within the Department of Health and Human Services. Information about the CICP is available at the toll free number 1–855–266–2427 or *http://www.hrsa.gov/cicp/.*

XV. Amendments

42 U.S.C. 247d–6d(b)(4)

Any amendments to this declaration will be published in the **Federal Register**.

Authority: 42 U.S.C. 247d–6d.

Dated: December 3, 2014.

Sylvia M. Burwell,

Secretary.

[FR Doc. 2014–28856 Filed 12–9–14; 8:45 am]

BILLING CODE 4150–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the National Coordinator for Health Information Technology; Federal Health IT Strategic Plan: 2015–2020 Open Comment Period

AGENCY: ONC, HHS.

ACTION: Notice.

Authority: Section 3001(c)(3) of the Public Health Service Act.

SUMMARY: Section 3001(c)(3) of the Public Health Service Act, as added by the Health Information Technology for Economic and Clinical Health (HITECH) Act, requires the National Coordinator for Health Information Technology (ONC) to update the Federal Health IT Strategic Plan (developed June 3, 2008; last updated on September 15, 2011) in consultation with other appropriate federal agencies and in collaboration with private and public entities. The Plan was developed in collaboration across multiple federal agencies, and ONC will seek input on the draft Plan from the private sector through the Health IT Policy Committee. This notice serves to announce that the public comment period for the Federal Health IT Strategic Plan is open through Tuesday, February 6 at 5:00 p.m. (Eastern). ONC welcomes and encourages all comments from the public regarding the Plan.

In order for your comments to be read and considered, you must submit your comment via *http://www.healthit.gov/policy-researchers-implementers/strategic-plan-public-comments.*

FOR FURTHER INFORMATION CONTACT: Matthew Swain, Program Analyst in the Office of Planning, Evaluation, and Analysis, *matthew.swain@hhs.gov,* 202.205.3754.

Dated: December 4, 2014.

Matthew Swain,

Program Analyst, Office of Planning, Evaluation, and Analysis, Office of the National Coordinator for Health Information Technology (ONC), Office of the Secretary (OS).

[FR Doc. 2014–28855 Filed 12–9–14; 8:45 am]

BILLING CODE 4150–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Mine Safety and Health Research Advisory Committee; Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the Mine Safety and Health Research Advisory Committee, Centers for Disease Control and Prevention, Department of Health and Human Services, has been renewed for a 2-year period through November 30, 2016.

For information, contact Jeffrey H. Welsh, B.A., Designated Federal Officer, Mine Safety and Health Research Advisory Committee, Centers for Disease Control and Prevention, Department of Health and Human Services, 626 Cochrans Mill Road, Mailstop P05, Pittsburgh, Pennsylvania 15236, Telephone (412) 386–4040 or fax (412) 386–6614.

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.

Claudette Grant,

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

[FR Doc. 2014–28933 Filed 12–9–14; 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, 2016.

Contact Person for More Information: Johnathan Mermin, M.D., M.P.H., 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.

Claudette Grant,

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

[FR Doc. 2014–28932 Filed 12–9–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Head Start Impact Study Participants Beyond 8th Grade.

OMB No.: 0970–0229.

Description: The Administration for Children and Families (ACF) within the Department of Health and Human Services (HHS) will collect follow-up information from children and families in the Head Start Impact Study. In anticipation of conducting a future follow-up for the study, ACF will collect information necessary to identify

respondents' current location and follow-up with respondents in the future.

The Head Start Impact Study is a longitudinal study involving 4,667 first-time enrolled three- and four-year-old preschool children across 84 nationally representative grantee/delegate agencies (in communities where there were more eligible children and families than can be served by the program). Participants were randomly assigned to either a Head Start group (that could enroll in Head Start services) or a control group (that could not enroll in Head services) or a control group (that could not enroll in Head Start services but could enroll in other available services selected by their parents). Data collection for the study began in fall of 2002 and has continued through late spring 2008 to include the participants' 3rd grade year. Location and contact information for participants has continued every spring beginning in 2009 and continued through spring 2014.

ACF will continue to collect a small amount of information for the sample through the spring of the participant's 12th grade year. To maintain adequate sample size, telephone interviews (with in-person follow-up as necessary) will be conducted in order to update the children's status and their location and contact information. Additionally, the parent interview will include a small set of items on children's special education needs, grade retention, school safety, school engagement, and parental monitoring to provide information on factors during adolescence that may influence long-term impacts of Head Start examined in a potential follow-up study. This information will be collected from parents or guardians in the spring of 2015 and 2016. Updates will take about 20 minutes to complete.

Respondents: The original sample of 4,667 treatment and control group members in the Head Start Impact Study, less 432 families that have given a ''hard'' refusal to participate in the study (*e.g.*, refused to participate if they were contacted again). The number of respondents for this requested data collection is 4,235.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
Parent Interview	8470	4235	1	1/3	1412

Estimated Total Annual Burden Hours: 1412.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: OPRE Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *OPREinfocollection@acf.hhs.gov*.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV*, Attn: Desk Officer for the Administration, for Children and Families.

Naomi Goldstein,

Director, Office of Planning, Research and Evaluation; Administration for Children and Families.

[FR Doc. 2014–28843 Filed 12–9–14; 8:45 am]

BILLING CODE 4184–22–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–2029]

Agency Information Collection Activities; Proposed Collection; Comment Request; Administrative Practices and Procedures; Formal Evidentiary Public Hearing

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on reporting requirements contained in current FDA regulations: Administrative Practices and Procedures; Formal Evidentiary Public Hearing.

DATES: Submit either electronic or written comments on the collection of information by February 9, 2015.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov.* Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.