• Documentation Requirements for Teaching Physicians

• Advanced Beneficiary Notice (provider education materials/carrier instructions).

For additional information and clarification on the topics listed, call the contact person in the ''For Further Information Contact'' section of this notice.

Individual physicians or medical organizations that represent physicians wishing to make 5-minute oral presentations on agenda issues should contact the Executive Director by 12 noon, *September 7, 2001,* to be scheduled. Testimony is limited to listed agenda issues only. The number of oral presentations may be limited by the time available. A written copy of the presenter's oral remarks should be submitted to the Executive Director no later than 12 noon, *September 7, 2001,* for distribution to Council members for review prior to the meeting. Physicians and organizations not scheduled to speak may also submit written comments to the Executive Director and Council members. The meeting is open to the public, but attendance is limited to the space available. Individuals requiring sign language interpretation for the hearing impaired or other special accommodation should contact John Lanigan at (202) 690–7418 at least 10 days before the meeting.

(Section 1868 of the Social Security Act (42 U.S.C. 1395ee) and section 10(a) of Public Law 92–463 (5 U.S.C. App. 2, section 10(a)); 45 C.F.R. part 11)

(Catalog of Federal Domestic Assistance Program No. 93.773, Medicare—Hospital Insurance; and Program No. 93.774, Medicare—Supplementary Medical Insurance Program)

Dated: August 28, 2001.

Thomas A. Scully,

Administrator, Centers for Medicare and Medicaid Services.

[FR Doc. 01–22065 Filed 8–30–01; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N–0581]

Agency Information Collection Activities; Announcement of OMB Approval; Requirements for Testing Human Blood Donors for Evidence of Infection Due to Communicable Disease Agents

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Requirements for Testing Human Blood Donors for Evidence of Infection Due to Communicable Disease Agents'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: JonnaLynn P. Capezzuto, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of June 11, 2001 (66 FR 31146), 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 0910–0472. The approval expires on August 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: August 24, 2001.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 01–22012 Filed 8–30–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01N–0231]

Agency Information Collection Activities; Proposed Collection; Comment Request; Veterinary Adverse Drug Reaction, Lack of Effectiveness, Product Defect Report; Reopening of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; reopening of comment period.

SUMMARY: The Food and Drug Administration (FDA) is reopening the comment period on a proposed collection of certain information by the agency until October 15, 2001. Under the Paperwork Reduction Act of 1995, Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information and to allow 60 days for public comment in response to the notice. This notice is reopening the comment period for a data collection effort to solicit comments on reporting and recordkeeping requirements obligating holders of approved new animal drug applications (NADAs) and abbreviated new animal drug applications (ANADAs) to submit information on adverse drug reactions, lack of effectiveness, and product defects.

DATES: Submit written or electronic comments on the collection of information by October 15, 2001.

ADDRESSES: Submit electronic comments on the collection of information via the internet at: http://www.accessdata.fda.gov/scripts/oc/dockets/edockethome.cfm. 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: Denver Presley, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1472.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of May 29, 2001 (66 FR 29141), FDA published a notice soliciting comments on reporting and recordkeeping requirements associated with 21 CFR part 510. To give interested persons additional time to submit comments on the proposed data collection, the agency is reopening the comment period until October 15, 2001.

Interested persons may submit to the Dockets Management Branch (address above) written comments by October 15, 2001. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

Dated: August 24, 2001.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 01–21967 Filed 8–30–01; 8:45 am]

BILLING CODE 4160–01–S