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
9. World Medical Association (WMA). Declaration of Helsinki – Ethical Principles for Medical Research Involving Human Subjects

20. U.S. Department of Health and Human Services, Food and Drug Administration. Comparison of FDA and HHS Human Subject Protection Regulations


