

that strive to augment the informed consent document and increase a subject's understanding of research participation. The meeting will conclude with a panel of speakers focusing on the regulatory barriers that may be associated with Community Based and Participatory Research. Public comment will be heard on both days.

Public attendance at the meeting is limited to space available. Individuals who plan to attend the meeting and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact persons. Members of the public will have the opportunity to provide comments on both days of the meeting. Public comment will be limited to five minutes per speaker. Any members of the public who wish to have printed materials distributed to SACHRP members for this scheduled meeting should submit materials to the Executive Director, SACHRP, prior to the close of business Thursday, October 22, 2009. Information about SACHRP and the draft meeting agenda will be posted on the SACHRP Web site at: <http://www.hhs.gov/ohrp/sachrp/index.html>.

Dated: October 6, 2009.

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

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Meeting of the Advisory Committee on Blood Safety and Availability

**AGENCY:** Department of Health and Human Services, Office of the Secretary.  
**ACTION:** Notice.

**SUMMARY:** As stipulated by the Federal Advisory Committee Act, the U.S. Department of Health and Human Services is hereby giving notice that the Advisory Committee on Blood Safety and Availability (ACBSA) will hold a meeting. The meeting will be open to the public.

**DATES:** The meeting will take place Thursday, November 19 and Friday, November 20, 2009 from 8:30 a.m. to 5 p.m.

**ADDRESSES:** The Universities at Shady Grove, 9630 Gudelsky Drive, Rockville, MD 20850, Phone: 301-738-6000.

**FOR FURTHER INFORMATION CONTACT:** Jerry A. Holmberg, PhD, Executive Secretary,

Advisory Committee on Blood Safety and Availability, Office of Public Health and Science, Department of Health and Human Services, 1101 Wootton Parkway, Suite 250, Rockville, MD 20852, (240) 453-8803, FAX (240) 453-8456, e-mail [ACBSA@hhs.gov](mailto:ACBSA@hhs.gov).

**SUPPLEMENTARY INFORMATION:** The Advisory Committee on Blood Safety and Availability provides advice to the Secretary and the Assistant Secretary for Health on a range of policy issues that impact (1) Definition of public health parameters around safety and availability of the blood supply and blood products, (2) broad public health, ethical and legal issues related to transfusion and transplantation safety, and (3) the implications for safety and the availability of various economic factors affecting product cost and supply.

In keeping with its established mission, the ACBSA has been asked to review and comment on the current processes and parameters which should be used in the decision-making process for transplantation safety policy. At the November 19 and 20, 2009 meeting, the Committee will be asked to comment and make recommendations on current safety decision making processes within the Department of Health and Human Services while considering those same processes within the Private Sector.

During the meeting, the ACBSA will be provided a briefing on Biovigilance (surveillance of blood, organs, and tissues safety). Specifically, the committee will be asked to comment on the white paper entitled, "Biovigilance in the United States: Efforts to Bridge a Critical Gap in Patient Safety and Donor Health."

The public will have opportunity to present their views to the Committee on both meeting days. A public comment session has been scheduled for November 19 and 20, 2009. 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 Executive Secretary at his/her earliest convenience. It is requested that those who wish to have printed material distributed to the Committee provide thirty (30) copies of the document to be distributed to the Executive Secretary, ACBSA, prior to close of business November 16, 2009. If it is not possible to provide 30 copies of the material to be distributed, then individuals are requested to provide at a minimum one (1) copy of the document(s) to be

distributed prior to the close of business November 16, 2009. It also is 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 Executive Secretary prior to close of business November 16, 2009.

Dated: September 25, 2009.

**Jerry A. Holmberg,**  
Executive Secretary, Advisory Committee on Blood Safety and Availability.

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[60Day-10-0612]

#### Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404-639-5960 and send comments to Maryam Daneshvar, CDC Acting Reports Clearance Officer, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an e-mail to [omb@cdc.gov](mailto:omb@cdc.gov).

Comments are invited 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) ways to enhance 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. Written comments should be received within 60 days of this notice.

#### Proposed Project

Well-Integrated Screening and Evaluation for Women Across the Nation (WISEWOMAN) Reporting System (OMB #0920-0612, exp. 1/31/2010)—Revision—National Center