

REGULATIONS, DIRECTIVES, INSTRUCTIONS AND POLICIES FOR THE
PROTECTION OF HUMAN SUBJECTS IN RESEARCH

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- b. Department of Health and Human Services (DHHS) Regulation, Title 45 CFR Public Welfare, Part 46: “Protection of Human Subjects”, 23 June 2005. (the “Common Rule”)
<http://www.hhs.gov/ohrp/humansubjects/guidance/45cfr46.htm>
- c. Department of Defense (DoD) Regulation, Title 32 CFR Part 219: “Protection of Human Subjects”, 19 August 1991. <http://www.dtic.mil/biosys/downloads/32cfr219.pdf>
- d. Federal Drug Administration (FDA) Regulation, Title 21 CFR Part 50: “Protection of Human Subjects”, 1 April 2006.
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- e. Federal Drug Administration (FDA) Regulation, Title 21 CFR Part 56: “Institutional Review Boards”, 1 April 2006.
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- j. Department of Defense (DoD) Directive 3210.7, “Research Integrity and Misconduct”, 14 May 2004. http://www.dtic.mil/whs/directives/corres/pdf/i32107_051404/i32107p.pdf
- k. U.S. Naval Academy Academic Dean and Provost Instruction, ACDEANINST 3920.3, “Integrity in Research and Scholarly Activity”, 4 February 2000.
- l. Secretary of the Navy Instruction, SECNAVINST 5210.8D, “Department of the Navy

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- n. Federal Privacy Act (5 USC 301, 552a (1994)).
[http://www.law.cornell.edu/uscode/search/display.html?terms=552\(a\)&url=/uscode/html/uscode05/uscode05_sec_05_00000552---a000-.html](http://www.law.cornell.edu/uscode/search/display.html?terms=552(a)&url=/uscode/html/uscode05/uscode05_sec_05_00000552---a000-.html)
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