

Tissue Safety and Availability (ACBTSA) will hold a meeting. The meeting will be open to the public.

**DATES:** The meeting will take place Tuesday April 7, 2105, from 8:00 a.m.–4:00 p.m. and Wednesday April 8, 2015, from 8:00 a.m.–3:30 p.m.

**ADDRESSES:** NIH Conference Room, 5635 Fishers Lane, Rockville, MD 20892.

**FOR FURTHER INFORMATION CONTACT:** Mr. James Berger, Designated Federal Officer for the ACBTSA, Senior Advisor for Blood and Tissue Policy, Office of the Assistant Secretary for Health, Department of Health and Human Services, 1101 Wootton Parkway, Suite 250, Rockville, MD 20852. Phone: (240) 453–8803; Fax (240) 453–8456; Email [ACBTSA@hhs.gov](mailto:ACBTSA@hhs.gov).

**SUPPLEMENTARY INFORMATION:** The ACBTSA provides advice to the Secretary through the Assistant Secretary for Health. The Committee advises on a range of policy issues to include: (1) Identification of public health issues through surveillance of blood and tissue safety issues with national biovigilance data tools; (2) identification of public health issues that affect availability of blood, blood products, and tissues; (3) broad public health, ethical and legal issues related to the safety of blood, blood products, and tissues; (4) the impact of various economic factors (*e.g.*, product cost and supply) on safety and availability of blood, blood products, and tissues; (5) risk communications related to blood transfusion and tissue transplantation; and (6) identification of infectious disease transmission issues for blood, organs, blood stem cells and tissues. The Committee has met regularly since its establishment in 1997.

The ACBTSA has made previous recommendations on the need to improve tissue tracking and traceability. These recommendations focused on tracking adverse events, creating a unique identifier for each donor, and improving patient outcomes. Past recommendations made by the ACBTSA may be viewed at [www.hhs.gov/bloodsafety](http://www.hhs.gov/bloodsafety).

The focus of the meeting will be to address current issues in tracking and traceability of tissue recovered from deceased donors. The discussion will focus on pertinent federal and state regulations, other mechanisms that impact tissue tracking and traceability, and current gaps in this area. Presenters will represent a wide range of government and non-government stakeholders, including federal agencies, accreditation organizations, tissue and eye banks, healthcare facilities, and medical practitioners.

The public will have an opportunity to present their views to the Committee during a public comment session scheduled for April 8, 2015. Comments will be limited to five minutes per speaker and must be pertinent to the discussion. Pre-registration is required for participation in the public comment session. Any member of the public who would like to participate in this session is encouraged to contact the Designated Federal Officer at his/her earliest convenience to register for time (limited to 5 minutes); registration must be completed prior to close of business on April 1, 2015. If it is not possible to provide 30 copies of the material to be distributed at the meeting, then individuals are requested to provide a minimum of one (1) copy of the document(s) to be distributed prior to the close of business on April 1, 2015. It is also requested that any member of the public who wishes to provide comments to the Committee utilizing electronic data projection submit the necessary material to the Designated Federal Officer prior to the close of business on April 1, 2015.

Dated: January 22, 2015.

**James J. Berger,**

*Senior Advisor for Blood and Tissue Safety Policy.*

[FR Doc. 2015–01680 Filed 1–28–15; 8:45 am]

**BILLING CODE 4150–41–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Nominations to the Advisory Committee on Blood and Tissue Safety and Availability

**AGENCY:** Office of the Secretary, Office of the Assistant Secretary for Health, Department of Health and Human Services.

**ACTION:** Notice.

**SUMMARY:** The Office of Assistant Secretary for Health (OASH) is seeking nominations of qualified individuals to be considered for appointment as members of the Advisory Committee on Blood and Tissue Safety and Availability (ACBTSA). ACBTSA is a Federal advisory committee within the Department of Health and Human Services. Management support for the activities of this Committee is the responsibility of the OASH. The qualified individuals will be nominated to the Secretary of Health and Human Services for consideration of appointment as members of the ACBTSA. Members of the Committee, including the Chair, are appointed by the Secretary. Members are invited to

serve on the Committee for up to four-year terms.

**DATES:** All nominations must be received no later than 4:00 p.m. EDT on March 2, 2015, at the address listed below.

**ADDRESSES:** All nominations should be mailed or delivered to Mr. James Berger, Senior Advisor for Blood and Tissue Policy; Office of Assistant Secretary for Health; Department of Health and Human Services; 1101 Wootton Parkway, Suite 250; Rockville, MD 20852. Telephone: (240) 453–8803; Fax (240) 453–8456; Email [ACBTSA@hhs.gov](mailto:ACBTSA@hhs.gov).

**FOR FURTHER INFORMATION CONTACT:** Mr. James Berger, Senior Advisor for Blood and Tissue Policy. Contact information for Mr. Berger is provided above.

A copy of the Committee charter and roster of the current membership can be obtained by contacting Mr. Berger or by accessing the ACBTSA Web site at <http://www.hhs.gov/bloodsafety>.

**SUPPLEMENTARY INFORMATION:** The ACBTSA shall provide advice to the Secretary through the Assistant Secretary for Health. The Committee shall advise on a range of policy issues to include: (1) Identification of public health issues through surveillance of blood and tissue safety issues with national biovigilance data tools; (2) identification of public health issues that affect availability of blood, blood products, and tissues; (3) broad public health, ethical and legal issues related to the safety of blood, blood products, and tissues; (4) the impact of various economic factors (*e.g.*, product cost and supply) on safety and availability of blood, blood products, and tissues; (5) risk communications related to blood transfusion and tissue transplantation; and (6) identification of infectious disease transmission issues for blood, organs, blood stem cells and tissues.

The Committee consists of 23 voting members. The Committee composition includes 14 public members, including the Chair, and nine (9) individuals designated to serve as official representative members. The public members are selected from state and local organizations, patient advocacy groups, provider organizations, academic researchers, ethicists, physicians, surgeons, scientists, risk communication experts, consumer advocates, and from among communities of persons who are frequent recipients of blood or blood products or who have received tissues or organs. The nine individuals who are appointed as official representatives are

selected to serve the interests of the blood, blood products, tissue and organ professional organizations or business sectors. The representative members are selected from the following groups: The AABB (formerly the American Association of Blood Banks); American Association of Tissue Banks; Eye Bank Association of America; Association of Organ Procurement Organizations; and one of either the American National Red Cross or America's Blood Centers on a rotating basis. The Committee composition can include additional representation from either the plasma protein fraction community or a trade organization; a manufacturer of blood, plasma, or other tissue/organ test kits; a manufacturer of blood, plasma or other tissue/organ equipment; a major hospital organization; or a major hospital accreditation organization. Where more than one company produces a specified product or process, representatives from those companies shall rotate on the same schedule as public members.

All ACBTSA members are authorized to receive the prescribed per diem allowance and reimbursement for travel expenses that are incurred to attend meetings and conduct Committee-related business, in accordance with Standard Government Travel Regulations. Individuals who are appointed to serve as public members are authorized also to receive a stipend for attending Committee meetings and to carry out other Committee-related business. Individuals who are appointed to serve as representative members for a particular interest group or industry are not authorized to receive a stipend for the performance of these duties.

This announcement is to solicit nominations of qualified candidates to fill nine public member positions that are scheduled to be vacated on the ACBTSA.

### Nominations

In accordance with the charter, persons nominated for appointment as members of the ACBTSA should be among authorities knowledgeable in blood banking, tissue banking, transfusion medicine, organ or tissue transplantation, plasma therapies, transfusion and transplantation safety, bioethics, and/or related disciplines. Nominations should be typewritten. The following information should be included in the package of material submitted for each individual being nominated for consideration of appointment: (a) The name, return address, daytime telephone number and affiliation(s) of the individual being nominated, the basis for the individual's

nomination, the category for which the individual is being nominated, and a statement bearing an original signature of the nominated individual that, if appointed, he or she is willing to serve as a member of the Committee; (b) the name, return address, and daytime telephone number at which the nominator may be contacted. Organizational nominators must identify a principal contact person in addition to the contact; and (c) a copy of a current curriculum vitae or resume for the nominated individual.

Individuals can nominate themselves for consideration of appointment to the Committee. All nominations must include the required information. Incomplete nominations will not be processed for consideration. The letter from the nominator and certification of the nominated individual must bear original signatures; reproduced copies of these signatures are not acceptable.

The Department is legally required to ensure that the membership of HHS Federal advisory committees is fairly balanced in terms of points of view represented and the functions to be performed by the advisory committee. Every effort is made to ensure that the views of women, all ethnic and racial groups, and people with disabilities are represented on HHS Federal Advisory committees and, therefore, the Department encourages nominations of qualified candidates from these groups. The Department also encourages geographic diversity in the composition of the committee. 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.

The Standards of Ethical Conduct for Employees of the Executive Branch are applicable to individuals who are appointed as public members of Federal advisory committees. Individuals appointed to serve as public members of Federal advisory committees are classified as special government employees (SGEs). The federal conflict of interest laws are applicable to SGEs. Therefore, individuals appointed to serve as public members of the ACBTSA are subject to an ethics review. The ethics review is conducted to determine if the individual has any interests and/or activities in the private sector that may conflict with performance of their official duties as a member of the Committee. Individuals appointed to serve as public members of the committee will be required to disclose information regarding financial holdings, consultancies, and research grants and/or contracts.

Dated: January 22, 2015.

**James J. Berger,**

*Senior Advisor for Blood and Tissue Policy.*

[FR Doc. 2015-01682 Filed 1-28-15; 8:45 am]

**BILLING CODE 4150-41-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[30-Day-15-0931]

### Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639-7570 or send an email to [omb@cdc.gov](mailto:omb@cdc.gov). Written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395-5806. Written comments should be received within 30 days of this notice.