number, OMB number, and OS document identifier, to *naomi.cook@hhs.gov*, or call the Reports Clearance Office on (202) 690–6162. Written comments and recommendations for the proposed information collections must be received within 30 days, and directed to the Desk Officer at the address below: OMB Desk Officer: John Kraemer, OMB Human Resources and Housing Branch, Attention: (OMB (0990–0269), New Executive Office Building, Room 10235, Washington DC 20503.

Dated: April 21, 2006.

Robert E. Polson,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.

[FR Doc. E6–6688 Filed 5–2–06; 8:45 am]

BILLING CODE 4153–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: OS–0990–New; 30-day notice]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Office of the Secretary, Office of Women's Health.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Type of Information Collection Request: Regular Clearance, New collection;

Title of Information Collection: National Women's Health Week (NWHW) Survey;

Form/OMB No.: OS–0990–New;

Use: The Office of Women's Health will evaluate how National Women's Health Week (NWHW) is implemented across the country, and to assess whether or not NWHW is reaching its program objectives, with an emphasis on messages, delivery mechanisms, levels of outreach and contact, and partnership strategies.

Frequency: Reporting, on occasion;

Affected Public: Business or other for-profit, not-for-profit institutions, Federal, State, local or tribal government;

Annual Number of Respondents: 1,400;

Total Annual Responses: 2,800;

Average Burden Per Response: 1 hour;

Total Annual Hours: 700.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access the HHS Web site address at *http://www.hhs.gov/ocio/infocollect/pending/* or e-mail your request, including your address, phone number, OMB number, and OS document identifier, to *naomi.cook@hhs.gov*, or call the Reports Clearance Office on (202) 690–6162. Written comments and recommendations for the proposed information collections must be received within 30 days, of this notice directly to the Desk Officer at the address below: OMB Desk Officer: John Kraemer, OMB Human Resources and Housing Branch, Attention: (OMB #0990–New), New Executive Office Building, Room 10235, Washington DC 20503.

Dated: April 21, 2006.

Robert E. Polson,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.

[FR Doc. E6–6689 Filed 5–2–06; 8:45 am]

BILLING CODE 4150–33–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Industry Exchange Workshop to Celebrate Food and Drug Administration Centennial: Past, Present, and Future of Regulated Food, Drugs, Nutritional Supplements, and Medical Devices; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

SUMMARY: The Food and Drug Administration (FDA) Philadelphia District, in cooperation with the Chemical Heritage Foundation (CHF), is announcing a workshop on past, current, and future issues and challenges in FDA regulation as part of the celebration of FDA's 100-year anniversary. Topics for discussion include: Turning points in FDA history; the impact of changes in science and technology on FDA regulation, regulation in the globalized economy, consumer access to information in the regulatory environment; and a risk-based approach to regulation as a model for the future. The purpose of this 1-day workshop for consumers, industry, academia, and regulators is to promote dialogue among regulators and these stakeholders.

Date and Time: The public workshop will be held on Tuesday, May 16, 2006, from 9 a.m. to 5 p.m.

Location: The public workshop will be held at the Chemical Heritage Foundation, 315 Chestnut St., Philadelphia, PA 19106, 215–873–8214, FAX: 215–629–5249.

Contact: Marie Falcone, Food and Drug Administration, U.S. Customhouse, 200 Chestnut St., rm. 900, Philadelphia, PA 19106, 215–717–3703, FAX: 215–597–5798, e-mail: *Marie.Falcone@fda.hhs.gov*.

Registration: Send registration information (including name, title, firm name, address, telephone, fax number) and the registration fee of $20.00 payable to the Chemical Heritage Foundation, 315 Chestnut St., Philadelphia, PA 19106. To register via the Internet go to *http://www.chemheritage.org/events/fda/index.html*. (FDA has verified the Web site address, but is not responsible for subsequent changes to the Web site after this document publishes in the **Federal Register**.) The registrar will also accept payment by major credit cards. The registration fee for FDA Philadelphia District employees is waived.

For more information on the meeting, or for questions about registration, contact the Chemical Heritage Foundation (CHF) at 215–873–8214, FAX: 215–629–5249, or via e-mail: *arthurd@chemheritage.org*.

Attendees are responsible for their own accommodations.

The registration fee will be used to offset the expenses of hosting the conference, including meals, refreshments, meeting rooms, and materials. Space is limited, therefore interested parties are encouraged to register early. Limited onsite registration may be available. Please arrive early to ensure prompt registration.

If you need special accommodations due to a disability, please contact Marie Falcone (see *Contact*) at least 7 days in advance of the workshop.

SUPPLEMENTARY INFORMATION: The ''FDA Past, Present, and Future of Regulating Food, Drugs, Medical Devices, and

Nutritional Supplements'' workshop helps fulfill the Department of Health and Human Services' and FDA's important mission to protect the public health by encouraging informed dialogue on the future direction of FDA regulation in the context of its historical accomplishments.

The workshop will also help to implement the objectives of section 406 of the FDA Modernization Act (21 U.S.C. 393) and the FDA Plan for Statutory Compliance, which include working more closely with stakeholders and providing access to scientific and technical expertise. Finally, the workshop furthers the goals of the Small Business Regulatory Enforcement Fairness Act (Public Law 104–121) by providing outreach activities by Government agencies directed to small businesses.

Dated: April 28, 2006.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 06–4185 Filed 5–1–06; 10:37 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[FDA 225–06–8400]

Memorandum of Understanding Between the Food and Drug Administration, United States Department of Health and Human Services, the Animal and Plant Health Inspection Service, the United States Department of Agriculture, and The National Institutes of Health, United States Department of Health and Human Services Concerning Laboratory Animal Welfare

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The purpose of this Memorandum of Understanding (MOU) is to set forth an agreement between the Animal and Plant Health Inspection Service, U.S. Department of Agriculture, and the National Institutes of Health, U.S. Department of Health and Human Services concerning Laboratory Animal Welfare and FDA (collectively ''the Parties'', or individually as a ''Party'') regarding the framework for reciprocal cooperation which will assist each agency in meeting its responsibilities in promoting proper laboratory animal care and welfare. This MOU replaces 225–83–8400.

DATES: The agreement became effective February 14, 2006.

FOR FURTHER INFORMATION CONTACT:

For FDA: Rodney T. Allnut, Office of Enforcement, Office of Regulatory Affairs, Food and Drug Administration, HFC–230, rm. 126, 15800 Crabbs Branch, Rockville, MD 20855, 240–632–6848, FAX: 240–632–6861.

For USDA: Chester Gipson, Animal and Plant Health Inspection Service, U.S. Department of Agriculture, 4700 River Rd., Unit 97, rm. 6A–16, Riverdale, MD 20737–1234, 301–734–4980, FAX: 301–734–4993.

For NIH: Carol Wigglesworth, Office of Laboratory Animal Welfare, Office of Extramural Research, National Institutes of Health, Rockwall I, suite 1050, MSC 7982, 6705 Rockledge Dr., Bethesda, MD 20892–7982, 301–496–7163, FAX: 301–402–2803.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 20.108(c), which states that all written agreements and MOUs between FDA and others shall be published in the **Federal Register**, the agency is publishing notice of this MOU.

Dated: April 26, 2006.

Jeffrey Shuren,

Assistant Commissioner for Policy.

BILLING CODE 4160–01–S