

Dated: May 29, 2024.

Miguelina Perez,

Program Analyst, Office of Federal Advisory Committee Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Aging; Notice of Closed Meeting

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute on Aging Special Emphasis Panel; Program Project.

Date: July 11, 2024.

Time: 12:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute on Aging, 5601 Fishers Lane, Rockville, MD 20852 (Virtual Meeting).

Contact Person: Dario Dieguez, Ph.D., Scientific Review Officer, Scientific Review Branch, National Institutes of Health, National Institute on Aging, 5601 Fishers Lane, Rockville, MD 20852, (301) 827–3101, dario.dieguez@nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.866, Aging Research, National Institutes of Health, HHS)

Dated: May 30, 2024.

Miguelina Perez,

Program Analyst, Office of Federal Advisory Committee Policy.

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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 on (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.

Project: Medications for the Treatment of Opioid Use Disorder—42 CFR Part 8 (OMB No. 0930–0206) and Opioid Treatment Programs (OTPs)—Extension

42 CFR part 8 establishes a certification program managed by SAMHSA's Center for Substance Abuse Treatment (CSAT). The regulation requires that opioid treatment programs (OTPs) be certified. "Certification" is the process by which SAMHSA determines that an OTP is qualified to provide opioid use disorder treatment under the federal opioid use disorder treatment standards established by the Secretary of Health and Human Services. To become certified, an OTP must be accredited by a SAMHSA-approved accreditation body. The

regulation also provides standards for such services as individualized treatment planning, medical care, and assessment of patient outcomes. This submission seeks continued approval of the information collection requirements in the regulation and of the forms used in implementing the regulation.

SAMHSA currently has approval for the *Application for Certification to Use Medications for the Treatment of Opioid Use Disorder in a Treatment Program Under 42 CFR 8.11* (Form SMA–162); the *Application for Approval as Accreditation Body Under 42 CFR 8.3(b)* (Form SMA–163); and the *Exception Request and Record of Justification Under 42 CFR 8.12* (Form SMA–168), which may be used on a voluntary basis by OTP practitioners when there is a patient care situation in which the OTP practitioner must make a treatment decision that falls outside of the required standards delineated in the regulation. Form SMA–168 is a simplified, standardized form to facilitate the documentation, request, and approval process for exceptions.

SAMHSA believes that the recordkeeping requirements in the regulation are customary and usual practices within the medical and rehabilitative communities and has not calculated a response burden for them. The recordkeeping requirements set forth in 42 CFR 8.4, 8.11 and 8.12 include maintenance of the following: 5-year retention by accreditation bodies of certain records pertaining to accreditation; documentation by an OTP of the following: a patient's medical examination when admitted to treatment, a patient's medical history, a care plan, any prenatal support provided the patient if applicable, the medical rationale for initial starting doses above 50mg, the medical rationale for a patient's dosage schedule, and care decisions made as a result of follow-up medical examinations.

The tables that follow summarize the annual reporting burden associated with the regulation, including burden associated with the forms. There are minor changes to these forms to improve data collection, remove unnecessary questions, and align terms with the final 42 CFR part 8 rule released February 2, 2024.

Form	Number of respondents	Responses/ respondent	Total responses	Hours/ response	Total hours
Estimated Annual Reporting Requirement Burden for Accreditation Bodies					
SMA-163	54	26.055	1,407	0.28	394
Estimated Annual Reporting Requirement Burden for Opioid Treatment Programs					
SMA-162	751.33	17.976	13,506	0.08	1,081
SMA-168	1,302.67	17.977	23,418	0.08	1,873
Subtotal	2,054	17.977	36,925	0.08	2,954
Total			38,332		3,348

Send comments to SAMHSA Reports Clearance Officer, 5600 Fisher Lane, Room 15E45, Rockville, MD 20852 OR email a copy to samhsapra@samhsa.hhs.gov. Written comments should be received by August 5, 2024.

Alicia Broadus,
Public Health Advisor.

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DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Automated Commercial Environment (ACE) Export Manifest for Air Cargo Test: Renewal of Test

AGENCY: U.S. Customs and Border Protection; Department of Homeland Security.

ACTION: General notice.

SUMMARY: This notice announces that CBP is renewing U.S. Customs and Border Protection’s (CBP’s) Automated Commercial Environment (ACE) Export Manifest for Air Cargo Test, a National Customs Automation Program (NCAP) test concerning ACE export manifest capability.

DATES: The voluntary pilot initially began on August 10, 2015, was modified and extended on August 14, 2017, and was further extended on December 22, 2021. This renewal is effective June 4, 2024. The extended test will run for an additional two years from the date of publication of this notice in the **Federal Register**.

ADDRESSES: Applications to participate in the ACE Export Manifest for Air Cargo Test must be submitted via email to CBP Export Manifest at cbpexportmanifest@cbp.dhs.gov. In the subject line of the email, please use “ACE Export Manifest for Air Cargo Test Application”. Applications will be accepted at any time during the test period. Written comments concerning

program, policy, and technical issues may also be submitted via email to CBP Export Manifest at cbpexportmanifest@cbp.dhs.gov. In the subject line of the email, please use “Comment on ACE Export Manifest for Air Cargo Test”. Comments may be submitted at any time during the test period.

FOR FURTHER INFORMATION CONTACT: Thomas J. Pagano, Branch Chief, or David Garcia, Program Manager, Outbound Enforcement and Policy Branch, Office of Field Operations, U.S. Customs and Border Protection, via email at cbpexportmanifest@cbp.dhs.gov.

SUPPLEMENTARY INFORMATION:

I. Background

The Automated Commercial Environment (ACE) Export Manifest for Air Cargo Test is a voluntary test in which participants agree to submit export manifest data to U.S. Customs and Border Protection (CBP) electronically, at least four hours prior to loading of the cargo onto the aircraft in preparation for departure from the United States. The ACE Export Manifest for Air Cargo Test is authorized under § 101.9(b) of title 19 of the Code of Federal Regulations (19 CFR 101.9(b)), which provides for the testing of National Customs Automation Program (NCAP) programs or procedures.

The ACE Export Manifest for Air Cargo Test examines the functionality regarding the filing of export manifest data for air cargo electronically in ACE. The ACE system creates a single automated export processing platform for certain export manifest, commodity, licensing, export control, and export targeting transactions. This will reduce costs for CBP, partner government agencies, and the trade community, as well as improve facilitation of export shipments through the supply chain.

The ACE Export Manifest for Air Cargo Test will also assess the feasibility of requiring the manifest information to be filed electronically in ACE within a specified time before the cargo is loaded

on the aircraft. This capability will enable CBP to calculate the risk and effectively identify and inspect shipments prior to loading of cargo to ensure compliance with all U.S. export laws.

CBP announced the procedures and criteria related to participation in the ACE Export Manifest for Air Cargo Test in a notice published in the **Federal Register** on July 10, 2015 (80 FR 39790). This test was originally set to run for approximately two years. On August 14, 2017, CBP extended the test period for one additional year (82 FR 37888). At that time, CBP also modified the original notice, making certain of the data elements optional, and opened the test to accept additional applications for participation from all parties who met the eligibility requirements.

The data elements, unless noted otherwise, are mandatory. Data elements which are “mandatory” must be provided to CBP for every shipment. Data elements which are “conditional” must be provided to CBP only if the particular information pertains to the cargo. Data elements which are “optional” may be provided to CBP but are not required. The data elements are set forth below:

- (1) Exporting Carrier
- (2) Marks of nationality and registration
- (3) Flight number
- (4) Port of lading
- (5) Port of unloading
- (6) Scheduled date of departure
- (7) Consolidator (conditional)
- (8) De-consolidator (conditional)
- (9) Air waybill type (Master, House, Simple or Sub)
- (10) Air waybill number
- (11) Number of pieces and unit of measure (optional)
- (12) Weight (kg./lb.)
- (13) Number of house air waybills (optional)
- (14) Shipper name and address
- (15) Consignee name and address
- (16) Cargo description
- (17) AES Internal Transaction Number (ITN) or AES Exemption Statement/ Exception Classification (per shipment)