

in this time period to request written confirmation from these companies of their commitment to pay these fees; if companies do not agree to make this commitment, FDA will request that they withdraw their submission(s), and such submissions will not be reviewed. For further information, contact Wayne Amchin (see **FOR FURTHER INFORMATION CONTACT**).

For information on how FDA will treat DTC television advertisement advisory review submissions not identified in response to the participation notice that are submitted after the 30-calendar-day time period for responding to that notice has elapsed, see sections II.A "Basis for the Fee" and II.B "Operating Reserves" of this document.

Dated: December 5, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7-24000 Filed 12-10-07; 8:45 am]

BILLING CODE 4160-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Drug Safety and Risk Management 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). The meeting will be open to the public.

Name of Committee: Drug Safety and Risk Management Advisory Committee.

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

Date and Time: The meeting will be held on February 1, 2008, from 8 a.m. to 5 p.m.

Location: Hilton Washington DC/ Silver Spring, Maryland Ballroom, 8727 Colesville Rd., Silver Spring, MD. The hotel phone number is 301-589-5200.

Contact Person: Teresa Watkins, Center for Drug Evaluation and Research (HFD-21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1093), Rockville, MD 20857, 301-827-7001, FAX: 301-827-6776, e-mail: Teresa.Watkins@fda.hhs.gov, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 3014512535. Please call the Information

Line for up-to-date information on this meeting. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the agency's Web site and call the appropriate advisory committee hot line/phone line to learn about possible modifications before coming to the meeting.

Agenda: The committee will discuss the efficacy and safety of new drug application (NDA) 22-054, INJECTAFER (ferric carboxymaltose injection), Luitpold Pharmaceuticals Incorporated, used for the treatment of iron deficiency anemia in patients with postpartum hemorrhage or heavy uterine bleeding.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at <http://www.fda.gov/ohrms/dockets/ac/acmenu.htm>, click on the year 2008 and scroll down to the appropriate advisory committee link.

Procedure: 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 on or before January 17, 2008. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Those desiring to make formal oral presentations should notify the contact person 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 on or before January 9, 2008. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by January 10, 2008.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee

meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Teresa Watkins at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at <http://www.fda.gov/oc/advisory/default.htm> for procedures on public conduct during advisory committee meetings.

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

Dated: December 4, 2007.

Randall W. Lutter,

Deputy Commissioner for Policy.

[FR Doc. E7-24003 Filed 12-10-07; 8:45 am]

BILLING CODE 4160-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committees; Filing of Closed Meeting Reports

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that, as required by the Federal Advisory Committee Act, the agency has filed with the Library of Congress the annual reports of those FDA advisory committees that held closed meetings during fiscal year 2007.

ADDRESSES: Copies are available from the Dockets Management Branch (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, 301-827-6860.

FOR FURTHER INFORMATION CONTACT:

Theresa L. Green, Committee Management Officer, Advisory Committee and Oversight Management Staff (HF-4), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1220.

SUPPLEMENTARY INFORMATION: Under section 10(d) of the Federal Advisory Committee Act (5 U.S.C. app.1) and 21 CFR 14.60(d), FDA has filed with the Library of Congress the annual reports for the following FDA advisory committees that held closed meetings during the period October 1, 2006 through September 30, 2007: *Center for Biologics Evaluation and Research:*

Cellular, Tissue and Gene Therapies

Advisory Committee,
Vaccines and Related Biological
Products Advisory Committee,
Center for Drugs Evaluation and
Research:

Antiviral Drugs Advisory Committee
Center for Devices and Radiological
Health:

Medical Devices Advisory Committee
(consisting of reports for Dental
Products Panel; Circulatory Devices
Panel)

Annual Reports are available for
public inspections between 9 a.m. and
4 p.m., Monday through Friday.

(1) The Library of Congress, Madison
Bldg., Newspaper and Current
Periodical Reading Room, 101
Independence Ave. SE, rm. 133,
Washington, DC; and (2) The
Dockets Management Branch (HFA-
305), Food and Drug
Administration, 5630 Fishers Lane,
rm. 1061, Rockville, MD 20852.

Dated: December 4, 2007.

Randall W. Lutter,

Deputy Commissioner for Policy.

[FR Doc. E7-23986 Filed 12-10-07; 8:45 am]

BILLING CODE 4160-01-S

DEPARTMENT OF HOMELAND SECURITY

Bureau of Immigration and Customs Enforcement

Agency Information Collection Activities: Extension of a Currently Approved Information Collection; Comment Request

ACTION: 60-Day Notice of Information
Collection Under Review; Data Relating
to Beneficiary of Private Bill, Form G-
79A.

The Department of Homeland
Security, Bureau of Immigration and
Customs Enforcement (ICE) 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 February 11, 2008.

Written comments and suggestions
from the public and affected agencies
concerning the collection of information
should address one or more of the
following four points:

(1) Evaluate whether the 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
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
collection.

(2) *Title of the Form/Collection:* Data
Relating to Beneficiary of Private Bill.

(3) *Agency form number, if any, and
the applicable component of the
Department of Homeland Security
sponsoring the collection:* Form G-79A.
Bureau of Immigration and Customs
Enforcement.

(4) *Affected public who will be asked
or required to respond, as well as a brief
abstract:* Primary: Individuals or
Households. The information is needed
to report on Private Bills to Congress
when requested.

(5) *An estimate of the total number of
respondents and the amount of time
estimated for an average respondent to
respond:* 100 responses at 1 hour per
response.

(6) *An estimate of the total public
burden (in hours) associated with the
collection:* 100 annual burden hours.

Comments and/or questions; requests
for a copy of the proposed information
collection instrument, with instructions;
or inquiries for additional information
should be directed to: Lee Shirkey,
Acting Chief, Records Management
Branch; U.S. Immigration and Customs
Enforcement, 425 I Street, NW., Room
1122, Washington, DC 20536; (202) 616-
2266.

Dated: December 6, 2007.

Lee Shirkey,

*Acting Branch Chief, Records Management
Branch, Bureau of Immigration and Customs
Enforcement, Department of Homeland
Security.*

[FR Doc. E7-23979 Filed 12-10-07; 8:45 am]

BILLING CODE 9111-28-P

DEPARTMENT OF HOMELAND SECURITY

Bureau of Immigration and Customs Enforcement

Agency Information Collection Activities: Extension of a Currently Approved Information Collection, Comment Request

ACTION: Request OMB Emergency
Approval and 60-Day Notice;
Immigration Bond; Form I-352, OMB
Control No. 1653-0022.

The Department of Homeland
Security, Bureau of Immigration and
Customs Enforcement has submitted the
following information collection request
for review and clearance in accordance
with the Paperwork Reduction Act of
1995. The proposed information
collection is published to obtain
comments from the public and affected
agencies. Comments are encouraged and
will be accepted for sixty days until
February 11, 2008.

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:*
Immigration Bond.

(3) *Agency form number, if any, and
the applicable component of the
Department of Homeland Security
sponsoring the collection:* Form I-352.
Bureau of Immigration and Customs
Enforcement.