DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Notice of public workshop.

Dated: May 24, 2016.

Jill Hartzler Warner
Associate Commissioner for Special Medical Programs.

[FR Doc. 2016–12684 Filed 5–27–16; 8:45 am]
BILLING CODE 4164–01–P

Supplementary Information: FDA is announcing a public workshop regarding antibacterial drug development for patients with unmet need and developing antibacterial drugs that target a single species. Discussions will focus on potential development pathways, aspects of clinical trials including patient population, trial designs, and endpoints, and the role of clinical trial networks in antibacterial drug development.

Registration: Registration is free for the public workshop. Interested parties are encouraged to register early. Seating will be available on a first-come, first-served basis. To register electronically, email registration information (including name, title, firm name, address, telephone, and fax number) to unmetneed2016@fda.hhs.gov. Persons without access to the Internet can call 301–796–1300 to register.

If you need special accommodations due to a disability, please contact Jessica Barnes or Lori Benner (see FOR FURTHER INFORMATION CONTACT) at least 7 days in advance.

Agenda: The workshop draft Agenda will be made available at: http://wwwfda.gov/Drugs/NewsEvents/ucm497650.htm at least 2 days prior to the meeting. The Agency encourages individuals, industry, health professionals, researchers, public health organizations and other interested persons to attend this public workshop.

Transcripts: Please be advised that as soon as a transcript is available, it will be accessible at http://www.regulations.gov. It may be viewed at the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1611, Rockville, MD 20852. A transcript will also be available in either hardcopy or on CD–ROM, after submission of a Freedom of Information request. The Freedom of Information office address is available on the Agency’s Web site at http://www.fda.gov. Transcripts will also be available on the Internet at: http://wwwfda.gov/Drugs/NewsEvents/ucm497650.htm approximately 45 days after the workshop.

Dated: May 24, 2016.
Leslie Kux,
Associate Commissioner for Policy.

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For Further Information Contact: Lori Benner and/or Jessica Barnes, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 22, Rm. 6221 Silver Spring, MD 20993–0002, 301–796–1300.

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