thereafter until the required divestiture is completed.

The purpose of this analysis is to facilitate public comment on the proposed Order, and it is not intended to constitute interpretation of the agreement and proposed Order or to modify in any way their terms.

Donald S. Clark,

Secretary.

[FR Doc. 97–33439 Filed 12–22–97; 8:45 am]

BILLING CODE 6750–01–M

GENERAL SERVICES ADMINISTRATION

Federal Acquisition Policy Division, FAR Secretariat Revision of Standard Forms

AGENCY: General Services Administration.

ACTION: Notice.

SUMMARY: The General Services Administration/FAR Secretariat has revised SF 1423, Inventory Verification Survey, SF 1426, Inventory Schedule A-Metals in Mill Product Form; SF 1428, Inventory Schedule B; SF 1430, Inventory Schedule C—(Work-In-Process); SF 1432, Inventory Schedule D—(Special Tooling and Special Test Equipment); SF 1434, Termination Inventory Schedule E (Short Form for Use With SF 1438 Only) to remove the need for particular certification requirements, and update the burden statement.

Since these forms are authorized for local reproduction, you can obtain new camera copy in three ways:

On the U.S. Government Management Policy CD–ROM;

On the internet. Address: http://www.gsa.gov/forms, or;

From CARM, Attn.: Barbara Williams, (202) 501–0581.

FOR FURTHER INFORMATION CONTACT:

FAR Secretariat, (202) 501–4755. This contact is for information on completing the form and interpreting the FAR only.

DATES: Effective December 23, 1997.

Dated: December 16, 1997.

Barbara M. Williams,

Deputy Standard and Optional Forms Management Officer.

[FR Doc. 97–33472 Filed 12–22–97; 8:45 am]

BILLING CODE 6820–34–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Safety and Occupational Health Study Section; NIOSH Meetings

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following committee meeting:

Name: Safety and Occupational Health Study Section, National Institute for Occupational Safety and Health (NIOSH).

Times and Dates: 8 a.m.–5:30 p.m., February 12, 1998. 8 a.m.–5:30 p.m., February 13, 1998.

Place: Old Town Alexandria Holiday Inn, 480 King Street, Alexandria, Virginia, 22314.

Status: Open business session, 8 a.m.–8:30 a.m., February 12, 1998; Closed evaluation sessions 8:30 a.m.–5:30 p.m., February 12, 1998; and 8 a.m.–5:30 p.m., February 13, 1998.

Purpose: The Safety and Occupational Health Study Section will review, discuss, and evaluate grant application(s) in response to the Institute's standard grants review and funding cycles pertaining to research issues in occupational safety and health and allied areas. It is the intent of NIOSH to support broad-based research endeavors in keeping with the Institute's program goals which will lead to improved understanding and appreciation for the magnitude of the aggregate health burden associated with occupational injuries and illnesses, as well as to support more focused research projects which will lead to improvements in the delivery of occupational safety and health services and the prevention of work-related injury and illness. It is anticipated that research funded will promote these program goals.

Matters to be discussed: The meeting will convene in open session from 8 a.m.–8:30 a.m. on February 12, 1998, to address matters related to the conduct of Study Section business. The meeting will proceed in closed session from 8:30 a.m. until scheduled adjournment (5:30 p.m.) on February 12, 1998. The meeting will continue in closed session from 8 a.m. until scheduled adjournment (5:30 p.m.) or earlier on February 13, 1998. The purpose of the closed sessions is for the Safety and Occupational Health Study Section to consider safety and occupational health related grant applications. These portions of the meeting will be closed to the public in accordance with provisions set forth in section 552(c) (4) and (6) title 5 U.S.C., and the Determination of the Associate Director for Management and Operations, CDC, pursuant to Pub. L. 92–463.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Pervis C. Major, Ph.D., Scientific Review Administrator, Office of Extramural Coordination and Special Projects, Office of the Director, NIOSH, 1095 Willowdale Road, Morgantown, West Virginia 26505. Telephone 304/285–5979.

Dated: December 17, 1997.

Carolyn J. Russell,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 97–33412 Filed 12–22–97; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97N–0510]

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 reinstatement of an existing collection of information, and allow 60 days for public comment in response to the notice. This notice solicits comments on the recordkeeping requirements for manufacturers of medicated animal feeds.

DATES: Submit written comments on the collection of information by February 23, 1998.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. 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: 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