

TABLE 1—ESTIMATES OF ANNUALIZED HOUR BURDEN—Continued

Type of respondent activity	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Adult questionnaire	10,000	2	20,000	.20	4,000
Youth questionnaire	2,500	2	5,000	.20	1000
Total	12,697	25,732	7,928

Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function.

Carlos Graham,
Reports Clearance Officer.

[FR Doc. 2021–23185 Filed 10–22–21; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

[Docket ID FEMA–2020–0016]

Meetings To Implement Pandemic Response Voluntary Agreement Under Section 708 of the Defense Production Act

AGENCY: Federal Emergency Management Agency, Department of Homeland Security.

ACTION: Announcement of meetings.

SUMMARY: The Federal Emergency Management Agency (FEMA) is holding a series of meetings to implement the Voluntary Agreement for the Manufacture and Distribution of Critical Healthcare Resources Necessary to Respond to a Pandemic.

DATES: The first meeting took place on Tuesday, October 12, 2021, from 10:00 a.m. to 12 p.m. Eastern Time (ET). The second meeting took place on Thursday, October 14, 2021, from 10:30 a.m. to 11 a.m. ET. The third meeting took place on Thursday, October 21, 2021, from 10:30 a.m. to 11 a.m. ET. The fourth meeting will take place on Thursday, October 28, 2021, from 10:30 a.m. to 11 a.m. ET. The fifth meeting will take place on Thursday, November 4, 2021, from 10:30 a.m. to 11 a.m. ET.

FOR FURTHER INFORMATION CONTACT: Robert Glenn, Office of Business, Industry, Infrastructure Integration, via

email at OB3I@fema.dhs.gov or via phone at (202) 212–1666.

SUPPLEMENTARY INFORMATION: Notice of these meetings is provided as required by section 708(h)(8) of the Defense Production Act (DPA), 50 U.S.C. 4558(h)(8), and consistent with 44 CFR part 332.

The DPA authorizes the making of “voluntary agreements and plans of action” with representatives of industry, business, and other interests to help provide for the national defense.¹ The President’s authority to facilitate voluntary agreements with respect to responding to the spread of COVID–19 within the United States was delegated to the Secretary of Homeland Security in Executive Order 13911.² The Secretary of Homeland Security further delegated this authority to the FEMA Administrator.³

On August 17, 2020, after the appropriate consultations with the Attorney General and the Chairman of the Federal Trade Commission, FEMA completed and published in the **Federal Register** a “Voluntary Agreement, Manufacture and Distribution of Critical Healthcare Resources Necessary to Respond to a Pandemic” (Voluntary Agreement).⁴ Unless terminated earlier, the Voluntary Agreement is effective until August 17, 2025, and may be extended subject to additional approval by the Attorney General after consultation with the Chairman of the Federal Trade Commission. The Agreement may be used to prepare for or respond to any pandemic, including COVID–19, during that time.

On December 7, 2020, the first plan of action under the Voluntary Agreement—the Plan of Action to Establish a National Strategy for the

Manufacture, Allocation, and Distribution of Personal Protective Equipment (PPE) to Respond to COVID–19 (PPE Plan of Action)—was finalized.⁵ The PPE Plan of Action established several sub-committees under the Voluntary Agreement, focusing on different aspects of the PPE Plan of Action.

On May 24, 2021, four additional plans of action under the Voluntary Agreement—the Plan of Action to Establish a National Strategy for the Manufacture, Allocation, and Distribution of Diagnostic Test Kits and other Testing Components to respond to COVID–19, the Plan of Action to Establish a National Strategy for the Manufacture, Allocation, and Distribution of Drug Products, Drug Substances, and Associated Medical Devices to respond to COVID–19, the Plan of Action to Establish a National Strategy for the Manufacture, Allocation, and Distribution of Medical Devices to respond to COVID–19, and the Plan of Action to Establish a National Strategy for the Manufacture, Allocation, and Distribution of Medical Gases to respond to COVID–19—were finalized.⁶ These plans of action established several sub-committees under the Voluntary Agreement, focusing on different aspects of each plan of action.

The meetings were chaired by the FEMA Administrator’s delegates from the Office of Response and Recovery (ORR) and Office of Policy and Program Analysis (OPPA), attended by the Attorney General’s delegates from the U.S. Department of Justice, and attended by the Chairman of the Federal Trade Commission’s delegates. In implementing the Voluntary Agreement, FEMA adheres to all procedural requirements of 50 U.S.C. 4558 and 44 CFR part 332.

Meeting Objectives: The objectives of the meetings are as follows:

1. Meet the Sub-Committee for Oxygen under the Medical Gases Plan of Action to establish priorities related to the COVID–19 response under the Voluntary Agreement.

⁵ See 85 FR 78869 (Dec. 7, 2020). See also 85 FR 79020 (Dec. 8, 2020).

⁶ See 86 FR 27894 (May 24, 2021). See also 86 FR 28851 (May 28, 2021).

¹ 50 U.S.C. 4558(c)(1).

² 85 FR 18403 (Apr. 1, 2020).

³ DHS Delegation 09052, Rev. 00.1 (Apr. 1, 2020); DHS Delegation Number 09052 Rev. 00 (Jan. 3, 2017).

⁴ 85 FR 50035 (Aug. 17, 2020). The Attorney General, in consultation with the Chairman of the Federal Trade Commission, made the required finding that the purpose of the voluntary agreement may not reasonably be achieved through an agreement having less anticompetitive effects or without any voluntary agreement and published the finding in the **Federal Register** on the same day. 85 FR 50049 (Aug. 17, 2020).

2. Gather Sub-Committee Participants and Attendees to ask targeted questions for situational awareness related to the Sub-Committee for Oxygen.

3. Identify potential Objectives and Actions that should be completed under the Sub-Committee for Oxygen.

4. Identify pandemic-related information gaps and areas that merit sharing by holding recurring meetings of the Sub-Committee for Oxygen with key stakeholders.

Meetings Closed to the Public: By default, the DPA requires meetings held to implement a voluntary agreement or plan of action be open to the public.⁷ However, attendance may be limited if the Sponsor⁸ of the voluntary agreement finds that the matter to be discussed at a meeting falls within the purview of matters described in 5 U.S.C. 552b(c), such as trade secrets and commercial or financial information.

The Sponsor of the Voluntary Agreement, the FEMA Administrator, found that these meetings to implement the Voluntary Agreement involved matters which fall within the purview of matters described in 5 U.S.C. 552b(c) and the meetings are therefore closed to the public.

Specifically, these meetings may require participants to disclose trade secrets or commercial or financial information that is privileged or confidential. Disclosure of such information allows for meetings to be closed to the public pursuant to 5 U.S.C. 552b(c)(4).

The success of the Voluntary Agreement depends wholly on the willing participation of the private sector participants. Failure to close these meetings to the public could reduce active participation by the signatories due to a perceived risk that sensitive company information could be prematurely released to the public. A premature public disclosure of a private sector participant's information could reduce trust and support for the Voluntary Agreement.

A resulting loss of support by the participants for the Voluntary Agreement would significantly frustrate the implementation of the Agency's objectives. Thus, these meeting closures are permitted pursuant to 5 U.S.C. 552b(c)(9)(B).

Deanne Criswell,

Administrator, Federal Emergency Management Agency.

[FR Doc. 2021-23226 Filed 10-22-21; 8:45 am]

BILLING CODE 9111-19-P

⁷ See 50 U.S.C. 4558(h)(7).

⁸ “[T]he individual designated by the President in subsection (c)(2) [of section 708 of the DPA] to administer the voluntary agreement, or plan of action.” 50 U.S.C. 4558(h)(7).

DEPARTMENT OF HOMELAND SECURITY

[Docket Number—DHS—2021—0037]

Agency Information Collection Activities: Office of the Immigration Detention Ombudsman Intake Form

AGENCY: Department of Homeland Security (DHS).

ACTION: 30-Day notice and request for comments; Office of the Immigration Detention Ombudsman Intake Form, 1601-0030, extension without change.

SUMMARY: The Department of Homeland Security, will submit the following Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and clearance in accordance with the Paperwork Reduction Act of 1995.

DATES: Comments are encouraged and will be accepted until November 24, 2021. This process is conducted in accordance with 5 CFR 1320.1.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this specific information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function.

SUPPLEMENTARY INFORMATION: The Department of Homeland Security's (DHS) Office of the Immigration Detention Ombudsman (OIDO) is an independent office tasked with resolving individual complaints from or about individuals in immigration detention regarding the potential violation of immigration detention standards or other potential misconduct. OIDO was established by Congress (Sec. 106 of the Consolidated Appropriations Act, 2020, Pub. L. 116-93). Its intake form is intended for use by individuals wishing to submit a complaint to OIDO. Information collected will provide the office with details about the allegations the submitter seeks to have OIDO address.

The information collected on this form will allow OIDO to identify: (1) The individual submitting the complaint and their contact information; (2) the detained individual who is the subject of the complaint; (3) the government-owned or contracted facility where the individual is or was detained and for how long; and (4) relevant details about the complaint. All of this information will be used by OIDO to investigate, resolve, and if appropriate, provide redress.

The use of this form is the most efficient means for collecting and processing the required data. Initially, collection will be via a paper form, which may be obtained from OIDO staff conducting routine visits in detention facilities. The form will also be available for download from the OIDO website. The PDF form will be able to be completed online, printed out, and submitted to OIDO by email, mail, or fax, or handed to a staff member in a detention facility.

After approval of the form described in this supporting statement, an electronic version will be developed so that submitters may complete and file via the OIDO website. The paper version will continue to be available; it will be noted on the form that using the paper method may result in processing delays for OIDO to complete data entry.

This information collection does not have an impact on small businesses or other small entities.

If this information is not collected, OIDO will not be able to accomplish its Congressional mandate to provide assistance to individuals who may be affected by misconduct, excessive force, or other violations of law or detention standards.

The assurance of confidentiality provided to the respondents for this information collection is based on the forthcoming Privacy Impact Assessment for the Immigration Detention Ombudsman Case Management System (ID-CMS) (June 21, 2021). Additionally, the information collected is covered by DHS/ALL-020 Department of Homeland Security Internal Affairs, April 28, 2014, 79 FR 23361 and DHS/ALL-025 Law Enforcement Authority in Support of the Protection of Property Owned, Occupied, or Secured by the Department of Homeland Security System of Records, June 14, 2017, 82 FR 27274.

This information collection was constructed in compliance with regulations and authorities under the purview of the DHS Privacy Office, DHS OCIO, DHS Records Management, and OMB regulations regarding data collection, use, sharing, storage, information security, and retrieval of information.

There are no changes to the information being collected and there is no change to the estimated burden associated with this collection.

The Office of Management and Budget is particularly interested in comments which:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including