Article 8
Duration

This Arrangement commences upon signature by all participants for ten (10) years. It may be extended for an additional ten-year period, after evaluation by the participants.

The participants may amend this document, by mutual written consent, specifying the date the amended Arrangement commences.

This Arrangement may be terminated by any participant upon thirty days advance written notice to the other participants.

Termination of this Arrangement does not affect the completion of cooperation activities that may have been formalized prior to termination.

SIGNED at Washington, D.C., this fourth day of September 2001, in quadruplicate, in the Spanish and English languages.

FOR THE DEPARTMENT OF HEALTH AND HUMAN SERVICES OF THE UNITED STATES OF AMERICA

FOR THE SECRETARIAT OF HEALTH OF THE UNITED MEXICAN STATES

FOR THE DEPARTMENT OF AGRICULTURE OF THE UNITED STATES OF AMERICA

FOR THE SECRETARIAT OF AGRICULTURE, LIVESTOCK, RURAL DEVELOPMENT, FISH, AND FOOD OF THE UNITED MEXICAN STATES

[FR Doc. 01–31576 Filed 12–21–01; 8:45 am]
BILLING CODE 4160–01–C

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Orthopaedic and Rehabilitation Devices Panel of the Medical Devices 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: Orthopaedic and Rehabilitation Devices Panel of the Medical Devices 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 January 10, 2002, from 9:30 a.m. to 5 p.m.

Location: Holiday Inn, Walker/Whetstone Rooms, Two Montgomery Village Ave., Gaithersburg, MD.

Contact: Hany W. Demian, Center for Devices and Radiological Health (HFZ–410), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–2036, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12521. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will discuss, make recommendations, and vote on premarket approval application (PMA) for a spinal fusion cage with a growth

factor soaked in a collagen sponge intended for use to treat lumbar degenerative disc disease. Background information, including the agenda and questions for the committee, will be available to the public on January 9, 2002, on the Internet at http://www.fda.gov/cdrh/panelmtg.html.

Procedure: On January 10, 2002, from 9:30 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 January 4, 2002. Oral presentations from the public will be scheduled between approximately 9:30 a.m. and 10 a.m. on January 10, 2002. Near the end of the committee deliberations for the PMA, a 30-minute open public session will be conducted for interested persons to address issues specific to the submission before the committee. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before January 4, 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 Presentation of Data: On January 10, 2002, from 1 p.m. to 2 p.m., the meeting will be closed to the public to permit the committee to discuss and review trade secret and/or confidential commercial information presented by a sponsor (5 U.S.C. 552b(c)(4)).

FDA regrets that it was unable to publish this notice 15 days prior to the January 10, 2002, Orthopaedic and Rehabilitation Devices Panel of the Medical Devices Advisory Committee meeting. Because the agency believes there is some urgency to bring this issue to public discussion and qualified members of the Orthopaedic and Rehabilitation Devices Panel of the Medical Devices 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).

Dated: December 18, 2001.

Linda A. Suydam,

Senior Associate Commissioner.

[FR Doc. 01–31578 Filed 12–21–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF THE INTERIOR

Office of the Secretary

Delaware and Lehigh Heritage Corridor Commission Meeting

AGENCY: Department of Interior, Office of the Secretary.

ACTION: Notice of meeting.

SUMMARY: This notice announces an upcoming meeting of the Delaware & Lehigh National Heritage Corridor Commission. Notice of this meeting is required under the Federal Advisory Committee Act (Public Law 92–463).

Meeting Date and Time: Friday, January 11, 2002, Time 1:30 p.m. to 4 p.m.

Address: Bethlehem Township Municipal Building, 4225 Easton Avenue, Bethlehem PA 18020.

The agenda for the meeting will focus on implementation of the Management Action Plan for the Delaware and Lehigh National Heritage Corridor and State Heritage Park. The Commission was established to assist the Commonwealth of Pennsylvania and its political subdivisions in planning and implementing an integrated strategy for protecting and promoting cultural, historic and natural resources. The Commission reports to the Secretary of the Interior and to Congress.

FOR FURTHER INFORMATION CONTACT: C. Allen Sachse, Executive Director, Delaware & Lehigh National Heritage Corridor Commission, 10 E. Church Street, Room A–208, Bethlehem, PA 18018, (610) 861–9345.

SUPPLEMENTARY INFORMATION: The Delaware & Lehigh National Heritage Corridor Commission was established by Public Law 100–692, November 18, 1988 and extended through Public Law 105–355, November 13, 1998.

Dated: December 18, 2001.

C. Allen Sachse,

Executive Director, Delaware & Lehigh National Heritage Corridor Commission.

[FR Doc. 01–31585 Filed 12–21–01; 8:45 am]

BILLING CODE 6820–PE–M

INTERNATIONAL TRADE COMMISSION

[Inv. No. 337–TA–459]

In the Matter of Certain Garage Door Operators Including Components Thereof; Notice of Decision to Modify an Initial Determination Granting a Motion to Intervene

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has determined to modify an initial determination (ID) (Order No. 5), as supplemented by Order No. 7, issued by the presiding administrative law judge (ALJ) in the above-captioned investigation. The ID is modified to the extent that the restrictions placed an intervenor's participation in the investigation are lifted.

FOR FURTHER INFORMATION CONTACT: Jean Jackson, Esq., Office of the General Counsel, U.S. International Trade Commission, 500 E Street, SW., Washington, DC 20436, telephone 202–205–3104. Copies of all nonconfidential documents filed in connection with this investigation are or will be available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street SW., Washington, DC 20436, telephone 202–205–2000. General information concerning the Commission may also be obtained by accessing its Internet server (http://www.usitc.gov). The public record for this investigation may be viewed on the Commission's electronic docket (EDIS–ON–LINE) at *http://dockets.usitc.gov/eol.public.* Hearing-impaired persons are advised that information on the matter can be obtained by contacting the Commission's TDD terminal on 202–205–1810.

SUPPLEMENTARY INFORMATION: The Commission instituted this investigation on July 16, 2001, based on a complaint filed by The Chamberlain Group, Inc. (''Chamberlain'') against six entities, not including Microchip Technology Inc. (Microchip). 66 FR 37704. Chamberlain's complaint alleges violations of section 337 in the importation into the United States, sale for importation, and/or sale within the United States after importation of certain garage door operators by reason of infringement of certain claims of Chamberlain's U.S. Letters Patents Nos. Re. 35,364 and Re. 36,703. On August 6, 2001, Microchip filed a motion to intervene in this investigation.

On October 1, 2001, the ALJ issued an ID (Order No. 5) granting Microchip's motion. The ID allowed Microchip to become an ''intervenor'' in the present investigation, but placed restrictions on Microchip's participation. Under the ID, Microchip was allowed to participate in the discovery phase of the investigation, but was not allowed to notice depositions during discovery or participate as a party at the hearing. On October 30, 2001, the Commission