analysis, and data management issues; (3) perspectives in developing multiplex devices for donor screening; and (4) workshop summary and conclusions.

Transcripts: Please be advised that as soon as possible after a transcript of the public workshop is available, it will be accessible at: http://www.fda.gov/BiologicsBloodVaccines/NewsEvents/WorkshopsMeetingsConferences/TranscriptsMinutes/default.htm.

Transcripts of the public workshop may also be requested in writing from the Division of Freedom of Information (ELEM-1029), Food and Drug Administration, 12420 Parklawn Dr., Rockville, MD 20857.

Dated: March 11, 2013.

Leslie Kux,

Assistant Commissioner for Policy.
[FR Doc. 2013–05987 Filed 3–14–13; 8:45 am]
BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Secretary's Advisory Committee on Heritable Disorders in Newborns and Children; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463, codified at 5 U.S.C. App. 2), notice is hereby given of the following meeting:

Name: Secretary's Advisory Committee on Heritable Disorders in Newborns and Children (SACHDNC).

Date and Time: April 19, 2013, 9:30 a.m. to 3:00 p.m.

Place: Virtual via Webinar.

Status: The meeting is open to the public. Pre-registration is required. For more information on registration and webinar details, please visit the SACHDNC Web site: http://www.hrsa.gov/advisorycommittees/mchbadvisory/heritabledisorders.

Purpose: The Secretary's Advisory Committee on Heritable Disorders in Newborns and Children (SACHDNC), as authorized by Public Law 106-310, which added section 1111 of the Public Health Service Act, codified at 42 U.S.C. 300b-10, was established by Congress to advise the Secretary of the Department of Health and Human Services regarding the development of newborn screening activities, technologies, policies, guidelines, and programs for effectively reducing morbidity and mortality in newborns and children having, or at risk for, heritable disorders. The SACHDNC's recommendations regarding additional conditions/inherited disorders for screening that have been adopted by the Secretary are included in the Recommended Uniform Screening Panel (RUSP) that constitutes part of the comprehensive guidelines supported by the Health Resources and Services

Administration. Pursuant to section 2713 of the Public Health Service Act, codified at 42 U.S.C. 300gg–13, non-grandfathered health plans are required to cover screenings included in the comprehensive guidelines without charging a co-payment, co-insurance, or deductible for plan years (i.e., policy years) beginning on or after the date that is one year from the Secretary's adoption of the condition for screening. The SACHDNC also provides advice and recommendations concerning grants and projects authorized under section 1109 of the Public Health Service Act (42 U.S.C. 300b–8).

Agenda: The meeting will include: (1) A policy paper report on the impact of recommendations related to sickle cell trait testing; (2) a presentation on the Affordable Care Act and the impact on individuals with heritable disorders; (3) a presentation by the Agency for Healthcare Research and Quality regarding the processes behind the U.S. Preventive Services Task Force review process; and (4) project reports on screening for Tyrosinemia Type I and Point of Care Screening and Lessons Learned.

Proposed agenda items are subject to change as priorities dictate. The agenda, webinar information, Committee Roster, Charter, presentations, and meeting materials are located on the Advisory Committee's Web site at http://www.hrsa.gov/advisorycommittees/mchbadvisory/heritabledisorders.

Public Comments: Members of the public can submit written comments and/or register to present oral comments. All comments, whether oral or written, are part of the official Committee record and will be available for public inspection and copying. Individuals who wish to make public comments are required to register for the webinar and email Lisa Vasquez (lvasquez@hrsa.gov) by April 10, 2013. The public comment period is scheduled for the morning of April 19, 2013. Written comments should be emailed to Lisa Vasquez (lvasquez@hrsa.gov) by April 10, 2013.

Written comments should identify the individual's name, address, email, telephone number, professional or business affiliation, type of expertise (i.e., parent, researcher, clinician, public health, etc.) and the topic/subject matter of comment. To ensure that all individuals who have registered to make oral comments can be accommodated, the allocated time may be limited. Individuals who are associated with groups or have similar interests may be requested to combine their comments and present them through a single representative. No audiovisual presentations are permitted.

Contact Person: Anyone interested in obtaining other relevant information should contact the designated federal officer (DFO), Debi Sarkar, Maternal and Child Health Bureau, Health Resources and Services Administration, Room 18A–19, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857; telephone: 301–443–1080; email: dsarkar@hrsa.gov.

More information on the Advisory Committee is available at http:// www.hrsa.gov/advisorycommittees/ mchbadvisory/heritabledisorders. Dated: March 11, 2013.

Bahar Niakan,

Director, Division of Policy and Information Coordination.

[FR Doc. 2013–06042 Filed 3–14–13; 8:45 am] BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Statement of Organization, Functions and Delegations of Authority

This notice amends Part R of the Statement of Organization, Functions and Delegations of Authority of the Department of Health and Human Services (HHS), Health Resources and Services Administration (HRSA) (60 FR 56605, as amended November 6, 1995; as last amended at 78 FR 14311–14312, dated March 5, 2013).

This notice reflects organizational changes to the Health Resources and Services Administration. This notice updates the functional statement for the Bureau of Primary Health Care (RC). Specifically, this notice: (1) Establishes the Office of National Assistance and Special Populations (RCE); (2) abolishes the Office of Training and Technical Assistance Coordination (RCS) and the Office of Special Population Health (RCG); and (3) updates the functional statement for the Office of the Associate Administrator (RC), the Office of Administrative Management (RCM), the Office of Policy and Program Development (RCH), and the Office of Quality and Data (RCK).

Chapter RC—Bureau of Primary Health Care

Section RC-10, Organization

Delete in its entirety and replace with the following:

The Bureau of Primary Health Care (RC) is headed by the Associate Administrator, who reports directly to the Administrator, Health Resources and Services Administration. The Bureau of Primary Health Care includes the following components:

- Office of the Associate Administrator (RC);
- (2) Office of Administrative Management (RCM);
- (3) Office of Policy and Program Development (RCH);
 - (4) Office of Quality and Data (RCK);
- (5) Office of National Assistance and Special Populations (RCE);
 - (6) Northeast Division (RCU);
 - (7) Central Southeast Division (RCV);
 - (8) North Central Division (RCT); and