#### Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2014–21379 Filed 9–8–14; 8:45 am] BILLING CODE 4163–18–P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

## Centers for Disease Control and Prevention

## Georgia Tuberculosis Outbreak Among Homeless

**AGENCY:** Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

**ACTION:** Notice of award.

SUMMARY: The Centers for Disease Control and Prevention (CDC) located within the United States Department of Health and Human Services (HHS) announces a notice of award to the Georgia Department of Public Health, Tuberculosis (TB) Program. This award will be in the amount of \$419,095.00.

The purpose of this award is to halt the further spread of a drug-resistant strain of tuberculosis associated with multiple homeless shelters in Fulton County, Georgia. **DATES:** It is expected the notice of award will begin on or about September 3, 2014. The project period will be for one year.

FOR FURTHER INFORMATION CONTACT: Gail Burns-Grant, Division of Tuberculosis Elimination, Field Services and Evaluation Branch, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS E–10, Atlanta, GA 30333; phone: 404–639–5344; email: *GAB2@cdc.gov*.

**SUPPLEMENTARY INFORMATION:** Currently, the state of Georgia is experiencing a public health emergency in Fulton County where there has been extensive transmission of a drug-resistant strain of tuberculosis (TB) associated with multiple homeless shelters in the county. The Georgia Department of Public Health asked CDC to provide emergency funding for the immediate implementation of CDC recommendations provided as a result of a May 2014 outbreak investigation to prevent further transmission of this drug-resistant strain of tuberculosis and to prevent further deaths associated with this outbreak. Project number is CDC-RFA-PS14-1416.

Dated: September 4, 2014.

### Ron A. Otten,

Acting Deputy Associate Director for Science, Centers for Disease Control and Prevention. [FR Doc. 2014–21455 Filed 9–4–14; 4:15 pm]

BILLING CODE 4163-18-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

### **Food and Drug Administration**

[Docket No. FDA-2014-N-0001]

## Advisory Committee Renewals; Correction

AGENCY: Food and Drug Administration,

HHS.

**ACTION:** Notice: correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice entitled "Advisory Committee Renewals" that appeared in the Federal Register of August 25, 2014 (79 FR 50658). The document announced the renewal of certain FDA advisory committees by the Commissioner of Food and Drugs. The table in the document contained several errors. This document corrects those errors.

FOR FURTHER INFORMATION CONTACT: Lisa Granger, Office of Policy, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 3330, Silver Spring, MD 20993–0002. 301–796–9115.

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of Monday, August 25, 2014, in FR Doc. 2014–20017, on page 50659 the table is corrected to read:

Name of committee	Date of expiration
Advisory Committee for Pharmaceutical Science and Clinical Pharmacology	January 22, 2016. March 3, 2016. March 23, 2016. April 5, 2016. April 25, 2016. May 1, 2016. May 13, 2016.
Pulmonary-Allergy Drugs Advisory Committee	May 30, 2016. May 31, 2016. June 2, 2016. June 4, 2016. June 4, 2016. June 9, 2016. June 26, 2016. July 9, 2016.

Dated: September 3, 2014.

### Jill Hartzler Warner,

Associate Commissioner for Special Medical Programs.

[FR Doc. 2014-21369 Filed 9-8-14; 8:45 am]

BILLING CODE 4164-01-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

## **Food and Drug Administration**

[Docket No. FDA-2014-N-1049]

Exploring the Expansion of Conditional Approval to Appropriate Categories of New Animal Drugs

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

SUMMARY: The Food and Drug
Administration (FDA) is announcing
that it is beginning the exploration
process described in a stated
performance goal in the Animal Drug
User Fee Amendments of 2013 (ADUFA
III) goals letter. Consistent with the
performance goal, the FDA is inviting
comments in regard to the Agency
exploring the use of statutory changes to
expand the use of conditional approval