confirmation when they have been accepted. If time and space permit, onsite registration on the day of the public workshop will be provided beginning at 7:30 a.m. We will let registrants know if registration closes before the day of the public workshop.

If you need special accommodations due to a disability, please contact Susan Monahan, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, Rm. 5231, Silver Spring, MD 20993–0002, 301–796–5661, email: *Susan.Monahan@fda.hhs.gov,* no later than April 9, 2018.

Streaming Webcast of the Public Workshop: This public workshop will also be webcast. The webcast link will be available on the registration web page after April 9, 2018. Organizations are requested to register all participants, but to view using one connection per location.

If you have never attended a Connect Pro event before, test your connection at *https://collaboration.fda.gov/common/help/en/support/meeting_test.htm.* To get a quick overview of the Connect Pro program, visit *https://www.adobe.com/go/connectpro_overview.* FDA has verified the website addresses in this document, as of the date this document publishes in the **Federal Register**, but websites are subject to change over time.

Transcripts: Please be advised that as soon as a transcript of the public workshop is available, it will be accessible at *https://www.regulations.gov.* It may be viewed at the Dockets Management Staff (see **ADDRESSES**). A link to the transcript will also be available approximately 45 days after the public workshop on the internet at *https://www.fda.gov/MedicalDevices/NewsEvents/WorkshopsConferences/default.htm.* (Select this public workshop from the posted events list.)

Dated: March 2, 2018.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2018–04629 Filed 3–7–18; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Information Technology Advisory Committee 2018 Schedule

AGENCY: Office of the National Coordinator for Health Information Technology (ONC), HHS.

ACTION: Notice of the Health Information Technology Advisory Committee 2018 schedule.

SUMMARY: This notice fulfills obligations under section 3002 of the Public Health Service Act (PHSA), as amended by the 21st Century Cures Act. Section 3002(b)(5) of the PHSA, as amended, mandates that the Health Information Technology Advisory Committee shall develop a schedule for the assessment of policy recommendations and the Secretary shall publish such schedule in the **Federal Register**.

FOR FURTHER INFORMATION CONTACT: Lauren Richie Designated Federal Officer, at *Lauren.Richie@hhs.gov.*

SUPPLEMENTARY INFORMATION: Section 3002 of the Public Health Service Act (PHSA), as amended by the 21st Century Cures Act (Pub. L. 114–255), establishes the Health Information Technology Advisory Committee (HITAC). The HITAC will be governed by the provisions of the Federal Advisory Committee Act (Pub. L. 92–463), as amended (5 U.S.C. App.), which sets forth standards for the formation and use of federal advisory committees. The HITAC, among other things, shall identify priorities for standards adoption and make recommendations to the National Coordinator for Health Information Technology (National Coordinator) on a policy framework to advance an interoperable health information technology infrastructure.

Health Information Technology Advisory Committee Schedule

Section 3002(b)(5) of the PHSA, as amended, provides that the HITAC shall develop a schedule for the assessment of policy recommendations developed by the HITAC and publish the schedule in the **Federal Register**. This schedule addresses the assessment of recommendations outlined in the policy framework recommended by the HITAC to the National Coordinator.

Accordingly, the schedule for the HITAC's assessment of policy recommendations is as follows:

1. Within 90 days of a charge by the National Coordinator for recommendations on a matter, identify the best mechanism to organize itself to develop recommendations, and at a minimum, will:

a. Develop an assessment of what policies, standards, implementation specifications, and certification criteria are currently available to be considered as part of the request;

b. Consider where gaps exist and identify potential organizations that have the capability to address those gaps (*i.e.,* no policy or standard is available or harmonization is required because more than one standard exists) related to the request; and

c. Create a timeline, which may also account for the National Institute of Standards and Technology (NIST) testing, where appropriate, and include dates when the HITAC is expected to issue the recommendation to the National Coordinator.

d. Include an opportunity for public comment during the consideration by the HITAC of the request by the National Coordinator for recommendations on a matter.

2. In responding to the National Coordinator:

a. Approve a timeline to deliver recommendations to the National Coordinator; and

b. Establish a task force to conduct analysis and solicit input, where appropriate, and develop draft recommendations to be considered by the full committee in a timely manner.

3. In collaboration with NIST, annually and through the use of public input, review and publish priorities for the use of health information technology, standards, and implementation specifications to support those priorities.

4. Recommend to the National Coordinator for purposes of adoption under section 3004, standards, implementation specifications, and certification criteria and an order of priority for the development, harmonization, and recognition of such standards, specifications, and certification criteria. Such recommendations shall include recommended standards, architectures, and software schemes for access to electronic individually identifiable health information across disparate systems including user vetting, authentication, privilege management, and access control.

The topics in which the HITAC is expected to address in FY2018 include, but may not be limited to the target areas as defined in section 3002 of the PHSA, as amended by the 21st Century Cures Act (Pub. L. 114–255), and they include:

1. Achieving a health information technology infrastructure that allows for the electronic access, exchange, and use of health information ;

2. The promotion and protection of privacy and security of health information in health information technology;

3. The facilitation of secure access by an individual to such individual's protected health information; and

4. Any other target area that the HITAC identifies as an appropriate target area to be considered. [42USC § 300jj (b)(2)(B)]

Notice of this meeting is given under section 3002(b)(5) of the PHSA, as amended.

Dated: March 1, 2018.

Lauren Richie,

Branch Chief, Office of Policy, Office of the National Coordinator for Health Information Technology.

[FR Doc. 2018–04543 Filed 3–7–18; 8:45 am]

BILLING CODE 4150–45–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Citizenship and Immigration Services

[OMB Control Number 1615–0122]

Agency Information Collection Activities; Revision of a Currently Approved Collection: Identity, Credential, and Access Management (ICAM)

AGENCY: U.S. Citizenship and Immigration Services, Department of Homeland Security.

ACTION: 60-Day notice.

SUMMARY: The Department of Homeland Security (DHS), U.S. Citizenship and Immigration (USCIS) invites the general public and other Federal agencies to comment upon this proposed revision of a currently approved collection of information or new collection of information. In accordance with the Paperwork Reduction Act (PRA) of 1995, the information collection notice is published in the **Federal Register** to obtain comments regarding the nature of the information collection, the categories of respondents, the estimated burden (*i.e.* the time, effort, and resources used by the respondents to respond), the estimated cost to the respondent, and the actual information collection instruments.

DATES: Comments are encouraged and will be accepted for 60 days until May 7, 2018.

ADDRESSES: All submissions received must include the OMB Control Number 1615–0122 in the body of the letter, the agency name and Docket ID USCIS–2011–0015. To avoid duplicate submissions, please use only *one* of the following methods to submit comments:

(1) *Online.* Submit comments via the Federal eRulemaking Portal website at *http://www.regulations.gov* under e-Docket ID number USCIS–2011–0015;

(2) *Mail.* Submit written comments to DHS, USCIS, Office of Policy and Strategy, Chief, Regulatory Coordination Division, 20 Massachusetts Avenue NW, Washington, DC 20529–2140.

FOR FURTHER INFORMATION CONTACT: USCIS, Office of Policy and Strategy, Regulatory Coordination Division, Samantha Deshommes, Chief, 20 Massachusetts Avenue NW, Washington, DC 20529–2140, telephone number 202–272–8377 (This is not a toll-free number. Comments are not accepted via telephone message). Please note contact information provided here is solely for questions regarding this notice. It is not for individual case status inquiries. Applicants seeking information about the status of their individual cases can check Case Status Online, available at the USCIS website at *http://www.uscis.gov,* or call the USCIS National Customer Service Center at 800–375–5283 (TTY 800–767–1833).

SUPPLEMENTARY INFORMATION:

Comments

You may access the information collection instrument with instructions, or additional information by visiting the Federal eRulemaking Portal site at: *http://www.regulations.gov* and enter USCIS–2011–0015 in the search box. Regardless of the method used for submitting comments or material, all submissions will be posted, without change, to the Federal eRulemaking Portal at *http://www.regulations.gov,* and will include any personal information you provide. Therefore, submitting this information makes it public. You may wish to consider limiting the amount of personal information that you provide in any voluntary submission you make to DHS. DHS may withhold information provided in comments from public viewing that it determines may impact the privacy of an individual or is offensive. For additional information, please read the Privacy Act notice that is available via the link in the footer of *http://www.regulations.gov.*

Written comments and suggestions from the public and affected agencies should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses.

Overview of This Information Collection

(1) *Type of Information Collection:* Revision of a Currently Approved Collection.

(2) *Title of the Form/Collection:* Identity, Credential, and Access Management (ICAM).

(3) *Agency form number, if any, and the applicable component of the DHS sponsoring the collection:* ICAM; USCIS.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract: Primary:* Individuals or households. In order to interact with USCIS electronic systems accessible through the USCIS ICAM portal, a first time user must establish an account. The account creation process requires the user to submit a valid email address; create a password; select their preference for receiving a one-time password (via email address, mobile phone, or two-factor authentication application on a mobile device); select five password reset questions and responses; and indicate the account type they want to set up (customer or legal representative). The account creation and the account login processes both require the user to receive and submit a one-time password. The one-time password can be provided either as an email to an email address or to a mobile phone via text message. The customer also has the option of receiving a one-time password readable by a two-factor authentication application on a mobile device. If the authentication application option is selected, the customer can either scan a QR code or enter a text code.

USCIS ICAM currently grants access to myUSCIS and the USCIS information collections available for e-filing.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* The estimated total number of respondents for the information collection ICAM is 1,772,600 and the estimated hour burden per response is 0.167 hours.

(6) *An estimate of the total public burden (in hours) associated with the collection:* The total estimated annual hour burden associated with this collection is 296,024 hours.

(7) *An estimate of the total public burden (in cost) associated with the collection:* The estimated total annual