TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN—Continued

Activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours	Total capital costs
Supplement to approved plan	10	1	10	30	300	120
Total					540	168

FDA estimates a total of 4 respondents will submit a new original warning plan and take 60 hours to complete a rotational warning plan for a total of 240 burden hours. In addition, 10 respondents will submit a supplement to an approved warning plan at 30 hours per response for a total of 300 hours. The total burden for this collection is estimated to be 540 hours.

Capital costs are based on 14 respondents mailing in their submission at a postage rate of $12 for a 5-pound parcel (business parcel post mail delivered from the furthest delivery zone). Therefore, FDA estimates that the total postage cost for mailing the rotational warning plans FDA to be $168.

We have adjusted our burden estimate, which has resulted in a decrease of 5,460 hours and 86 respondents to the currently approved burden. We received a total number of 44 original smokeless warning plans, and a total of 17 supplements. After receiving the initial influx of original warnings plans, FDA does not expect to receive as many original warning plans annually. We expect that a few supplements will continue to be received as new products are marketed or as warning plans are revised. We anticipate a total number of 10 supplements submitted annually and 4 original smokeless warning plans.

Dated: June 7, 2019.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2019–12472 Filed 6–12–19; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Assistant Secretary for Health

Meeting of the Pain Management Best Practices Inter-Agency Task Force; Correction

AGENCY: Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services.

ACTION: Notice; correction.

SUMMARY: The Office of the Assistant Secretary for Health published a document in the **Federal Register** of June 3, 2019, announcing the Pain Management Best Practices Inter-Agency Task Force's virtual public meeting. This document is announcing a change in the meeting date.

FOR FURTHER INFORMATION CONTACT: Ms. Alicia Richmond Scott, 240–453–2816; *paintaskforce@hhs.gov.*

SUPPLEMENTARY INFORMATION:

Correction

In the **Federal Register** of June 3, 2019, in FR Doc. 2019–11473, on page 25548, in the first column, correct the **DATES** caption to read:

DATES: The Task Force meeting will be held on Wednesday, June 26, 2019 from 5:00 p.m. to 6:30 p.m. Eastern Time (ET). The agenda will be posted on the Task Force website at *https://www.hhs.gov/ash/advisory-committees/pain/index.html.*

Dated: June 6, 2019.

Vanila M. Singh,

Chief Medical Officer, Chair, Pain Management Best Practices Inter-Agency Task Force, Office of the Assistant Secretary for Health.

[FR Doc. 2019–12482 Filed 6–12–19; 8:45 am]

BILLING CODE 4150–28–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Center for Scientific Review Special Emphasis Panel; PAR17–094: Maximizing Investigators' Research Award (R35).

Date: July 9, 2019.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Bethesda North Marriott Hotel Conference Center, Montgomery County Conference Center Facility, 5701 Marinelli Road, North Bethesda, MD 20852.

Contact Person: Baishali Maskeri, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2022, Bethesda, MD 20892, 301–827–2864, *maskerib@mail.nih.gov*

Name of Committee: Center for Scientific Review Special Emphasis Panel; RFA Panel: Healthy Brain and Child Development Study (Healthy BCD).

Date: July 11, 2019.

Time: 8:30 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Residence Inn Capital View, 2850 South Potomac Avenue, Arlington, VA 22202

Contact Person: Heidi B. Friedman, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 1012A, MSC 7770, Bethesda, MD 20892, 301–379–5632, *hfriedman@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; PAR Panel: Chronic Disease and the Reduction of Health Disparities.

Date: July 12, 2019.

Time: 10:30 a.m. to 3:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Karen Nieves Lugo, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, Bethesda, MD 20892, *karen.nieveslugo@nih.gov.*

Name of Committee: AIDS and Related Research Integrated Review Group; Population and Public Health Approaches to HIV/AIDS Study Section

Date: July 15–16, 2019.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications,

Place: Hotel Nikko San Francisco, 222 Mason Street, San Francisco, CA 94102

Contact Person: Jose H. Guerrier, Ph.D., Scientific Review Officer, Center for