DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the National Coordinator for Health Information Technology, American Health Information Community Population Health and Clinical Care Connections Workgroup Meeting

ACTION: Announcement of meeting.

SUMMARY: This notice announces the 14th meeting of the American Health Information Community Population Care and Clinical Care Connections Workgroup [formerly Biosurveillance Workgroup] in accordance with the Federal Advisory Committee Act (Pub. L. 92–463, 5 U.S.C., App.)

DATES: March 2, 2007, from 1 p.m. to 4 p.m.

ADDRESSES: Mary C. Switzer Building (330 C Street, SW, Washington, DC 20201), Conference Room 4090 (please bring photo ID for entry to a Federal building)

FOR FURTHER INFORMATION CONTACT: *http://www.hhs.gov/healthit/ahic/population/*

SUPPLEMENTARY INFORMATION: The Workgroup will discuss the priority area of Response Management.

The meeting will be available via internet access. For additional information, go to *http://www.hhs.gov/healthit/ahic/population/pop_instructhtml.*

Dated: February 7, 2007.

Judith Sparrow,

Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.

[FR Doc. 07–707 Filed 2–14–06; 8:45 am]

BILLING CODE 4150–24–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the National Coordinator for Health Information Technology; American Health Information Community Meeting

ACTION: Announcement of meeting.

SUMMARY: This notice announces the 12th meeting of the American Health Information Community in accordance with the Federal Advisory Committee Act (Pub. L. 92–463, 5 U.S.C., App.) The American Health Information Community will advise the Secretary and recommend specific actions to achieve a common interoperability framework for health information technology (IT).

DATES: March 13, 2007, from 8:30 a.m. to 3 p.m.

ADDRESSES: Hubert H. Humphrey building (200 Independence Avenue, SW., Washington, DC 20201), Conference Room 800.

FOR FURTHER INFORMATION CONTACT: Visit *http://www.hhs.gov/healthit/ahic.html.*

SUPPLEMENTARY INFORMATION: The meeting will include presentations by the Quality, Population Health and Clinical Care Connections, Consumer Empowerment, and Confidentiality, Privacy and Security Workgroups on their Recommendations; an update on the Certification Commission for Healthcare Information Technology (CCHIT); and a panel presentation on Privacy and Security issues.

A Web cast of the Community meeting will be available on the NIH Web site at: *http://www.videocast.nih.gov/.*

If you have special needs for the meeting, please contact (202) 690–7151.

Dated: February 7, 2007.

Judith Sparrow,

Director, American Health Information Community, Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.

[FR Doc. 07–708 Filed 2–14–07; 8:45 am]

BILLING CODE 4150–24–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day–07–05CO]

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

The Centers for Disease Control and Prevention's Consumer Response Services Center (CDC–INFO) Evaluation-New-National Center for Health Marketing (NCHM), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC is launching an integrated ''one face to the public'' approach across all communication channels to handle inquiries concerning a broad spectrum of public health topics. The overall objective is to ensure consistent, timely, reliable health information for dissemination to a variety of consumers (public, health professionals, researchers, etc.) and to address variations in inquiry volumes related to public health emergencies, news events, and dynamic, shifting public health priorities. The CDC has integrated over 40 hotlines into one Consumer Response Services Center CDC–INFO. CDC–INFO has an exceptionally wide scope because content currently divided between over 40 hotlines handling nearly 2,000,000 telephone contacts annually will be consolidated under CDC–INFO. All CDC hotlines were consolidated in one center beginning in February 2005, with all CDC program areas transitioning into CDC–INFO through a phased approach during the next three years. CDC–INFO itself will be operational for at least the next seven years. The primary objectives of the national evaluation are to (1) Proactively evaluate customer interactions and service effectiveness by employing assessment measures and data collection mechanisms to support performance management, gathering insights and understandings for improving service levels, and implementing effective measures to meet customer satisfaction goals; (2) develop an ongoing understanding of customer requirements and satisfaction trends to achieve best of practice quality standards and to provide qualitative assessments, quantitative data, and cost factors to drive improvement and reinforce operational objectives; (3) measure CDC–INFO contractor service performance to assist in determining whether performance incentives have been achieved; and (4) to collect data in order to address public concern and response to emergencies, outbreaks, and media events.

Sample size, respondent burden, and intrusiveness have been minimized to be consistent with national evaluation objectives. Procedures will be employed to safeguard the privacy and confidentiality of participants. Pilot tests assisted in controlling burden and ensuring the user-relevance of questions. The following table shows the estimated annualized burden for data collection. There are no respondent

costs other than the amount of time required to respond to the survey.

ESTIMATED ANNUALIZED BURDEN HOURS

Data collection instrument	Number of respondents	Responses/ respondent	Average burden per response (in hrs)	Average annual burden hours
Satisfaction survey (callers)	25,000	1	3/60	1,250
Satisfaction survey (e-mail inquiries)	330	1	3/60	17
Follow up survey	3,125	1	7/60	365
Key informant survey	100	1	7/60	12
Postcard survey for bulk mailing	950	1	1/60	16
Postcard survey for individual publications	2,100	1	1/60	35
Web survey for e-mail publication orders	1,000	1	1/60	17
Web survey for internet publications	950	1	1/60	16
Special event/Outreach survey—General Public	25,600	1	5/60	2,133
Special event/Outreach survey—Professionals	10,400	1	5/60	867
Emergency response survey—Level 1 emergency—General Public	31,151	1	5/60	2596
Emergency response survey—Level 1 emergency—Professionals	7,459	1	5/60	622
Emergency response survey—Level 2 emergency—General Public	57,579	1	5/60	4798
Emergency response survey—Level 2 emergency—Professionals	51,821	1	5/60	4318
Emergency response survey—Level 3 emergency—General Public	351,863	1	5/60	29,322
Emergency response survey—Level 3 emergency—Professional	316,678	1	5/60	26,390
Emergency response survey—Level 4 emergency—General Public	645,630	1	5/60	53,803
Emergency response survey—Level 4 emergency—Professional	596,504	1	5/60	49,709
Total Burden Hours				176,286

Dated: February 6, 2007.

Joan F. Karr,

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

[FR Doc. E7–2637 Filed 2–14–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0430]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; General Licensing Provisions: Biologics License Application, Changes to an Approved Application, Labeling, Revocation and Suspension, Postmarketing Studies Status Reports, and Forms FDA 356h and 2567; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of February 2, 2007 (72 FR 5057). The document announced that an opportunity for public comment on a proposed collection of information had been submitted to the Office of Management and Budget for review and clearance under the Paperwork Reduction Act of 1995. The notice published with an error in titles referring to an FDA form number in two places in the document. This document corrects those errors.

DATES: February 15, 2007.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of Friday, February 2, 2007, the following corrections are made on page 5057:

1. In the first column, in the ninth line of the title of the document, the phrase ''Forms FDA 456h'' is corrected to read ''Forms FDA 356h''.

2. In the second column, in the **SUPPLEMENTARY INFORMATION** section of the document, in the sixth line of the title, the phrase ''Forms FDA 456h'' is corrected to read ''Forms FDA 356h''.

Dated: February 8, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–2576 Filed 2–14–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0436]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on How To Use E-Mail To Submit a Study Protocol

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by March 19, 2007.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–6974.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Jr., Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers