

**REGISTRATION:** There is no registration fee, however, space is limited. Persons will be registered in the order in which calls are received. Please call Betty B. Palsgrove at 301-827-6618 to register. Registrations also may be transmitted by fax to 1-800-344-3332 or 301-443-2446. Please include the name and title of the person attending and the name of the organization.

**FOR FURTHER INFORMATION CONTACT:** Peter H. Rheinwein, M.D., J.D., Office of Health Affairs (HFY-40), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-6630.

**SUPPLEMENTARY INFORMATION:**

The purpose of the meeting is to provide an opportunity for representatives of health professional organizations and other interested persons to be briefed by senior FDA staff. It will also provide an opportunity for informal discussion on these topics of particular interest to health professional organizations.

This public meeting is free of charge; however, space is limited. Registration for the meeting will be accepted in the order received and should be sent to the contact person. Registration should include the name and title of the person attending and the name of the organization being represented, if any.

Dated: January 20, 1998.

**William K. Hubbard,**

*Associate Commissioner for Policy Coordination.*

[FR Doc. 98-1850 Filed 1-26-98; 8:45 am]

BILLING CODE 4160-01-F

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**Medical Imaging 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). At least one portion of the meeting will be closed to the public.

*Name of Committee:* Medical Imaging Drugs Advisory Committee.

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

*Date and Time:* The meeting will be held on February 9, 1998, 8 a.m. to 5 p.m.

*Location:* Holiday Inn, Versailles Ballrooms I and II, 8120 Wisconsin Ave., Bethesda, MD.

*Contact Person:* Leander B. Madoo, Center for Evaluation and Research (HFD-21), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-5455, or FDA Advisory Committee Information Line, 1-80-741-8138 (301-443-0572 in the Washington, DC area), code 12540. Please call the Information Line for up-to-date information on this meeting.

*Agenda:* The committee will discuss the safety and efficacy of new drug application (NDA) 20-887 AcuTect™, Diatide, Inc., a radiopharmaceutical agent for the detection and localization of acute venous thrombosis.

*Procedure:* On February 9, 1998, from 8 a.m. to 1 p.m. and from 2 p.m. to 5 p.m. the meeting is open to the public. 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 by February 2, 1998. Oral presentations from the public will be scheduled between approximately 8:30 a.m. and 9:30 a.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before February 2, 1998, 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.

*Closed Committee Deliberations:* On February 9, 1998, from 1 p.m. to 2 p.m., the meeting will be closed to permit discussion and review of trade secret and/or confidential information relating to NDA 20-887 AcuTect™ (5 U.S.C. 552b(c)(4)).

FDA regrets that it was unable to publish this notice 15 days prior to the February 9, 1998, Medical Imaging Drugs Advisory Committee meeting. Because the agency believes there is some urgency to bring this issue to public discussion and qualified members of the Medical Imaging Drugs Advisory Committee were available at this time, the Commissioner concluded that it was in the public interest to hold this meeting even if there was not sufficient time for the customary 15-day public notice.

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

Dated: January 22, 1998.

**Michael A. Friedman,**

*Deputy Commissioner for Operations.*

[FR Doc. 98-2022 Filed 1-23-98; 11:47 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. 98D-0017]

**International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products (VICH); Draft Guidance on Validation of Analytical Procedures: Definition and Terminology (#63), and Validation of Analytical Procedures: Methodology (#64); Availability; Request for Comments**

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

**ACTION:** Notice; request for comments.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing the availability for comment of two draft guidance for industry (GFI) documents entitled "Validation of Analytical Procedures: Definition and Terminology" (number 63) and "Validation of Analytical Procedures: Methodology" (number 64). These related draft GFI documents have been adapted for veterinary use by the International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products (VICH) from two guidelines, Q2A and Q2B, that were adopted by the International Conference on Harmonisation (ICH) of Technical Requirements for Registration of Pharmaceuticals for Human Use. The draft guidance is intended to provide guidance on characteristics that should be considered during the validation of analytical procedures included as part of registration applications for approval of veterinary medicinal products submitted to the European Union, Japan, and the United States.

**DATES:** Submit written comments on these draft GFI documents by March 30, 1998.

**ADDRESSES:** Submit written comments on the two draft GFI documents to the Dockets Management Branch (HFA-305), Food and Drug Administration, 12420 Parklawn Dr., rm 1-23, Rockville, MD 20857. Comments should be identified with the full title of the draft GFI document and the docket number found in the heading of this document.