comments and ATSDR responses, is available on the ATSDR Web site at: *http://www.atsdr.cdc.gov/substances/dioxin/policy/index.html.*

DATES: Comments concerning this document must be received by February 27, 2007.

ADDRESSES: Public comments should be forwarded to Ms. Athena Gemella, ATSDR, Office of Science, 1600 Clifton Road, N.E., Mail stop E–28. Atlanta, GA. 30333, or e-mail at *AGemella@cdc.gov.*

FOR FURTHER INFORMATION CONTACT: Ms. Athena Gemella, Office of Science, telephone (404) 498–0621.

Dated: December 22, 2006.

Kenneth Rose,

Acting Director, Office of Policy, Planning and Evaluation, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry.

[FR Doc. E6–22388 Filed 12–28–06; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-07–06AI]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an e-mail to *omb@cdc.gov.* Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–6974. Written comments should be received within 30 days of this notice.

Proposed Project

Metropolitan Atlanta Stillbirth Management Survey: Knowledge, Attitudes and Practice Patterns from Obstetricians,—New—National Center on Birth Defects and Developmental Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The U.S. Congress House Report 108–792 (joint conference report for the Fiscal Year 2005 omnibus appropriations bill) provides specific funding to devise a comprehensive strategy for expanding existing birth defects surveillance systems to incorporate surveillance data on all intrauterine fetal deaths of 20 or more week's gestation into the Metropolitan Atlanta Congenital Defects Program (MACDP). Stillbirth is largely an understudied adverse pregnancy outcome even though it accounts for nearly one half of all perinatal mortality. There is currently no nationally accepted definition of what constitutes a stillbirth, and there are no universally recommended, standardized stillbirth evaluation protocols in use for the evaluation of fetal deaths. The proposed survey has been designed to evaluate and assess the knowledge, attitudes and practice management patterns of obstetricians in the metropolitan Atlanta area regarding stillbirths in general, as well as in their medical practice. This information will be used to identify prevailing deficiencies leading to incomplete and inaccurate reporting of data relative to stillbirths, and to develop targeted awareness and educational strategies for participating MACDP facilities. Ongoing, accurate and reliable population-based registries of stillbirths are essential for conducting epidemiologic studies on the causes of and risk factors for this pregnancy outcome. This survey will be mailed to randomly selected obstetricians whose practices serve residents of the 5 counties comprising metropolitan Atlanta. This survey will be conducted once and will take approximately 2–3 months to collect the data. NCBDDD is requesting OMB clearance for 1 (one) year. There is no cost to the survey respondents except for the time necessary to complete the survey. The total annual burden hours are 122

ESTIMATE OF ANNUALIZED BURDEN HOURS

Respondents	Participant status	Number of respondents	Number of reponses per respondent	Average burden per response (in hours)
Obstetricians	Non-Participant	120	1	1/60
	Participant	480	1	15/60

Dated: December 22, 2006.

Joan F. Karr,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E6–22381 Filed 12–28–06; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–R–284]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid Services (CMS), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the Agency's function; (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.

1. *Type of Information Collection Request:* Revision of a currently approved collection;.

Title of Information Collection: Medicaid Statistical Information System.

Use: State data are reported by the Federally mandated electronic process, known as Medicaid Statistical Information System (MSIS). These data are the basis of actuarial forecasts for Medicaid service utilization and costs; of analysis and cost savings estimates required for legislative initiatives relating to Medicaid; and for responding to requests for information from CMS components, the Department, Congress and other customers. Form Number: CMS–R–284 (OMB#: 0938–0345).

Frequency: Quarterly.

Affected Public: State, Local or Tribal Government.

Number of Respondents: 53.

Total Annual Responses: 212.

Total Annual Hours: 3,392.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS Web Site address at *http://www.cms.hhs.gov/PaperworkReductionActof1995*, or E-mail your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov*, or call the Reports Clearance Office on (410) 786–1326.

Written comments and recommendations for the proposed information collections must be mailed or faxed within 30 days of this notice directly to the OMB desk officer: OMB Human Resources and Housing Branch, Attention: Katherine Astrich, New Executive Office Building, Room 10235, Washington, DC 20503, Fax Number: (202) 395–6974.

Dated: December 21, 2006.

Michelle Shortt,

Director, Regulations Development Group, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. E6–22233 Filed 12–28–06; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committees; Tentative Schedule of Meetings for 2007

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing a tentative schedule of forthcoming meetings of its public advisory committees for 2007. During 1991, at the request of the Commissioner of Food and Drugs (the Commissioner), the Institute of Medicine (the IOM) conducted a study of the use of FDA's advisory committees. In its final report, one of the IOM's recommendations was for the agency to publish an annual tentative schedule of its meetings in the **Federal Register**. This publication implements the IOM's recommendation.

FOR FURTHER INFORMATION CONTACT: Theresa L. Green, Advisory Committee Oversight and Management Staff (HF–4), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1220.

SUPPLEMENTARY INFORMATION: The IOM, at the request of the Commissioner, undertook a study of the use of FDA's advisory committees. In its final report in 1992, one of the IOM's recommendations was for FDA to adopt a policy of publishing an advance yearly schedule of its upcoming public advisory committee meetings in the **Federal Register**; FDA has implemented this recommendation. The annual publication of tentatively scheduled advisory committee meetings will provide both advisory committee members and the public with the opportunity, in advance, to schedule attendance at FDA's upcoming advisory committee meetings. Because the schedule is tentative amendments to this notice will not be published in the **Federal Register**. However, changes to the schedule will be posted on the FDA advisory committees' Internet site located at *http://www.fda.gov/oc/advisory/default.htm*. FDA will continue to publish a **Federal Register** notice 15 days in advance of each upcoming advisory committee meeting, to announce the meeting (21 CFR 14.20).

The following list announces FDA's tentatively scheduled advisory committee meetings for 2007. You may also obtain up-to-date information by calling the Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area).

TABLE 1.

Committee Name	Tentative Date(s) of Meeting(s)	Advisory Committee 10–Digit Information Line Code
OFFICE OF THE COMMISSIONER		
Pediatric Advisory Committee	April day(s) to be announced.	8732310001
Science Board to the Food and Drug Administration	June day(s) to be announced.	3014512603
CENTER FOR BIOLOGICS EVALUATION AND RESEARCH		
Allergenic Products Advisory Committee	April 18, October 19.	3014512388
Blood Products Advisory Committee	April 26–27, August 16–17, December 13–14.	3014519516
Cellular Tissue and Gene Therapies Advisory Committee	March 29–30, July 26–27, November 15–16.	3014512389
Transmissible Spongiform	Encephalopathies Advisory Committee to be announced.	3014512392
Vaccines and Related Biological Advisory Committee	February 27–28, May 16–17, September 19–20, November 14–15.	3014512391
CENTER FOR DRUG EVALUATION AND RESEARCH		
Anesthetic and Life Support Drugs Advisory Committee	March 29.	3014512529