delivering behavioral health resources and TA to support those implementing CDSME programs in the aging and disability network. LECOM has already established relationships with CDSME grantees with a behavioral health focus, which will allow them to seamlessly expand the TA and training they can offer to ensure CDSME grantees' success in meeting their goals and deliverables.

Additionally, the grantee has worked diligently to develop relationships with stakeholders through outreach and conference presentations, and to ensure that an inclusive range of partners are in place, engaged in the work, and committed to the success of CDSME.

LECOM is successfully meeting all programmatic goals under the current *Expanding Outreach and Professional Training to Engage Older Adults with Behavioral Health Conditions in Evidence-Based Health Promotion Programs* cooperative agreement.

Authority: Older Americans Act, Title IV; Patient Protection and Affordable Care Act, 42 U.S.C. 300U–11 (Prevention and Public Health Fund).

Dated: September 9, 2026.

Richard Nicholls,

Deputy Administrator and Chief of Staff of the Administration for Community Living.

[FR Doc. 2026–18729 Filed 9–11–26; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2026–N–2740]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Customer/Partner Customer Service Satisfaction Surveys

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: Submit written comments (including recommendations) on the collection of information by October 14, 2026.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to *https://www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under Review—Open for Public Comments'' or by using the search function. The OMB control number for this information collection is 0910–0360. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Kelly Covington, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 240–402–5661, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Customer/Partner Customer Service Satisfaction Surveys

OMB Control Number 0910–0360—Extension

Under section 1003 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 393), FDA is authorized to conduct research and public information programs about regulated products and responsibilities of the Agency. Executive Order 12862, entitled ''Setting Customer Service Standard,'' directs Federal Agencies that ''provide significant services directly to the public'' to ''survey customers to determine the kind and quality of services they want and their level of satisfaction with existing services.'' FDA is seeking to extend OMB approval to conduct customer service satisfaction surveys to implement Executive Order 12862. Participation in the surveys is voluntary. This request covers customer/partner (including State and local governments) service satisfaction surveys of regulated entities, such as food processors; cosmetic, drug, biologic, and medical device manufacturers; animal drugs, animal food and feed; tobacco products; and consumers and health professionals.

FDA will use the information from these surveys to identify strengths and weaknesses in service to customers/partners and to make improvements. The surveys will measure timeliness, appropriateness, clarity, and accuracy of information, courtesy, and problem resolution in the context of individual programs.

FDA estimates conducting approximately 20 customer/partner service satisfaction surveys per year, each requiring an average of 25 minutes for review and completion. We estimate respondents to these surveys to be between 100 and 20,000 customers/partners. Some of these surveys will be repeats of earlier surveys for purposes of monitoring customer/partner service and developing long-term data. Respondents to this collection of information cover a broad range of stakeholders who have experience with certain products regulated by or services provided by FDA.

In the **Federal Register** of April 3, 2026 (91 FR 16951), FDA published a 60-day notice requesting public comment on the proposed collection of information. One comment was received; however, it did not respond to the questions posed in § 1320.8(d).

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN[1]

Activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Mail, telephone, web-based survey	85,000	1	85,000	.42 (25 minutes)	35,700

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

Based on a review of the information collection since our last request for OMB approval, we have made no adjustments to our burden estimate.

Grace R. Graham,

Deputy Commissioner for Policy, Legislation, and International Affairs.

[FR Doc. 2026–18691 Filed 9–11–26; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the National Coordinator for Health Information Technology; Health Information Technology Advisory Committee; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the Health Information Technology Advisory Committee (HITAC), originally scheduled for September 24, 2026, from 10:00 a.m. to 3:00 p.m. ET, which was published in the **Federal Register** on January 21, 2026 (91 FR 2542).

The meeting will be held on October 8, 2026, from 10:00 a.m. to 3:00 p.m. ET. The meeting is open to the public and will be held virtually. For web conference instructions and the most up-to-date meeting information, please visit the HITAC calendar at *www.healthit.gov/topic/federal-advisory-committees/hitac-calendar.*

Dated: September 9, 2026.

Tara Porter,

Designated Federal Officer, HITAC.

[FR Doc. 2026–18682 Filed 9–11–26; 8:45 am]

BILLING CODE 4150–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; NIH Electronic Application System for NIH Certificates of Confidentiality (CoC)

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the Office of Science Policy (OSP), in the Office of the Director (OD), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

DATES: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, Email your comment or request, including your address to: *NIH-CoC-Coordinator@mail.nih.gov* or contact: Dr. Adam Berger, Division of Clinical and Healthcare Research Policy, Office of Science Policy, OD, NIH, 6705 Rockledge Drive, Suite 630, Bethesda, MD 20892, or call non-toll-free number (301) 496–9838. Formal requests for additional plans and instruments must be requested in writing.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: written comments and/or suggestions from the public and affected agencies are invited to address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) 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) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Proposed Collection Title: Electronic Application for NIH Certificates of Confidentiality (CoC E-application System), 0925–0689, REVISION, exp., date 08/31/2028. Office of Science Policy (OSP), National Institutes of Health (NIH).

Need and Use of Information Collection: The purpose of this electronic system is to submit and process requests for NIH to issue discretionary Certificates of Confidentiality (CoC) to research organizations that request a CoC from NIH. As described in the authorizing legislation (Section 301(d) of the Public Health Service Act, 42 U.S.C. 241(d)), CoCs are issued by specific agencies of the Department of Health and Human Services (DHHS), including NIH, to authorize researchers to protect the privacy of human research subjects by prohibiting them from releasing names and identifying characteristics of research participants to anyone not connected with the research, except in limited circumstances specified in statute. At NIH, the issuance of CoCs has been delegated to the NIH Office of Science Policy (OSP) in the NIH Office of the Director. The NIH has been using an online CoC system to review requests and issue CoCs since 2015. The current CoC request form includes 27 questions to collect information from research organizations and six Institutional Assurance statements to be affirmed by the Institutional Official. The information provided is used to determine eligibility for a CoC and to issue the CoC to the requesting organization. Eligible requesting organizations that provide legally binding affirmations that they will abide by the terms of the CoC are issued a Certificate of Confidentiality. This system has increased efficiency and reduced burden for both requesters and NIH staff who currently process these requests. NIH received 518 requests for CoCs from January 202*5* through December 202*5* and expects to receive approximately the same number of requests in subsequent years.

OMB approval is requested for three years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 777.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
Individuals	518	1	90/60	777
Total		518		777