SUMMARY: This Government-wide Travel Advisory Committee (GTAC) (the Committee) is a Federal Advisory Committee established in accordance with the Federal Advisory Committee Act (FACA), 5 U.S.C., App 2. This notice announces the next two meetings, which are open to the public via teleconference and webinar.

DATES: The upcoming March 26, 2014 and April 30, 2014 meetings will begin at 9:00 a.m. Eastern Standard Time and end no later than 4:00 p.m. Eastern Standard Time. February 24, 2014.

FOR FURTHER INFORMATION CONTACT: Ms. Marcerto Barr, Designated Federal Officer (DFO), Government-wide Travel Advisory Committee (GTAC), Office of Government-Wide Policy, General Services Administration, 1800 F Street NW., Washington, DC 20405, 202–208–7654 or by email to: gtac@gsa.gov.

SUPPLEMENTARY INFORMATION: The purpose of the GTAC is to conduct public meetings, submit reports and to make recommendations to existing travel policies, processes and procedures, including the per diem methodology to assure that official travel is conducted in a responsible manner with the need to minimize costs.

Authority: The GSA Office of Asset and Transportation Management, Travel and Relocation Division, establishes policy that governs travel by Federal civilian employees and others authorized to travel at Government expense on temporary duty travel through the Federal Travel Regulation (FTR).

Agenda: The March meeting will include a follow-up discussion of previous topics, including Data and Meals and Incidental Expenditure Allowances. The April meeting will discuss Managed Lodging, Long-term stay, and reduced per diem.

Meeting Access: The meeting is open to the public via teleconference and webinar. Members of the public wishing to listen in on the GTAC discussion are recommended to visit the GTAC Web site at: www.gsa.gov/gtac to obtain registration details. Members of the public will not have the opportunity to ask questions or otherwise participate in the meeting. However, members of the public wishing to comment on the discussion or topics outlined in the agenda should follow the steps detailed in Procedures for Providing Public Comments.

Availability of Materials for the Meeting: Please see the GTAC Web site www.gsa.gov/gtac for any available materials and detailed meeting notes after the meeting.

Procedures for Providing Public Comments: In general, public comments will be posted to www.gsa.gov/gtac. Non-electronic documents will be made available for public inspection and copying at GSA, 1800 F Street NW., Washington, DC 20405, on official business days between the hours of 10:00 a.m. Eastern Standard Time and 4:00 p.m. Eastern Standard Time. The public can make an appointment to inspect comments by telephoning the DFO at 202–208–7654. All comments, including attachments and other supporting materials received, are part of the public record and subject to public disclosure. Any comments submitted in connection with the GTAC meeting will be made available to the public under the provisions of the Federal Advisory Committee Act.

The public is invited to submit written comments within 7 business days after each meeting by either of the following methods and cite Meeting Notice–GTAC–2014–01.

Electronic or Paper Comments: (1) Submit electronic comments to gtac@ gsa.gov; or (2) submit paper comments to the attention of Ms. Marcerto Barr at GSA, 1800 F Street NW., Washington, DC 20405.

Dated: February 18, 2014.

Carolyn Austin-Diggs,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: HHS-OS-21431-60D]

Agency Information Collection Activities; Proposed Collection; Public Comment Request

AGENCY: Office of the Secretary, HHS. **ACTION:** Notice.

SUMMARY: In compliance with section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). The ICR is for reinstatement of a previously-approved information collection assigned OMB control number 0990–0313, which expired on October 31, 2013. Prior to submitting

that ICR to OMB, OS seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on the ICR must be received on or before April 25, 2014.

ADDRESSES: Submit your comments to *Information.CollectionClearance*@ *hhs.gov* or by calling (202) 690–6162.

FOR FURTHER INFORMATION CONTACT:

Information Collection Clearance staff, *Information.CollectionClearance@ hhs.gov* or (202) 690–6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the document identifier HHS-OS-21431-60D for reference.

Information Collection Request Title: National Blood Collection and

Utilization Survey.

Abstract: The National Blood Collection & Utilization Survey (NBCUS) is a biennial survey of the blood collection and utilization community (industry) to produce reliable and accurate estimates of national and regional collections, utilization, safety, and availability of all blood products, some cellular therapeutic products, as well as information on bacterial testing and human tissue transplantation that are of interest to the transfusion medicine community. The 2013 NBCUS shall be funded by the U.S. Department of Health and Human Services (DHHS) and performed by (contactor, to be determined). In previous years, the NBCUS program was performed under the auspices of the National Blood Data Resource Center (NBDRC), a private subsidiary of AABB (formerly known as the American Association of Blood Banks), with private funding.

The survey includes a core of standard questions on blood collection, processing, and utilization practices to allow for comparison with data from previous surveys; additionally, questions to specifically address emerging and developing issues and technologies in blood collection and utilization are included. Biovigilance remains a key theme for the 2013 survey, as continued from the 2007, 2009, and 2011 iterations. To that end, questions on transfusion transmitted infections, transfusion associated circulatory overload, acute hemolysis, delayed hemolysis, and severe allergic reactions are included in the survey.

Need and Proposed Use of the Information: Under the authority of Section 301 of the Public Health Service Act (42 U.S.C.241), as identified in the 1997 HHS Blood Action Plan, and twice in the Advisory Committee on Blood & Tissue Safety & Availability's (ACBTSA) recommendations to the Secretary, there is a need to provide national policy makers with current supply and demand data.

Likely Respondents: Respondents will include approximately 3,000 institutions that include U.S. blood collection and processing facilities, hospital-based transfusion blood banks, and cord blood banks. Participating institutions will be selected from the

American Hospital Association (AHA) annual survey database and AABB member list of blood collection facilities.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions, to develop, acquire, install and utilize technology and systems for the purpose

of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information, to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information, and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
National Blood Collection and Utilization Survey	3,000	1	1	3,000
Total	3,000	1	1	3,000

OS specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Darius Taylor,

Deputy, Information Collection Clearance Officer.

[FR Doc. 2014–03829 Filed 2–21–14; 8:45 am]

BILLING CODE 4150-41-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: HHS-OS-21435-60D]

Agency Information Collection Activities; Proposed Collection; Public Comment Request

AGENCY: HHS, Office of the Secretary. **ACTION:** Notice.

SUMMARY: In compliance with section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, announces plans to submit a new Information Collection Request (ICR), described below, to the Office of Management and Budget

(OMB). Prior to submitting that ICR to OMB, OS seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR. **DATES:** Comments on the ICR must be received on or before April 25, 2014. **ADDRESSES:** Submit your comments to Information.CollectionClearance@ hhs.gov or by calling (202) 690–6162.

FOR FURTHER INFORMATION CONTACT: Information Collection Clearance staff, Information.CollectionClearance@ hhs.gov or (202) 690–6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the document identifier HHS OS-21435-60D for reference.

Information Collection Request Title: HIPAA Covered Entity and Business Associate Pre-Audit Survey.

Abstract: This information collection consists of a survey of up to 1200 Health Insurance Portability and Accountability Act of 1996 (HIPAA) covered entities (health plans, health care clearinghouses, and certain health care providers) and business associates (entities that provider certain services to a HIPAA covered entity) to determine suitability for the Office for Civil Rights (OCR) HIPAA Audit Program. The survey will gather information about respondents to enable OCR to assess the size, complexity, and fitness of a respondent for an audit. Information collected includes, among other things, recent data about the number of patient

visits or insured lives, use of electronic information, revenue, and business locations.

Need and Proposed Use of the Information: The Office for Civil Rights (OCR) is mandated to conduct periodic audits to assess the compliance of covered entities and business associates with the HIPAA Privacy, Security, and Breach Notification Rules. This information collection will enable OCR to assess the suitability of respondent covered entities and business associates for audits.

Likely Respondents: Respondents will include both HIPAA covered entities and business associates.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions, to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information, to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information, and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.