Secretary, Federal Communications Commission.

Parties are strongly encouraged to file comments electronically using the Commission's Electronic Comment Filing System (ECFS). Parties are also requested to send a courtesy copy of their comments via email to *jennifer.mckee@fcc.gov.*

Two (2) copies of the comments and reply comments should also be sent to Chief, Pricing Policy Division, Wireline Competition Bureau, Federal Communications Commission, 445 12th Street, SW., Room 5–A221, Washington, DC 20554. Parties shall also serve one copy with Qualex International, Portals II, 445 12th Street, SW., Room CY–B402, Washington, DC 20554, (202) 863–2893, or via email to <*qualexint@aol.com*>.

Documents in WC Docket No. 03–266, including the Level 3 Petition, are available for public inspection and copying during business hours at the FCC Reference Information Center, Portals II, 445 12th St. SW., Room CY–A257, Washington, DC 20554. The documents may also be purchased from Qualex International, telephone (202) 863–2893, facsimile (202) 863–2898.

This matter shall be treated as a ''permit-but-disclose'' proceeding in accordance with the Commission's *ex parte* rules. *See* 47 CFR 1.1200, 1.1206. Persons making oral *ex parte* presentations are reminded that memoranda summarizing the presentations must contain summaries of the substance of the presentations and not merely a listing of the subjects discussed. More than a one or two sentence description of the views and arguments presented generally is required. *See* 47 CFR 1.1206(b). Other rules pertaining to oral and written *ex parte* presentations in permit-but-disclose proceedings are set forth in section 1.1206(b) of the Commission's rules.

Federal Communications Commission.

William F. Maher, Jr.,

Chief, Wireline Competition Bureau.

[FR Doc. 04–666 Filed 1–12–04; 8:45 am]

BILLING CODE 6712–01–U

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N–0187]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Postmarket Surveillance of Medical Devices

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Postmarket Surveillance of Medical Devices'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Peggy Robbins, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of August 20, 2003 (68 FR 58690), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0449. The approval expires on December 31, 2006. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets.*

Dated: January 6, 2004.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–599 Filed 1–12–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N–0318]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Procedures for the Safe and Sanitary Processing and Importing of Fish and Fishery Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Procedures for the Safe and Sanitary Processing and Importing of Fish and Fishery Products'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Peggy Robbins, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of November 24, 2003 (68 FR 65937), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0354. The approval expires on December 31, 2006. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets.*

Dated: January 6, 2004.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–602 Filed 1–12–04; 8:45 am]

BILLING CODE 4160–01–S