Carriage (CS Docket No. 97–248 and RM–9097). Summary: The Commission will consider action concerning amendments to the program access rules.

6—International—Title: Biennial Regulatory Review -- Reform of the International Settlements Policy and Associated Filing Requirements. Summary: The Commission will consider allowing U.S. international telecommunications carriers greater flexibility in negotiating arrangements with foreign telecommunications carriers for the exchange of international telecommunications traffic.

Additional information concerning this meeting may be obtained from Maureen Peratino or David Fiske, Office of Public Affairs, telephone number (202) 418–0500; TTY (202) 418–2555.

Copies of materials adopted at this meeting can be purchased from the FCC's duplicating contractor, International Transcription Services, Inc. (ITS, Inc.) at (202) 857–3800; fax (202) 857–3805 and 857–3184; or TTY (202) 293–8810. These copies are available in paper format and alternative media, including large print/type; digital disk; and audio tape. ITS may be reached by e-mail: its—inc@ix.netcom.com. Their Internet address is http://www.itsi.com.

This meeting can be viewed over George Mason University's Capitol Connection. For information on this service call (703) 993–3100. The audio portion of the meeting will be broadcast live on the Internet via the FCC's Internet audio broadcast page at <http://www.fcc.gov/realaudio/>. The meeting can also be heard via telephone, for a fee, from National Narrowcast Network, telephone (202) 966–2211 or fax (202) 966–1770; and from Conference Call USA (available only outside the Washington, DC metropolitan area), telephone 1–800–962–0044. Audio and video tapes of this meeting can be purchased from Infocus, 341 Victory Drive, Herndon, VA 20170, telephone (703) 834–0100; fax number (703) 834–0111.

Dated July 30, 1998.

Federal Communications Commission.

Magalie Roman Salas,

Secretary.

[FR Doc. 98–20867 Filed 7–31–98; 11:36 am]

BILLING CODE 6712–01–F

FEDERAL COMMUNICATIONS COMMISSION

Public Information Collection Approved by Office of Management and Budget

July 28, 1998.

The Federal Communications Commission (FCC) has received Office of Management and Budget (OMB) approval for the following public information collection pursuant to the Paperwork Reduction Act of 1995, Pub. L. 96–511. An agency may not conduct or sponsor a collection of information unless it displays a currently valid control number. Not withstanding any other provisions of law, no person shall be subject to any penalty for failing to comply with a collection of information subject to the Paperwork Reduction Act (PRA) that does not display a valid control number. Questions concerning the OMB control numbers and expiration dates should be directed to Judy Boley, Federal Communications Commission, (202) 418–0214.

Federal Communications Commission

OMB Control No.: 3060–0834.

Expiration Date: 12/31/1998.

Title: Reconsideration of Rules and Policies for the 220–222 MHz Radio Service—PR 98–552, GN 92–252, PR 93–253.

Form No.: N/A.

Estimated Annual Burden: 44,850 annual hours; 30 minutes—12 hours per response; 18,400 responses. This collection contains various burdens including coordinating actions with other licensees, submitting certifications with applications for modification of authorization, and seeking a waiver of section 90.729(b).

Description: The information collected will be used by the Commission to verify licensee compliance with Commission rules and regulations and to ensure the integrity of the 220 MHz service, and to ensure that licensees continue to fulfill their statutory responsibilities in accordance with the Communications Act of 1934.

Federal Communications Commission.

Magalie Roman Salas,

Secretary.

[FR Doc. 98–20747 Filed 8–3–98; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

AGENCY HOLDING THE MEETING: Board of Governors of the Federal Reserve System.

TIME AND DATE: 11:00 a.m., Monday, August 10, 1998.

PLACE: Marriner S. Eccles Federal Reserve Board Building, 20th and C Streets, N.W., Washington, D.C. 20551.

STATUS: Closed.

MATTERS TO BE CONSIDERED:

1. Personnel actions (appointments, promotions, assignments, reassignments, and salary actions) involving individual Federal Reserve System employees.

2. Any items carried forward from a previously announced meeting.

CONTACT PERSON FOR MORE INFORMATION: Lynn S. Fox, Assistant to the Board; 202–452–3204.

SUPPLEMENTARY INFORMATION: You may call 202–452–3206 beginning at approximately 5 p.m. two business days before the meeting for a recorded announcement of bank and bank holding company applications scheduled for the meeting; or you may contact the Board's Web site at http://www.bog.frb.fed.us for an electronic announcement that not only lists applications, but also indicates procedural and other information about the meeting.

Dated: July 31, 1998

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 98–20915 Filed 7–31–98; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[INFO–98–24]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of section 3506(c) (2) (A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer at (404) 639–7090.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the

agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Send comments to Seleda Perryman, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice. Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Proposed Projects

1. Requirement for a Special Permit to Import Cynomolgus-African Green or Rhesus Monkeys—(0920–0263)—Extension—National Center for Infectious Disease (NCID) Division of Quarantine—A registered importer nonhuman primates must submit to the Director, CDC, a written plan which specifies the steps that will be taken to prevent exposure of persons and animals during the entire importation and quarantine process for the arriving nonhuman primates. Under the special permit arrangement, registered importers must submit a plan to CDC for the importation and quarantine if they wish to import the specific monkeys covered. The plan must address disease prevention procedures to be carried out in every step of the chain of custody of such monkeys, from embarkation in the country of origin to release from quarantine. Information such as species, origin and intended use for monkeys, transit information, isolation and quarantine procedures, and procedures for testing of quarantined animals is necessary for CDC to make public health decisions. This information enables CDC to evaluate compliance with the standards and determine whether the measures being taken to prevent exposure of persons and animals during importation are adequate. Once CDC is assured, through the monitoring of shipments (normally no more than 2), that the provisions of a special permit plan are being followed by a new permit holder, and that the use of adequate disease control practices is being demonstrated, the special permit is extended to cover the receipt of additional shipments under the same plan for a period of 180 days, and may be renewed upon request. This eliminates the burden on importers to repeatedly report identical information, requiring only that specific shipment itineraries and information on changes to the plan which require approval be submitted.

The respondents are commercial or not-for-profit importers of nonhuman primates. We are requesting clearance for 3 years. Total cost to Respondents: $350 (14 x $25)

Respondents	Number of respondents	Number of responses/respondents	Avg. burden/responses (in hrs.)	Total burden (in hrs.)
Businesses	5	5	2@0.5; 3@0.1	6.5
Organizations	15	5	0.1	7.5
Total				14

Dated: July 29, 1998.

Charles W. Gollmar,

Acting Associate Director for Policy, Planning and Evaluation Centers for Disease Control and Prevention (CDC).

[FR Doc. 98–20719 Filed 8–3–98; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N–0453]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection and information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the submission of applications to recognized accreditation bodies that will assess potential U.S. Conformity Assessment Bodies (CAB's) seeking to be designated under the United States (U.S.)/European Community (EC) Mutual Recognition Agreement (MRA) to assess medical devices produced for the EC market. Elsewhere in this issue of the **Federal Register**, FDA is publishing a notice announcing OMB's approval of this collection of information (OMB control number 0910–0378). Since this was an emergency approval that expires on January 31, 1999, FDA is following the normal PRA clearance procedures by issuing this notice.

DATES: Submit written comments on the collection of information by October 5, 1998.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Margaret R. Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth below.