

course sequence, additional related courses open to students, time devoted to lecture, and clinical and research experience addressing the relationship with didactic programs in the educational process.

e. The extent to which the program effort is capable of supporting the number and type of students proposed.

f. Extent to which the program has initiated collaborative relationships with external agencies and institutions to expand and strengthen its research capabilities by providing student and faculty research opportunities.

g. Evidence of previous record of training in occupational injury prevention, including placement of graduates and employment history.

h. The extent to which the applicant documents methods in use or proposed methods for evaluating the effectiveness of the training, including the use of feedback mechanisms from graduates and employers, placement of graduates in research positions, research accomplishments of graduates and reports from consultations and cooperative activities with other universities, professional associations, and other outside agencies.

i. Competence, experience and training of the Program Director, faculty and advisors in relation to the type and scope of research training involved.

j. Degree of institutional commitment to Program goals. An example of institutional commitment to the long-term stability of academic programs is the commitment of tenured or tenure-track faculty positions to each participating academic program.

k. Adequacy of the academic and physical environment in which the training will be conducted, including access to appropriate occupational injury prevention research resources.

l. The extent to which the budget is adequate, justified, and consistent with the intended use of the grant funds.

m. Evidence of a plan for establishment of an Advisory Committee, including meeting times, roles and responsibilities.

I. Other Requirements

Technical Reporting Requirements

Provide CDC with original plus two copies of:

1. The initial interim progress report is due December 1, 2004. This report is required on December 1, on an annual basis. The progress report will serve as your non-competing continuation application, and must contain the following elements:

a. Current Budget Period Objectives and Activities.

b. Current Budget Period Financial Progress.

c. New Budget Period Program Proposed Activities and Objectives.

d. Detailed Line-Item Budget and Justification.

e. Additional Requested Information.

2. Financial status report, no more than 90 days after the end of each budget period. The initial report is due September 30, 2005.

3. Final financial and performance reports, no more than 90 days after the end of the project period.

Send all reports to the Grants Management Specialist identified in the "Where to Obtain Additional Information" section of this announcement.

Additional Requirements:

The following additional requirements are applicable to this program. For a complete description of each, see Attachment I of the program announcement, as posted on the CDC Web site.

AR-2* Requirements for Inclusion of Women and Racial and Ethnic Minorities in Research

AR-3* Animal Subjects Requirements

AR-10 Smoke-Free Workplace Requirements

AR-11 Healthy People 2010

AR-12 Lobbying Restrictions

*Applies only to ERC Pilot Project Research Training Program applications.

Executive Order 12372 does not apply to this program.

J. Where To Obtain Additional Information

This and other CDC announcements, the necessary applications, and associated forms can be found on the CDC Web site, Internet address: <http://www.cdc.gov>. Click on "Funding" then "Grants and Cooperative Agreements".

For general questions about this announcement, contact: Technical Information Management, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341-4146, Telephone: 770-488-2700.

For business management and budget assistance, contact: Cynthia Y. Mitchell, Grants Management Specialist, Procurement and Grants Office, Program Announcement 04001, Centers for Disease Control and Prevention (CDC), 626 Cochran Mill Rd., Mailstop P05, Pittsburgh, PA 15236, Telephone: (412) 386-6434, e-mail address: CMitchell@cdc.gov.

For program technical assistance, contact: John T. Talty, Principal Engineer, Office of Extramural Programs, National Institute for

Occupational Safety and Health, Centers for Disease Control and Prevention (CDC), 4676 Columbia Parkway, Mailstop C-7, Cincinnati, OH 45226-1998, Telephone (513) 533-8241, e-mail address: jtt2@cdc.gov.

Dated: May 9, 2003.

Sandra R. Manning,

Director, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC).

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02N-0077]

Agency Information Collection Activities; Announcement of OMB Approval; Emergency Medical Device Shortage Program Survey

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Emergency Medical Device Shortage Program Survey" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Peggy Robbins, Office of Information Resources Management (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1223.

SUPPLEMENTARY INFORMATION: In the *Federal Register* of March 17, 2003 (68 FR 12705), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910-0491. The approval expires on January 31, 2004. A copy of the supporting statement for this information collection is available on the Internet at <http://www.fda.gov/ohrms/dockets>.

Dated: May 9, 2003.

Jeffrey Shuren,

Assistant Commissioner for Policy.

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