

Websites		
<u>Category</u>	<u>Homepage</u>	<u>Subcategory Sites</u>
Advisory Committee on Human Radiation Experiments	http://www.gwu.edu/~nsarchiv/radiation/	
American Society for Bioethics and Humanities	http://www.asbh.org	
Association of American Universities (AAU)	http://www.aau.edu/	
Belmont Report (<i>Ethical Principles and Guidelines</i>)	https://www.hhs.gov/ohrp/regulations-and-policy/belmont-report/index.html	
Bioresearch Monitoring (<i>FDA Program</i>)	https://www.fda.gov/science-research/clinical-trials-and-human-subject-protection/bioresearch-monitoring-program-bimo	
CBER (<i>Center for Biologics Evaluation and Research, FDA</i>)	https://www.fda.gov/about-fda/office-medical-products-and-tobacco/about-center-biologics-evaluation-and-research-cber	
CDC (<i>Center for Disease Control and Prevention, DHHS</i>)	http://www.cdc.gov/	
CDER (<i>Center for Drug Evaluation and Research, FDA</i>)	https://www.fda.gov/about-fda/center-drug-evaluation-and-research/center-drug-evaluation-and-research	
CDRH (<i>Center for Devices and Radiological Health, FDA</i>)	https://www.fda.gov/about-fda/office-medical-products-and-tobacco/center-devices-and-radiological-health	
Council for International Organizations of Medical Sciences (CIOMS)	http://www.cioms.ch/	
Declaration of Helsinki (<i>Biomedical research recommendations</i>)	https://www.wma.net/policies-post/wma-declaration-of-helsinki-ethical-principles-for-medical-research-involving-human-subjects/	

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Department of Education <i>(Protection of Human Subjects in Research)</i>	https://www2.ed.gov/about/offices/list/ocfo/humansub.html	<i>PPRA for Parents =</i> https://www2.ed.gov/policy/gen/guid/fpco/ppra/parents.html
DOE <i>(Department of Energy)</i>	http://energy.gov/	<i>Human Genome Project Information=</i> https://web.ornl.gov/sci/techresources/Human_Genome/
DHHS <i>(Dept. of Health and Human Services)</i>	http://www.hhs.gov/	
eCFR <i>(Electronic Code of Federal Regulations)</i>	https://gov.ecfr.io/cgi-bin/ECFR	
FDA <i>(Food and Drug Administration)</i>	http://www.fda.gov	<i>FDA Bioresearch Monitoring Information =</i> https://www.fda.gov/science-research/clinical-trials-and-human-subject-protection/bioresearch-monitoring-program-bimo <i>Title 21 CFR 50 =</i> https://gov.ecfr.io/cgi-bin/text-idx?SID=6da9926849ad0af5a3275a02bd8bbb4f&mc=true&node=pt21.1.50&rgn=div5 <i>Title 21 CFR 50.24 Consent Waiver =</i> https://gov.ecfr.io/cgi-bin/text-idx?SID=6da9926849ad0af5a3275a02bd8bbb4f&mc=true&node=pt21.1.50&rgn=div5#se21.1.50_124 <i>MedWatch =</i> http://www.fda.gov/Safety/MedWatch/default.htm
GPO <i>(Government Publishing Office)</i>	https://www.gpo.gov/	<i>Code of Federal Regulations & Federal Register =</i> https://www.govinfo.gov/bulkdata/CFR/#default
Guide to Good Clinical Practice <i>(Ordering information)</i>	http://fda.thompson.com/Guide-to-Good-Clinical-Practice	
ICH <i>(International Conference on Harmonization)</i>	http://www.ich.org/home.html	

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IRB Forum (formerly MCWIRB now maintained by PRIM&R)	http://www.irbforum.org/	
MedWatch (FDA Medical Products Reporting Program)	http://www.fda.gov/Safety/MedWatch/default.htm	MedWatch (FDA Medical Products Reporting Program)
NCTR (National Center for Toxicological Research)	https://www.fda.gov/about-fda/office-chief-scientist/national-center-toxicological-research	
NHGRI (National Human Genome Research Institute)	http://www.genome.gov/	
NIH (National Institutes of Health)	http://www.nih.gov/	Intramural Program: Office of Human Subjects Research = http://ohsr.od.nih.gov/ Office for Human Research Protections (OHRP) = http://www.hhs.gov/ohrp/ Division of Extramural Research: Ethical, Legal, and Social Implications Research Program = http://www.genome.gov/10001618 Human Genome Project = http://www.genome.gov/10001772
NSF (National Science Foundation)	http://www.nsf.gov/	
Nuremberg Code (Directives for Human Experimentation)	https://history.nih.gov/research/downloads/nuremberg.pdf	
Office of Good Clinical Practice (formerly Office of Human Research Trial) FDA	https://www.fda.gov/science-research/clinical-trials-and-human-subject-protection/good-clinical-practice-contacts	
OHRP (Office of Human Research Protections)	http://www.hhs.gov/ohrp/	OHRP IRB Guidebook = http://wayback.archive-it.org/org-745/20150930181805/http://www.hhs.gov/ohrp/archive/irb/irb_guidebook.htm

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<i>ORA (Office of Regulatory Affairs) FDA</i>	https://www.fda.gov/about-fda/office-global-regulatory-operations-and-policy/office-regulatory-affairs	
<i>ORI (Office of Research Integrity) Department of Health and Human Resources</i>	https://ori.hhs.gov/	
<i>ORO (Office of Research Oversight)</i>	http://www.va.gov/oro/	
Presidential Commission for the Study of Bioethical Issues (formerly National Bioethics Advisory Commission)	https://bioethicsarchive.georgetown.edu/pcsbi/node/851.html	
<i>PRIM&R (Public Responsibility in Medicine and Research)</i>	http://www.primr.org/	
<i>PHS (Public Health Service)</i>	http://www.usphs.gov/	<i>Policy on Humane Care & Use of Laboratory Animals =</i> http://grants.nih.gov/grants/olaw/references/phspol.htm
<i>R&D (VA Research & Development)</i>	http://www.research.va.gov/default.cfm	
<i>UK Homepage (University of Kentucky)</i>	http://www.uky.edu/	
<i>USDA (U.S. Dept. of Agriculture)</i>	http://www.usda.gov/wps/portal/usda/usdahome	