FEDERAL RETIREMENT THRIFT INVESTMENT BOARD

Sunshine Act; Notice of Meeting July 24, 2018, Telephonic, 12:00 p.m., 10th Floor Board Meeting Room, 77 K Street NE, Washington, DC 20002

Agenda

FEDERAL RETIREMENT THRIFT INVESTMENT BOARD MEMBER MEETING STATUS: All parts will be open to the public.

MATTERS TO BE CONSIDERED:

Open Session

1. Approval of the Minutes for the June 25, 2018 Board Member Meeting
2. Monthly Reports
   (a) Participant Activity
   (b) Legislative Report
3. Quarterly Reports
   (c) Investment Performance
   (d) Budget Review
   (e) Audit Status
4. Enterprise Risk Management Update
5. IT Update

CONTACT PERSON FOR MORE INFORMATION: Kimberly Weaver, Director, Office of External Affairs, (202) 942–1640.

Dated: July 16, 2018.

Dharmesh Vashee,

Deputy General Counsel, Federal Retirement Thrift Investment Board.

[FR Doc. 2018–15433 Filed 7–16–18; 4:15 pm]

BILLING CODE 6760–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB NO.: 0970–0383]

Submission for OMB Review; Comment Request; Evaluation of the Transitional Living Program (TLP)—Extension

Description:

The Family and Youth Services Bureau (FYSB) and the Office of Planning, Research, Evaluation (OPRE) in the Administration for Children and Families (ACF) are requesting to continue collecting data as part of a currently approved information collection (OMB No. 0970–0383). The purpose is to continue baseline data collection at study enrollment and follow-up data collection for the Evaluation of the Transitional Living Program (TLP). The TLP evaluation was designed to examine the effects of FYSB's Transitional Living Program on runaway and homeless youth, focusing on such outcomes as housing and homelessness, education or training, employment, social connections, socio-emotional well-being, and risk behaviors.

Data collection will include three primary surveys: (1) A survey administered at the time of TLP enrollment (baseline), (2) a survey administered 6 months after enrollment, which will collect information on short-terms outcomes; and (3) a survey administered at 12 months, which will collect information on longer-term outcomes.'' Participants will be enrolled through the TLP study sites.

Respondents: Runaway and homeless youth ages 16 to 22 who agree to participate in the study upon enrollment into one of the TLP study sites.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
Young Adult Baseline Survey	600	200	1	0.62	124
Young Adult 6-Month Follow Up Survey	600	200	1	0.61	122
Young Adult 12-Month Follow Up Survey	600	200	1	0.61	122
Estimated Total Burden Hours					368

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, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *OPREinfocollection@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

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2018–15307 Filed 7–17–18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0742]

Agency Information Collection Activities; Proposed Collection; Comment Request; Registration of Producers of Drugs and Listing of Drugs in Commercial Distribution

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each 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 for drug establishment registration and product listing.

DATES: Submit either electronic or written comments on the collection of information by September 17, 2018.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before September 17,