proper care instructions may vary greatly among firms, primarily based on the number of different lines of textile garments introduced per year that require new or revised care instructions. Staff estimates the burden of determining care instructions to be 100 hours each year per firm, for a cumulative total of 2,264,200 hours. Staff further estimates that the burden of drafting and ordering labels is 80 hours each year per firm, for a total of 1,811,360 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 18.4 billion garments that are required to have care instructions on permanent labels.[12] For the remaining 11.04 billion items (60 percent of 18.4 billion), the process is semi-automated and requires an average of approximately ten seconds per item, for a total of 30,666,667 hours per year. Thus, the total estimated annual burden for all firms is 34,742,226 hours (2,264,200 hours to determine care instructions + 1,811,360 hours to draft and order labels + 30,666,666 hours to attach labels).

Estimated annual cost burden: $258,329,000, rounded to the nearest thousand (solely relating to labor costs). The chart below summarizes the total estimated costs.

Task	Hourly rate	Burden hours	Labor cost
Determine care instructions	$ 26.00	2,264,200	$58,869,200
Draft and order labels	17.00	1,811,360	30,793,120
Attach labels	[13] 5.50	30,666,667	168,666,669
Total			258,328,989

Staff believes that there are no current start-up costs or other capital costs associated with the Care Labeling 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.

Request for Comments

You can file a comment online or on paper. Write ''Apparel Rules: FTC File No. P074201'' on your comment. Your comment—including your name and your state—will be placed on the public record of this proceeding, including, to the extent practicable, on the public Commission Web site, at *http://www.ftc.gov/os/publiccomments.shtm.* As a matter of discretion, the Commission tries to remove individuals' home contact information from comments before placing them on the Commission Web site. Because your comment will be made public, you are solely responsible for making sure that your comment does not include any sensitive personal information, such as a Social Security number, date of birth, driver's license number or other state identification number or foreign country equivalent, passport number, financial account number, or credit or debit card number. You are also solely responsible for making sure that your comment does not include any sensitive health information, such as medical records or other individually identifiable health information. In addition, do not include any ''[t]rade secret or any commercial or financial information which is . . . privileged or confidential,'' as discussed in Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2). In particular, do not include competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

If you want the Commission to give your comment confidential treatment, you must file it in paper form, with a request for confidential treatment, and you must follow the procedure explained in FTC Rule 4.9(c), 16 CFR 4.9(c). Your comment will be kept confidential only if the FTC General Counsel, in his or her sole discretion, grants your request in accordance with the law and the public interest. Postal mail addressed to the Commission is subject to delay due to heightened security screening. As a result, the Commission encourages you to submit your comments online. To make sure that the Commission considers your online comment, you must file it at *https://ftcpublic.commentworks.com/ftc/apparelrulespra* by following the instructions on the Web-based form. If this Notice appears at *http://www.regulations.gov,* you also may file a comment through that Web site.

If you file your comment on paper, write ''Apparel Rules: FTC File No. P074201'' on your comment and on the envelope, and mail it to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW., Suite CC–5610, (Annex J), Washington, DC 20580, or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW., 5th Floor, Suite 5610, (Annex J), Washington, DC 20024. If possible, submit your paper comment to the Commission by courier or overnight service.

The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before March 10, 2015. You can find more information, including routine uses permitted by the Privacy Act, in the Commission's privacy policy, at *http://www.ftc.gov/ftc/privacy.htm.*

David C. Shonka,

Principal Deputy General Counsel.

[FR Doc. 2015–00166 Filed 1–8–15; 8:45 am]

BILLING CODE 6750–01–P

[12] About 1 billion of the 19.4 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] *See* note 3.

GENERAL SERVICES ADMINISTRATION

[Notice–FAS–2014–02; Docket: 2014–0002; Sequence: 39]

Notice of Public Meeting for the Supplemental Draft Environmental Impact Statement for the U.S. Department of State Foreign Affairs Security Training Center in Nottoway County, Virginia

AGENCY: U.S. General Services Administration (GSA).

ACTION: Meeting Notice.

SUMMARY: Pursuant to the Council on Environmental Quality regulations implementing the procedural provisions of the National Environmental Policy Act, the U.S. General Services Administration (GSA) has prepared and filed with the U.S. Environmental Protection Agency (EPA) a Supplement to the 2012 Draft Environmental Impact Statement (EIS) for the U.S. Department of State (DOS), Bureau of Diplomatic Security, Foreign Affairs Security Training Center (FASTC). GSA is the lead agency; cooperating agencies are DOS, U.S. Army Corps of Engineers (ACE), EPA, and National Guard Bureau (NGB). The Supplemental Draft EIS was prepared to evaluate the environmental impacts of site acquisition and development of FASTC at the Virginia Army National Guard Maneuver Training Center (VA ANG MTC) at Fort Pickett and Nottoway County's Local Redevelopment Authority area in Nottoway County, Virginia.

DATES: *Comment date:* The public may submit comments on the Supplemental Draft EIS during a 45-day public review and comment period beginning January 9, 2015 with publication of this notice and ending on February 23, 2015. Instructions for submitting comments may be found under the heading **SUPPLEMENTARY INFORMATION** in this notice.

Public Meeting: A public information meeting is scheduled for January 26, 2015, between 7:00 p.m. and 8:00 p.m., Eastern Standard Time, in the Auditorium at the Blackstone Conference and Retreat Center located at 707 Fourth Street, Blackstone VA 23824.

ADDRESSES: Send written comments by email to *FASTC.info@gsa.gov,* or U.S. Postal Service to: Ms. Abigail Low, GSA Project Manager, 20 N. 8th Street, Philadelphia, PA 19107.

FOR FURTHER INFORMATION CONTACT: Abigail Low, GSA Project Manager; 20 N. 8th Street, Philadelphia, PA 19107 215–446–4815, or email at *FASTC.info@gsa.gov.*

SUPPLEMENTARY INFORMATION:

Background: The purpose of the proposed FASTC in Nottoway County is to consolidate existing dispersed hard skills training functions into a single suitable location to improve training efficiency and enhance training operations. The proposed FASTC is needed to meet the increased demand for well-trained security personnel. A Notice of Intent to prepare a Supplemental Draft EIS was published in the **Federal Register** at 79 FR 52336 on September 3, 2014. A public scoping period and public scoping meeting for the proposed action were held in October 2011 in relation to the 2012 Draft EIS. Additionally, the public was invited to submit comments concerning the proposal for 30 days from publication of the September 3, 2014 Notice of Intent to prepare a Supplemental Draft EIS.

The alternatives fully evaluated in the Supplemental Draft EIS include the No Action Alternative and Build Alternative 3. Build Alternative 3 was developed based on the 2014 Master Plan Update that incorporates the adjustments in the FASTC program. The Preferred Alternative is Build Alternative 3.

The Supplemental Draft EIS incorporates by reference and builds upon the analyses presented in the 2012 Draft EIS and documents the Section 106 process under the National Historic Preservation Act of 1966, as amended (36 CFR part 800). The Supplemental Draft EIS addresses substantial changes to the proposed action that are relevant to environmental concerns and assesses any new circumstances or information relevant to potential environmental impacts.

In early 2013, all efforts on the Final EIS and work on the proposed site at Fort Pickett and Nottoway County's LRA area were put on hold pending additional due diligence and reviews at an existing federal training site in Georgia. As part of this due diligence effort, DOS conducted site visits to the Federal Law Enforcement Training Center (FLETC) in Glynco, Georgia. During this time period, DOS also assessed the scope and size of the FASTC project and determined a smaller platform was more fiscally prudent.

In April 2014, the earlier DOS selection of the proposed site for FASTC at Fort Pickett and Nottoway County was reaffirmed. Planning for the site resumed based on a reduced scope of requirements compared with the 2012 plan. The project would now proceed as a hard skills only facility, which consists of driving tracks, firing ranges, a mock urban environment, explosives ranges, and associated classrooms and administrative functions. Several hard skills training venues have been consolidated. Soft skills training, such as simulation labs, and life support functions, such as dormitories and dining facilities have been eliminated from the program. More information on the proposed FASTC program is available at *http://www.state.gov/recovery/fastc.*

The Supplemental Draft EIS has been distributed to various federal, state, and local agencies. The Supplemental Draft EIS is available for review on the project Web site *http://www.state.gov/recovery/fastc.* A printed copy of the Supplemental Draft EIS is available for viewing at the following libraries:

- Nottoway County Library—Louis Spencer Epes Memorial Library, 415 South Main St., Blackstone, VA
- Amelia County—James L. Hamner Public Library, 16351 Dunn Street, Amelia, VA
- Brunswick County—Brunswick County Library, 133 W. Hicks Street, Lawrenceville, VA
- Dinwiddie County—Dinwiddie Library, 14103 Boydton Plank Road, Dinwiddie, VA
- Lunenburg County—Ripberger Library, 117 South Broad St., Kenbridge, VA
- Prince Edward County—Farmville—Prince Edward Community Library, 1303 West 3rd Street, Farmville, VA
- Chesterfield County—Central Library, 9501 Lori Road, Chesterfield, VA
- Mecklenburg County—Southside Regional Library, 316 Washington Street, Boydton, VA

Federal, state, and local agencies, and other interested parties, are invited and encouraged to be present or represented at the public meeting on January 26, 2015. All formal comments will become part of the public record and substantive comments will be responded to in the Final EIS.

Public Comments: Comments on the Supplemental Draft EIS can be submitted three ways: (1) Submit comments via the FASTC email address: *FASTC.info@gsa.gov,* (2) provide written comments during the public meeting, or (3) mail a comment form or letter to: Ms. Abigail Low, GSA Project Manager, 20 N. 8th Street, Philadelphia, PA 19107. Written comments postmarked by February 23, 2015 will become part of the official public record.

Public Meeting: An informational presentation will be followed by an informal question and answer session. Informational posters will be on display, and representatives from GSA and DOS will be available to explain the proposed project, answer questions, and receive comments from the public.

Comment forms will be available for the public to provide formal written comments.

Dated: December 24, 2014.

Myles Vaughan,

NEPA Program Manager, Facilities Management & Services Programs Division, U.S. GSA, Mid-Atlantic Region.

[FR Doc. 2014–30791 Filed 1–8–15; 8:45 am]

BILLING CODE 6820–89–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2015–0001]

Proposed Revised Vaccine Information Materials for Multiple Pediatric Vaccines (''Your Baby's First Vaccines'')

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: Under the National Childhood Vaccine Injury Act (NCVIA) (42 U.S.C. 300aa–26), the Centers for Disease Control and Prevention (CDC) within the Department of Health and Human Services (HHS) develops vaccine information materials that all health care providers are required to give to patients/parents prior to administration of specific vaccines. HHS/CDC seeks written comment on the proposed updated vaccine information statement for multiple pediatric vaccines.

DATES: Written comments must be received on or before March 10, 2015.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2015–0001, by any of the following methods:

- Federal eRulemaking Portal: *http://www.regulations.gov*. Follow the instructions for submitting comments.
- Mail: Written comments should be addressed to Suzanne Johnson-DeLeon (*msj1@cdc.gov*), National Center for Immunization and Respiratory Diseases, Centers for Disease Control and Prevention, Mailstop A–19, 1600 Clifton Road NE., Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and docket number. All relevant comments received will be posted without change to *http://www.regulations.gov,* including any personal information provided. For access to the docket to read background documents or comments received, go to *http://www.regulations.gov*.

FOR FURTHER INFORMATION CONTACT: Skip Wolfe (*crw4@cdc.gov*), National Center for Immunization and Respiratory Diseases, Centers for Disease Control and Prevention, Mailstop A–19, 1600 Clifton Road NE., Atlanta, Georgia 30329.

SUPPLEMENTARY INFORMATION: The National Childhood Vaccine Injury Act of 1986 (Pub. L. 99–660), as amended by section 708 of Public Law 103–183, added section 2126 to the Public Health Service Act. Section 2126, codified at 42 U.S.C. 300aa–26, requires the Secretary of Health and Human Services to develop and disseminate vaccine information materials for distribution by all health care providers in the United States to any patient (or to the parent or legal representative in the case of a child) receiving vaccines covered under the National Vaccine Injury Compensation Program (VICP).

Development and revision of the vaccine information materials, also known as Vaccine Information Statements (VIS), have been delegated by the Secretary to the Centers for Disease Control and Prevention (CDC). Section 2126 requires that the materials be developed, or revised, after notice to the public, with a 60-day comment period, and in consultation with the Advisory Commission on Childhood Vaccines, appropriate health care provider and parent organizations, and the Food and Drug Administration. The law also requires that the information contained in the materials be based on available data and information, be presented in understandable terms, and include:

(1) A concise description of the benefits of the vaccine,

(2) A concise description of the risks associated with the vaccine,

(3) A statement of the availability of the National Vaccine Injury Compensation Program, and

(4) Such other relevant information as may be determined by the Secretary.

The vaccines initially covered under the National Vaccine Injury Compensation Program were diphtheria, tetanus, pertussis, measles, mumps, rubella and poliomyelitis vaccines. Since April 15, 1992, any health care provider in the United States who intends to administer one of these covered vaccines is required to provide copies of the relevant vaccine information materials prior to administration of any of these vaccines. Since then, the following vaccines have been added to the National Vaccine Injury Compensation Program, requiring use of vaccine information materials for them as well: Hepatitis B, *Haemophilus influenzae* type b (Hib), varicella (chickenpox), pneumococcal conjugate, rotavirus, hepatitis A, meningococcal, human papillomavirus (HPV), and seasonal influenza vaccines. Instructions for use of the vaccine information materials are found on the CDC Web site at: *http://www.cdc.gov/vaccines/hcp/vis/index.html.* Copies of the Vaccine Information Statements are found in the docket at *www.regulations.gov* (Docket CDC–2015–0001) under ''Supporting and Related Materials.''

HHS/CDC is proposing an updated version of the multiple pediatric vaccines (''Your Baby's First Vaccines'') vaccine information statement.

The vaccine information materials referenced in this notice are being developed in consultation with the Advisory Commission on Childhood Vaccines, the Food and Drug Administration, and parent and health care provider groups.

We invite written comment on the proposed vaccine information material entitled ''Your Baby's First Vaccines: What You Need to Know.'' A copy of the proposed vaccine information statement is available at *www.regulations.gov* (see Docket ID CDC–2015–0001). Comments submitted will be considered in finalizing these materials. When the final materials are published in the **Federal Register**, the notice will include an effective date for their mandatory use.

Dated: January 5, 2015.

Ron A. Otten,

Acting Deputy Associate Director for Science, Centers for Disease Control and Prevention.

[FR Doc. 2015–00171 Filed 1–8–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Statement of Organization, Functions, and Delegations of Authority

Part C (Centers for Disease Control and Prevention) of the Statement of Organization, Functions, and Delegations of Authority of the Department of Health and Human Services (45 FR 67772–76, dated October 14, 1980, and corrected at 45 FR 69296, October 20, 1980, as amended most recently at 79 FR 21760–21763, dated April 17, 2014) is amended to reflect the reorganization of the Office of Safety, Security and Asset Management.

Section C–B, Organization and Functions, is hereby amended as follows:

Delete items (1), (2) and (3) of the functional statement for the *Office of Operations (CAJ13), Office of the*