DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committee for Pharmaceutical Science and Clinical Pharmacology; 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: Advisory Committee for Pharmaceutical Science and Clinical Pharmacology.

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 March 18, 2008, from 8:30 a.m. to 5 p.m. and on March 19, 2008, from 8:30 a.m. to 12:30 p.m.

Location: Food and Drug Administration, Center for Drug Evaluation and Research Advisory Committee Conference Room, rm. 1066, 5630 Fishers Lane, Rockville, MD.

Contact Person: Mimi Phan, 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: *mimi.phan@fda.hhs.gov*, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in Washington, DC area), code 3014512539. 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: On March 18, 2008, the committee will: (1) Discuss and provide comments on three new topics of this meeting; first new topic: The new clinical pharmacogenomics (PGx) concept paper. Key issues in the concept paper include an industry survey on the collection of PGx samples, and the applications of PGx in clinical development will be presented and (2) discuss and provide comments on the second new topic: Quantitative clinical pharmacology: Critical path opportunities. An example of a disease model and its applications will be presented. The regulatory experience, designs, and implications of pediatric studies will be discussed. On March 19, 2008, the committee will consider the third new topic: Renal impairment concept paper. The effects of renal impairment on Cytochrom P (CYP)/transporter, methods of evaluation of renal function, and the effects of hemodialysis on drug clearance will be discussed.

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 March 4, 2008. Oral presentations from the public will be scheduled between approximately 10:30 a.m. and 11 a.m. each day. 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 February 27, 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 February 28, 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 Mimi Phan 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: February 4, 2008.

Randall W. Lutter,

Deputy Commissioner for Policy.

[FR Doc. E8–2540 Filed 2–11–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

[USCG–2008–0075]

Commercial Fishing Industry Vessel Safety Advisory Committee

AGENCY: Coast Guard, DHS.

ACTION: Notice of Open Teleconference Meeting.

SUMMARY: This notice announces a teleconference of the Commercial Fishing Industry Vessel Safety Advisory Committee (CFIVSAC). The purpose of the teleconference is for CFIVSAC to discuss and prepare recommendations for the Coast Guard concerning the work of the Communications Subcommittee and the Risk Management Subcommittee and to discuss other CFIVSAC actions resulting from its last public meeting on November 13 and 14, 2007.

DATES: The teleconference call will take place on Wednesday, February 27, 2008, from 1:30 p.m. until approximately 3 p.m. Eastern Standard Time.

ADDRESSES: Committee members and members of the public may participate by dialing 1–877–451–9782 on a touch-tone phone. You will then be prompted to enter your ''participant code number,'' which is 9559674#. Please ensure that you enter the # mark after the participant code. Public participation is welcomed; however, the number of teleconference lines is limited, and lines are available first-come, first-served. Members of the public may also participate by coming to Room 1116 U.S. Coast Guard Headquarters; 2100 Second Street, SW., Washington, DC 20593–0001. We request that members of the public who plan to attend this meeting notify Mr. Mike Rosecrans at 202–372–1245 so that

he may notify building security officials. You may also gain access to this docket at *http://dms.dot.gov/search/searchFormSimple.cfm*. Background information is available at *http://www.fishsafe.info*.

FOR FURTHER INFORMATION CONTACT: Mr. Mike Rosecrans, Assistant Executive Director of CFIVSAC, telephone 202–372–1245, fax 202–372–1917.

SUPPLEMENTARY INFORMATION: The Federal Advisory Committee Act requires that public notice of these meetings be announced in the **Federal Register** [5 U.S.C. App. (Pub. L. 92–463)]. CFIVSAC is chartered under that Act. It provides advice and makes recommendations to the Secretary on issues regarding safety of commercial fishing industry vessels.

Tentative Agenda: Wednesday, February 27, 2008 1:30 p.m.: Welcome, introduction of new members and Opening Remarks—CFIVSAC Chairman Mr. Jerry Dzugan.

Open discussion concerning the work of the Communications Subcommittee and Task 07–01 Completion of Fishing Vessel Digest.

Discussion of the work of the Risk Management Subcommittee and Task 07–02—Roles and Mission and Risk Management Best Practices.

Discussion of Task 07–03—Long Range Goals for the Committee.

Public comment period.

Discussion of plans for next meeting.

3 p.m.: Adjourn.

This tentative agenda is subject to change and the meeting may adjourn early if all

Committee business has been completed.

Public Participation

The Chairman of CFIVSAC is empowered to conduct the teleconference in a way that will, in his judgment, facilitate the orderly conduct of business. During its teleconference, the Committee welcomes public comment. The Committee will make every effort to hear the views of all interested parties, including the public. Written comments may be submitted to Mr. Mike Rosecrans, Assistant Executive Director, CFIVSAC; Commandant (CG–5433); 2100 Second Street, SW., Washington, DC 20593–0001.

Information on Services for Individuals With Disabilities

For information on facilities or services for individuals with disabilities or to request special assistance at the meeting, contact Mr. Rosecrans as soon as possible.

Dated: February 7, 2008.

H.L. Hime,

Acting Director of Commercial Standards and Regulations.

[FR Doc. 08–656 Filed 2–8–08; 1:54 pm]

BILLING CODE 4910–15–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Accreditation and Approval of SGS North America, Inc., as a Commercial Gauger and Laboratory

AGENCY: U.S. Customs and Border Protection, Department of Homeland Security.

ACTION: Notice of accreditation and approval of SGS North America, Inc., as a commercial gauger and laboratory.

SUMMARY: Notice is hereby given that, pursuant to 19 CFR 151.12 and 19 CFR 151.13, SGS North America, Inc., 2 Avenue J, Bayonne, NJ 07002, has been approved to gauge and accredited to test petroleum and petroleum products, organic chemicals and vegetable oils for customs purposes, in accordance with the provisions of 19 CFR 151.12 and 19 CFR 151.13. anyone wishing to employ this entity to conduct laboratory analyses and gauger services should request and receive written assurances from the entity that it is accredited or approved by the U.S. Customs and Border Protection to conduct the specific test or gauger service requested. Alternatively, inquiries regarding the specific test or gauger service this entity is accredited or approved to perform may be directed to the U.S. Customs and Border Protection by calling (202) 344–1060. The inquiry may also be sent to *cbp.labhq@dhs.gov.* Please reference the Web site listed below for a complete listing of CBP approved gaugers and accredited laboratories. *http://cbp.gov/xp/cgov/import/operations_support/labs_scientific_svcs/commercial_gaugers/.*

DATES: The accreditation and approval of SGS North America, Inc., as commercial gauger and laboratory became effective on September 20, 2007. The next triennial inspection date will be scheduled for September 2010.

FOR FURTHER INFORMATION CONTACT: Commercial Gauger Laboratory Program Manager, Laboratories and Scientific Services, U.S. Customs and Border Protection, 1300 Pennsylvania Avenue, NW., Suite 1500N, Washington, DC 20229, 202–344–1060.

Dated: January 31, 2008.

Ira S. Reese,

Executive Director, Laboratories and Scientific Services.

[FR Doc. 08–589 Filed 2–11–08; 8:45 am]

BILLING CODE 9111–14–M

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Accreditation and Approval of Inspectorate America Corporation, as a Commercial Gauger and Laboratory

AGENCY: U.S. Customs and Border Protection, Department of Homeland Security.

ACTION: Notice of accreditation and approval of Inspectorate America Corporation, as a commercial gauger and laboratory.

SUMMARY: Notice is hereby given that, pursuant to 19 CFR 151.12 and 19 CFR 151.13, Inspectorate America Corporation, 1404 Joliet Road, Suite G, Romeoville, IL 60446, has been approved to gauge and accredited to test petroleum and petroleum products, organic chemicals and vegetable oils for customs purposes, in accordance with the provisions of 19 CFR 151.12 and 19 CFR 151.13. Anyone wishing to employ this entity to conduct laboratory analyses and gauger services should request and receive written assurances from the entity that it is accredited or approved by the U.S. Customs and Border Protection to conduct the specific test or gauger service requested. Alternatively, inquiries regarding the specific test or gauger service this entity is accredited or approved to perform may be directed to the U.S. Customs and Border Protection by calling (202) 344–1060. The inquiry may also be sent to *cbp.labhq@dhs.gov*. Please reference the Web site listed below for a complete listing of CBP approved gaugers and accredited laboratories. *http://cbp.gov/xp/cgov/import/operations_support/labs_scientific_svcs/commercial_gaugers/*.

DATES: The accreditation and approval of Inspectorate America Corporation, as commercial gauger and laboratory became effective on August 08, 2007. The next triennial inspection date will be scheduled for August 2010.

FOR FURTHER INFORMATION CONTACT: Commercial Gauger Laboratory Program Manager, Laboratories and Scientific Services, U.S. Customs and Border Protection, 1300 Pennsylvania Avenue, NW., Suite 1500N, Washington, DC 20229, 202–344–1060.