a member of the Committee; (2) the nominator's name, address, daytime telephone number, and the home and/or work address, telephone number, and email address of the individual being nominated; and (3) a current copy of the nominee's curriculum vitae. Federal employees should not be nominated for consideration of appointment to this Committee.

The Department makes every effort to ensure that the membership of HHS Federal advisory committees is fairly balanced in terms of points of view represented and the committee's function. Every effort is made to ensure that individuals from a broad representation of geographic areas, women and men, ethnic and minority groups, and the disabled are given consideration for membership on HHS Federal advisory committees. Appointment to this Committee shall be made without discrimination on the basis of age, race, ethnicity, gender, sexual orientation, disability, and cultural, religious, or socioeconomic status.

Individuals who are selected to be considered for appointment will be required to provide detailed information regarding their financial holdings, consultancies, and research grants or contracts. Disclosure of this information is necessary in order to determine if the selected candidate is involved in any activity that may pose a potential conflict with the official duties to be performed as a member of SACHRP.

Dated: December 22, 2014.

Jerry Menikoff,

Director, Office for Human Research Protections, Executive Secretary, Secretary's Advisory Committee, on Human Research Protections.

[FR Doc. 2014–30400 Filed 12–24–14; 8:45 am]

BILLING CODE 4150–36–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Request for Information

AGENCY: Office of Child Support Enforcement (OCSE), Administration for Children and Families, HHS.

ACTION: Notice of request for information.

SUMMARY: The Administration for Children and Families (ACF) published a notice in the **Federal Register** on October 23, 2014, (79 FR 63406) requesting public comments to inform its upcoming Report to Congress. The Report to Congress is required to be submitted no later than June 30, 2015, under title III, section 305 of H.R. 4980 (Pub. L. 113–183), Preventing Sex Trafficking and Strengthening Families Act of 2014. ACF stated in the notice that the request for information would remain open until December 22, 2014, for the receipt of public comments. To provide the public with more time to comment, ACF extends the period of time for which the comments will remain open.

To provide clarification on the first bullet point under the Background Section, which was truncated in the first **Federal Register** Notice, please consider the following: A review of the effectiveness of state child support programs and collection practices and an analysis of the extent to which the practices result in unintended consequences or performance issues.

DATES: Comments must be received by 11:59 p.m. on February 27, 2015, to be considered.

FOR FURTHER INFORMATION CONTACT: The Office of Child Support Enforcement at *OCSEreport@acf.hhs.gov*.

Dated: December 19, 2014.

Donna Bonar,

Deputy Comissioner, Office of Child Support Enforcement.

[FR Doc. 2014–30285 Filed 12–24–14; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–2214]

Next Generation Sequencing Diagnostic Tests; Public Workshop; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop; request for comments.

SUMMARY: The Food and Drug Administration (FDA) is announcing the following public workshop entitled ''Optimizing FDA's Regulatory Oversight of Next Generation Sequencing Diagnostic Tests.'' The purpose of this workshop is to discuss and receive feedback from the community on the questions in the discussion paper on diagnostic tests for human genetics or genomics using next generation sequencing (NGS) technology.

DATES: The public workshop will be held on February 20, 2015, from 8:30 a.m. to 5 p.m.

ADDRESSES: The public workshop will be held at the Natcher Center at the National Institutes of Health Campus, 9000 Rockville Pike, Bldg. 45 Auditorium, Bethesda, MD 20814. For parking and security information, please refer to *http://www.nih.gov/about/visitor/*.

FOR FURTHER INFORMATION CONTACT: David Litwack, Office of In Vitro Diagnostics and Radiological Health, Center for Devices and Radiological Health, Food and Drug Administration, Bldg. 66, Rm. 5544, 10903 New Hampshire Ave., Silver Spring, MD 20993, 301–796–6697, email: *ernest.litwack@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Registration: Registration is free and available on a first-come, first-served basis. Persons interested in attending this public workshop must register online by 4 p.m. February 12, 2015. Early registration is recommended because facilities are limited and, therefore, FDA may limit the number of participants from each organization. If time and space permits, onsite registration on the day of the public workshop will be provided beginning at 7:30 a.m.

If you need special accommodations due to a disability, please contact Susan Monahan, 301–796–5661, email: *Susan.Monahan@fda.hhs.gov* no later than February 6, 2015.

To register for the public workshop, please visit FDA's Medical Devices News & Events—Workshops & Conferences calendar at *http://www.fda.gov/MedicalDevices/NewsEvents/WorkshopsConferences/default.htm.* (Select this public workshop from the posted events list.) Please provide complete contact information for each attendee, including name, title, affiliation, address, email, and telephone number. If you are unable to register online, please contact Susan Monahan (see *Registration*.) Registrants will receive confirmation after they have been accepted. You will be notified if you are on a waiting list.

Streaming Webcast of the Public Workshop: This public workshop will also be Webcast. Persons interested in viewing the Webcast must register online by February 12, 2015. Early registration is recommended because Webcast connections are limited. Organizations are requested to register all participants, but to view using one connection per location. Webcast participants will be sent technical system requirements after registration and will be sent connection access information after February 13, 2015. If you have never attended a Connect Pro