1. Food and Drugs Act
2. Food and Drug Regulations: Part C, Division 5, Drugs for 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
7. ICH GCP International Conference on Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use – ICH Harmonised Tripartite Guideline – Guidelines for Good Clinical Practice E6(R1)
10. World Medical Association (WMA). Declaration of Helsinki – Ethical Principles for Medical Research Involving Human Subjects


