

20892. The meeting will be open to the public with attendance limited to space available. The meeting will also be Web cast.

The main agenda item will be a review of the revised draft report on genetics education and training and discussion of the final draft recommendations. The meeting will also include sessions on genomic data sharing and the implications of affordable whole-genome sequencing, an update on the implementation of the Genetic Information Nondiscrimination Act, and a briefing from the Food and Drug Administration on activities related to genetic testing.

As always, the Committee welcomes hearing from anyone wishing to provide public comment on any issue related to genetics, health and society. Please note that because SACGHS operates under the provisions of the Federal Advisory Committee Act, all public comments will be made available to the public. Individuals who would like to provide public comment should notify the SACGHS Executive Secretary, Ms. Sarah Carr, by telephone at 301-496-9838 or e-mail at carrs@od.nih.gov. The SACGHS office is located at 6705 Rockledge Drive, Suite 750, Bethesda, MD 20892. Anyone planning to attend the meeting who needs special assistance, such as sign language interpretation or other reasonable accommodations, is also asked to contact the Executive Secretary.

Under authority of 42 U.S.C. 217a, Section 222 of the Public Health Service Act, as amended, the Department of Health and Human Services established SACGHS to serve as a public forum for deliberations on the broad range of human health and societal issues raised by the development and use of genetic and genomic technologies and, as warranted, to provide advice on these issues. The draft meeting agenda and other information about SACGHS, including information about access to the Web cast, will be available at the following Web site: http://oba.od.nih.gov/SACGHS/sacghs_meetings.html.

Dated: August 24, 2010.

Jennifer Spaeth,

Director, NIH Office of Federal Advisory Committee Policy.

[FR Doc. 2010-21532 Filed 8-27-10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Committee on Vital and Health Statistics: Meeting

Pursuant to the Federal Advisory Committee Act, the Department of Health and Human Services (HHS) announces the following advisory committee meeting.

Name: National Committee on Vital and Health Statistics (NCVHS), Full Committee Meeting.

Time and Date: September 15, 2010 9 a.m.–2 p.m.; September 16, 2010 8:30 a.m.–12:30 p.m.

Place: Embassy Suites Crystal City Hotel, 1300 Jefferson Davis Highway, Arlington, VA 22202, (703) 979-9799.

Status: Open.

Purpose: At this meeting the Committee will hear presentations and hold discussions on several health data policy topics. On the morning of the first day the Committee will hear updates from the Department, the Center for Medicare and Medicaid Services, and the Office of the National Coordinator for Health Information Technology. Draft letters to the HHS Secretary regarding the HIPAA national health plan identifier and operating rules on eligibility and claim status will also be discussed. In the afternoon there will be a discussion about a letter to the HHS Secretary regarding sensitive information in medical records.

On the morning of the second day there will be a review of the final letters regarding the national health plan identifier, operating rules on eligibility and claim status, and sensitive information in medical records. Subcommittees will also present their reports. The afternoon of the second day will conclude with a discussion of the 60th Anniversary Symposium that was held in June 2010.

The times shown above are for the full Committee meeting. Subcommittee breakout sessions can be scheduled for late in the afternoon of the first day and second day and in the morning prior to the full Committee meeting on the second day. Agendas for these breakout sessions will be posted on the NCVHS Web site (URL below) when available.

Contact Person for More Information: Substantive program information as well as summaries of meetings and a roster of committee members may be obtained from Marjorie S. Greenberg, Executive Secretary, NCVHS, National Center for Health Statistics, Centers for Disease Control and Prevention, 3311 Toledo Road, Room 2402, Hyattsville, Maryland 20782, telephone (301) 458-4245. Information also is available on the NCVHS home page of the HHS Web site: <http://www.ncvhs.hhs.gov/>, where further information including an agenda will be posted when available.

Should you require reasonable accommodation, please contact the CDC Office of Equal Employment Opportunity on (301) 458-4EEO (4336) as soon as possible.

Dated: August 24, 2010.

James Scanlon,

Deputy Assistant Secretary for Planning and Evaluation (Science and Data Policy), Office of the Assistant Secretary for Planning and Evaluation.

[FR Doc. 2010-21516 Filed 8-27-10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the National Biodefense Science Board

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 National Biodefense Science Board (NBSB) will be holding a public meeting. The meeting is open to the public.

DATES: The NBSB will hold a public meeting on September 22, 2010 from 8 a.m. to 5 p.m. ET. The agenda is subject to change as priorities dictate.

ADDRESSES: Washington, DC Metro Area. The venue details will be posted on the NBSB webpage at <http://www.phe.gov/Preparedness/legal/boards/nbsb/Pages/default.aspx> as they become available.

FOR FURTHER INFORMATION CONTACT: 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.

Background: A portion of this public meeting will be dedicated to a report and presentation by the Disaster Mental Health Subcommittee to the NBSB on their assessment of the Department of Health and Human Services' progress to better integrate behavioral health into emergency preparedness and response

activities. Subsequent agenda topics will be added as priorities dictate.

Availability of Materials: The meeting agenda and materials will be posted on the NBSB Web site at <http://www.phe.gov/Preparedness/legal/boards/nbsb/Pages/default.aspx> prior to the meeting.

Procedures for Providing Public Input: Any member of the public providing oral comments at the meeting must sign in at the registration desk and provide his/her name, address, and affiliation. All written comments must be received prior to September 21, and should be sent by e-mail to NBSB@HHS.GOV with "NBSB Public Comment" as the subject line. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should e-mail NBSB@HHS.GOV.

Dated: August 10, 2010.

Nicole Lurie,

Assistant Secretary for Preparedness and Response.

[FR Doc. 2010-21504 Filed 8-27-10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) approve the proposed information collection project: "Understanding Patients' Knowledge and Use of Acetaminophen—Phase 2." In accordance with the Paperwork Reduction Act, 44 U.S.C. 3501-3520, AHRQ invites the public to comment on this proposed information collection.

DATES: Comments on this notice must be received by October 29, 2010.

ADDRESSES: Written comments should be submitted to: Doris Lefkowitz, Reports Clearance Officer, AHRQ, by e-mail at doris.lefkowitz@AHRQ.hhs.gov. Copies of the proposed collection plans, data collection instruments, and specific details on the estimated burden can be obtained from the AHRQ Reports Clearance Officer.

FOR FURTHER INFORMATION CONTACT: Doris Lefkowitz, AHRQ Reports Clearance Officer, (301) 427-1477, or by e-mail at doris.lefkowitz@AHRQ.hhs.gov.

SUPPLEMENTARY INFORMATION:

Proposed Project

Understanding Patients' Knowledge and Use of Acetaminophen—Phase 2

AHRQ proposes a cross-sectional prospective survey to identify issues that relate to the misuse and overdosing of over-the-counter (OTC) acetaminophen. The survey was developed based on results from a previous data collection (OMB control number 0935-0154, approved on 10/13/2009). Acetaminophen is the most widely used analgesic and antipyretic drug in the U.S. When appropriately used, it is a very safe agent. However, a single large overdose, or several supratherapeutic dosages in a short period of time, has been associated with acute liver failure, which can occur with dosages over 250 mg/kg over a 24-hour period, or > 12 g in an adult. Toxicity from acetaminophen has been on the rise in the past 3 decades, and is now the most common cause of acute liver failure in the U.S., surpassing viral hepatitis.

This project has the following aims:

- (1) To estimate frequency of use, knowledge, and practices regarding use of OTC acetaminophen, and
 - (2) To evaluate potential determinants of misuse in community-based samples.
- This information will be useful for policy makers to consider and to evaluate regulations and legislation with respect to the distribution, dispensing and sales of OTC acetaminophen.

This study is being conducted by AHRQ through its contractor, the University of Texas. This project

supports AHRQ's Centers for Education and Research on Therapeutics initiative to promote the safe and effective use of therapeutics. See 42 U.S.C. 299b-1(b). It also supports AHRQ's mandate for the inclusion of priority populations. See 42 U.S.C. 299(c).

Method of Collection

To achieve the projects' aims the following data collections will be implemented:

(1) Surveys with parents of young children (age < 8 years). The purpose of this survey is to learn how parents administer acetaminophen to their children and to identify determinants of misuse of acetaminophen;

(2) Surveys with adolescents (ages 13 to 20). The purpose of this survey is to learn how adolescents use acetaminophen and to identify determinants of misuse of acetaminophen;

(3) Surveys with adults (21 to 65 years of age). The purpose of this survey is to learn how adults use acetaminophen and to identify determinants of misuse of acetaminophen;

(4) Surveys with adults (greater than 65 years of age). The purpose of this survey is to learn how older adults use acetaminophen and to identify determinants of misuse of acetaminophen, particularly in regards to age-related factors.

Data will be collected in person using paper questionnaires administered by the project personnel.

Estimated Annual Respondent Burden

Exhibit 1 shows the estimated annualized burden hours for the respondent's time to participate in this project. Each of the four questionnaires used in the planned face-to-face surveys will require approximately 30 minutes to complete. The total annualized burden for all participants is estimated to be 400 hours.

Exhibit 2 shows the estimated annualized cost burden for the respondent's time to participate in the project. The total annualized cost burden is estimated to be \$8,361.

EXHIBIT 1—ESTIMATED ANNUALIZED BURDEN HOURS

Data collection mode	Number of respondents	Number of responses per respondent	Hours per response	Total burden hours
Surveys with Parents of Children < 8 years of age	300	1	30/60	150
Surveys with Adolescents (13 to 20 years of age)	200	1	30/60	100
Surveys with Adults (20 to 65 years)	150	1	30/60	75
Surveys with Adults (greater than 65 years)	150	1	30/60	75