in this notice, became members of the SEC.

John Howard,

Director, National Institute for Occupational Safety and Health.

[FR Doc. 2013–28452 Filed 11–26–13; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: 45 CFR 303.7—Provision of Services in Intergovernmental IV–D; Federally Approved Forms.

OMB No.: 0970–0085.

Description: The Intergovernmental forms were initially approved by OMB in 1988; 45 CFR 303.7 requires child support programs to use the OMB federally-approved forms in intergovernmental IV–D cases unless a country has provided alternative forms as a part of its chapter in a Caseworker's Guide to Processing Cases with Foreign Reciprocating Countries. Additionally Public Law 104–193, the Personal Responsibility and Work Opportunity Reconciliation Act of 1996, amended 42 U.S.C. 666 to require State Child Support Enforcement (CSE) agencies to enact the Uniform Interstate Family Support Act (UIFSA) into State law by January 1, 1998. Section 311(b) of UIFSA requires the States to use forms mandated by Federal law.

Based on the comments we received in response to the 60-day notice in the **Federal Register** (Volume 78, Number 126, page 39298), we have determined we need to address several issues, particularly relating to the protection of Personal Identifiable Information (PII), and may need to restructure the intergovernmental forms. Because multiple changes to the forms may impact state and federal procedures and systems, we need to thoroughly analyze the options for revising the forms, and provide states and the public the opportunity to comment on any changes and associated burden.

Therefore, at this time, we are requesting an extension of the current forms without any changes. Once we complete the analysis of the issues raised in response to the recent 60 day notice, we will propose changes to the forms and associated burden and request a new round of comments under the Paperwork Reduction Act. The changes will be based on the state's needs and the best interest of the program.

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 respondent	Average burden hours per response	Total burden hours
Transmittal 1	54	19,392	0.25	261,790.25
Transmittal 2	54	14,544	0.08	62,829.66
Transmittal 3	54	970	0.08	4,188.64
Uniform Petition	54	11,635	0.08	50,263.73
General Testimony	54	11,635	0.33	207,337.88
Affidavit Paternity	54	5,818	0.17	53,405.21
Locate Data Sheet	54	388	0.08	1,675.46
Notice of Controlling Order	54	388	0.08	1,675.46
Registration Statement	54	7,757	0.08	33,509.15
Estimated Total Annual Burden Hours				676,683.20

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. 2013–28448 Filed 11–26–13; 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: National Extranet Optimized Runaway and Homeless Youth Management Information System (NEORHYMIS) Version 2.1.

OMB No.: 0970–0123.

Description: The Runaway and Homeless Youth Act, as amended by Public Law 106–71 (42 U.S.C. 5701 et seq.), mandates that the Department of Health and Human Services (HHS) report regularly to Congress on the status of HHS-funded programs serving runaway and homeless youth. Such reporting is similarly mandated by the Government Performance and Results Act. Organizations funded under the Runaway and Homeless Youth program are required by statute (42 U.S.C. 5712, 42 U.S.C. 5714–2) to meet certain data collection and reporting requirements. These requirements include maintenance of client statistical records on the number and the characteristics of the runaway and homeless youth, and youth at risk of family separation, who participate in the project, and the services provided to such youth by the project.

Respondents: States localities, private entities and coordinated networks of

such entities. Typical respondents are non-profit community based organizations who are reporting on the youth that they serve through their Basic Center, Transitional Living and Street Outreach programs.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Youth Profile: BCP Entrance Report	321	118	.125	4, 735
Youth Profile: TLP Entrance Report	205	19	.125	487
Youth Profile: BCP Exit Report	321	118	.125	4,735
Youth Profile: TLP Exit Report	205	19	.125	487
Brief Contacts	526	153	.05	4, 024
BCP Turn-a-ways	321	9	.05	144
TLP Turn-a-ways	205	24	.05	246
Street Outreach Report	138	5,660	.02	15,622
Data Transfer	664	2	.50	664

Estimated Total Annual Burden Hours: 31, 441.

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, Email: *OIRA_SUBMISSION@OMB.EOP.GOV*. Attn: Desk Officer for the Administration for Children and Families

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2013–28430 Filed 11–26–13; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–1423]

Agency Information Collection Activities; Proposed Collection; Comment Request; Importer's Entry Notice

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 FDA's Importer's Entry Notice.

DATES: Submit either electronic or written comments on the collection of information by January 27, 2014.

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: FDA PRA Staff, Office of Operations, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, *PRAStaff@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.

Information Request Regarding Importer's Entry Notice—(OMB Control Number 0910–0046)—Extension

Section 801 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (U.S.C. 381) charges the Secretary of Health and Human Services (HHS), through FDA, with the responsibility of assuring foreign origin FDA regulated foods, drugs, cosmetics, medical devices, radiological health, and tobacco products offered for import into the United States meet the same requirements of the FD&C Act as do domestic products, and for preventing products from entering the country if they are not in compliance. The discharge of this responsibility involves close coordination and cooperation