R. Record Source Categories

Information in this system is obtained from many sources, including the following: (1) Directly from the respondent or complainant or his/her representative; (2) derived from materials supplied by the respondent or complainant or his/her representative; (3) from information supplied by the institutions, witnesses, scientific publications, and other nongovernmental sources; (4) from observation and analysis made by FDA and ORI staff and scientific experts; (5) from departmental and other Federal, State, and local government records; (6) from hearings and other administrative proceedings; and (7) from any other relevant source.

S. Records Exempted From Certain Provisions of the Privacy Act

FDA records related to research misconduct proceedings will be exempt from the Privacy Act requirements pertaining to providing an accounting of disclosures, access and amendment, notification, and Agency procedures and rules under sections 552a(k)(2) and (k)(5) of the Privacy Act.

Elsewhere in this issue of the **Federal Register**, FDA is publishing a notice of proposed rulemaking and direct final rule to apply these exemptions to records in this system related to ongoing investigations or that would reveal a confidential source. These exemptions are necessary to safeguard the integrity of the research misconduct proceedings and to ensure that FDA's efforts to obtain accurate and objective information will not be hindered. In the course of investigations of allegations of research misconduct, it is often necessary to give an express promise to withhold the identity of an individual who has provided relevant information. Sources of information necessary to complete an effective investigation may be reluctant to provide sensitive information unless they can be assured that their identities will not be revealed. The proposed exemptions will ensure that the records related to ongoing investigations will not be disclosed inappropriately and that the identities of confidential sources will be protected.

The notice of proposed rulemaking and direct final rule provide additional detail regarding the bases for these exemptions.

II. Comments

FDA invites comments on all parts of the systems notice. Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) either electronic or written comments regarding this document. It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Dated: June 12, 2012.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2012–20888 Filed 8–27–12; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Commission on Childhood Vaccines, 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 meeting:

Name: Advisory Commission on Childhood Vaccines (ACCV).

Date and Time: September 06, 2012, 1:00 p.m. to 5:15 p.m. EDT.

Place: Parklawn Building (and via audio conference call), Conference Room 10–65, 5600 Fishers Lane, Rockville, MD 20857.

The ACCV will meet on Thursday, September 06 from 1:00 p.m. to 5:15 p.m. (EDT). The public can join the meeting via audio conference call by dialing 1–800–369–3104 on September 06, and providing the following information:

Leader's Name: Dr. Geoffrey Evans
Password: ACCV

Agenda: The agenda items for the September meeting will include, but are not limited to: Updates from the Division of Vaccine Injury Compensation (DVIC), Department of Justice (DOJ), National Vaccine Program Office (NVPO), Immunization Safety Office (Centers for Disease Control and Prevention), National Institute of Allergy and Infectious Diseases (National Institutes of Health) and Center for Biologics Evaluation and Research (Food and Drug Administration). A draft agenda and additional meeting materials will be posted on the ACCV web site (*http://www.hrsa.gov/vaccinecompensation/accv.htm*) prior to the meeting. Agenda items are subject to change as priorities dictate.

Public Comment: Persons interested in attending the meeting in person or providing an oral presentation should submit a written request, along with a copy of their presentation to: Annie Herzog, DVIC, Healthcare Systems Bureau (HSB), Health Resources and Services Administration (HRSA), Room 11C–26, 5600 Fishers Lane, Rockville, Maryland 20857 or email: *aherzog@hrsa.gov*. Requests should contain the name, address, telephone number, email address, and any business or professional affiliation of the person desiring to make an oral presentation. Groups having similar interests are requested to combine their comments and present them through a single representative. The allocation of time may be adjusted to accommodate the level of expressed interest. DVIC will notify each presenter by email, mail, or telephone of their assigned presentation time. Persons who do not file an advance request for a presentation, but desire to make an oral statement, may announce it at the time of the public comment period. Public participation and ability to comment will be limited to space and time as available.

FOR FURTHER INFORMATION CONTACT: Anyone requiring information regarding the ACCV should contact Annie Herzog, DVIC, HSB, HRSA, Room 11C–26, 5600 Fishers Lane, Rockville, MD 20857; telephone (301) 443–6593, or email: *aherzog@hrsa.gov*.

Dated: August 22, 2012.

Bahar Niakan,

Director, Division of Policy and Information Coordination.

[FR Doc. 2012–21093 Filed 8–27–12; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; 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.