DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Division of Unaccompanied Children's Services (DUCS) Request for Specific Consent.

OMB No.: New Collection.

Description: The William Wilberforce Trafficking Victims Protection Reauthorization Act of 2008 (TVPRA of 2008), Public Law 110–457 was enacted into law December 23, 2008. Section 235(d) directs the Secretary of HHS to grant or deny requests for specific consent for unaccompanied alien children in HHS custody who seek to invoke the jurisdiction of a state court for a dependency order and who also seek to invoke the jurisdiction of a state court to determine or alter his or her custody status or release from ORR. These requests can be extremely time sensitive since a child must ask a state court for dependency before turning 18 years old.

In developing procedures for collecting the necessary information from unaccompanied alien children, their attorneys, or other representatives to allow HHS to approve or deny consent requests, ORR/DUCS devised a form. Specifically, the form asks the requestor for his/her identifying information, basic identifying information on the unaccompanied alien child, the name of the HHS-funded facility where the child is in HHS custody and care, the name of the court and its location, and the kind of request (*e.g.,* for a change in custody, etc.). The form also asks that the unaccompanied alien child's attorney or authorized representative attach a Notice of Representation, which is an approved federal government agency form used for immigration procedures that authorizes the attorney to act on behalf of the child (*i.e.,* G–28, EOIR–28, EOIR–29), or any other form of authorization to act on behalf of the unaccompanied alien child.

Respondents: Attorneys, accredited legal representatives, or others authorized to act on behalf of a unaccompanied alien child.

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Request for Specific Consent to Juvenile Court Jurisdiction (ORR–0132)	72	1	0.3333	24

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: *infocollection@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, Fax: 202–395–7285, E-mail: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2010–32504 Filed 12–27–10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Notice of Interstate Lien.

OMB No.: 0970–0153.

Description: Section 452(a)(11) of the Social Security Act requires the Secretary of Health and Human Services to promulgate a form for imposition of liens to be used by the State child support enforcement (Title IV–D) agencies in interstate cases. Section 454(9)(E) of the Social Security Act requires each State to cooperate with any other State in using the Federal form for imposition of liens in interstate child support cases. Tribal IV–D agencies are not required to use this form but may choose to do so. OMB approval of this form is expiring in February 2011 and the Administration for Children and Families is requesting an extension of this form.

Respondents: State, local or Tribal agencies administering a child support enforcement program under title IV–D of the Social Security Act.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondents	Average burden hours per responses	Total burden hours
Notice of Lien	1,832,384	1	0.25	458,096

Estimated Total Annual Burden Hours: 458,096.

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: infocollection@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. Fax: 202–395–7285. *E-mail: OIRA_SUBMISSION@OMB.EOP.GOV. Attn:* Desk Officer for the Administration for Children and Families.

Dated: December 22, 2010.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2010–32592 Filed 12–27–10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0198]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Premarket Notification

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Premarket Notification" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Daniel Gittleson, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–5156, e-mail: *Daniel.Gittleson@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In the **Federal Register** of August 11, 2010 (75 FR 48696), the Agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0120. The approval expires on December 31, 2013. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: December 21, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010–32508 Filed 12–27–10; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0447]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medical Devices; Third Party Review Program Under the Food and Drug Administration Modernization Act

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (the PRA).

DATES: Fax written comments on the collection of information by January 27, 2011.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or e-mailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0375. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Daniel Gittleson, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–5156, e-mail: *Daniel.Gittleson@fda.hhs.gov.*

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.

Medical Devices; Third Party Review Program Under the Food and Drug Administration Modernization Act—(OMB Control Number 0910–0375)—Extension

Section 210 of the Food and Drug Administration Modernization Act (FDAMA) established section 523 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 360m), directing FDA to accredit persons in the private sector to review certain premarket notifications (510(k)s). Participation in this third-party review program by accredited persons is entirely voluntary. A third party wishing to participate will submit a request for accreditation to FDA. Accredited third-party reviewers have the ability to review a manufacturer's 510(k) of the FD&C Act (21 U.S.C. 360) submission for selected devices. After reviewing a submission, the reviewer will forward a copy of the 510(k) submission, along with the reviewer's documented review and recommendation to FDA. Third-party reviewers should maintain records of their 510(k) reviews and a copy of the 510(k) for a reasonable period of time, usually a period of 3 years.

This information collection will allow FDA to continue to implement the accredited person review program established by FDAMA and improve the efficiency of 510(k) review for low- to moderate-risk devices.

Respondents to this information collection are businesses or other for-profit organizations.

I. Reporting

510(k) Reviews Conducted by Accredited Third Parties

According to FDA's data in 2009, the Agency has experienced that the number of 510(k)s submitted for third-party review is approximately 260 annually, which is 26 annual reviews per each of the 10 accredited reviewers.

II. Recordkeeping

Third party reviewers are required to keep records of their review of each submission. According to FDA's in 2009, the Agency anticipates approximately 260 submissions of 510(k)s for third-party review per year.

In the **Federal Register** of September 22, 2010 (75 FR 57801), FDA published a 60-day notice requesting public comment on the proposed collection of information. FDA received one comment; however, it was not PRA related.

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