Estimated Total Annual Burden Hours: 9,256.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Fax: 202–395–7285, email: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2012–15389 Filed 6–22–12; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0001]

Oncologic Drugs Advisory Committee; Amendment of Notice

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

The Food and Drug Administration (FDA) is announcing an amendment to the notice of meeting of the Oncologic Drugs Advisory Committee. This meeting was announced in the **Federal Register** of May 31, 2012 (77 FR 32125–32126). The amendment is being made to reflect a change in the *Date and Time,* and *Procedure* portions of the document. The *Date and Time* of the meeting will change to July 24, 2012, from 8 a.m. to 6 p.m. The *Procedure* portion of the document has changed to reflect an updated public participation time of 10:30 a.m. to 11 a.m., and 4:30 p.m. to 5 p.m. There are no other changes.

FOR FURTHER INFORMATION CONTACT: Caleb Briggs, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993–0002, 301–796–9001, FAX: 301–847–8533, *ODAC@fda.hhs.gov,* or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), and follow the prompts to the desired center or product area. Please call the Information Line for up-to-date information on this meeting.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of May 31, 2012, FDA announced that a meeting of the Oncologic Drugs Advisory Committee would be held on July 24, 2012. On page 32125, in the third column, the *Date and Time* portion of the document is changed to read as follows:

Date and Time: The meeting will be held on July 24, 2012 from 8 a.m. to 6 p.m.

On page 32126, in the first column, the third sentence in the *Procedure* portion of the document is changed to read as follows:

Procedure: Oral presentations from the public will be scheduled between approximately 10:30 a.m. to 11 a.m., and 4:30 p.m. to 5 p.m.

This notice is issued under the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR part 14, relating to the advisory committees.

Dated: June 20, 2012.

Jill Hartzler Warner,

Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2012–15393 Filed 6–22–12; 8:45 am]

BILLING CODE 4160–01–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 (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.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Cardiovascular Sciences.

Date: July 17–18, 2012.

Time: 9:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Maqsood A Wani, DVM, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2114, MSC 7814, Bethesda, MD 20892, 301–435–2270, *wanimaqs@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Fellowship: Endocrinology, Metabolism, Nutrition and Reproductive Sciences.

Date: July 18, 2012.

Time: 11:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Dianne Camp, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, ROOM # 6164, Bethesda, MD 20892, 301–435–1044, *campdm@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: June 19, 2012.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2012–15473 Filed 6–22–12; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute; Notice of Closed Meeting

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 meeting.

The meeting 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: National Heart, Lung, and Blood Institute Special Emphasis Panel; Phase II Trials in Lung Disease.

Date: July 12, 2012.

Time: 8:30 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Renaissance Washington DC, Dupont Circle, 1143 New Hampshire Avenue NW., Washington, DC 20037.

Contact Person: Stephanie L Constant, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7189, Bethesda, MD 20892, 301–443–8784, *constantsl@nhlbi.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: June 14, 2012.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2012–15472 Filed 6–22–12; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish a summary of information collection requests under OMB review, in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these documents, call the SAMHSA Reports Clearance Officer on (240) 276–1243.

Project: Enhancing Substance Abuse Treatment Services To Address Hepatitis Infection Among Intravenous Drug Users Hepatitis Testing and Vaccine Tracking Form (OMB No. 0930–0300)—Reinstatement and Extension

The Substance Abuse and Mental Health Services Administration's (SAMHSA) Center Substance Abuse Treatment (CSAT), is responsible for the Hepatitis Testing and Vaccine Tracking Form for the prevention of Viral Hepatitis in patients in designated Opioid Treatment Programs (OTPs). There are no changes to the form or added burden.

This form allows SAMHSA/CSAT to collect essential Clinical information that will be used for quality assurance, quality performance and product monitoring on approximately 264 Rapid Hepatitis C Test kits and 10,628 doses of hepatitis vaccine (Twinrix, HAV, or HBV). The above kits and vaccines will be provided to designated OTPs serving the minority population in their communities. The information collected on the Form solicits and reflect the following information:

- Demographics (age, gender, ethnicity) of designated OTP site
- History (Screening) of Hepatitis C exposure
- Results of Rapid Hepatitis C Testing (Kit) and Follow-up information
- Service Provided (type of vaccine given) Divalent vaccine (Twinrix-combination HAV and HBV) or Monovalent vaccine (HAV and/or HBV)
- Substance Abuse Treatment Outcomes (Information regarding the beginning, continuing or completion of vaccination series)
- Type of Referral Services Indicated (i.e., Gastroenterology, TB; Mental Health, Counseling, Reproductive/ Prenatal, etc.)

This program is authorized under Section 509 of the Public Health Service (PHS) Act [42 U.S.C. 290bb–2].

The form increases the screening and reporting of viral hepatitis in high risk minorities in OTPs. The information collected allows SAMHSA to address the increased morbidity and mortality of hepatitis in minorities being treated for drug addiction.

The SAMHSA/CSAT Hepatitis Testing and Vaccine Tracking Form supports quality of care, provide minimum but adequate clinical and product monitoring, and provide appropriate safeguards against fraud, waste and abuse of Federal funds.

The table below reflects the annualized hourly burden.

Number of respondents screened	Responses/ respondent	Burden hours	Total burden hours
50,000	1	0.05	2,500

Written comments and recommendations concerning the proposed information collection should be sent by July 25, 2012 to the SAMHSA Desk Officer at the Office of Information and Regulatory Affairs, Office of Management and Budget (OMB). To ensure timely receipt of comments, and to avoid potential delays in OMB's receipt and processing of mail sent through the U.S. Postal Service, commenters are encouraged to submit their comments to OMB via email to: *OIRA_Submission@omb.eop.gov.* Although commenters are encouraged to send their comments via email, commenters may also fax their comments to: 202–395–7285. Commenters may also mail them to: Office of Management and Budget, Office of Information and Regulatory Affairs, New Executive Office Building, Room 10102, Washington, DC 20503.

Summer King,

Statistician.

[FR Doc. 2012–15414 Filed 6–22–12; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HOMELAND SECURITY

[Docket No. DHS–2012–0026]

Committee Name: Homeland Security Academic Advisory Council

AGENCY: Department of Homeland Security.

ACTION: Committee Management; Notice of Federal Advisory Committee Meeting.

SUMMARY: The Homeland Security Academic Advisory Council (HSAAC) will meet on July 10, 2012 in Washington, DC. The meeting will be open to the public.