

1. Quarterly data progress reports. These quarterly reports are required by the Office of Management and Budget (OMB) authorizing legislation (OMB Form 0920-0282.) The reports are due 30 days after the end of each quarter.

2. An interim progress narrative report, due no less than 90 days before the end of the budget period. This progress report will serve as your non-competing continuation application, and must contain the following elements:

a. Progress on Current Budget Period Objectives and Activities.

b. Current Budget Period Financial Progress.

c. New Budget Period Proposed Program Objectives and Activities.

d. Detailed Line-Item Budget and Justification.

3. Calendar-year surveillance data, submitted annually in the approved OMB format, no later than April 30. In addition, a written surveillance summary must be disseminated to state and local public health officials, policy makers, the CDC project officer, and others.

4. Financial Status Reports, due within 90 days of the end of the budget period.

5. Final financial reports and performance reports, due within 90 days after the end of the project period.

6. Projects that involve the collection of information from 10 or more individuals, and are funded by a cooperative agreement will be subject to review by OMB under the Paperwork Reduction Act. Data collection initiated under this cooperative agreement program has been approved by OMB under OMB number 0920-0337, "National Childhood Blood Lead Surveillance System", Expiration Date: 6/30/2004.

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-9, Paperwork Reduction Act Requirements

AR-10, Smoke-Free Workplace Requirements

AR-11, Healthy People 2010

AR-12, Lobbying Restrictions

AR-21, Small, Minority & Women-Owned Businesses

Executive Order 12372 does not apply to this program.

J. Where To Obtain Additional Information

Two telephone conference calls for application technical assistance will be held during the application period. For further information, please contact Rob Henry at (770) 488-4024. This, and other CDC announcements, necessary applications, and associated forms can be found on the CDC home page Internet address: <http://www.cdc.gov>. Click on "Funding", then "Grants and Cooperative Agreements."

For general questions regarding this announcement, contact: Technical Information Management, CDC Procurement and Grants Office, 2920 Brandywine Road, Room 3000, Atlanta, GA 30341-4146, Telephone: (770) 488-2700.

For business management and budget assistance, contact: Mildred Garner, Grants Management Officer, CDC Procurement and Grants Office, 2920 Brandywine Road, Room 3000, Atlanta, GA 30341-4146, Telephone: (770) 488-2745, E-mail address: mgarner@cdc.gov.

For business management and budget assistance in the territories, contact: Charlotte Flitcraft, Grants Management Officer, CDC Procurement and Grants Office, 2020 Brandywine Rd., Room 3000, Atlanta, GA 30319, Telephone: 770-488-2632, E-mail address: caf5@cdc.gov.

For program technical assistance, contact: Rob Henry, Acting Team Leader, Program Services Section, Lead Poisoning Prevention Branch, Centers for Disease Control and Prevention, 1600 Clifton Rd, NE, MS E-25, Atlanta, GA 30333, Telephone: (770) 488-4024, E-mail address: rhenry@cdc.gov.

Dated: December 31, 2002.

Sandra R. Manning,

CGFM, Director, Procurement and Grants Office, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 03N-0009]

Agency Information Collection Activities; Proposed Collection; Comment Request; Application for Exemption From Federal Preemption of State and Local Medical Device Requirements

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on FDA's requirements for State and local governments' applications for exemption from preemption for medical device requirements.

DATES: Submit written or electronic comments on the collection of information by March 24, 2003.

ADDRESSES: Submit electronic comments on the collection of information to <http://www.accessdata.fda.gov/scripts/oc/dockets/edockethome.cfm>. Submit written comments on the collection of information to the Dockets Management Branch (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

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: Under the PRA (44 U.S.C. 3501-3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. "Collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or