FEDERAL COMMUNICATIONS COMMISSION

Notice of Public Information Collection(s) Being Reviewed by the Federal Communications Commission for Extension Under Delegated Authority


SUMMARY: The Federal Communications Commission, as part of its continuing effort to reduce paperwork burden invites the general public and other Federal agencies to take this opportunity to comment on the following information collection(s), as required by the Paperwork Reduction Act (PRA) of 1995, Public Law 104-13. An agency may not conduct or sponsor a collection of information unless it displays a currently valid control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the Paperwork Reduction Act that does not display a valid control number. Comments are requested concerning (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; (b) the accuracy of the Commission’s burden estimate; (c) ways to enhance the quality, utility, and clarity of the information collected; and (d) ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology.

DATES: Written Paperwork Reduction Act (PRA) comments should be submitted on or before January 31, 2005. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

ADDRESSES: Direct all Paperwork Reduction Act (PRA) comments to Judith B. Herman, Federal Communications Commission, Room 1-A, C804, 445 12th Street, SW., Washington, DC 20554 or via the Internet to Judith-B.Herman@fcc.gov.

FOR FURTHER INFORMATION CONTACT: For additional information or copies of the information collection(s), contact Judith B. Herman at 202-418-0214 or via the Internet at Judith-B.Herman@fcc.gov.

SUPPLEMENTARY INFORMATION: OMB Control No.: 3060-0513.
Number of Respondents: 83.
Estimated Time Per response: 50 hours.
Frequency of Response: Annual reporting requirement.
Total Annual Burden: 4,150 hours.
Total Annual Cost: Not Applicable.
Privacy Act Impact Assessment: Not Applicable.
Needs and Uses: The Joint Cost Report is needed to administer our joint cost rules (Part 64) and to analyze data in order to prevent cross-subsidization of non-regulated operations by the regulated operations of Tier 1 carriers. The information contained in the ARMS Report 43-03 provides the necessary detail to enable the Commission to fulfill its regulatory responsibilities. Automated reporting of these data greatly enhances the Commission’s ability to process and analyze the extensive amounts of data that it needs to administer its rules. ARMS facilitates the timely and efficient analysis of revenue requirements, rates of return and price caps, and provides an improved basis for auditing and other oversight functions. It also enhances the Commission’s ability to quantify the effects of policy proposals.

FOR FURTHER INFORMATION CONTACT:
Michelle A. Smith, Director, Office of Board Members; 202-452-2955.

SUPPLEMENTARY INFORMATION: You may call 202-452-3206 beginning at approximately 5 p.m. two business days before the meeting for a recorded announcement of bank and bank holding company applications scheduled for the meeting; or you may contact the Board’s Web site at http://www.federalreserve.gov for an electronic announcement that not only lists applications, but also indicates procedural and other information about the meeting.


Robert deV. Frierson, Deputy Secretary of the Board.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Information Collections Related to Reunification Procedures for Unaccompanied Alien Children. OMB No: New Collection.

Description: Following the passage of the 2002 Homeland Security Act (Pub. L. 107-296), the Administration for Children and Families (ACF), Office of Refugee Resettlement (ORR), is charged with the care and placement of unaccompanied alien children in Federal custody, and implementing a policy for the release of these children, when appropriate, upon the request of suitable sponsors while awaiting immigration proceedings. In order for ORR to make determinations regarding the release of these children, the potential sponsors must meet certain conditions pursuant to section 462 of the Homeland Security Act and the Flores v. Reno settlement agreement, No. CV85-4544-RJK (C.D. Cal. 1997). ORR considers the suitability of a sponsor based on the sponsor’s ability...
and agreement to provide for the physical, mental and financial well-being of an unaccompanied minor and the sponsor’s assurance to appear before immigration courts. To ensure the safety of the children, sponsors must undergo a background check. Suitable sponsors may be parents, close relatives, friends or entities concerned with the child’s welfare. In this Notice, ACF announces that it proposes to employ the use of several information collections for recording: (1) The Sponsor’s Agreement to Conditions of Release, which collects the sponsor’s affirmation to the terms of the release; (2) the Verification of Release, which collects the children’s affirmation to the terms of their release; (3) the Family Reunification Packet, which collects information related to the sponsor’s ability to provide for the physical, mental and financial well-being of the child(ren); and (4) the Authorization for Release of Information, which collects information to be utilized for a background check.


### ANNUAL BURDEN ESTIMATES

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Total burden hours</th>
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<td>Family Reunification Packet</td>
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<td>3,000</td>
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<tr>
<td>Authorization for Release of Information</td>
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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**[Docket No. 2004N–0498]**

**Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Devices; Device Tracking**

**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 information 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 information collection requirements for tracking of medical devices.

**DATES:** Submit written or electronic comments on the collection of information by January 31, 2005.

**ADDRESSES:** Submit electronic comments on the collection of information to: http://www.fda.gov/dockets/ecomments. 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.

**FOR FURTHER INFORMATION CONTACT:** Peggy Robbins, Office of Management Programs (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 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, including each proposed extension of an existing 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 these topics: (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