

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Substance Abuse and Mental Health Services Administration****Agency Information Collection Activities: Proposed Collection; Comment Request**

In compliance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning opportunity for public comment on proposed collections of information, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information collection plans, call the SAMHSA Reports Clearance Officer at (240) 276-0361.

Comments are invited on: (a) Whether the proposed collections of information are necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection

of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

Proposed Project: Regulations To Implement SAMHSA's Charitable Choice Statutory Provisions—42 CFR Parts 54 and 54a (OMB No. 0930-0242)—Extension

Section 1955 of the Public Health Service Act (42 U.S.C. 300x-65), as amended by the Children's Health Act of 2000 (Pub. L. 106-310) and Sections 581-584 of the Public Health Service Act (42 U.S.C. 290kk *et seq.*, as added by the Consolidated Appropriations Act (Pub. L. 106-554)), set forth various provisions which aim to ensure that religious organizations are able to compete on an equal footing for federal funds to provide substance use services. These provisions allow religious organizations to offer substance use services to individuals without impairing the religious character of the organizations or the religious freedom of the individuals who receive the

services. The provisions apply to the Substance Abuse Prevention and Treatment Block Grant (SABG), to the Projects for Assistance in Transition from Homelessness (PATH) formula grant program, and to certain Substance Abuse and Mental Health Services Administration (SAMHSA) discretionary grant programs (programs that pay for substance use treatment and prevention services, not for certain infrastructure and technical assistance activities). Every effort has been made to assure that the reporting, recordkeeping and disclosure requirements of the proposed regulations allow maximum flexibility in implementation and impose minimum burden.

No changes are being made to the regulations or the burden hours. This information collection has been approved without changes since 2010.

Information on how states comply with the requirements of 42 CFR part 54 was approved by the Office of Management and Budget (OMB) as part of the Substance Abuse Prevention and Treatment Block Grant FY 2019-2021 annual application and reporting requirements approved under OMB control number 0930-0168.

42 CFR Citation and purpose	Number of respondents	Responses per respondent	Total responses	Hours per response	Total hours
Part 54—States Receiving SA Block Grants and/or Projects for Assistance in Transition from Homelessness (PATH)					
Reporting:					
96.122(f)(5) Annual report of activities the state undertook to comply 42 CFR part 54 (SABG)	60	1	60	1	60
54.8(c)(4) Total number of referrals to alternative service providers reported by program participants to States (respondents).					
SABG	6	(avg.) 23	135	1	135
PATH	10	5	50	1	50
54.8 (e) Annual report by PATH grantees on activities undertaken to comply with 42 CFR part 54	56	1	56	1	56
Disclosure:					
54.8(b) State requires program participants to provide notice to program beneficiaries of their right to referral to an alternative service provider.					
SABG	60	1	60	.05	3
PATH	56	1	56	.05	3
Recordkeeping:					
54.6(b) Documentation must be maintained to demonstrate significant burden for program participants under 42 U.S.C. 300x-57 or 42 U.S.C. 290cc-33(a)(2) and under 42 U.S.C. 290cc-21 to 290cc-35	60	1	60	1	60
Part 54—Subtotal	115	477	367

Part 54a—States, local governments and religious organizations receiving funding under Title V of the PHS Act for substance abuse prevention and treatment services

Reporting:					
54a.8(c)(1)(iv) Total number of referrals to alternative service providers reported by program participants to states when they are the responsible unit of government	25	4	100	.083	8

42 CFR Citation and purpose	Number of respondents	Responses per respondent	Total responses	Hours per response	Total hours
54a(8)(d) Total number of referrals reported to SAMHSA when it is the responsible unit of government. (Note: This notification will occur during the course of the regular reports that may be required under the terms of the funding award.)	20	2	40	.25	10
Disclosure: 54a.8(b) Program participant notice to program beneficiaries of rights to referral to an alternative service provider	1,460	1	1,460	1	1,460
Part 54a—Subtotal	1,505	1,600	1,478
Total	1,620	2,077	1,845

Send comments to Carlos Graham, SAMHSA Reports Clearance Officer, 5600 Fishers Lane, Room 15E57–A, Rockville, Maryland 20857, or email a copy to Carlos.Graham@samhsa.hhs.gov. Written comments should be received by March 21, 2022.

Carlos Graham,

Reports Clearance Officer.

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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, under the Plan of Action to Establish a National Strategy for the Manufacture, Allocation, and Distribution of Medical Gases to Respond to COVID–19, to implement the Voluntary Agreement for the Manufacture and Distribution of Critical Healthcare Resources Necessary to Respond to a Pandemic.

DATES:

- Thursday, January 6, 2022, from 2 p.m. to 3 p.m. Eastern Time (ET).
- Thursday, January 13, 2022, from 2 p.m. to 3 p.m. ET.
- Thursday, January 20, 2022, from 2 p.m. to 3 p.m. ET.
- Thursday, January 27, 2022, from 2 p.m. to 3 p.m. ET.

FOR FURTHER INFORMATION CONTACT:

Robert Glenn, FEMA Office of Response and Recovery’s 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

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).

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.

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).