procedures under which CMS will periodically disclose the identity of Medicare enrollees whose records have been inactive for a specific period of time (not to be less than 1 year). SSA will use the selected data as an indicator of cases that should be reviewed in order to determine continued eligibility to SSA-administered programs. Cases may be selected for review directly from the match of non-utilization of Medicare or may be used as a factor in a system to prioritize reviews. SSA's Office of Inspector General (OIG) will investigate individual cases alleging fraud, waste, and/or abuse referred to the OIG.

CATEGORIES OF RECORDS AND INDIVIDUALS COVERED BY THE MATCH:

SSA will furnish CMS with an electronic file containing Title II Claim Account Number and Title II Beneficiary Identification Code and any other information needed to accurately match the records. CMS will match the SSA file against its ''Enrollment Database'' system of records (formerly known as the Health Insurance Master Record), system No. 09–70–0502, and will disclose information on non-utilization of Medicare benefits by SSA recipients.

INCLUSIVE DATES OF THE MATCH:

The Computer Matching Program shall become effective no sooner than 40 days after the report of the matching program is sent to OMB and Congress, or 30 days after publication in the **Federal Register**, whichever is later. The matching program will continue for 18 months from the effective date and may be extended for an additional 12 months thereafter, if certain conditions are met.

[FR Doc. 01–26491 Filed 10–19–01; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01N–0175]

Agency Information Collection Activities; Announcement of OMB Approval; Survey of Single-Use Medical Device Reuse and Reprocessing in Hospitals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Survey of Single-Use Medical Device Reuse and Reprocessing in Hospitals'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

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

SUPPLEMENTARY INFORMATION: In the **Federal Register** of July 25, 2001 (66 FR 38713), 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 0910–0477. The approval expires on October 31, 2004. A copy of the supporting statement for this information collection is available on the Internet at http://www.fda.gov/ohrms/dockets.

Dated: October 12, 2001.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 01–26554 Filed 10–19–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N–1637]

Agency Information Collection Activities; Announcement of OMB Approval; Transmittal of Advertising and Promotional Labeling for Drugs and Biologics for Human Use

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Transmittal of Advertising and Promotional Labeling for Drugs and Biologics for Human Use'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Karen L. Nelson, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of December 21, 2000 (65 FR 80437), 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 0910–0376. The approval expires on October 31, 2004. A copy of the supporting statement for this information collection is available on the Internet at http://www.fda.gov/ohrms/dockets.

Dated: October 12, 2001.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 01–26555 Filed 10–19–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N–1682]

Agency Information Collection Activities; Announcement of OMB Approval; Radioactive Drug Research Committee

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

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

FOR FURTHER INFORMATION CONTACT: Karen L. Nelson, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of January 5, 2001 (66 FR 1137), 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 0910–0053. The approval expires on October 31, 2004. A copy of the supporting statement for this information collection is available on