

1. Food and Drugs Act
2. Food and Drug Regulations: Part C, Division 5, Drugs for Clinical Trials Involving Human Subjects
3. Natural Health Products Regulations: Part 4, Clinical Trials Involving Human Subjects
4. Medical Devices Regulations: Part 3, Medical Devices for Investigational Testing Involving Human Subjects
5. Personal Information Protection and Electronic Documents Act
6. United States Code of Federal Regulations: 21 CFR 50, 56, 312, 812 and 45 CFR 46
7. ICH GCP International Council on Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use – ICH Harmonised Tripartite Guideline – Guidelines for Good Clinical Practice E6(R2)
8. Canadian Institutes of Health Research, Natural Sciences and Engineering Research Council of Canada, and Social Sciences and Humanities Research Council of Canada, Tri-Council Policy Statement: Ethical Conduct for Research Involving Humans, TCPS2 2018
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