ensuring Sabal Trail's ability to negotiate additional Transco expansions. First, the proposed Order incorporates the capacity lease agreement between Transco and Sabal Trail, which reflects terms Transco and Sabal Trail reached when an independent and motivated commercial partner owned Transco. The proposed Order gives Sabal Trail additional flexibility and optionality in obtaining the phased capacity expansions already contemplated by the capacity lease agreement. The proposed Order terminates twelve years after it issues, in order to cover the entirety of ETE's obligations for the expansions currently outlined in the capacity lease agreement.

Second, the Order requires that, within one year of the closing of the Acquisition, ETE offer to amend the capacity lease agreement to allow Sabal Trail to request expansions for as long as an additional eight years after the last expansion currently in the capacity lease agreement. These provisions ensure that Sabal Trail has the same future expansion opportunities as would have existed if an independent Williams continued to own Transco.

ETE must offer future expansions on the same terms and conditions that Transco negotiated as an independent entity. For each requested expansion, ETE must inform Sabal Trail of the estimated expansion cost, using the same methodology for each that Transco uses in its normal course of business. ETE then is obligated to expand Transco as requested by Sabal Trail. However, to prevent Sabal Trail from requesting cost-prohibitive expansions—expansions that an independent Williams would not have agreed to—ETE retains the right to require Sabal Trail to pay for the capital costs of the expansion, in which case ETE would not charge Sabal Trail a lease fee for that particular expanded capacity.

The proposed Order does not obligate ETE to expand Transco if Sabal Trail does not have (or has not secured pre-construction commitments from shippers for) sufficient capacity to use the expansion to serve Florida. The Acquisition does not change the incentives of Transco's owner to deny capacity expansions to serve areas outside of Florida. Thus, without this limitation, the proposed Order could give Sabal Trail expansion rights it would have been unable to negotiate from an independent Transco.

The Commission does not intend this analysis to constitute an official interpretation of the proposed Order or to modify its terms in any way.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 2016–14092 Filed 6–14–16; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0056; Docket 2016–0053; Sequence 23]

Information Collection; Report of Shipment

AGENCY: Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

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

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement concerning report of shipment.

DATES: Submit comments on or before August 15, 2016.

ADDRESSES: Submit comments identified by Information Collection 9000–0056, Report of Shipment, by any of the following methods:

- *Regulations.gov: http://www.regulations.gov.*

Submit comments via the Federal eRulemaking portal by searching the OMB control number. Select the link ''Submit a Comment'' that corresponds with ''Information Collection 9000–0056, Report of Shipment''. Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 9000–0056, Report of Shipment'' on your attached document.

- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0056, Report of Shipment.

Instructions: Please submit comments only and cite Information Collection 9000–0056, Report of Shipment, in all correspondence related to this collection. Comments received generally will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *www.regulations.gov,* approximately two to three days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

FOR FURTHER INFORMATION CONTACT: Mr. Curtis E. Glover, Sr., Procurement Analyst, Office of Acquisition Policy, by telephone at 202–501–1448 or *curtis.glover@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

Per FAR 47.208, military (and, as required, civilian agency) storage and distribution points, depots, and other receiving activities require advance notice of shipments en-route from contractors' plants. Generally, this notification is required only for classified material; sensitive, controlled, and certain other protected material; explosives, and some other hazardous materials; selected shipments requiring movement control; or minimum carload or truckload shipments. It facilitates arrangements for transportation control, labor, space, and use of materials handling equipment at destination. Also, timely receipt of notices by the consignee transportation office precludes the incurring of demurrage and vehicle detention charges. Unless otherwise directed by a contracting officer, a contractor shall send the notice to the consignee transportation office at least twenty-four hours before the arrival of the shipment.

B. Annual Reporting Burden

Respondents: 33.
Responses per Respondent: 303.
Annual Responses: 9,999.
Hours per Response: .167.
Total Burden Hours: 1,670.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary; 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; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street, Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0056, Report of Shipment, in all correspondence.

Dated: June 9, 2016.

Lorin S. Curit,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2016–14119 Filed 6–14–16; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0853]

Tobacco Product Manufacturing Facility Visits

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA), Center for Tobacco Products (CTP) is announcing an invitation for participation in its Tobacco Product Manufacturing Facility Visits. This program is intended to give FDA staff an opportunity to visit facilities involved in the manufacturing of newly deemed tobacco products and their components and parts, including any related laboratory testing, and to observe the manufacturing operations of the tobacco industry. The purpose of this document is to invite parties interested in participating in Tobacco Product Manufacturing Facility Visits to submit requests to CTP.

DATES: Submit either an electronic or written request for participation by August 15, 2016. See section IV of this document for information on requests for participation.

ADDRESSES: If your facility is interested in participating in Tobacco Product Manufacturing Facility Visits, please submit a request either electronically to *http://www.regulations.gov* or in writing to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Matthew Brenner, Center for Tobacco Products, Food and Drug Administration, Document Control Center, Bldg. 71, Rm. G335, 10903 New Hampshire Ave., Silver Spring, MD 20993–0002, 877–287–1373, email: *CTPRegulations@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

On June 22, 2009, the Family Smoking Prevention and Tobacco Control Act (Pub. L. 111–31; 123 Stat. 1776) was signed into law, amending the Federal Food, Drug, and Cosmetic Act (the FD&C Act) and giving FDA authority to regulate tobacco product manufacturing, distribution, and marketing. The new provisions include, among other things, the authority to issue regulations related to tobacco product manufacturing practice in order to protect the public health and to assure that tobacco products are in compliance with the FD&C Act. Specifically, section 906(e) of the FD&C Act (21 U.S.C. 387f(e)) provides that in applying manufacturing restrictions to tobacco, the Secretary shall prescribe regulations (which may differ based on the type of tobacco product involved) requiring that the methods used in, and the facilities and controls used for, the manufacture, preproduction design validation (including a process to assess the performance of a tobacco product), packing, and storage of a tobacco product conform to current good manufacturing practice, or hazard analysis and critical control point methodology.

CTP is instituting Tobacco Product Manufacturing Facility Visits to provide FDA staff with the opportunity to:

- Observe tobacco product manufacturing operations—from the receipt of raw materials to the distribution of newly deemed tobacco products, and
- Learn about the manufacturing practices and processes unique to your facility and newly deemed tobacco products.

This program will also inform FDA staff as they implement the tobacco provisions of the FD&C Act.

II. Description of the Tobacco Product Manufacturing Facility Visits

In this program, groups of FDA staff plan to observe the following facilities and their operations:

- Manufacturing facilities, including establishments that process, package, label, and distribute different types of newly deemed tobacco products (*e.g.,* dissolvable products, gels, cigars, pipe tobacco, waterpipe tobacco products, and electronic nicotine delivery systems (ENDS) (including e-cigarettes, e-hookah, e-cigars, vape pens, advanced refillable personal vaporizers, and electronic pipes) and liquid nicotine and flavors) (see 81 FR 28973, May 10, 2016),
- Laboratory facilities that perform tobacco testing (whether third-party or in-house), and
- Manufacturing facilities for tobacco products for further manufacturing into finished tobacco products (including, but not limited to, components, parts, flavors, casings, e-liquids).

Please note that Tobacco Product Manufacturing Facility Visits are not intended to include or replace official FDA inspections of facilities to determine compliance with the FD&C Act; rather, these facility visits are meant to educate FDA staff and improve their understanding of the tobacco industry and its manufacturing operations.

III. Site Selection

CTP plans to select sites from one or more of each of the following categories:

- Dissolvable products,
- Gels,
- Cigars,
- Pipe tobacco,
- Waterpipe tobacco products,
- ENDS (including e-cigarettes, e-hookah, e-cigars, vape pens, advanced refillable personal vaporizers, and electronic pipes) and liquid nicotine and flavors,
- Tobacco laboratories,
- Importers of finished tobacco products,
- Distributors and wholesalers of regulated tobacco products, and/or
- Manufacturers of tobacco products for further manufacturing into finished tobacco products (including, but not limited to, components, parts, flavors, casings, e-liquids).

Final site selections will be based on the availability of CTP funds and resources for the relevant fiscal year, as well as the following factors, as applicable: (1) Compliance status of the requesting facility and affiliated firm; (2) whether the requesting facility or affiliated firm, if applicable, has a significant request or marketing application or submission pending with FDA; and (3) whether the requesting facility will be engaged in active manufacturing or processing during the proposed time of the visit. All travel expenses associated with Tobacco Product Manufacturer Facility Visits will be the responsibility of CTP.

IV. Requests for Participation

The request for participation should include the following identification information:

- The name and contact information (including address, phone number, and email) of your point of contact for the request;