SYSTEM MANAGER(S) AND ADDRESS:

Director, Division of Data Liaison and Distribution, Enterprise Databases Group, Office of Information Services, CMS, Room N2–04–27, 7500 Security Boulevard, Baltimore, Maryland, 21244–1850.

NOTIFICATION PROCEDURE:

For purpose of access, the subject individual should write to the system manager, who will require the system name, the subject individual's name (woman's maiden name, if applicable), social security number (SSN) (furnishing the SSN is voluntary, but it may make searching for a record easier and prevent delay), address, date of correspondence and control number.

RECORD ACCESS PROCEDURE:

For purpose of access, use the same procedures outlined in Notification Procedures above. Requestors should also reasonably specify the record contents being sought. (These procedures are in accordance with Department regulation 45 CFR 5b.5(a)(2).)

CONTESTING RECORD PROCEDURES:

The subject individual should contact the system manager named above, and reasonably identify the record and specify the information to be contested. State the corrective action sought and the reasons for the correction with supporting justification. (These procedures are in accordance with Department regulation 45 CFR 5b.7.)

RECORD SOURCE CATEGORIES:

CMS's National Claims History system of records, Enrollment Database system of records, Medicare Beneficiary Database system of records, and Medicaid Statistical Information System of records.

SYSTEMS EXEMPTED FROM CERTAIN PROVISIONS OF THE ACT:

None.

[FR Doc. 02–25427 Filed 10–4–02; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Online Interstate Referral Guide (IRG).

OMB No.: 0970–0209.

Description: The IRG is an essential reference maintained by OCSE that provides States with an effective and efficient way of viewing and updating State profile, address, and FIPS code information by consolidation data available through numerous discrete sources into a single centralized, automated repository.

Respondents: State IV–D Child Support Programs.

Annual Burden Estimates:

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Online IRG	54	18	.3	292

Estimated Total Annual Burden Hours: 292.

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 Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 02447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection.

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: October 1, 2002.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 02–25424 Filed 10–4–02; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: OCSE–369A: Financial Report; and OCSE–34A: Quarterly Report of Collections.

OMB No.: 0970–0181.

Description: Each State agency administering the Child Support Enforcement Program under Title IV–D of the Social Security Act is required to provide information to the Office of Child Support Enforcement concerning its administrative expenditures and its receipt and disposition of child support payments from non-custodial parents. These quarterly reporting forms enable each State to provide that information, which is used to compute both the quarterly grants awarded to each State and the annual incentive payments earned by each State. This information is also included in a published annual statistical and financial report, available to the general public.

Comments sent to the Office of Child Support Enforcement, both directly and in response to an earlier **Federal Register** Notice (67 FR 39727, *et seq.*), provided many useful recommendations to update and correct these financial reporting forms. However, several comments strongly indicated that State agencies would have inadequate time to incorporate these revisions in time to meet the reporting requirements for the fiscal year beginning October 1, 2002. In addition, legislation has been introduced in Congress that, if enacted, may require additional revisions to these forms.

For these reasons, we have decided to request that the expiration date of the existing forms be extended, without change, through September 30, 2004.

Respondents: State agencies administering the Child Support Enforcement Program.

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
OCSE–396A	54	4	8	1,728
OCSE–34A	54	4	8	1,728

Estimated Total Annual Burden Hours: 3,456.

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.

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, 725 17th Street, NW., Washington, DC 20503, Attn: Desk Officer for ACF.

Dated: October 1, 2002.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 02–25425 Filed 10–4–02; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Los Angeles District Office; Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of meeting.

The Food and Drug Administration (FDA) is announcing a meeting which is intended to give the drugs, devices, and biologics industries and consumers an opportunity to exchange information with the FDA Los Angeles District staff. The main focus of the meeting is to provide an opportunity for the Los Angeles District leadership to interact with industry and the public, and to discuss regulatory affairs, plans, and future programs. The open house is sponsored by the Orange County Regulatory Affairs Discussion Group (OCRA).

Date and Time: The open house will be held on Tuesday, October 22, 2002, from 6 p.m. to 9 p.m.

Location: The open house will be held at FDA Los Angeles District, 19900 MacArthur Boulevard, suite 300, Irvine, CA 92612.

Contact: Ramlah Oma, Food and Drug Administration, 19900 MacArthur Blvd., suite 300, Irvine, CA 92612, 949–798–7611, FAX: 949–798–7656, or Jack Dhuwalia, OCRA, PMB 624, 5405 Alton Pkwy, suite 5A, Irvine, CA 92604, 888–532–4357, FAX: 949–854–2672, Internet: www.ocra-dg.org.

Registration and open house Information: For registration information, including registration form and electronic payment, see the OCRA Internet site at www.ocra-dg.org (click on '' OCRA meetings'').

Registrations fees are $40.00 for members of OCRA, Southern California Pharmaceutical Discussion Group (SCPDG), and Parenteral Drug Association (PDA), and $45.00 for nonmembers. The cost includes hot and cold hors d'oeuvres, dessert and nonalcoholic beverages, but excludes parking fees.

If you need special accommodations due to a disability, please contact Ramlah Oma (see *Contact*) at least 7 days in advance.

Dated: October 1, 2002.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 02–25392 Filed 10–4–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02D–0385]

Guidance on the Petition Process to Request Approval of Labeling for Foods That Have Been Treated By Irradiation; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a guidance document entitled ''Guidance; Implementation of Section 10809 of the Farm Security and Rural Investment Act of 2002, Pub. L. No. 107–171, § 10809 (2002) Regarding the Petition Process to Request Approval of Labeling for Foods That Have Been Treated By Irradiation,'' which explains the recommended process for petitioning the agency for approval of labeling, which is not false or misleading in any material respect, of a food that has been treated by irradiation.

DATES: Submit written or electronic comments at any time.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to http://www.fda.gov/dockets/ecomments.

FOR FURTHER INFORMATION CONTACT: Loretta A. Carey, Center for Food Safety and Applied Nutrition (HFS–822), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 301–436–2371.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a guidance document implementing the part of section 10809 of the Farm Security and Rural Investment Act of 2002 (Public Law 107–171, § 10809 (2002)), that states that ''[p]ending promulgation of the final rule * * *, any person may petition the Secretary [FDA] for approval of labeling, which is not false or misleading in any material respect, of a food which has been treated by irradiation using radioactive isotope, electronic beam, or x-ray.'' Section 10809 of the Farm Security and Rural Investment Act of 2002 also requires that, pending promulgation of the final rule, ''[t]he Secretary [FDA] shall approve or deny such a petition within 180 days of receipt of the petition, or the petition shall be deemed denied, except to the extent additional agency review is mutually agreed upon by the Secretary [FDA] and the petitioner.''

FDA is issuing this guidance to interested parties who wish to petition the agency for approval of the labeling of a food treated by irradiation. As explained in the guidance, FDA recommends that interested parties who wish to petition the agency use the procedures set forth in § 10.30 (21 CFR 10.30), except that § 10.30(e)(2)(iii), regarding 180-day tentative responses,