valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

OBTAINING COPIES OF PROPOSALS: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW, Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0035, Claims and Appeals, in all correspondence.

Dated: May 10, 2018.

Lorin Curit,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2018–10408 Filed 5–15–18; 8:45 am]

BILLING CODE 6820–EP–P

GENERAL SERVICES ADMINISTRATION

[Notice–MA–2018–03; Docket No. 2018–0002, Sequence No. 7]

Rescission of FTR Bulletins

AGENCY: Office of Government-wide Policy (OGP), General Services Administration (GSA).

ACTION: Notice of Federal Travel Regulation (FTR) Bulletin 18–04, rescission of FTR Bulletins.

SUMMARY: GSA is officially rescinding various FTR bulletins to ensure the Travel/Per Diem Bulletin section on the agency's FTR website displays only current information. Agencies' policies should be updated as warranted.

DATES: The rescission is as of May 16, 2018.

FOR FURTHER INFORMATION CONTACT: For clarification of content, please contact Mr. Cy Greenidge, Office of Government-wide Policy, Office of Asset and Transportation Management, at 202–219–2349, or by email at *travelpolicy@gsa.gov.* Please cite Notice of FTR Bulletin 18–04.

SUPPLEMENTARY INFORMATION: Executive Order 13777, Enforcing the Regulatory Reform Agenda, Section 3, paragraph (d)(ii), states in part, the Regulatory Reform Task Force shall attempt to identify regulations that are outdated, unnecessary, or ineffective. GSA has conducted a thorough review of all FTR bulletins on the FTR Travel/Per Diem Bulletins website (*https://www.gsa.gov/policy-regulations/regulations/federal-travel-regulation/federal-travel-regulation-and-related-files#TravelPerDiemBulletins*) and determined that some of the Bulletins contain outdated, duplicative, expired, or inapplicable content. FTR Bulletin 18–04 lists all rescinded bulletins meeting one of the aforementioned criterion.

Dated: May 11, 2018.

Alexander Kurien,

Deputy Associate Administrator, Office of Asset and Transportation Management, Office of Government-wide Policy.

[FR Doc. 2018–10436 Filed 5–15–18; 8:45 am]

BILLING CODE 6820–14–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Evaluation of the Transitional Living Program (TLP).

OMB No.: 0970–0383.

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, previously approved by OMB: (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.

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, 330 C Street SW, Washington, DC 20201, Attn: OPRE 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.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2018–10461 Filed 5–15–18; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2018–N–1773]

Blood Products Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) announces a forthcoming public advisory committee meeting of the Blood Products Advisory Committee. The general function of the Blood Products Advisory Committee is to provide advice and recommendations to the Agency on regulatory issues related to blood and products derived from blood. On the first day of the meeting, the Committee will provide advice regarding bacterial risk control strategies to enhance the safety and availability of platelets for transfusion. On the second day of the meeting, the Committee, supplemented with members from the Microbiology Devices Panel of the Medical Devices Advisory Committee, will function as a medical device panel to provide advice and recommendations to the Agency on classification of devices. The meeting will be open to the public.

DATES: The meeting will be held on July 18, 2018, from 8 a.m. to 5 p.m. and July 19, 2018, from 8 a.m. to 3 p.m.

ADDRESSES: FDA White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503, sections B and C), Silver Spring, MD 20993–0002. Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.*

FOR FURTHER INFORMATION CONTACT: Bryan Emery, Division of Scientific Advisors and Consultants, Center for Biologics Evaluation and Research (CBER), Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 6268, Silver Spring, MD 20993–0002, 240–402–8054, *bryan.emery@fda.hhs.gov;* or Joanne Lipkind, Division of Scientific Advisors and Consultants, CBER, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 6270, Silver Spring, MD 20993–0002, 240–402–8106, *joanne.lipkind@fda.hhs.gov;* or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's website at *https://www.fda.gov/AdvisoryCommittees/default.htm* and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting. For those unable to attend in person, the meeting will be also be available via webcast. The webcast will be available at the following link on both days: *https://collaboration.fda.gov/bpac0718/.*

SUPPLEMENTARY INFORMATION:

Agenda: On July 18, 2018, the Blood Products Advisory Committee will meet in open session to discuss and provide advice regarding bacterial risk control strategies for blood collection establishments and transfusion services to enhance the safety and availability of platelets for transfusion. The Committee will discuss the available strategies to control the risk of bacterial contamination of platelets with 5-day and 7-day dating, including bacterial testing using culture-based devices and rapid bacterial detection devices and implementation of pathogen reduction technology.

On July 19, 2018, the Committee will function as a medical device panel. The Committee will meet in open session to discuss and provide advice regarding the device reclassification from class III to class II of nucleic acid and serology-based point-of-care and laboratory-based in vitro diagnostic devices indicated for use as aids in the diagnosis of human immunodeficiency virus (HIV) infection. The devices that will be discussed by the Committee during the meeting are post-amendment devices that currently are classified into class III under section 513(f)(1) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360c(f)(1)).

FDA intends to make background material available to the public approximately 2 weeks and no later than 2 business days before the meeting. If FDA is unable to post the background material on its website prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's website after the meeting. Background material is available at *https://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee meeting link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before July 11, 2018. Oral presentations from the public will be scheduled between approximately 2:15 p.m. and 3:15 p.m. on July 18, 2018, and between 12:30 p.m. and 1:30 p.m. on July 19, 2018. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before July 5, 2018. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by July 6, 2018.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with disabilities. If you require accommodations due to a disability, please contact Bryan Emery