[FR Doc. 2012–20074 Filed 8–16–12; 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: Federal Child Support Services Portal Registration.

OMB No.: 0970–0370.

The purpose of the Federal Child Support Services Portal Registration is to collect information from an authorized individual registering to use the Federal Parent Locator Service (FPLS) Child Support Services Portal. This information collection is necessary to authenticate the individual's identity and comply with the statutory requirement that federal Office of Child Support Enforcement (OCSE) establish and implement safeguards to restrict access to confidential information in the FPLS to authorized persons. 42 U.S.C. 653(m)(2).

After identity is authenticated, secure accounts will be created for authorized users to view data for their respective applications.

Respondents: Employers, Financial Institutions, Insurers, State Agencies, Local Access and Visitation Providers.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Registration Screen	588	1	0.1	58.8

Estimated Total Annual Burden Hours: 58.8

Additional Information: Copies of the proposed collection may be obtained 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.

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.

Bob Sargis,

Reports Clearance, Officer.

[FR Doc. 2012–20164 Filed 8–16–12; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: DHHS/ACF/OPRE Head Start Classroom-based Approaches and Resources for Emotion and Social skill promotion (CARES) project: Tracking Participants.

OMB No.: 0970–0364.

Description: The Head Start Classroom-based Approaches and Resources for Emotion and Social skill promotion (CARES) project is an evaluation of three social emotional program enhancements within Head Start settings serving three- and four-year-old children. This project focuses on identifying the central features of effective programs to provide the information federal policy makers and Head Start providers will need if they are to increase Head Start's capacity to improve the social and emotional skills and school readiness of preschool age children. The Head Start CARES project completed data collection for cohort (1) 4-year-olds and cohort (2) 3-year-olds in spring of 2011 and cohort (2) 4-year-olds in the spring of 2012.

ACF is proposing to collect information necessary to identify CARES study respondents' current location and follow-up with respondents until the children reach third grade. In support of an examination of third grade outcomes, information must be collected from parents or guardians until the third grade year. Therefore, in the spring of 2013 tracking of all children will be necessary, in the spring of 2014 for the three- and four-year-old children in cohort 2 only, and in the spring of 2015 the three-year-olds in cohort 2 only. To enable the opportunity to conduct data collection in 3rd grade, complete tracking information on the full sample, both ages and cohorts, for all years until third grade is necessary. In addition to location and contact information, a small set of additional items will provide information on the parents' perception of the children's well-being.

Respondents: The respondents to the tracking phone calls will be low-income parents and their Head Start children. This is a three-year information collection request.

ANNUAL BURDEN ESTIMATES

Instrument	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Estimated annual burden hours
Parent Survey Cohort 1–4 year olds	201	1	0.33	66
Parent Survey Cohort 2–4 year olds	690	2	0.33	1380
Parent Survey Cohort 2–3 year olds	320	3	0.33	106
Total				1552

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 Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email address: *OPREinfocollection@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,

ACF Reports Clearance Officer.

[FR Doc. 2012–20207 Filed 8–16–12; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0873]

Agency Information Collection Activities; Proposed Collection; Comment Request; Bar Code Label Requirement for Human Drug and Biological Products

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, 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 the bar code label requirements for human drug and biological products.

DATES: Submit either electronic or written comments on the collection of information by October 16, 2012.

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.

FOR FURTHER INFORMATION CONTACT: Ila Mizrachi, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–7726, *ila.mizrachi@fda.hhs.gov.*

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 collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Bar Code Label Requirement for Human Drug and Biological Products—(OMB Control Number 0910–0537)—Extension

In the **Federal Register** of February 26, 2004 (69 FR 9120), we issued regulations that required human drug product and biological product labels to have bar codes. The rule required bar codes on most human prescription drug products and on over-the-counter (OTC) drug products that are dispensed under an order and commonly used in health care facilities. The rule also required machine-readable information on blood and blood components. For human prescription drug products and OTC drug products that are dispensed under an order and commonly used in health care facilities, the bar code must contain the National Drug Code number for the product. For blood and blood components, the rule specifies the minimum contents of the machine-readable information in a format approved by the Center for Biologics Evaluation and Research Director as blood centers have generally agreed upon the information to be encoded on the label. The rule is intended to help reduce the number of medication errors in hospitals and other health care settings by allowing health care professionals to use bar code scanning equipment to verify that the right drug (in the right dose and right route of administration) is being given to the right patient at the right time.

Most of the information collection burden resulting from the final rule, as calculated in table 1 of the final rule (69 FR at 9149), was a one-time burden that does not occur after the rule's compliance date of April 26, 2006. In addition, some of the information collection burden estimated in the final rule is now covered in other OMB-approved information collection packages for FDA. However, parties may continue to seek an exemption from the bar code requirement under certain, limited circumstances. Section 201.25(d) (21 CFR 201.25(d)) requires submission of a written request for an exemption and describes the contents of such requests. Based on the number of exemption requests we have received, we estimate that approximately 2 exemption requests may be submitted annually, and that each exemption request will require 24 hours to complete. This would result in an annual reporting burden of 48 hours.