the use of photographic images, the ONC invites suggestions for extending the HL7 DAM to accommodate images.

Ownership of intellectual property is determined by the following:

- Each entrant retains title and full ownership in and to their submission. Entrants expressly reserve all intellectual property rights not expressly granted under the challenge agreement.
- By participating in the challenge, each entrant hereby irrevocably grants to Sponsor and Administrator a limited, non-exclusive, royalty free, worldwide, license and right to reproduce, publically perform, publically display, and use the Submission to the extent necessary to administer the challenge, and to publically perform and publically display the Submission, including, without limitation, for advertising and promotional purposes relating to the challenge.

Authority: 15 U.S.C. 3719.

Dated: November 20, 2012.

Farzad Mostashari,

National Coordinator for Health Information Technology.

[FR Doc. 2012–29524 Filed 12–5–12; 8:45 am]

BILLING CODE 4150–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–13–0214]

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 (404) 639–7570 or send an email to *omb@cdc.gov*. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

National Health Interview Survey (NHIS), (OMB No. 0920–0214 expiration 08/31/2014)—Revision—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Section 306 of the Public Health Service (PHS) Act (42 U.S.C. 242k), as amended, authorizes that the Secretary of Health and Human Services (DHHS), acting through NCHS, shall collect statistics on the extent and nature of illness and disability of the population of the United States.

The annual National Health Interview Survey is a major source of general statistics on the health of the U.S. population and has been in the field continuously since 1957. Clearance is sought for three years, to collect data for 2013, 2014, and 2015. This voluntary household-based survey collects demographic and health-related information on a nationally representative sample of persons and households throughout the country. Information is collected using computer assisted personal interviews (CAPI). A core set of data is collected each year while sponsored supplements vary from year to year. For 2013, there are supplementary questions on cancer screening, asthma, immune suppression, hepatitis, epilepsy, HIV testing, neighborhood characteristics, financial worries, sleep issues, and sexual identity.

Cases in a 5,000 case test were randomly assigned to receive questions on HIV testing, neighborhood characteristics, financial worries, sleep issues, and sexual identity in either CAPI or ACASI. Prevalence estimates for the sexual identity questions were compared by mode of administration. Since a documented advantage of ACASI is the enhanced level of privacy it affords, we anticipated higher prevalence estimates from this mode of administration. Estimates were similar for the two modes of administration. Therefore, the questions will be administered in CAPI, the more cost efficient mode.

In accordance with the 1995 initiative to increase the integration of surveys within the Department of Health and Human Services, respondents to the NHIS serve as the sampling frame for the Medical Expenditure Panel Survey conducted by the Agency for Healthcare Research and Quality. The NHIS has long been used by government, university, and private researchers to evaluate both general health and specific issues, such as cancer, diabetes, and access to health care. It is a leading source of data for the Congressionally-mandated ''Health US'' and related publications, as well as the single most important source of statistics to track progress toward the National Health Promotion and Disease Prevention Objectives.

There is no cost to the respondents other than their time. As shown below, the estimated overall average annual burden for the 2013, 2014, and 2015 surveys is 57,099 hours.

ANNUALIZED BURDEN TABLE

Questionnaire (respondent)	Number of respondents	Number of responses per respondent	Average burden per respondent in hours
Screener Questionnaire	12,000	1	5/60
Family Core (adult family member)	55,000	1	23/60
Adult Core (sample adult)	44,000	1	15/60
Child Core (adult family member)	17,000	1	10/60
Child/Teen Record Check (medical provider)	10,000	1	5/60
Supplements (adult family member)	60,000	1	12/60
Sexual Identity Module (adult family member)	44,000	1	4/60
Multi-mode study (adult family member)	5,000	1	30/60
Reinterview Survey	5,000	1	5/60
Sample Frame Test (adult family member)	5,000	1	30/60

Kimberly S. Lane,

Deputy Director, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2012–29474 Filed 12–5–12; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

The Centers for Disease Control (CDC)/ Health Resources and Services Administration (HRSA) Advisory Committee on HIV, Viral Hepatitis and STD Prevention and Treatment Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the CDC/HRSA Advisory Committee on HIV, Viral Hepatitis and STD Prevention and Treatment, Department of Health and Human Services, has been renewed for a 2-year period through November 25, 2014.

Contact Person for More Information: Kevin Fenton, M.D., Ph.D., Designated Federal Officer, CDC/HRSA Advisory Committee on HIV, Viral Hepatitis and STD Prevention and Treatment, Department of Health and Human Services, CDC, 1600 Clifton Road, NE., Mailstop E07, Atlanta, Georgia 30333, telephone (404) 639–8000 or fax (404) 639–8600.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: November 29, 2012.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2012–29471 Filed 12–5–12; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Request for Nominations for Candidates To Serve on the National Public Health Surveillance and Biosurveillance Advisory Committee (NPHSBAC)

Correction: This notice was published in the **Federal Register** on November 1, 2012 Volume 77, Number 215, page 66620. This notice is to announce the extension of submission for potential nominees.

Nominations should be sent, in writing, and postmarked by December 21, 2012: Vernellia Johnson, Management and Program Analyst, Public Health Surveillance and Informatics Program Office, Centers for Disease Control and Prevention, Office of Surveillance, Epidemiology and Laboratory Services Century, 1600 Clifton Road NE., MS E–97, Atlanta, GA 30333 or via email to *hft9@cdc.gov.* Telephone and facsimile submissions cannot be accepted.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: November 30, 2012.

Cathy Ramadei,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 2012–29478 Filed 12–5–12; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–D–1086]

Compliance Guidance for Small Business Entities on Labeling and Effectiveness Testing; Sunscreen Drug Products for Over-the-Counter Human Use; Notice of Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a compliance guidance for small business entities entitled ''Labeling and Effectiveness Testing: Sunscreen Drug Products for Over-the-Counter Human Use; Small Entity Compliance Guide.'' This guidance is intended to help small businesses understand and comply with the requirements of the final rule addressing labeling and effectiveness testing requirements for over-the counter (OTC) sunscreen drug products. The guidance describes the requirements of the final rule in plain language and provides answers to common questions on how to comply with the rule. This guidance was prepared in accordance with the Small Business Regulatory Enforcement Fairness Act.

DATES: Submit either electronic or written comments on Agency guidances at any time.

ADDRESSES: Submit written requests for single copies of this guidance to the Division of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 2201, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your requests. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance document.

Submit electronic comments on the guidance to *http://www.regulations.gov.* Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Reynold Tan, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 22, Rm. 5493, Silver Spring, MD 20993–0002, 301–796–1009.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a compliance guidance for small business entities entitled ''Labeling and Effectiveness Testing: Sunscreen Drug Products for Over-the-Counter Human Use; Small Entity Compliance Guide.'' This guidance summarizes the June 17, 2011, final rule (76 FR 35620) regarding labeling and testing requirements for OTC sunscreen drug products. Under the 2011 sunscreen final rule, required and permitted labeling is based upon the results of effectiveness testing. The effectiveness testing consists of a sun protection factor (SPF) Test and a Broad Spectrum (ultraviolet A (UVA) and ultraviolet B (UVB) protection) Test. In addition, a test demonstrating water resistance that accompanies the SPF Test to ensure retention of SPF