

data on positions in to-be-issued Treasury coupon securities, mainly the trading on a when-issued delivery basis.

Current Actions: On November 2, 2009, the Federal Reserve published a notice in the **Federal Register** (74 FR 56633) requesting public comment for 60 days on the extension, with revision, of the FR 2004. The comment period for this notice expired on January 4, 2010. The Federal Reserve did not receive any comments. The revisions will be implemented as proposed.

Final approval under OMB delegated authority of the implementation of the following survey:

Report title: Central Bank Survey of Foreign Exchange and Derivatives Market Activity.

Agency form number: FR 3036.

OMB control number: 7100-0285.

Frequency: One-time.

Reporters: Financial institutions that serve as intermediaries in the wholesale foreign exchange and derivatives market and dealers.

Estimated annual reporting hours: 2,165 hours.

Estimated average hours per response: Turnover survey, 55 hours; outstandings survey, 60 hours.

Number of respondents: Turnover survey, 35; outstandings survey, 4.

General description of report: This information collection is voluntary (12 U.S.C. 225a and 263) and is given confidential treatment (5 U.S.C. 552(b)(4)).

Abstract: The FR 3036 is the U.S. part of a global data collection that is conducted by central banks once every three years. More than 50 central banks plan to conduct the survey in 2010. The Bank for International Settlements compiles national data from each central bank to produce global market statistics.

The Federal Reserve System and other government agencies use the survey to monitor activity in the foreign exchange and derivatives markets. Respondents use the published data to gauge their market share.

Current Actions: On November 2, 2009, the Federal Reserve published a notice in the **Federal Register** (74 FR 56633) requesting public comment for 60 days on the implementation of the FR 3036. The comment period for this notice expired on January 4, 2010. The Federal Reserve did not receive any comments. The survey will be implemented as proposed.

Board of Governors of the Federal Reserve System, January 12, 2010.

Jennifer J. Johnson,
Secretary of the Board.

[FR Doc. 2010-751 Filed 1-15-10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Biodefense Science Board: Notification of Public Teleconference

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

ACTION: Notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the Department of Health and Human Services (HHS) is hereby giving notice that the National Biodefense Science Board (NBSB) will hold a teleconference meeting. The meeting is open to the public. Pre-registration is NOT required, however, individuals who wish to participate in the public comment session should e-mail NBSB@HHS.GOV to RSVP.

DATES: The meeting will be held on February 10, 2010 from 2 p.m. to 4 p.m. ET.

ADDRESSES: The meeting will occur by teleconference. To attend, call 1-866-395-4129, pass-code "ASPR." Please call 15 minutes prior to the beginning of the conference call to facilitate attendance.

FOR FURTHER INFORMATION: E-mail: NBSB@HHS.GOV

SUPPLEMENTARY INFORMATION: Pursuant to section 319M of the Public Health Service Act (42 U.S.C. 247d-7f) and section 222 of the Public Health Service Act (42 U.S.C. 217a), the Department of Health and Human Services established the National Biodefense Science Board. The Board shall provide expert advice and guidance to the Secretary on scientific, technical, and other matters of special interest to the Department of Health and Human Services regarding current and future chemical, biological, nuclear, and radiological agents, whether naturally occurring, accidental, or deliberate. The Board may also provide advice and guidance to the Secretary on other matters related to public health emergency preparedness and response.

The Board will discuss and consider recommendations from the National Biodefense Science Board's Medical Countermeasure Markets and Sustainability Working Group report titled "Inventory of Issues Constraining or Enabling Industry Involvement in Medical Countermeasure Efforts."

Members of the public are invited to attend by teleconference via a toll-free call-in phone number. The teleconference will be operator assisted to allow the public the opportunity to provide comments to the Board. Public participation will be limited to time and space available. Public comments will

be limited to no more than 3 minutes per speaker. To be placed on the public comment list, notify the operator when you join the teleconference.

Public comments received by close of business one week prior to each teleconference will be distributed to the NBSB in advance. Submit comments via e-mail to NBSB@HHS.GOV, with "NBSB Public Comment" as the subject line.

A draft agenda and any additional materials/agendas will be posted on the NBSB Web site (<http://www.hhs.gov/aspr/omsph/nbsb/>) prior to the meeting.

Dated: January 11, 2010.

Nicole Lurie,

Assistant Secretary for Preparedness and Response.

[FR Doc. 2010-778 Filed 1-15-10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2010-N-0001]

Advisory Committees; Filing of Closed Meeting Reports

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that, as required by the Federal Advisory Committee Act, the agency has filed with the Library of Congress the annual reports of those FDA advisory committees that held closed meetings during fiscal year 2009.

ADDRESSES: Copies are available from the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, 301-827-6860.

FOR FURTHER INFORMATION CONTACT:

Teresa L. Hays, Committee Management Officer, Advisory Committee Oversight and Management Staff (HF-4), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1220.

SUPPLEMENTARY INFORMATION: Under section 10(d) of the Federal Advisory Committee Act (5 U.S.C. app. 1) and 21 CFR 14.60(d), FDA has filed with the Library of Congress the annual reports for the following FDA advisory committees that held closed meetings during the period October 1, 2008, through September 30, 2009: *Center for Biologics Evaluation and Research;*

Blood Products Advisory Committee,