

for information on where and how to submit applications.

To be effective, the size of the committee will be limited. Each organization affected by emergency alarm systems on passenger vessels need not have its own representative on the committee. Rather, each interest must be adequately represented and the membership must be fairly balanced.

After reviewing the comments received in response to this notice and any applications for membership, the Board will issue a notice in the **Federal Register** announcing the establishment of the committee and the committee membership, unless it is determined based on comments that the establishment of the committee would be inappropriate. The first committee meeting is tentatively scheduled for August 15 and 16, 2007 at the Access Board offices in Washington, DC.

The Board expects the committee to hold no more than three meetings and all meetings will be in the Washington, DC area. The meetings will be open to the public. Future committee meetings will be announced in the **Federal Register**.

The Board will provide staff support to the committee. Members of the committee will not be compensated for their service. The Board may pay travel expenses for a limited number of persons who would otherwise be unable to serve on the committee. Members will not be considered special government employees since they will serve as representatives of their organizations and will not be required to file confidential financial disclosure reports.

Availability of Copies and Electronic Access

Single copies of this publication may be obtained at no cost by calling the Access Board's automated publications order line (202) 272-0080, by pressing 2 on the telephone keypad and then please record your name, address, city, State, zip code, telephone number and request the emergency alarms advisory committee notice. Persons using a TTY should call (202) 272-0082. This document is available in alternate formats upon request. Persons who want this publication in an alternate format should specify the type of format (cassette tape, Braille, large print, or ASCII disk). This document is also

available on the Board's Web site (<http://www.access-board.gov>).

Tricia Mason,

Chair, Architectural and Transportation Barriers Compliance Board.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

42 CFR Part 52

RIN 0925-AA42

Grants for Research Projects

AGENCY: National Institutes of Health, Department of Health and Human Services.

ACTION: Notice of proposed rulemaking.

SUMMARY: The National Institutes of Health (NIH) proposes to amend the existing regulations governing grants for research projects by revising the definition of Principal Investigator to mean one or more individuals designated by the grantee in the grant application and approved by the Secretary, who is or are responsible for the scientific and technical direction of the project, rather than limiting the role of principal investigator to one single individual, and the conditions for multiple or concurrent awards permitting the Secretary to evaluate, approve and make one or more awards pursuant to one or more applications. **DATES:** Comments must be received on or before August 24, 2007 in order to assure that NIH will be able to consider the comments in preparing the final rule.

ADDRESSES: Persons and organizations interested in submitting comments, identified by RIN 0925-AA42, may do so by any of the following methods:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the instructions for submitting comments.
- *E-mail:* jm40z@nih.gov. Include RIN number 0925-AA42 in the subject line of the message.
- *Fax:* 301-402-0169.
- *Mail:* Jerry Moore, NIH Regulations Officer, Office of Management Assessment, National Institutes of Health, 6011 Executive Boulevard, Suite 601, MSC 7669, Rockville, MD 20892.
- *Hand Delivery/Courier:* 6011 Executive Boulevard, Suite 601, Rockville, MD 20892.

FOR FURTHER INFORMATION CONTACT: Jerry Moore at the address above, or

telephone 301-496-4607 (not a toll-free number).

SUPPLEMENTARY INFORMATION: On September 30, 2003, NIH Director Elias A. Zerhouni announced a series of far-reaching strategic initiatives known collectively as the NIH Roadmap for Medical Research (NIH Roadmap). The NIH Roadmap is an innovative approach designed to transform the nation's medical research capabilities and accelerate fundamental research discovery and translation of that knowledge into effective prevention strategies and new treatments. One of the NIH Roadmap initiatives encourages interdisciplinary research and team science and includes a recommendation to modify grant and research contract applications to allow proposing of more than one Principal Investigator when appropriate. This is congruent with the January 4, 2005, directive issued by the President's Office of Science and Technology Policy (OSTP) to all Federal research agency heads instructing the heads to accommodate the recognition of two or more Principal Investigators on research projects (grants and contracts). While this new OSTP policy does not prohibit the use of a single Principal Investigator when that is most appropriate for a particular research project, it simply permits the designation of more than one Principal Investigator when that more accurately reflects the management needs of a research project.

For the purpose of implementing the NIH Roadmap initiatives, the NIH plans to modify research grant and contract applications to request information on more than one Principal Investigator, consistent with the new OSTP policy establishing the appropriateness of multiple Principal Investigators. Accordingly, we propose to revise the definition of the term Principal Investigator set forth in § 52.2 of the Grants for Research Projects regulations codified at 42 CFR Part 52, and the conditions for multiple or concurrent awards permitting the Secretary to evaluate, approve and make one or more awards pursuant to one or more applications.

Specifically, in this Notice of Proposed Rulemaking (NPRM) we propose to amend the existing regulations governing grants for research projects by revising the definition of Principal Investigator so that it does not limit the role of Principal Investigator to one single individual.

As announced in NIH notice number NOT-OD-07-017 (<http://grants.nih.gov/grants/guide/notice-files/NOT-OD-07-017.html>), these individual(s) must be

judged by the applicant organization to have the appropriate level of authority and responsibility to direct the project or program supported by the grant in order to be considered Principal Investigator(s). While this rule would permit the applicant organization to designate multiple individuals as Principal Investigators who share the authority and responsibility for leading and directing the project, intellectually and logistically, each Principal Investigator is responsible and accountable to the applicant organization (or, as appropriate, to a collaborating organization), for the proper conduct of the project or program, including the submission of all required reports. In other words, the presence of more than one identified Principal Investigator on an application or award diminishes neither the responsibility nor the accountability of any individual Principal Investigator.

Additionally, we propose to amend § 52.6 by revising paragraph (d) permitting the Secretary to evaluate, approve and make one or more awards pursuant to one or more applications.

Under current regulations, the Secretary is permitted to evaluate, approve and make more than one award pursuant to two or more applications. In some cases, however, it may be desirable to disaggregate a single application to make more than one award. For example, in the case of an application for support of a project that involves more than one Principal Investigator affiliated with more than one institution, it may be desirable to administer the project with more than one award. In addition, applications that involve subprojects may be disaggregated into separate awards to improve scientific management. The revised regulatory language clarifies options and provides an opportunity to contemplate more than one award that may involve more than one institution in response to a single application. In some of these cases separate records will be associated in the NIH data system so that the components can be managed as a single project to promote close collaboration with their counterparts. Actual awards also will be associated through special terms of award to clearly note collaborations and any special requirements resulting from such collaborations. In other cases, it may be appropriate to consider multiple applications from more than one institutions that are managed as a single unit with multiple awards to the different institutions to facilitate collaboration. This change will foster interdisciplinary and collaborative research and will improve management

flexibility even when components of such collaborative research programs are administered by different NIH awarding components.

The purpose of this NPRM is to invite public comment on the proposed regulation. The following is provided as public information.

Executive Order 12866

Executive Order 12866 requires that all regulatory actions reflect consideration of the costs and benefits they generate, and that they meet certain standards, such as avoiding the imposition of unnecessary burdens on the affected public. If a regulatory action is deemed to fall within the scope of the definition of the term "significant regulatory action" contained in § 3(f) of the Order, prepublication review by the Office of Management and Budget's Office of Information and Regulatory Affairs (OIRA) is necessary. This proposed rule was reviewed under Executive Order 12866 by OIRA and was deemed significant.

Executive Order 12866 also requires each agency to write all rules in plain language. In addition to your substantive comments on this proposed rule, we invite comments on how to make this proposed rule easier to understand. For example:

- Have we organized the material to suit your needs?
- Are the requirements in the rule clearly stated?
- Does the rule contain technical language or jargon that is not clear?
- Would a different format (grouping and order of sections, use of headings, paragraphing) make the rule easier to understand?
- Could we improve clarity by adding illustrative examples, tables, lists, or diagrams?
- What else could we do to make the rule easier to understand?

Regulatory Flexibility Act

The Regulatory Flexibility Act of 1980 (5 U.S.C. chapter 6) requires that regulatory proposals be analyzed to determine whether they create a significant impact on a substantial number of small entities. The Director, NIH, certifies that any final rule resulting from this proposed rule will not have any such impact.

Executive Order 13132

Executive Order 13132, Federalism, requires that federal agencies consult with State and local government officials in the development of regulatory policies with federalism implications. The Director, NIH, reviewed the proposed rule as required

under the Executive Order and determined that it does not have any federalism implications. The Director, NIH, certifies that the proposed rule will not have an effect on the States, or on the distribution of power and responsibilities among the various levels of government.

Paperwork Reduction Act

This proposed rule does not contain any information collection requirements which are subject to Office of Management and Budget (OMB) approval under the Paperwork Reduction Act of 1995 (44 U.S.C. chapter 35).

Catalog of Federal Domestic Assistance

The Catalog of Federal Domestic Assistance numbered programs affected by the proposed regulations are:

- 93.113—Biological Response to Environmental Health Hazards
- 93.114—Applied Toxicological Research and Testing
- 93.115—Biometry and Risk Estimation—Health Risks from Environmental Exposures
- 93.118—Acquired Immunodeficiency Syndrome (AIDS) Activity
- 93.121—Oral Diseases and Disorders Research
- 93.135—Centers for Research and Demonstration for Health Promotion and Disease Prevention
- 93.136—Injury Prevention and Control Research and State and Community Based Programs
- 93.172—Human Genome Research
- 93.173—Research Related to Deafness and Communication Disorders
- 93.184—Disabilities Prevention
- 93.213—Research and Training in Complementary and Alternative Medicine
- 93.242—Mental Health Research Grants
- 93.262—Occupational Safety and Health Program
- 93.271—Alcohol Research Career Development Awards for Scientists and Clinicians
- 93.273—Alcohol Research Programs
- 93.279—Drug Abuse and Addiction Research Programs
- 93.281—Mental Health Research Career/Scientist Development Awards
- 93.283—Centers for Disease Control and Prevention—Investigations and Technical Assistance
- 93.361—Nursing Research
- 93.389—National Center for Research Sources
- 93.390—Academic Research Enhancement Award
- 93.393—Cancer Cause and Prevention Research
- 93.394—Cancer Detection and Diagnosis Research

93.395—Cancer Treatment Research
 93.396—Cancer Biology Research
 93.821—Biophysics and Physiological Sciences Research
 93.837—Heart and Vascular Diseases Research
 93.838—Lung Diseases Research
 93.839—Blood Diseases and Resources Research
 93.846—Arthritis, Musculoskeletal and Skin Diseases Research
 93.847—Diabetes, Endocrinology and Metabolism Research
 93.848—Digestive Diseases and Nutrition Research
 93.849—Kidney Diseases, Urology and Hematology Research
 93.853—Clinical Research Related to Neurological Disorders
 93.855—Allergy, Immunology and Transplantation Research
 93.856—Microbiology and Infectious Diseases Research
 93.859—Biomedical Research and Research Training
 93.865—Research for Mothers and Children
 93.866—Aging Research
 93.867—Vision Research
 93.879—Medical Library Assistance
 93.929—Center for Medical Rehabilitation Research
 93.934—Fogarty International Center Research Collaboration Award
 93.939—Blood Diseases and Resources Research
 93.941—HIV Demonstration, Research, Public and Professional Education Projects
 93.942—Research, Treatment and Education Programs on Lyme Disease in the United States
 93.943—Epidemiologic Research Studies of Acquired Immunodeficiency Syndrome (AIDS) and Human Immunodeficiency Virus (HIV) Infection in Selected Population Groups
 93.947—Tuberculosis Demonstration, Research, Public and Professional Education

List of Subjects in 42 CFR Part 52

Grant programs—Health; Medical research; Occupational safety and health.

Dated: May 11, 2006.

Elias A. Zerhouni,

Director, National Institutes of Health.

Approved: October 12, 2006.

Michael O. Leavitt,
 Secretary.

Editorial Note: This document was received by the Office of the Federal Register on June 20, 2007.

For reasons presented in the preamble, it is proposed to amend part

52 of title 42 of the Code of Federal Regulations as set forth below.

PART 52—GRANTS FOR RESEARCH PROJECTS

1. The authority citation for part 52 continues to read as follows:

Authority: 42 U.S.C. 216.

1A. We propose to amend § 52.2 by revising the definition of the term “Principal investigator” to read as follows:

§ 52.2 Definitions.

* * * * *

Principal investigator means the individual(s) judged by the applicant organization to have the appropriate level of authority and responsibility to direct the project or program supported by the grant and who is or are responsible for the scientific and technical direction of the project.

* * * * *

2. We propose to amend § 52.6 by revising paragraph (d) to read as follows:

§ 52.6 Grant awards.

* * * * *

(d) *Multiple or concurrent awards.* Whenever a research project involves a number of different but related problems, activities or disciplines which require evaluation by different groups, or whenever support for a project could be more effectively administered by separate handling of separate aspects of the project, the Secretary may evaluate, approve and make one or more awards pursuant to one or more applications.

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DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

50 CFR Part 17

Endangered and Threatened Wildlife and Plants; 12-Month Finding on a Petition To List the Sierra Nevada Distinct Population Segment of the Mountain Yellow-Legged Frog (*Rana muscosa*)

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Notice of an amended 12-month petition finding.

SUMMARY: We, the U.S. Fish and Wildlife Service (Service), announce an amended 12-month finding on a petition

to list the Sierra Nevada distinct population segment (DPS) of the mountain yellow-legged frog (*Rana muscosa*) under the Endangered Species Act of 1973, as amended (Act). We are amending our previous 12-month petition finding, which found that listing is warranted but precluded, by revising the preclusion and expeditious progress section of that finding.

DATES: The finding announced in this document was made on June 25, 2007.

ADDRESSES: Supporting documentation used in the development of this amended 12-month finding will be available for inspection, by appointment, during normal business hours at the Endangered Species Program, Division of Conservation and Classification, U.S. Fish and Wildlife Service, 4401 N. Fairfax Drive, Room 420, Arlington, VA 22203. Comments and materials received, as well as supporting documentation used in the development of the initial 12-month finding published on January 16, 2003 (68 FR 2283), are available for inspection, by appointment, during normal business hours at the Sacramento Fish and Wildlife Office, U.S. Fish and Wildlife Service, 2800 Cottage Way, Room W-2605, Sacramento, CA 95825.

FOR FURTHER INFORMATION CONTACT: Chris Nolin, Chief, Division of Conservation and Classification, Endangered Species Program (see **ADDRESSES** section) (telephone 703-358-2171). Persons who use a telecommunications device for the deaf (TDD) may call the Federal Information Relay Service (FIRS) at 1-800-877-8339, 24 hours a day, 7 days a week.

SUPPLEMENTARY INFORMATION:

Background

Section 4(b)(3)(B) of the Endangered Species Act of 1973, as amended (Act) (16 U.S.C. 1531 *et seq.*), requires that, for any petition to revise the Lists of Endangered and Threatened Wildlife and Plants that contains substantial scientific or commercial information that the petitioned action may be warranted, we make a finding within 12 months of the date of the receipt of the petition on whether the petitioned action is: (a) Not warranted, (b) warranted, or (c) warranted, but that the immediate proposal of a regulation implementing the petitioned action is precluded by other pending proposals to determine whether any species is threatened or endangered, and expeditious progress is being made to add or remove qualified species from the Lists of Endangered and Threatened Wildlife and Plants (Lists). Such 12-