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 3, 2015, 9:00 a.m. to 4:30 p.m. EDT.

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

The ACCV will meet on Thursday, September 3, 2015, from 9:00 a.m. to 4:30 p.m. (EDT). The public can join the meeting by:

1. (In Person) Persons interested in attending the meeting in person are encouraged to submit a written notification to: Annie Herzog, DICP, Healthcare Systems Bureau (HSB), Health Resources and Services Administration (HRSA), Room 11C–26, 5600 Fishers Lane, Rockville, Maryland 20857 or email: aherzog@hrsa.gov. Since this meeting is going to be held in a federal government building, attendees will need to go through a security check to enter the building and participate in the meeting. Written notification is encouraged so a list of attendees can be provided to Annie Herzog to make entry through security quicker.

2. (Audio Conference) The conference phone number is 877–917–4913. When calling, provide the following information:

   Leaders Name: Dr. A. Melissa Houston. Password: ACCV.

3. (Visual Portion) Connect to the ACCV Adobe Connect Pro meeting using the following URL: https://hrsa.connectsolutions.com/accv/ (copy and paste the link into your browser if it does not work directly, and enter as a guest). Participants should call and connect 15 minutes prior to the meeting in order for logistics to be set up. If you have never attended an Adobe Connect meeting, please test your connection using the following URL: https://hrsa.connectsolutions.com/common/help/en/support/meeting_test.htm and get a quick overview by following URL: http://www.adobe.com/go/connectpro_overview. Call (301) 443–6634 or send an email to aherzog@hrsa.gov if you are having trouble connecting to the meeting site.

Agenda: The agenda items for the September 2015 meeting will include, but are not limited to: updates from ACCV Adult Immunization Workgroup, the Division of Injury Compensation Programs (DICP), 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 providing an oral presentation should submit a written request, along with a copy of their presentation to: Annie Herzog, DICP, 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. DICP 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 it permits.

For Further Information Contact: Anyone requiring information regarding the ACCV should contact Annie Herzog, DICP, HSB, HRSA, Room 11C–26, 5600 Fishers Lane, Rockville, Maryland 20857; telephone (301) 443–6593, or email: aherzog@hrsa.gov.

Jackie Painter, Director, Division of the Executive Secretariat.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: In compliance with Section 507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Health Resources and Services Administration (HRSA) has submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB). The OMB is required to approve or reject the ICR within 30 days, or, if it chooses not to act on the ICR, to inform HRSA of its decision. In compliance with Section 3506(c)(4)(A) of the Paperwork Reduction Act of 1995, this notice is published to provide an opportunity for public comments on the content and proposed burden of the OMB-approved information collection instrument.

Requests for a public hearing should be submitted to the contact person below. Written comments and听证会请求 should be submitted to: Joanne Rodriguez, Office of Public Health Programs, Health Resources and Services Administration, 5600 Fishers Lane, Rockville, MD 20857. Comments may also be submitted electronically by accessing the following Web site: http://www.regulations.gov. Instructions for submitting comments are available at this URL. The collection of information in 21 CFR part 812 have been approved under OMB control number 0910–0485; and the collections of information in 21 CFR part 814 have been approved under OMB control number 0910–0231; the collections of information in 21 CFR parts 801 and 809 have been approved under OMB control number 0910–0485; and the collections of information in the guidance document entitled “Informed Consent For In Vitro Diagnostic Device Studies Using Leftover Human Specimens That Are Not Individually Identifiable” have been approved under OMB control number 0910–0582.

V. Comments

Interested persons may submit either electronic comments regarding this document to http://www.regulations.gov or written comments to the Division of Dockets Management (see ADDRESSES). 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, and will be posted to the docket at http://www.regulations.gov.

Dated: August 10, 2015.

Leslie Kux, Associate Commissioner for Policy.