

**ACTION:** Notice of public meeting; request for comments.

The Food and Drug Administration (FDA) is announcing a public meeting entitled "Town Hall Discussion With the Director of the Center for Devices and Radiological Health and Other Senior Center Management." The purpose of this meeting is to present the Center for Devices and Radiological Health (CDRH) fiscal year (FY) 2010 priorities. In addition, FDA is interested in engaging in discussions about issues that are of importance to the medical device industry.

**Date and Time:** The public meeting will be held on June 22, 2010, from 9 a.m. to 5 p.m.

**Location:** The public meeting will be held at the Hilton Boston/Woburn, Two Forbes Rd., Woburn, MA 01801. The meeting will not be videotaped or Web cast.

**Contact:** Heather Howell, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, rm. 4320, Silver Spring, MD 20993, 301-796-5718, e-mail:

[heather.howell@fda.hhs.gov](mailto:heather.howell@fda.hhs.gov).

**Registration and Requests for Oral Presentations:** If you wish to attend the public meeting, you must register online at: <http://www.fda.gov/MedicalDevices/NewsEvents/WorkshopsConferences/ucm206671.htm>. Provide complete contact information for each attendee, including name, title, company or organization, address, e-mail, and telephone number. Registration requests must be received by 5 p.m. on Wednesday, June 9, 2010.

If you wish to make an oral presentation during any of the sessions at the meeting (see section II of this document), you must indicate this at the time of registration. FDA will do its best to accommodate requests to speak. Individuals and organizations with common interests are urged to consolidate or coordinate their presentations, and to request time for a joint presentation. FDA will determine the amount of time allotted to each presenter and the approximate time that each oral presentation is scheduled to begin.

Registration is free and will be on a first-come, first-served basis. Early registration is recommended because seating is limited. FDA may limit the number of participants from each organization based on space limitations. Registrants will receive confirmation once they have been accepted. Onsite registration on the day of the public meeting will be provided on a space-available basis beginning at 8 a.m.

If you need special accommodations due to a disability, please contact Susan Monahan at 301-796-5661 or e-mail: [susan.monahan@fda.hhs.gov](mailto:susan.monahan@fda.hhs.gov) at least 7 days in advance of the meeting.

**Comments:** FDA is holding this public meeting to share information and discuss issues of importance to the medical device industry. CDRH is specifically interested in addressing the following question: What mechanism(s) would you prefer or suggest for FDA to engage with industry? The deadline for responding to this question and for submitting other comments related to this public meeting is Wednesday, June 9, 2010.

Regardless of attendance at the public meeting, interested persons may submit written or electronic comments. Submit written comments to the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to <http://www.regulations.gov>. Submit a single copy of electronic comments or two paper copies of any mailed comments, except that individuals may submit one paper copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

#### **SUPPLEMENTARY INFORMATION:**

##### **I. Background**

CDRH has announced four priority areas of activity for FY 2010, each of which presents significant opportunities to improve CDRH's effectiveness in fulfilling our public health mission. More information, including specific goals and actions associated with each priority, is available under "CDRH Strategic Planning" at <http://www.fda.gov/AboutFDA/CentersOffices/CDRH>.

##### **II. Public Meeting**

The objective of this public meeting is to present CDRH's FY 2010 priorities. In addition, FDA is interested in engaging in discussions about issues that are of importance to the medical device industry. CDRH wishes to obtain feedback/ideas for facilitating two-way communication between CDRH and the medical device industry.

The meeting will open with an introduction of CDRH Senior Staff in attendance. Following introductions, Dr. Jeffrey Shuren, the Director of CDRH, will present the FY 2010 CDRH priorities. Industry representatives and other members of the public will then be given the opportunity to present

comments to CDRH Senior Staff. Attendees from CDRH may respond to questions presented by industry and other members of the public.

In advance of the meeting, additional information, including a meeting agenda with a speakers' schedule, will be made available on the Internet. This information will be placed on file in the public docket (docket number found in brackets in the heading of this document), which is available at <http://www.regulations.gov>. This information will also be available at <http://www.fda.gov/MedicalDevices/NewsEvents/WorkshopsConferences/default.htm> (select the appropriate meeting from the list).

##### **III. 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. 1061, Rockville, MD. A transcript will also be available in either hardcopy or on CD-ROM, after submission of a Freedom of Information request. Written requests are to be sent to Division of Freedom of Information (HFI-35), Office of Management Programs, Food and Drug Administration, 5600 Fishers Lane, rm. 6-30, Rockville, MD 20857.

Dated: April 30, 2010.

**Leslie Kux,**

*Acting Assistant Commissioner for Policy.*

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#### **DEPARTMENT OF HEALTH AND HUMAN SERVICES**

##### **Food and Drug Administration**

[Docket No. FDA-2010-N-0002]

##### **Withdrawal of Approval of New Animal Drug Applications; Coumaphos; Novobiocin; Buquinolate and Lincomycin**

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is withdrawing approval of three new animal drug applications (NADAs). In a final rule published elsewhere in this issue of the **Federal Register**, FDA is amending the animal drug regulations to remove portions reflecting approval of the single NADA of the three that is codified.

**DATES:** Withdrawal of approval is effective May 17, 2010.

**FOR FURTHER INFORMATION CONTACT:** John Bartkowiak, Center for Veterinary Medicine (HFV-212), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240-276-9079, e-mail: [john.bartkowiak@fda.hhs.gov](mailto:john.bartkowiak@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:** The following sponsors have requested that FDA withdraw approval of the three NADAs listed in Table 1:

TABLE 1.

Sponsor	NADA No./Product (Established Name of Drug)	21 CFR Cite (Sponsor's Drug Labeler Code)
Pharmacia & Upjohn Co., a Division of Pfizer, Inc., 235 East 42d St., New York, NY 10017	NADA 13-467/ALBAMIX Susceptibility Disks (novobiocin)	Not codified
Pharmacia & Upjohn Co., a Division of Pfizer, Inc., 235 East 42d St., New York, NY 10017	NADA 45-738/LINCOMIX/BONAIID (lincomycin/buquinolate)	Not codified
Purina Mills, Inc., P.O. Box 66812, St. Louis, MO 63166-6812	NADA 42-117/Purina 6 Day Worm-Kill Concentrate (coumaphos)	558.185 (017800)

Therefore, under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, and in accordance with § 514.116 *Notice of withdrawal of approval of application* (21 CFR 514.116), notice is given that approval of NADAs 13-467, 42-117, and 45-738, and all supplements and amendments thereto, is hereby withdrawn, effective May 17, 2010.

In a final rule published elsewhere in this issue of the **Federal Register**, FDA is amending the animal drug regulations to reflect the withdrawal of approval of NADA 42-117.

Dated: April 30, 2010.

**Bernadette Dunham,**

*Director, Center for Veterinary Medicine.*

[FR Doc. 2010-10567 Filed 5-4-10; 8:45 am]

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## DEPARTMENT OF HOMELAND SECURITY

### United States Immigration and Customs Enforcement

#### Agency Information Collection Activities: Extension of an Existing Information Collection; Comment Request

**ACTION:** 60-Day Notice of Information Collection for Review; Form I-333, Obligor Change of Address.

The Department of Homeland Security, U.S. Immigration and Customs Enforcement (USICE), has submitted the following information collection request for review and clearance in accordance with the Paperwork Reduction Act of 1995. The information collection is published to obtain comments from the public and affected agencies. Comments

are encouraged and will be accepted for sixty days until July 6, 2010.

Written comments and suggestions regarding items contained in this notice, and especially with regard to the estimated public burden and associated response time should be directed to the Department of Homeland Security (DHS), Joseph M. Gerhart, Chief, Records Management Branch, U.S. Immigration and Customs Enforcement, 500 12th Street, SW., Room 3138, Washington, DC 20536; (202) 732-6337.

Comments are encouraged and will be accepted for sixty days until July 6, 2010. Written comments and suggestions from the public and affected agencies concerning the proposed collection of information should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

#### Overview of this information collection:

(1) *Type of Information Collection:* Extension of a currently approved information collection.

(2) *Title of the Form/Collection:* Obligor Change of Address.

(3) *Agency form number, if any, and the applicable component of the Department of Homeland Security sponsoring the collection:* Form I-333, U.S. Immigration and Customs Enforcement.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: Individual or Households, Business or other non-profit. The information collected on the Form I-333 is necessary for U.S. Immigration and Customs Enforcement (ICE) to provide immigration bond obligors a standardized method to notify ICE of address updates. Upon receipt of the formatted information records will then be updated to ensure accurate service of correspondence between ICE and the obligor.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* 12,000 responses at 15 minutes (.25 hours) per response.

(6) *An estimate of the total public burden (in hours) associated with the collection:* 3,000 annual burden hours.

Requests for a copy of the proposed information collection instrument, with instructions; or inquiries for additional information should be requested via e-mail to: [forms.ice@dhs.gov](mailto:forms.ice@dhs.gov) with "ICE Form I-333" in the subject line.

Dated: April 28, 2010.

**Joseph M. Gerhart,**

*Branch Chief, Records Management Branch, U.S. Immigration and Customs Enforcement, Department of Homeland Security.*

[FR Doc. 2010-10546 Filed 5-4-10; 8:45 am]

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