

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]

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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.

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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