

Governors not later than October 22, 2012.

A. Federal Reserve Bank of Chicago (Colette A. Fried, Assistant Vice President) 230 South LaSalle Street, Chicago, Illinois 60690-1414:

1. *United Community MHC*, Lawrenceburg, Indiana, proposes to convert to stock form and merge with United Community Bancorp, Lawrenceburg, Indiana, an existing savings and loan holding company. The existing United Community Bancorp will merge with a new company, also called United Community Bancorp, Lawrenceburg, Indiana, which will become a savings and loan holding company through the acquisition of 100 percent of the outstanding stock of United Community Bank, Lawrenceburg, Indiana, a federal savings bank.

Board of Governors of the Federal Reserve System, September 21, 2012.

**Robert deV. Frierson,**

*Secretary of the Board.*

[FR Doc. 2012-23706 Filed 9-25-12; 8:45 am]

**BILLING CODE 6210-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Meeting of the Presidential Advisory Council on HIV/AIDS

**AGENCY:** Department of Health and Human Services, Office of the Secretary, Office of the Assistant Secretary for Health.

**ACTION:** Notice.

**SUMMARY:** As stipulated by the Federal Advisory Committee Act, the U.S. Department of Health and Human Service (DHHS) is hereby giving notice that the Presidential Advisory Council on HIV/AIDS (PACHA) will hold a meeting. The meeting will be open to the public.

**DATES:** The meeting will be held October 25, 2012 and October 26, 2012 from 9 a.m. to approximately 5 p.m. (EDT).

**ADDRESSES:** Washington Marriott at Metro Center, 775 12th Street NW., Washington, DC 20005-3901.

**FOR FURTHER INFORMATION CONTACT:** Ms. Caroline Talev, Public Health Assistant, Presidential Advisory Council on HIV/AIDS, Department of Health and Human Services, 200 Independence Avenue SW., Room 443H, Hubert H. Humphrey Building, Washington, DC 20201; (202) 205-1178. More detailed information about PACHA can be obtained by accessing the Council's Web site [www.aids.gov/pacha](http://www.aids.gov/pacha).

**SUPPLEMENTARY INFORMATION:** PACHA was established by Executive Order 12963, dated June 14, 1995 as amended by Executive Order 13009, dated June 14, 1996. The Council was established to provide advice, information, and recommendations to the Secretary regarding programs and policies intended to promote effective prevention of HIV disease and AIDS. The functions of the Council are solely advisory in nature.

The Council consists of not more than 25 members. Council members are selected from prominent community leaders with particular expertise in, or knowledge of, matters concerning HIV and AIDS, public health, global health, philanthropy, marketing or business, as well as other national leaders held in high esteem from other sectors of society. Council members are appointed by the Secretary or designee, in consultation with the White House Office on National AIDS Policy. The agenda for the upcoming meeting will be posted on the Council's Web site at [www.aids.gov/pacha](http://www.aids.gov/pacha).

Public attendance at the meeting is limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact person. Pre-registration for public attendance is advisable and can be accomplished by contacting Caroline Talev at [caroline.talev@hhs.gov](mailto:caroline.talev@hhs.gov). Members of the public will have the opportunity to provide comments at the meeting. Any individual who wishes to participate in the public comment session must register with Caroline Talev at [caroline.talev@hhs.gov](mailto:caroline.talev@hhs.gov); registration for public comment will not be accepted by telephone. Public comment will be limited to two minutes per speaker. Any members of the public who wish to have printed material distributed to PACHA members at the meeting should submit, at a minimum, 1 copy of the materials to Caroline Talev, no later than close of business Thursday, October 18, 2012. Contact information for the PACHA contact person is listed above.

Dated: September 18, 2012.

**B. Kaye Hayes,**

*Executive Director, Presidential Advisory Council on HIV/AIDS.*

[FR Doc. 2012-23720 Filed 9-25-12; 8:45 am]

**BILLING CODE 4150-43-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2012-N-0012]

### Office of the Commissioner of Food and Drugs; Delegation of Authority

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

**ACTION:** Notice.

Notice is hereby given that I have delegated to the Commissioner of Food and Drugs (the Commissioner) certain authority added to the Public Health Service Act by section 801 of Public Law 110-85, the Food and Drug Administration Amendments Act of 2007 (42 U.S.C. 282(j)), pertaining to the expansion of the Clinical Trial Registry and Results Data Bank described therein. Specifically, the Commissioner is delegated the following authority:

- Section 402(j)(5)(C)(ii) of the Public Health Service Act (42 U.S.C. 282(j)(5)(C)(ii))—To determine that any clinical trial information was not submitted as required under 42 U.S.C. 282(j) or was submitted but is false or misleading in any particular and to notify the responsible party and give such party an opportunity to remedy non-compliance by submitting required revised clinical trial information not later than 30 days after such notification.

This authority may be redelegated. This delegation will be exercised in accordance with the Department of Health and Human Services' applicable policies, procedures, guidelines, and regulations.

I ratify and affirm any actions taken by the Commissioner or her subordinates that involved the exercise of the authority delegated herein prior to the effective date of this delegation. This delegation is effective upon date of signature.

Dated: September 5, 2012.

**Kathleen Sebelius,**

*Secretary of Health and Human Services.*

[FR Doc. 2012-23598 Filed 9-25-12; 8:45 am]

**BILLING CODE 4160-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2012-N-0001]

### Anti-Infective Drugs Advisory Committee; Notice of Meeting

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

**ACTION:** Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

*Name of Committee:* Anti-Infective Drugs Advisory Committee.

*General Function of the Committee:* To provide advice and recommendations to the Agency on FDA's regulatory issues.

*Date and Time:* The meeting will be held on November 28, 2012, from 8 a.m. to 5 p.m.

*Location:* DoubleTree by Hilton Hotel Washington DC—Silver Spring, The Ballroom, 8727 Colesville Rd., Silver Spring, MD. The hotel's phone number is 301-589-5200.

*Contact Person:* Diane Goyette, 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, email: [AIDAC@fda.hhs.gov](mailto:AIDAC@fda.hhs.gov), or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site at <http://www.fda.gov/AdvisoryCommittees/default.htm> and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

*Agenda:* The committee will discuss the safety and efficacy of new drug application (NDA) 204384, bedaquiline tablets, submitted by Janssen Therapeutics, Division of Janssen Products, LP. The proposed indication (use) for this product is for the treatment of patients with multi-drug resistant pulmonary tuberculosis.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at <http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm>. Scroll down to the

appropriate advisory committee meeting link.

*Procedure:* Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before November 13, 2012. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before November 2, 2012. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by November 5, 2012.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Diane Goyette at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at <http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm> for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: September 18, 2012.

**Jill Hartzler Warner,**

*Acting Associate Commissioner for Special Medical Programs.*

[FR Doc. 2012-23573 Filed 9-25-12; 8:45 am]

**BILLING CODE 4160-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES****Health Resources and Services Administration****Epilepsy Program**

**AGENCY:** Health Resources and Services Administration, HHS.

**ACTION:** Notice of Noncompetitive Program Expansion Supplemental Award to the Epilepsy Foundation of America.

**SUMMARY:** The Health Resources and Services Administration will be issuing noncompetitive supplemental funding under the Maternal and Child Health Bureau's Epilepsy Program to the Epilepsy Foundation of America (U23MC19824) to support additional evaluation activities.

**SUPPLEMENTARY INFORMATION:**

*Intended Recipient of the Award:* Epilepsy Foundation of America.

*Amount of the Noncompetitive Supplemental Funding:* \$250,700.

**Authority:** Section 501(a)(2) of the Social Security Act, as amended.

CFDA Number: 93.110.

*Project Period:* September 1, 2012, through August 31, 2013, for a total of 12 months.

*Justification for the Exception to Competition:* The Epilepsy Foundation of America currently contracts with Mathematica to evaluate the HRSA Epilepsy Program. There is a need to increase their allotted award through program expansion supplemental funds to support additional evaluation activities for Year 3. Mathematica will conduct a more comprehensive evaluation of the entire Epilepsy Program in order to determine the extent to which the program has addressed the legislative requirements, increased access to care in medically underserved areas, and developed/implemented evidence-based strategies to achieve the legislative purpose of the program. As a result of the evaluation, a quality improvement strategy for the Epilepsy Program will be developed and implemented by the Quality Improvement (QI) contractor. Results should provide a more comprehensive understanding of the impact across the entire program. Indicators and measures developed will also be used for the next competitive cycle to track performance, quality and outcomes within the context of the legislation and the newly published Institute of Medicine recommendations for the Epilepsies, focusing on evaluating prevention efforts.

Mathematica, John Snow Inc. (QI Contractor), and the Epilepsy