

Dated: December 26, 2001.

**Linda A. Suydam,**

*Senior Associate Commissioner.*

[FR Doc. 02-26 Filed 1-2-02; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

#### Joint Meeting of the Transmissible Spongiform Encephalopathies Advisory Committee and the Blood Products Advisory Committee; Notice of Meeting

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

**ACTION:** Notice.

This notice announces a forthcoming joint meeting of two public advisory committees of the Food and Drug Administration (FDA). At least one portion of the joint meeting will be closed to the public.

*Name of Committees:* Transmissible Spongiform Encephalopathies Advisory Committee and the Blood Products Advisory Committee.

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

*Date and Time:* The meeting will be held on January 16, 2002, from 1 p.m. to 4:30 p.m.; and on January 17, 2002, from 8 a.m. to 5:15 p.m.

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

*Contact:* William Freas or Sheila D. Langford, Center for Biologics Evaluation and Research (HFM-71), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852-1448, 301-827-0314, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 12392. Please call the Information Line for up-to-date information on this meeting.

*Agenda:* On January 17, 2002, the committees will listen to updates on the "Revised FDA Guidance on Preventive Measures to Reduce the Possible Risk of Transmission of Creutzfeldt-Jakob Disease (CJD) and Variant Creutzfeldt-Jakob Disease (vCJD) by Blood and Blood Products" document, and other related topics. For the purpose of further evaluating the adequacy of our present blood deferral recommendations, the committee will then discuss the effectiveness of measures taken to protect humans from foodborne exposure to the bovine spongiform

encephalopathy (BSE) agent in countries with BSE.

*Procedure:* On January 17, 2002, from 8 a.m. to 4 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 January 11, 2002. Oral presentations from the public will be scheduled between approximately 1:20 p.m. and 2:20 p.m. on January 17, 2002. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before January 11, 2002, 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 January 16, 2002, from 1 p.m. to 4 p.m., the meeting will be closed to permit discussion and review of trade secret and/or confidential information (5 U.S.C. 552b(c)(4)). This portion of the meeting will be closed to permit discussion of this material.

FDA regrets that it was unable to publish this notice 15 days prior to the January 16 and 17, 2002, Joint Meeting of the Transmissible Spongiform Encephalopathies Advisory Committee and the Blood Products Advisory Committee meeting. Because the agency believes there is some urgency to bring these issues to public discussion and qualified members of the Joint Meeting of the Transmissible Spongiform Encephalopathies Advisory Committee and the Blood Products Advisory Committee were available at this time, the Commissioner of Food and Drugs 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).

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**Linda A. Suydam,**

*Senior Associate Commissioner.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Government-Owned Inventions; Availability for Licensing

**AGENCY:** National Institutes of Health, DHHS.

**ACTION:** Notice.

**SUMMARY:** The inventions listed below are owned by agencies of the U.S. Government and are available for licensing in the U.S. in accordance with 35 U.S.C. 207 to achieve expeditious commercialization of results of federally-funded research and development. Foreign patent applications are filed on selected inventions to extend market coverage for companies and may also be available for licensing.

**ADDRESSES:** Licensing information and copies of the U.S. patent applications listed below may be obtained by writing to the indicated licensing contact at the Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, Maryland 20852-3804; telephone: 301/496-7057; fax: 301/402-0220. A signed Confidential Disclosure Agreement will be required to receive copies of the patent applications.

#### XAGE-1, A Gene Expressed in Multiple Cancers and Uses Thereof

Drs. Ira H. Pastan (NCI), Xiu F. Liu (NCI), Byungkook Lee (NCI) and Lee J. Helman (NCI)

DHHS Ref. No. E-161-00/0 (Provisional Application) filed September 1, 2000 and E-161-00/1 (PCT Application) filed August 31, 2001

*Licensing Contact:* Richard Rodriguez; 301/496-7056 ext. 287; e-mail: [rodrigur@od.nih.gov](mailto:rodrigur@od.nih.gov).

The XAGE-1 gene is a human X-linked gene that is strongly expressed in breast cancer, lung cancer and several other cancers as well as normal testes. The largest open reading frame of the XAGE-1 transcript encodes a putative protein of 16.3 kD (p16) with a potential transmembrane domain at the amino terminus. In addition, the XAGE-1 transcript contains a second ATG in the reading frame corresponding to residue 66, which would encode a 9 kD protein (p9). *In vitro* transfection experiments using 293 T cells have revealed a 9 kD protein. However, the size of the endogenously expressed protein is not yet known. XAGE-1 shares homology with GAGE/PAGE proteins in the C-terminal end.