

Public Health Service, HHS

§ 52.8

§ 52.7 Use of funds; changes.

(a) *Delegation of fiscal responsibility.* The grantee may not in whole or in part delegate or transfer to another person responsibility for the use or expenditure of grant funds.

(b) *Changes in project.* The permissible changes by the principal investigator in the approved project shall be limited to changes in methodology, approach or other aspects of the project to expedite achievement of the project's research objectives, including changes that grow out of the approved project and serve the best scientific strategy. If the grantee and the principal investigator are uncertain whether a change complies with this provision, the question must be referred to the Secretary for a final determination.

(c) *Changes in project period.* The project period determined pursuant to § 52.5(b) may be extended by the Secretary, with or without additional grant support, for such an additional period as the Secretary determines may be required to complete, or fulfill the purposes of, the approved project.

[45 FR 12240, Feb. 25, 1980]

§ 52.8 Other HHS regulations and policies that apply.

Several other HHS policies and regulations apply to grants under this part. These include, but are not necessarily limited to:

- 37 CFR part 401—Rights to inventions made by nonprofit organizations and small business firms under government grants, contracts, and cooperative agreements
- 42 CFR part 50, subpart A—Responsibility of PHS awardee and applicant institutions for dealing with and reporting possible misconduct in science
- 42 CFR part 50, subpart D—Public Health Service grant appeals procedure
- 42 CFR part 50, subpart F—Responsibility of applicants for promoting objectively in research for which PHS funding is sought
- 45 CFR part 16—Procedures of the Departmental Grant Appeals Board
- 45 CFR part 46—Protection of human subjects
- 45 CFR part 74—Administration of grants
- 45 CFR part 75—Informal grant appeals procedures
- 45 CFR part 76—Governmentwide debarment and suspension (nonprocurement) and governmentwide requirements for drug-free workplace (grants)

45 CFR part 80—Nondiscrimination under programs receiving Federal assistance through the Department of Health and Human Services—effectuation of title VI of the Civil Rights Act of 1964

45 CFR part 81—Practice and procedure for hearings under part 80 of this title

45 CFR part 84—Nondiscrimination on the basis of handicap in programs and activities receiving Federal financial assistance

45 CFR part 86—Nondiscrimination on the basis of sex in education programs and activities receiving or benefiting from Federal financial assistance

45 CFR part 91—Nondiscrimination on the basis of age in HHS programs or activities receiving Federal financial assistance

45 CFR part 92—Uniform administrative requirements for grants and cooperative agreements to State and local governments

45 CFR part 93—New restrictions on lobbying
59 FR 14508 (March 28, 1994)—NIH Guidelines on the Inclusion of Women and Minorities as Subjects in Clinical Research.

NOTE: This policy is subject to changes, and interested persons should contact the Office of Research on Women's Health, NIH, Room 201, Building 1, MSC 0161, BETHESDA, MD 20892-0161 (301-402-1770; not a toll-free number) to obtain references to the current version and any amendments.]

59 FR 34496 (July 5, 1994)—NIH Guidelines for Research Involving Recombinant DNA Molecules.

NOTE: This policy is subject to changes, and interested persons should contact the Office of Recombinant DNA Activities, NIH, Suite 323, 6000 Executive Boulevard, MSC 7010, Bethesda, MD 20892-7010 (301-496-9838; not a toll-free number) to obtain references to the current version and any amendments.]

“PHS Grants Policy Statement,” DHHS Publication No. (OASH) 94-50,000 (Rev.) April 1, 1994.

NOTE: This policy is subject to changes, and interested persons should contact the Grants Policy Branch, OASH, Room 17A45, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857 (301-443-1874; not a toll-free number) to obtain references to the current version and any amendments.]

“Public Health Service Policy on Humane Care and Use of Laboratory Animals,” Office for Protection from Research Risks, NIH (Revised September 1986).

NOTE: This policy is subject to changes, and interested persons should contact the Office for Protection from Research Risks, NIH, Suite 3B01, 6100 Executive Boulevard, MSC 7507, Rockville, MD 20852-7507 (301-496-

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7005; not a toll-free number) to obtain references to the current version and any amendments.]

[61 FR 55106 Oct. 24, 1996]

§ 52.9 Additional conditions.

The Secretary may with respect to any grant award or class of awards impose additional conditions prior to or at the time of any award when in the Secretary's judgment such conditions are necessary to assure or protect advancement of the approved project, the interests of the public health, or the conservation of grant funds.

[45 FR 12240, Feb. 25, 1980; 45 FR 20096, Mar. 27, 1980]

PART 52a—NATIONAL INSTITUTES OF HEALTH CENTER GRANTS

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SOURCE: 57 FR 61006, Dec. 23, 1992, unless otherwise noted.

§ 52a.1 To which programs do these regulations apply?

(a) The regulations of this part apply to grants by the National Institutes of Health and its organizational components to support the planning, establishment, expansion, and operation of research and demonstration and/or multipurpose centers in health fields described in this paragraph. Specifically, these regulations apply to:

(1) National Institute of Mental Health centers of excellence with respect to research on autism, as authorized by section 409C of the Act (42 U.S.C. 284g);

(2) National cancer research and demonstration centers (including payments for construction), as authorized by section 414 of the Act (42 U.S.C. 285a–3);

(3) National cancer research and demonstration centers with respect to breast cancer, as authorized by section 417 of the Act (42 U.S.C. 285a–6);

(4) National cancer and demonstration centers with respect to prostate cancer, as authorized by section 417A of the Act (42 U.S.C. 285a–7);

(5) National research and demonstration centers for heart, blood vessel, lung, and blood diseases, sickle cell anemia, blood resources, and pediatric cardiovascular diseases (including payments for construction), as authorized by section 422 of the Act (42 U.S.C. 485b–4);

(6) Research and training centers (including diabetes mellitus, and digestive, endocrine, metabolic, kidney and urologic diseases), as authorized by section 431 of the Act (42 U.S.C. 285c–5);

(7) Research and training centers regarding nutritional disorders, as authorized by section 434 of the Act (42 U.S.C. 285c–8);

(8) Multipurpose arthritis and musculoskeletal diseases centers (including payments for alteration, but not construction), as authorized by section 441 of the Act (42 U.S.C. 285d–6);

(9) Alzheimer's disease centers, as authorized by section 445 of the Act (42 U.S.C. 285e–2);

(10) Claude D. Peppers Older Americans Independence Centers, as authorized by section 445A of the Act (42 U.S.C. 285e–3);

(11) Centers of excellence in Alzheimer's disease research and treatment, as authorized by section 445I of the Act (42 U.S.C. 285e–10a);

(12) Research centers regarding chronic fatigue syndrome, as authorized by section 447 of the Act (42 U.S.C. 285f–1);

(13) Research centers with respect to contraception and infertility, as authorized by section 452A of the Act (42 U.S.C. 285g–5);

(14) Child health research centers, as authorized by section 452C of the Act (42 U.S.C. 285g–7);

(15) Fragile X research centers, as authorized by 452E of the Act (42 U.S.C. 285g–9);