the TCPS2: CORE Tutorial.
- Freedom of Information and Protection of Privacy Act (FIPPA) See Part 3 - Protection of Privacy: Division 2 - Use and Disclosure of Personal Information by Public Bodies - Articles 32 and 35.
- PIPA: Personal Information Protection Act (See Part 6 - Disclosure of Personal Information: Articles 21 and 22.
- Natural Health Products Regulations Health Canada: Food and Drug Act
- The Belmont Report: written by the U.S. National Commission for the Protection of Human Subjects of Biomedical and Behavioural Research

Contact us

- Contact the Research Ethics Board at researchethics@interiorhealth.ca or by phone at 250-870-4602.