Respondents: The respondents to these follow-up surveys will be low-income individuals from the five states represented by the four sites currently participating in the HtE Project: Kansas, Missouri, New York, Pennsylvania and Rhode Island. Many will be current or former TANF participants, and many will be current or former recipients of Medicaid. These populations are at heightened risk for all of the barriers that cause people to be hard-to-employ.

Prior to these follow-up surveys, basic demographic information for all survey respondents will have been obtained wherever possible from the existing automated systems or brief baseline information forms. In the Rhode Island site, respondents will have completed a more detailed baseline survey, which is required to establish baseline measures of depression and related conditions.

The annual burden estimates are detailed below, and the substantive content of each survey are detailed in the supporting statement.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Rhode Island, 6–month	734	1	38 minutes or .63 hrs	464.87
Rhode Island, 15–month	734	1	45 minutes or .75 hrs	550.50
New York City, 12–month	1,000	1	32 minutes or .53 hrs	533.33
Philadelphia, 12–month	750	1	25 minutes or .42 hrs	312.50
Kansas/Missouri, 12–month	680	1	45 minutes or .75 hrs	510.00

Estimated Total Annual Burden Hours. 2,371.20

Additional Information

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families. Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. E-mail address: *grjohnson@acf.hhs.gov.*

OMB Comment

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register.** Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Attn: Desk Officer for ACF, E-mail address: *Katherine_T._Astrich@omb.eop.gov.*

Dated: February 8, 2005

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 05–2825 Filed 2–14–05; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request Proposed Projects:

Title: Community-Based Child Abuse Prevention Program (CBCAP).

OMB No.: 0970–0155.

Description: The Program Instruction, prepared in response to the enactment of the Community-Based Grants for the Prevention of Child Abuse and Neglect (administratively known as the Community-Based Child Abuse Prevention Program (CBCAP)), as set forth in Title II of Pub. L. 108–36, Child Abuse Prevention and Treatment Act Amendments of 2003, provides direction to the States and Territories to accomplish the purposes of (1) supporting community-based efforts to develop, operate, expand and, where appropriate, to network initiatives aimed at the prevention of child abuse and neglect and to support networks of coordinated resources and activities to better strengthen and support families to reduce the incidence of child abuse and neglect; and (2) fostering an understanding, appreciation and knowledge of diverse populations in order to be effective in preventing and treating child abuse and neglect. This Program Instruction contains information collection requirements that are found in Pub. L. 108–36 at Sections 201, 202, 203, 205, 206, 207, and pursuant to receiving a grant award. The information submitted will be used by the agency to ensure compliance with the statute, complete the calculation of the grant award entitlement, and provide training and technical assistance to the grantee.

Respondents: State Government.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Application	52	1	40	2,080
Annual Report	52	1	24	1,248

Estimated Total Annual Burden Hours 3,328

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF

Reports Clearance Officer. All requests should be identified by the title of the information collection. E-mail: *grjohnson@acf.hhs.gov*.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility, (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: February 8, 2005.

Robert Sargis,

Reports Clearance, Officer.

[FR Doc. 05–2826 Filed 2–14–05; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005D–0042]

Draft Guidance on the Open Public Hearing; Food and Drug Administration Advisory Committee Meetings; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance document entitled ''The Open Public Hearing; FDA Advisory Committee Meetings.'' This draft guidance is for members of the public who choose to participate in the open public hearing (OPH) session of an FDA advisory committee meeting. The draft guidance is intended to answer more fully questions about how the public may participate at an OPH session, and it includes topics such as meeting logistics and administrative requirements.

DATES: Submit written or electronic comments on this draft guidance by June 15, 2005. General comments on agency guidance documents are welcome at any time.

ADDRESSES: Submit written requests for single copies of the draft guidance to Linda Ann Sherman, Advisory Committee Oversight and Management Staff (HF–4), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857. Submit written comments on the draft guidance to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.fda.gov.dockets/ecomments*. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.

FOR FURTHER INFORMATION CONTACT: Linda Ann Sherman, Office of the Commissioner (HF–4), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1220, e-mail: *disclosure@oc.fda.gov*.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a draft guidance document entitled ''The Open Public Hearing; FDA Advisory Committee Meetings.''

Guidance documents are prepared for FDA staff, applicants/sponsors, and the public that describe the agency's interpretation of, or policy on, a regulatory issue. Every committee meeting includes an OPH during which interested persons may present relevant information or views orally or in writing 21 CFR 14.25(a). The hearing is conducted in accordance with 21 CFR 14.29. FDA encourages the participation from all public speakers in its decisionmaking processes. The draft guidance is intended to answer more fully questions about how (including topics such as meeting logistics and administrative requirements) the public may participate at an OPH session. This includes, but is not limited to, general members of the public; individuals or spokespersons from the regulated industry; consumer advocacy groups; and professional organizations, societies, or associations.

This level 1 draft guidance is being issued consistent with FDA's good guidance practices (21 CFR 10.115). The draft guidance, when finalized will represent the agency's current thinking on an FDA advisory committee open public hearing. It does not create or confer any rights for or on any person and does not operate to bind FDA or the public. An alternative approach may be used if such approach satisfies the requirements of the applicable statute and regulations.

II. Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) written or electronic comments on the draft guidance. Two paper copies of mailed comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

III. Electronic Access

Persons with access to the Internet may obtain the document at *http://www.fda.gov/oc/advisory/default.htm* in the policy and guidance section of FDA's advisory committee Intranet Web site.

Dated: February 8, 2005.

William K. Hubbard,

Associate Commissioner for Policy and Planning.

[FR Doc. 05–2822 Filed 2–14–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005D–0033]

Draft Guidance for Industry on Internal Radioactive Contamination—Development of Decorporation Agents; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance for industry entitled ''Internal Radioactive Contamination—Development of Decorporation Agents.'' This draft document provides guidance to industry on the development of decorporation agents for the treatment of internal radioactive contamination when evidence is needed to demonstrate the effectiveness of the agents, but human efficacy studies are unethical or infeasible. In such instances, the Animal Efficacy Rule may be invoked to approve new medical products not previously marketed or new indications for previously marketed products. Specifically, this draft guidance addresses chemistry, manufacturing and controls (CMC) information; animal efficacy, safety pharmacology, and toxicology studies; clinical pharmacology, biopharmaceutics, and human safety studies; and postapproval commitments.

DATES: Submit written or electronic comments on the draft guidance by May