

STATUS: This meeting will be open to the public.

ITEMS TO BE DISCUSSED:

Correction and Approval of Minutes.
Draft Advisory Opinion 2000-24;
Alaska Democratic Party by counsel,
Neil Reiff.

Statements of Reasons—Requests to
Deny Certification of Public Funds to
Patrick J. Buchanan and Ezola Foster
(LRA#598/599).

Notice of Disposition of Petition for
Rulemaking Filed by the Project on
Government Oversight.

Administrative Matters.

PERSON TO CONTACT FOR INFORMATION:

Mr. Ron Harris, Press Officer,
Telephone: (202) 694-1220.

Mary W. Dove,

Acting Secretary of the Commission.

[FR Doc. 00-27654 Filed 10-24-00; 11:49
am]

BILLING CODE 6715-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Committee on Vital and Health Statistics; Meeting

Pursuant to the Federal Advisory
Committee Act, the Department of
Health and Human Services (HHS)
announces the following advisory
committee meeting.

Name: National Committee on Vital and
Health Statistics (NCVHS).

Time and Date: November 28, 2000—9
a.m.–5 p.m. EDT.

November 29, 2000—10:15 a.m.–3:30 p.m.
EDT.

Place: Hubert H. Humphrey Building, 200
Independence Avenue SW., Room 705A,
Washington, DC 20201.

Status: Open.

Purpose: At this meeting the Committee
will hear presentations and hold discussions
on several health data policy topics. On the
first day an update from HHS has been
scheduled on the implementation of the
administrative simplification provisions of
the Health Insurance Portability and
Accountability Act of 1996 (HIPAA). The
Committee will be briefed by the Director of
the National Center for Health Statistics on
several health data activities. In addition,
there may be a discussion of a possible draft
letter to the HHS Secretary regarding digital
signatures. The Committee will also discuss
action items reported in the summary from
its 50th Anniversary Symposium held earlier
in the year. There will also be a report on two
recent meetings of the World Health
Organization's (WHO) collaborating Center
for the Classification of Diseases. A panel
discussion has been scheduled on HIPAA
implementation issues. The first day will end
with breakout sessions for subcommittees
and workgroups. Day two will also begin
with breakout sessions and then the full

committee will be briefed on selected HHS
data policy initiatives and will hear an
analysis of State privacy laws. The afternoon
session will be devoted to hearing reports
from the subcommittees and workgroups and
the setting of future agendas.

Notice: In the interest of security, HHS has
instituted stringent procedures for entrance
to the Hubert H. Humphrey building by non-
government employees. Persons without a
government identification card may need to
have the guard call for an escort to the
meeting.

Contact Person for Core Information:

Substantive program information as well as
summaries of meetings and a roster of
committee members may be obtained from
Marjorie S. Greenberg, Executive Secretary,
NCVHS, National Center for Health Statistics,
Centers for Disease Control and Prevention,
Room 1100, Presidential Building, 6525
Belcrest Road, Hyattsville, Maryland 20782,
telephone (301) 458-4245. Information also
is available on the NCVHS home page of the
HHS website: <http://www.ncvhs.hhs.gov/>,
where further information including an
agenda will be posted when available.

Dated: October 18, 2000.

James Scanlon,

*Director, Division of Data Policy, Office of
the Assistant Secretary for Planning and
Evaluation.*

[FR Doc. 00-27462 Filed 10-25-00; 8:45 am]

BILLING CODE 4151-05-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N-1268]

Agency Information Collection Activities; Announcement of OMB Approval; Food Additives and Food Additive Petitions

AGENCY: Food and Drug Administration,
HHS.

ACTION: Notice.

SUMMARY: The Food and Drug
Administration (FDA) is announcing
that a collection of information entitled
“Food Additives and Food Additive
Petitions” 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 August 3, 2000 (65
FR 47736), 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 0910-0016. The
approval expires on October 31, 2003. A
copy of the supporting statement for this
information collection is available on
the Internet at <http://www.fda.gov/ohrms/dockets>.

Dated: October 20, 2000.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 00-27546 Filed 10-25-00; 8:45 am]

BILLING CODE 4160-01-F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N-1373]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Reporting and Recordkeeping Requirements for Mammography Facilities

AGENCY: Food and Drug Administration,
HHS.

ACTION: Notice.

SUMMARY: The Food and Drug
Administration (FDA) is announcing
that the proposed collection of
information listed below has been
submitted to the Office of Management
and Budget (OMB) for review and
clearance under the Paperwork
Reduction Act of 1995.

DATES: Submit written comments on the
collection of information by November
27, 2000.

ADDRESSES: Submit written comments
on the collection of information to the
Office of Information and Regulatory
Affairs, OMB, New Executive Office
Bldg., 725 17th St. NW., rm. 10235,
Washington, DC 20503, Attn: Wendy
Taylor, Desk Officer for FDA.

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
compliance with 44 U.S.C. 3507, FDA
has submitted the following proposed
collection of information to OMB for
review and clearance.