when the list of FCC ICRs currently under review appears, look for the OMB control number of this ICR and then click on the ICR Reference Number. A copy of the FCC submission to OMB will be displayed.

SUPPLEMENTARY INFORMATION:
OMB Control Number: 3060–0179.
Title: Section 73.1590, Equipment Performance Measurements.
Form Number: N/A.
Type of Review: Extension of a currently approved collection.
Respondents: Business or other for-profit entities; not-for-profit institutions.
Number of Respondents: 13,049.
Estimated Time per Response: 0.5–18 hours.
Frequency of Response: Recordkeeping requirement.
Total Annual Burden: 12,335 hours.
Total Annual Cost: None
Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this collection is contained in Section 154(i) of the Communications Act of 1934, as amended.
Nature and Extent of Confidentiality: There is no need for confidentiality with this collection of information.
Privacy Impact Assessment: No impact(s).
Needs and Uses: 47 CFR 73.1590(d) requires licensees of AM, FM and TV stations to make audio and video equipment performance measurements for each main transmitter. These measurements and a description of the equipment and procedures used in making the measurements must be kept on file at the transmitter or remote control point for 2 years. In addition, this information must be made available to the FCC upon request.
OMB Control Number: 3060–0500.
Title: Section 76.1713, Resolution of Complaints.
Form Number: N/A.
Type of Review: Extension of a currently approved collection.
Respondents: Business or other for-profit entities.
Number of Respondents and Responses: 10,750 respondents and 21,500 responses.
Estimated Hours per Response: 1–17 hours.
Frequency of Response: Recordkeeping and third party disclosure requirements; annual reporting requirement.
Total Annual Burden: 193,500 hours.
Total Annual Cost: None.
Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this collection is contained in Section 76.1713 of the Communications Act of 1934, as amended.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Decision To Evaluate a Petition To Designate a Class of Employees From the Kansas City Plant in Kansas City, Missouri, To Be Included in the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: NIOSH gives notice as required by 42 CFR 83.12(e) of a decision to evaluate a petition to designate a class of employees from the Kansas City Plant in Kansas City, Missouri, to be included in the Special Exposure Cohort under the Energy Employees Occupational Illness Compensation Program Act of 2000. The initial proposed definition for the class being evaluated, subject to revision as warranted by the evaluation, is as follows:

Facility: Kansas City Plant.
Location: Kansas City, Missouri.
Job Titles and/or Job Duties: All employees who worked in any area.

FOR FURTHER INFORMATION CONTACT: Stuart L. Hinnefeld, Director, Division of Compensation Analysis and Support,
In compliance with the requirements of Section 506(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 Planning, Research and Evaluation, 370 L’Enfant Promenade SW., Washington, DC 20447, Att: ACF Reports Clearance Officer. Email address: infocollection@acf.hhs.gov. 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.

Robert Sargis, Reports Clearance Officer.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0764]

Agency Information Collection Activity; Proposed Collection; Comment Request; Draft Animal Feed Regulatory Program Standards; Availability

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 the information collection associated with the draft Animal Feed Regulatory Program Standards (AFRPS). The draft feed standards are neither final nor intended for implementation at this time.

DATES: Submit either electronic or written comments on the collection of information by September 9, 2013.

ADDRESSES: Submit electronic comments on the collection of information to http://www.regulations.gov. Submit written comments on the collection of information to the Division of Dockets Management (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. Submit written requests for single copies of the draft feed standards to the U.S. Food and Drug Administration, Office of Regulatory Affairs, Office of Partnerships, 12420 Parklawn Dr., ELEM–3033, Rockville, MD 20857. Send one self-addressed adhesive label to assist the office in processing your request, or fax your request to 301–827–3588. See the SUPPLEMENTARY INFORMATION section for an electronic copy of the draft feed standards.

FOR FURTHER INFORMATION CONTACT: With regard to the information collection:

Ila S. Mizrahi, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850,