

proposes to carry out the second phase of a net-cost analysis of a program of prenatal and infancy home visiting.

This project is being funded noncompetitively, because of its uniqueness in examining cost savings to government resulting from an investment in a program of prenatal and early childhood home visitation. The duration of this project is two years, beginning April 1, 2000, for a total cost of \$312,766.

FOR FURTHER INFORMATION CONTACT: K.A. Jagannathan, Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade, SW, Washington, DC 20447, Phone: 202-205-4829.

Dated: March 31, 2000.

Howard Rolston,

Director, Office of Planning, Research and Evaluation.

[FR Doc. 00-8409 Filed 4-5-00; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Senior Executive Service; Performance Review Board Members

AGENCY: Administration for Children and Families.

ACTION: Announcing appointment of Performance Review Board members.

SUMMARY: 5 U.S.C. 4314(c)(4) of the Civil Service Reform Act of 1978, Pub. L. 95-454, requires that the appointment of Performance Review Board members be published in the **Federal Register**.

The following persons will serve on the Performance Review Boards or Panels which oversee the evaluation of performance appraisals of Senior Executive Service members of the Department of Health and Human Services, Administration for Children and Families: Diann Dawson, Leon R. McCowan, Madeline Mocko.

Dated: March 29, 2000.

Elizabeth M. James Duke,

Deputy Assistant Director for Administration, Office of Administration.

[FR Doc. 00-8413 Filed 4-5-00; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N-1060]

Agency Information Collection Activities; Proposed Collection; Comment Request; Adoption of the FDA Food Code by Local, State, and Tribal Governments

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, and to allow 60 days for public comment in response to the notice. This notice solicits comments on FDA's collection of information from local, State, and tribal agencies concerning their adoption of, or plans to adopt, all or portions of the FDA Food Code or its equivalent by regulation, law, or ordinance. The Association of Food and Drug Officials (AFDO) has been contracted by FDA to telephonically and/or electronically contact local, State, and tribal food program administrators to determine Food Code adoption in their respective jurisdictions.

DATES: Submit written comments on the collection of information by June 5, 2000.

ADDRESSES: 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: JonnaLynn P. Capezuto, Office of Information Resources Management (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-4659.

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 provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Adoption of the FDA Food Code by Local, State, and Tribal Governments

FDA has developed the model Food Code to assist and promote consistent implementation of national food safety regulatory policy among the several thousand local, State, and tribal jurisdictions that have primary responsibility for the regulation or oversight of retail level food operations. The FDA Food Code provides a scientifically sound technical and legal basis for regulating the retail segment of the food industry. Authority for providing such assistance is derived from section 311(a) of the Public Health Service Act (42 U.S.C. 243) and delegation of authority from the Public Health Service to the Commissioner of Food and Drugs relative to food protection is contained in 21 CFR 5.10(a)(2) and (a)(4). Under 31 U.S.C. 1535, FDA provides assistance to other Federal agencies such as the Indian Health Service.

Nationwide adoption of the model FDA Food Code is an important step to further the goals of the President's Council on Food Safety for consistent, scientifically sound, and risk-based food safety standards and practices and to work more effectively with partners in State, local, and tribal governments. FDA has established a site on the Internet at <http://www.cfsan.fda.gov> under "Federal/State Food Programs"

and "Retail Food Safety References" to list jurisdictions that have reported adoptions of the FDA Food Code. Because it is self-reported, the list is incomplete and has not been evaluated to determine whether all the adopted codes are equivalent to the model Food Code. It is important to FDA to have a comprehensive, accurate, and current inventory of Food Code adoptions to help achieve the aims of the President's Council on Food Safety and the agency's Food Safety Initiative goals.

FDA has obtained the services of AFDO to develop and implement an active surveillance system to track and report on the adoption of the FDA Food Code by State and local agencies and tribal nations of native Americans. The

contractor will develop and maintain an active data base to track adoptions of the Food Code; identify and periodically contact State, local, and tribal food safety program administrators to determine the current status of adoptions of the Food Code or its equivalent; evaluate the equivalency of the adopted codes with the FDA Food Code; and provide quarterly progress reports to FDA from the data base in tabular and graphic form. Reports may be placed on the Internet at <http://www.fda.gov>.

Initial contacts by AFDO to local, State, and tribal program administrators will be by telephone and/or e-mail to determine the Food Code status in their jurisdiction(s). Verbal responses to

questions will be acceptable as will electronic or facsimile information. Followup contacts to clarify responses will be by telephone or e-mail to minimize the burden on respondents.

The types of questions to be asked will be whether or not the FDA Food Code has been adopted in the respondent's jurisdiction, which version of the Food Code is in effect, and if not, which local jurisdictions need to be contacted for Food Code adoption status. AFDO will also determine with the local/State/tribal governments that it has the latest version of the code for analysis.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN¹

No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
500	2	1,000	1	1,000
Total Hours				1,000

¹ There are no capital costs or operating and maintenance costs associated with this collection of information.

FDA based its estimate on the number of State agencies (100) involved in Food Code-related regulatory programs, 300 local agencies with local ordinance authority that may consider Food Code adoption in any one year, and 100 tribal agencies. Estimating the number of local agencies is difficult before the start of this project because in some States, adoption by a State agency automatically applies to all local jurisdictions in that state. In other States, some metropolitan jurisdictions may adopt the FDA Food Code individually. Similar circumstances may apply to tribal nations' agencies that may be adopting the FDA Food Code. When the initial information gathering is completed, FDA will be able to identify more accurately the number of local and tribal agencies for which tracking adoption of the FDA Food Code will be necessary.

Frequency of reporting will range from once per year to quarterly for any one jurisdiction. This is because agencies that have already adopted the Food Code will require less frequent contact, perhaps only annually, than those that are in the process of adopting the Food Code. An average of two contacts in 1 year, therefore, was selected. Because most reporting will be done telephonically or electronically, reporting times often will be much less than 1 hour.

These estimates will fluctuate from year to year as agencies adopt, revise,

and consider adoption of the FDA Food Code. Over the next 3 years, the frequency of contacts should decrease as jurisdictions adopt the FDA Food Code. This project will take several years to complete because the adoption process in some States can extend to 2 years or more. For example, some States have biennial legislative sessions. Others have extensive notice-and-comment administrative rulemaking procedures that can extend well beyond 1 year.

Dated: March 30, 2000.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 00-8416 Filed 4-5-00; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99N-4166]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Electronic Records; Electronic Signatures

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the proposed collection of

information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments on the collection of information by May 8, 2000.

ADDRESSES: Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Wendy Taylor, Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: JonnaLynn P. Capezzuto, Office of Information Resources Management (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-4659. **SUPPLEMENTARY INFORMATION:** In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Electronic Records; Electronic Signatures—Part 11 (21 CFR Part 11) (OMB Control Number 0910-0303)—Extension

FDA regulations in part 11 (21 CFR part 11) provide criteria for acceptance by FDA of electronic records, electronic signatures, and handwritten signatures executed to electronic records as equivalent to paper records and handwritten signatures executed on