garments introduced per year that require new or revised care instructions. Staff estimates the burden of determining care instructions to be 43 hours each year per respondent, for a cumulative total of 1,145,821 hours. Staff further estimates that the burden of drafting and ordering labels is 2 hours each year per respondent, for a total of 53,294 hours. Staff believes that the process of attaching labels is fully automated and integrated into other production steps for about 40 percent of the approximately 19.1 billion garments that are required to have care instructions on permanent labels.[12] For the remaining 11.46 billion items (60 percent of 19.1 billion), the process is semi-automated and requires an average of approximately two seconds per item, for a total of 6,366,667 hours per year. Thus, the total estimated annual burden for all respondents is 7,565,782 hours (1,145,821 hours to determine care instructions + 53,294 hours to draft and order labels + 6,366,667 hours to attach labels).

Estimated annual cost burden: $61,410,000[13] , rounded to the nearest thousand (solely relating to labor costs).

Task	Hourly Rate	Burden Hours	Labor Cost
Determine care instructions	$22.00	1,145,821	$25,208,062
Draft and order labels	$16.27	53,294	$867,093
Attach labels	$5.55 [14]	6,366,667	$35,335,002
TOTAL			$61,410,157

[14] *See* note 6.

Staff believes that there are no current start-up costs or other capital costs associated with the Rule. Because the labeling of textile products has been an integral part of the manufacturing process for decades, manufacturers have in place the capital equipment necessary to comply with the Rule's labeling requirements. Based on knowledge of the industry, staff believes that much of the information required by the Rule would be included on the product label even absent those requirements.

David C. Shonka,

Acting General Counsel.

[FR Doc. E9–3056 Filed 2–11–09: 8:45 am]

[Billing code: 6750–01–S]

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0281]

National Capital Region (NCR), Office of Childcare Services; Information Collection; General Services Administration (GSA) Child Care Specialist Feedback Form

AGENCY: NCR Office of Childcare Services, Public Buildings Service (PBS), GSA.

ACTION: Notice of request for comments regarding an extension to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the General Services Administration will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a currently approved information collection requirement. This information will be used to assess satisfaction with services delivered by staff from the Office of Child Care Services. The respondents are current users of the Office of Child Care Services. The OMB clearance currently expires on April 30, 2009.

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate and based on valid assumptions and methodology; and ways to enhance the quality, utility, and clarity of the information to be collected.

DATES: Submit comments on or before: April 13, 2009.

FOR FURTHER INFORMATION CONTACT: Leo G. Bonner, Regional Child Care Coordinator, Office of Child Care Services, at telephone (202) 401–7403 or via e-mail to *leo.bonner@gsa.gov*.

ADDRESSES: Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to the Regulatory Secretariat (VPR), General Services Administration, Room 4041, 1800 F Street, NW., Washington, DC 20405. Please cite OMB Control No. 3090–0281, General Services Administration (GSA) Child Care Specialist Feedback Form, in all correspondence.

SUPPLEMENTARY INFORMATION:

A. Purpose

This information will be used to assess consumer satisfaction with services delivered by staff from the Office of Child Care services.

B. Annual Reporting Burden

Respondents: 144.
Responses Per Respondent: 1.
Hours Per Response: .083 (5 minutes).
Total Burden Hours: 12.
OBTAINING COPIES OF PROPOSALS: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (VPR), 1800 F Street, NW., Room 4041, Washington, DC 20405, telephone (202) 501–4755. Please cite OMB Control No. 3090–0281, General Services Administration (GSA) Child Care Specialist Feedback Form, in all correspondence.

Dated: January 30, 2009.

Casey Coleman,

Chief Information Officer.

[FR Doc. E9–2945 Filed 2–11–09; 8:45 am]

BILLING CODE 6820–A4–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Advisory Committee on Minority Health

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of Public Health and Science, Office of Minority Health.

ACTION: Notice: correction.

SUMMARY: The Department of Health and Human Services published a notice in the **Federal Register** of February 4, 2009 announcing a February 24, 2009 meeting of the Advisory Committee on Minority Health. It was announced that this meeting would be held at The Westin National Harbor, 171 Waterfront Street, Oxon Hill, MD. Due to unforseen

[12] About 1 billion of the 20.1 billion garments produced annually are either not covered by the Care Labeling Rule (gloves, hats, caps, and leather, fur, plastic, or leather garments) or are subject to an exemption that allows care instructions to appear on packaging (hosiery).

[13] We have corrected an error in this calculation that appeared in the prior 60-day Federal Register notice.

circumstances the location of the meeting has been changed.

FOR FURTHER INFORMATION CONTACT: Ms. Monica A. Baltimore, Tower Building, 1101 Wootton Parkway, Suite 600, Rockville, Maryland 20852. Phone: 240–453–2882 Fax: 240–453–2883.

Correction

In the **Federal Register** of February 4, 2009, Vol. 74, No. 22, on page 6041, in the 2nd column, correct the **ADDRESSES** caption to read:

The meeting will be held at The Gaylord National and Convention Center, Annapolis Rooms 1 & 2, 201 Waterfront Street (National Harbor), Oxon Hill, MD 20745.

Dated: February 9, 2009.

Mirtha R. Beadle,

Deputy Director, Office of Minority Health, Office of Public Health and Science, Office of the Secretary, U.S. Department of Health and Human Services.

[FR Doc. E9–3014 Filed 2–11–09; 8:45 am]

BILLING CODE 4150–29–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Secretary's Advisory Committee on Human Research Protections

AGENCY: Department of Health and Human Services, Office of the Secretary.

ACTION: Notice.

SUMMARY: Pursuant to Section 10(a) of the Federal Advisory Committee Act, U.S.C. Appendix 2, notice is hereby given that the Secretary's Advisory Committee on Human Research Protections (SACHRP) will hold its nineteenth meeting. The meeting will be open to the public.

DATES: The meeting will be held on Tuesday, March 3, 2009 from 8:30 a.m. until 5 p.m. and Wednesday, March 4, 2009 from 8:30 a.m. until 5 p.m.

ADDRESSES: The Sheraton National Hotel, 900 South Orme Street, Arlington, Virginia 22204. Phone: 703–521–1900.

FOR FURTHER INFORMATION CONTACT: Jerry Menikoff, J.D., M.D., Director, Office for Human Research Protections (OHRP), or Julia Gorey, J.D., Executive Director, SACHRP; U.S. Department of Health and Human Services, 1101 Wootton Parkway, Suite 200, Rockville, Maryland 20852; 240–453–8141; fax: 240–453–6909; e-mail address: *sachrp@osophs.dhhs.gov*.

SUPPLEMENTARY INFORMATION: Under the authority of 42 U.S.C. 217a, Section 222 of the Public Health Service Act, as amended, SACHRP was established to provide expert advice and recommendations to the Secretary of Health and Human Services and the Assistant Secretary for Health on issues and topics pertaining to or associated with the protection of human research subjects.

On March 3, 2009, SACHRP will receive and discuss a report from an internal task force charged with prioritizing SACHRP's existing recommendations to OHRP. The Committee will then hear a presentation of the recent National Academy of Sciences report entitled ''Health Research and the Privacy of Health Information—The HIPAA Privacy Rule,'' followed by a presentation of the Association of Academic Health Centers' recent survey on the impact of the HIPAA Privacy Rule on research. Lastly, SACHRP will hear a report from the Subpart A Subcommittee, which is charged with developing recommendations for consideration by SACHRP about the application of Subpart A of 45 CFR part 46 in the current research environment. This subcommittee was established by SACHRP at its October 4–5, 2004 meeting.

On March 4, 2009, the Committee will receive and discuss a report from the Subcommittee on Inclusion of Individuals with Impaired Decision-Making in Research. That subcommittee is charged with developing recommendations for consideration by SACHRP about whether guidance or additional regulations are needed for research involving individuals with impaired decision-making capacity. It was formed as a result of discussions during the July 31–August 1, 2006 SACHRP meeting. The day will conclude with a panel discussion addressing harmonization issues associated with the Common Rule and the FDA regulations.

Public attendance at the meeting is limited to space available. Individuals who plan to attend the meeting and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact persons. Members of the public will have the opportunity to provide comments on both days of the meeting. Public comment will be limited to five minutes per speaker. Any members of the public who wish to have printed materials distributed to SACHRP members for this scheduled meeting should submit materials to the Executive Director, SACHRP, prior to the close of business Friday, February 27, 2009. Information about SACHRP and the draft meeting agenda will be posted on the SACHRP Web site at: *http://www.hhs.gov/ohrp/sachrp/index.html*.

Dated: February 6, 2009.

Jerry Menikoff,

Director, Office for Human Research Protections, Executive Secretary, Secretary's Advisory Committee on Human Research Protections.

[FR Doc. E9–3015 Filed 2–11–09; 8:45 am]

BILLING CODE 4150–36–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–09–08BF]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an e-mail to *omb@cdc.gov*. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–6974. Written comments should be received within 30 days of this notice.

Proposed Project

Evaluation Models to Assess Patient Perspectives on Opt-out HIV Testing in Clinical Settings—New—National Center for HIV, Viral Hepatitis, STD and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In 2006, CDC published the *Revised Recommendations for HIV Testing of Adults, Adolescents and Pregnant Women in Health Care Settings* which recommends routine, opt-out HIV testing to persons 13–64 years of age in health care settings. The goal of this project is to develop evaluation models for health care providers in a variety of settings to independently assess the effect that expanded HIV screening activities have on patient attitudes toward and acceptance of HIV testing.

The evaluation models will be packaged into a toolkit containing educational materials, administrative tools and a model questionnaire to measure patients' perceptions of their ability to decline testing, the sufficiency and effectiveness of methods used to