

preparation and review process for 510(k).

Under § 807.90, submitters may request information on their 510(k) review status 90 days after the initial login date of the 510(k). Thereafter, the submitter may request status reports every 30 days following the initial status request. To obtain a 510(k) status report,

the submitter should complete the status request form, Form FDA 3541, and fax it to the Center for Devices and Radiological Health office identified on the form.

The most likely respondents to this information collection will be specification developers and medical device manufacturers.

In the **Federal Register** of July 23, 2013 (78 FR 44130), FDA published a 60-day 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

Activity/21 CFR part/Section/Form No.	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
510(k) submission (807 subpart E)	3,900	1	3,900	79	308,100
Summary cover sheet (807.87) and FDA 3514	1,956	1	1,956	0.5 (30 minutes)	978
Status request (807.90(a)(3)) and FDA 3541	218	1	218	0.25 (15 minutes)	55
Standards (807.87(d) and (f)); FDA 3654	2,700	1	2,700	10	27,000
510(k) summary and statement (807.92 and 807.93).	225	10	2,250	10	22,500
Total					358,633

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

Dated: October 22, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

CDC/HRSA Advisory Committee on HIV, Viral Hepatitis and STD Prevention and Treatment; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), notice is hereby given of the following virtual committee meeting.

Name: CDC/HRSA Advisory Committee on HIV, Viral Hepatitis and STD Prevention and Treatment

Dates and Times: November 13, 2013, 10:00 a.m.–4:30 p.m. November 14, 2013, 10:00 a.m.–12:30 p.m.

Place: This meeting is accessible via audio conference call and Adobe Connect Pro.

Status: This meeting is open to the public. The available lines will accommodate approximately 300 people.

Purpose: This Committee is charged with advising the Director, CDC, and the Administrator, HRSA, regarding activities related to prevention and control of HIV/AIDS, Viral Hepatitis and other STDs, the support of health care services to persons living with HIV/AIDS, and education of health

professionals and the public about HIV/AIDS, Viral Hepatitis, and other STDs.

Agenda: Agenda items include: (1) Affordable Care Act (ACA) Updates; (2) Clinical Workforce Issues; and (3) CHAC Workgroup Updates. Agenda items are subject to change as priorities dictate. To register for this event, please go to the following link: <https://hrsa.connectsolutions.com/advisorychac/event/registration.html>.

After you register, you should receive a confirmation email with a URL link for access to this event. You will also need to enter your Adobe Connect user name and password. If you've never used Adobe Connect, get a quick overview: http://www.adobe.com/go/connectpro_overview.

The public can join the meeting by:

1. (Audio Portion) Calling the Toll free Phone Number 1-888-324-9564 and providing the Participant Pass Code 4805129, and

2. (Visual Portion) Connecting to the Advisory Committee Adobe Connect Pro Meeting using the following URL: <https://hrsa.connectsolutions.com/advisorychac/event/login.html> (copy and paste the link into your browser if it does not work directly).

Participants should call and connect 15 minutes prior to the meeting in order for logistics to be set up. Call (301) 443-9684 or send an email to sgordon@hrsa.gov if you are having trouble registering for this meeting; or call (301) 443-2843 or send an email to lfores@hrsa.gov if you are having trouble connecting to the meeting site.

Public Comment: Persons who desire to make an oral statement may request it at the time of the public comment

period. Public participation and ability to comment will be limited to space and time as it permits.

FOR FURTHER INFORMATION CONTACT: Shelley B Gordon, Health Resources and Services Administration, HIV/AIDS Bureau, 5600 Fishers Lane, Rockville, Maryland 20857, telephone at (301) 443-9684.

Dated: October 21, 2013.

Bahar Niakan,

Director, Division of Policy and Information Coordination.

[FR Doc. 2013-25231 Filed 10-25-13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Environmental Health Sciences; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), 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.