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.

[FR Doc. 2014–17664 Filed 7–25–14; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects:

Title: State High Performance Bonus System (HPBS) Transmission File Layouts for HPBS Work Measures

OMB No.: 0970–0230

Description: There is no longer a High Performance Bonus associated with this information collection. The Deficit Reduction Act of 2005 (Pub. L. 109–171) eliminated the funding for the High Performance Bonus (HPB), but we are still requesting that States continue to submit data necessary to calculate the work measures previously reported under the HPB.

Specifically, The TANF program was reauthorized under the Deficit Reduction Act of 2005. The statute eliminated the funding for the HPB under section 403(a)(4). Nevertheless the Department is required under section 413(d) to annually rank State performance in moving TANF recipients into private sector employment. We are, therefore, requesting that States continue to transmit monthly files of adult TANF recipients necessary to calculate the work measures performance data. To the extent States do not provide the requested information, we will extract the matching information from the TANF Data Report. This may result in calculation of the work performance measures based on sample data, which would provide us less precise information on States' performance.

The Transmission File Layouts form provides the format that States will continue to use for the quarterly electronic transmission of monthly data on TANF adult recipients. States that have separate TANF–MOE files on these programs are also requested to transmit similar files. We are not requesting any changes to the Transmission File Layouts form.

Respondents: Respondents may include any of the 50 States, the District of Columbia, Guam, Puerto Rico, and the Virgin Islands.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
State High Performance Bonus System (HPBS) Transmission File Layouts for HPBS Work Measures	42	2	12	1,008

Estimated Total Annual Burden Hours: 1,008.

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, Attn: 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.

[FR Doc. 2014–17668 Filed 7–25–14; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Head Start Performance Information Report.

OMB No.: 0970–0427.

Description: The Office of Head Start within the Administration for Children and Families, United States Department of Health and Human Services, is proposing to renew authority to collect information using the Head Start Program Information Report, monthly enrollments, contacts, locations, and reportable conditions. All information is collected electronically through the Head Start Enterprise System (HSES). The PIR provides information about Head Start and Early Head Start services received by the children and families enrolled in Head Start programs. The information collected in the PIR is used to inform the public about these programs, to make periodic reports to Congress about the status of children in Head Start programs as required by the Head Start Act, and to assist the administration and training/technical assistance of Head Start programs.

Respondents: Head Start and Early Head Start program grant recipients.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Head Start Program Information Report	3,041	1	4	12,164
Grantee Monthly Enrollment Reporting	1,773	12	0.05	1,063.8
Contacts, Locations & Reportable Conditions	3,041	1	0.25	760.25

Estimated Total Annual Burden Hours: 13,988.05.

Additional Information:

Copies of the proposed collection may be obtained 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. All requests should be identified by the title of the information collection. Email 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, *Email: OIRA_SUBMISSION@OMB.EOP.GOV*, Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2014–17654 Filed 7–25–14; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0723]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Medical Devices; Reports of Corrections and Removals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Medical Devices; Reports of Corrections and Removals'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: On April 8, 2014, the Agency submitted a proposed collection of information entitled ''Medical Devices; Reports of Corrections and Removals'' 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 0910–0359. The approval expires on July 31, 2017. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: July 22, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–17633 Filed 7–25–14; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–0079]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Animal Generic Drug User Fee Cover Sheet

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Animal Generic Drug User Fee Cover Sheet'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: On April 28, 2014, the Agency submitted a proposed collection of information entitled ''Animal Generic Drug User Fee Cover Sheet'' 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 0910–0632. The approval expires on July 31, 2017. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: July 23, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–17712 Filed 7–25–14; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–1008]

Exploring the Possibility of Proprietary Name Reservation for Drug Products; Establishment of a Public Docket

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is establishing a public docket to discuss issues related to reserving proprietary names for drug products. During the negotiations for the 2007 reauthorization of the Prescription Drug User Fee Amendments Act (PDUFA IV), FDA agreed to several performance goals related to the review of drug and biological product proprietary names to reduce medication error. Among those goals, FDA and industry expressed an