

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](mailto: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]

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## 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.