increasing the probability that customers would pay higher prices for fibre channel switches and that innovation would be lessened.

### **Entry**

Entry into the worldwide fibre channel switch market is not likely to occur in a timely, likely, or sufficient magnitude, character and scope to deter or counteract any anticompetitive effects created by the proposed Acquisition. Entry is unlikely in light of slowly declining demand for fibre channel switches in a mature market, customers that tend to stay with one fibre channel switch manufacturer for extended periods of time, and the significant capital costs required for entry.

### The Consent Agreement

To remedy the alleged competitive concern stemming from Broadcom's access to Cisco's competitively sensitive confidential information, the consent decree prevents the Cisco information from being shared among Broadcom employees who could use such information to raise prices or lessen innovation.

Pursuant to the proposed Order, only authorized individuals will have access to Cisco's competitively sensitive confidential information that is given to the firewalled entity, which is defined as Broadcom's business group responsible for the development, production, sale, and marketing of fibre channel ASICs for Cisco. The firewalled entity will have separate facilities and a separate information technology system with security protocols assuring access only to the authorized individuals. Furthermore, Broadcom shall require all authorized individuals to sign a nondisclosure agreement, requiring compliance with the terms of the proposed Order. Additionally, the proposed Order provides for a cooling off period whereby any authorized individual who leaves his or her position at the firewalled entity will not work in the development, production, sale, or marketing of fibre channel ASICs for Brocade's business unit or in the development, production, sales, and marketing of fibre channel switches for twelve months.

The proposed Order also requires Broadcom to use Cisco's competitively sensitive confidential information only in furtherance of the design, manufacturing, and sale of fibre channel ASICs for Cisco. Moreover, Broadcom will be required to take all actions necessary to prevent access to, or the disclosure or use of Cisco's competitively sensitive confidential information by or to anyone who is not an authorized individual. The proposed Order also incorporates by reference non-disclosure provisions contained in four prior private Confidentiality Agreements that Broadcom, or its predecessor, signed with Cisco.

To ensure compliance with the proposed Order, the Commission will appoint a Monitor to oversee Broadcom's and Brocade's performance of their obligations pursuant to the Consent Agreement. The Monitor will be appointed to a five-year term, but the Commission may extend or modify the term as appropriate up to a ten-year period. Further, the Consent Agreement contains appropriate reporting requirements.

### **Opportunity for Public Comment**

The purpose of this analysis is to facilitate public comment on the proposed Consent Agreement to aid the Commission in determining whether it should make the proposed Consent Agreement final. This analysis is not an official interpretation of the proposed Consent Agreement and does not modify its terms in any way.

By direction of the Commission.

#### Donald S. Clark,

Secretary.

[FR Doc. 2017–14536 Filed 7–11–17; 8:45 am]
BILLING CODE 6750–01–P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Centers for Disease Control and Prevention** 

Sunshine Act Meeting: Board of Scientific Counselors NCEH/ATSDR; Cancelation

FEDERAL REGISTER CITATION OF PREVIOUS ANNOUNCEMENT: The original Federal Register Notice for this meeting was published in the Federal Register on May 22, 2017, Volume 82, Number 2017–10333, page/s/23250–23251.

PREVIOUSLY ANNOUNCED TIME AND DATE OF THE MEETING: 9:00 a.m.—noon, EDT, June 23, 2017.

**CHANGES IN THE MEETING:** This meeting is being canceled in its entirety.

CONTACT PERSON FOR MORE INFORMATION: Amanda Malasky, BS, ORISE Fellow, CDC, 4770 Buford Hwy., Atlanta, GA 30344, telephone 770–488–7699; yoo0@cdc.gov.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee

management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

### Claudette Grant,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2017–14745 Filed 7–10–17; 4:15 pm]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Food and Drug Administration** 

[Docket No. FDA-2017-N-0002]

New Animal Drugs; Withdrawal of Approval of a New Animal Drug Application

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

11101

**ACTION:** Notice of withdrawal.

**SUMMARY:** The Food and Drug Administration (FDA) is withdrawing approval of two new animal drug applications (NADAs). This action is being taken at the sponsor's request because these products are no longer manufactured or marketed.

**DATES:** Withdrawal of approval is effective July 24, 2017.

### FOR FURTHER INFORMATION CONTACT:

Sujaya Dessai, Center for Veterinary Medicine (HFV–212), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–402–5761, sujaya.dessai@fda.hhs.gov.

**SUPPLEMENTARY INFORMATION:** Zoetis, Inc., 333 Portage St., Kalamazoo, MI 49007 has requested that FDA withdraw approval of NADA 065–291 for bulk dihydrostreptomycin sulfate and NADA 065–324 for bulk streptomycin sulfate because the products are no longer manufactured or marketed.

Therefore, under authority delegated to the Commissioner of Food and Drugs, and in accordance with § 514.116 Notice of withdrawal of approval of application (21 CFR 514.116), notice is given that approval of NADA 065–291 and NADA 065–324, and all supplements and amendments thereto, is hereby withdrawn, effective July 24, 2017.

As neither of these NADAs was codified, the animal drug regulations do not require amendment to reflect the voluntary withdrawal of approval of these applications.