notice requesting public comment on the proposed collection of information. No comments were received.

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

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

21 CFR section	Number of respondents	Number of responses per respondent	Total number of annual responses	Average burden per response in hours	Total hours
1150.5(a), (b)(1) and (2), and Form FDA 3852; Identification and removal information (monthly)	820	12	9,840	3	29,520
1150.5(b)(3); Certified copies (monthly)	820	12	9,840	1	9,840
Voluntary premium cigar data submission (monthly)	50	12	600	1.5	900
1150.13; Payment of user fee assessment (quarterly)	319	4	1,276	1	1,276
1150.15(a); Submission of user fee dispute (at discretion of respondent)	2	1	2	10	20
1150.15(d); Submission of request for further review of dispute of user fee (at discretion of respondent)	1	1	1	5	5
Total			21,559		41,561

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

We have revised our burden estimates to this information collection request. 21 CFR 1150.5 is reflecting an increase in 120 respondents from 700 to 820. FDA requires the use of Form FDA 3852 to capture the monthly identification and removal information specified under § 1150.5(b)(1) and (b)(2). The form also directs manufacturers and importers to attach supporting documentation required by § 1150.5(b)(3). FDA considered the number of active Alcohol and Tobacco Tax and Trade Bureau (TTB) permits (based on TTB data) in FY23 for domestic manufacturers and importers of tobacco products subject to tobacco user fees.

Voluntary premium cigar data submission (monthly) is reflecting a reduction in 50 respondents from 100 to 50 and a reduction in average burden per response from 2.5 to 1.5 hours. FDA updated this data based on reasonable estimates of the burden of voluntary submissions in FY24. There may be some fluctuations in this number.

Section 1150.13 (21 CFR 1150.13) is reflecting a reduction in 57 respondents from 376 to 319. FDA considered the number of user fee assessments issued to domestic manufacturers and importers of tobacco products subject to tobacco user fees on average each quarter for FY23. Note, entities may have more than one TTB permit, however, tobacco user fee assessments are aggregated based on Employer Identification Number and not TTB permit number. Therefore, we expect the number of respondents to be lower for § 1150.13.

21 CFR 1150.15(a) is reflecting a reduction in 3 respondents from 5 to 2, and 21 CFR 1150.15(d) is reflecting a reduction in 2 respondents from 3 to 1 and a reduction in average burden per response from 10 to 5 hours. FDA considered the historical submission of tobacco user fee disputes and requests for additional Agency review.

The cumulative changes to the estimated burden for this information collection reflects an overall increase of 3,377 burden hours and a corresponding increase of 2,047 responses.

Dated: July 23, 2025.

Grace R. Graham,

Deputy Commissioner for Policy, Legislation, and International Affairs.

[FR Doc. 2025–14221 Filed 7–28–25; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Deafness and Other Communication Disorders; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the National Deafness and Other Communication Disorders Advisory Council, September 04, 2025, 01:00 p.m. to September 05, 2025, 01:05 p.m., National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852 which was published in the **Federal Register** on July 21, 2025, 90 FR 34281.

The meeting time has been changed from 9:00 a.m. to 9:40 a.m. to 9:30 a.m. to 10:00 a.m. The meeting is partially Closed to the public.

Dated: July 24, 2025.

Bruce A. George,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2025–14239 Filed 7–28–25; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Aging; Notice of Closed Meeting

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the Board of Scientific Counselors, NIA.

The meeting will be closed to the public as indicated below in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended for the review, discussion, and evaluation of individual grant applications conducted by the National Institute On Aging, including consideration of personnel qualifications and performance, and the competence of individual investigators, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Board of Scientific Counselors, NIA.

Date: May 19–21, 2026.

Time: May 19, 2026, 8:00 a.m. to 4:30 p.m.

Agenda: Executive Session; Board Business; Review of Labs and PI's; Adjournment.

Address: National Institute of Aging, 251 Bayview Blvd., Baltimore, MD 21224 (Virtual Meeting).

Time: May 20, 2026, 8:00 a.m. to 4:30 p.m.