

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice via www.reginfo.gov/public/do/PRAMain. Find this information collection by selecting “Currently under Review” and “Select Agency: Department of Health and Human Services”.

FOR FURTHER INFORMATION CONTACT: When requesting information, such as a free copy of the form to be used for the information collection, please include the document identifier OS–0945-New-30D and project title for reference, to Conner O’Brien by email at OCRPrivacy@hhs.gov, or by phone at (202) 240–3110.

SUPPLEMENTARY INFORMATION: Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency’s functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Title of the Collection: Confidentiality of Substance Use Disorder Patient Records Complaint Form.

Type of Collection: New Collection.

OMB No. 0945-New: Office for Civil Rights (OCR)—Health Information Privacy, Data, and Cybersecurity Division.

Abstract: OCR is requesting Office of Management and Budget (OMB) approval for a new information collection. OCR published a 60-day notice in the **Federal Register** on September 11, 2025,¹ and, following a government shutdown, OCR reopened the comment period for an additional 47 days on November 19, 2025.² No public comments were received. Under OCR’s delegated civil enforcement authority, a person may file a complaint with OCR for alleged noncompliance with 42 CFR part 2 (“Part 2”). OCR will collect information from individuals using the form, Confidentiality of Substance Use Disorder Patient Records Complaint, to allow OCR to collect the minimum information needed from individuals who file complaints with OCR to form the basis for the initial processing of such complaints to satisfy the right of an individual to file a complaint under 42 CFR 2.4(b).

I. Authority

OCR has delegated civil enforcement authority³ under 42 U.S.C. 290dd–2(f), as amended by section 3221 of the Coronavirus Aid, Relief, and Economic Security Act, which applies section 1176 of the Social Security Act (“the Act”), 45 U.S.C. 1320d–5, to a violation of Part 2 in the same manner as it applies to a violation of part C of title XI of the Act, Administrative Simplification. The Administrative Simplification provisions were added to the Act by the Health Insurance Portability and Accountability Act of 1996 (HIPAA). OCR currently exercises its enforcement authority under section 1176 of the Act to enforce the HIPAA

Privacy, Security, and Breach Notification Rules (collectively, “HIPAA Rules”) in accordance with 45 CFR part 160 (“Enforcement Rule”). To implement 42 U.S.C. 290dd–2, as amended, the Department published a final rule modifying Part 2, 89 FR 12472 (February 16, 2024) (“2024 Part 2 Final Rule”), which included, among other provisions, a new right to file complaints of noncompliance and applied 45 CFR part 160, subparts C, D, and E, as the enforcement regulation for Part 2.

II. Method of Collection

Individuals will be able to submit a Part 2 complaint using the HHS OCR online complaint portal or by filling out a copy of the form obtained from OCR’s website and submitting it to OCR electronically or by postal mail.

III. Estimated Burden

The estimated burden on individuals for gathering information and completing the Part 2 complaint form is 45 minutes per complaint based on the hourly burden for HIPAA complaints. Assuming that 1 in every 1,000 patients treated for SUD would file a Part 2 complaint, the Department estimates a total of 1,398 burden hours for 1,864 complaints annually (1,864,367 patients/1,000 × .75 hours = 1,398).

Likely Respondents: Patients of substance use disorder treatment programs who believe that a Part 2 program, covered entity, business associate, qualified service organization, or other lawful holder of Part 2 records is noncompliant with 42 CFR part 2.

Estimated Annualized Burden Table:

42 CFR	Type of respondent	Number of respondents	Number of responses per respondent	Total responses	Average burden hours per response	Total burden hours
2.4	Individuals filing a Part 2 complaint	1,864	1	1,864	0.75	1,398

Catherine Howard,
Paperwork Reduction Act Reports Clearance Officer, Department of Health and Human Services, Office of the Secretary.

[FR Doc. 2026–00499 Filed 1–12–26; 8:45 am]

BILLING CODE 4153–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

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

The meetings 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: Center for Scientific Review Special Emphasis Panel: Cell Structure Function.

Date: January 27, 2026.

Time: 2:00 p.m. to 3:30 p.m.

Agenda: To review and evaluate grant applications.

¹ 90 FR 44077 (September 11, 2025).

² 90 FR 52070 (November 19, 2025).

³ 90 FR 41833 (August 27, 2025).

Address: National Institutes of Health,
6701 Rockledge Drive, Bethesda, MD 20892.
Meeting Format: Virtual Meeting.
Contact Person: Jessica Smith, Ph.D.,
Scientific Review Officer, Center for
Scientific Review, National Institutes of
Health, 6701 Rockledge Drive, Bethesda, MD
20892, 301-402-3717, jessica.smith6@nih.gov.

This notice is being published less than 15 days from the meeting date due to exceptional circumstances. As a result of the 43-day government shutdown, due to lapsed appropriations, the above meeting was canceled. This meeting was to assess the scientific and technical merit of NIH grant applications, required by statute to disburse NIH funds. The meeting must take place urgently so that evaluations of biomedical research applications addressing multiple major public health priorities can be submitted to the national advisory councils for timely funding recommendations.

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393-93.396, 93.837-93.844, 93.846-93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: January 8, 2026.

Bruce A. George,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2026-00451 Filed 1-12-26; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

Notice of Issuance of Final Determination Concerning Lumens VC-TR60A Camera

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

ACTION: Notice of final determination.

SUMMARY: This document provides notice that U.S. Customs and Border Protection (CBP) has issued a final determination concerning the country of origin of the Lumens VC-TR60A Camera. Based upon the facts presented, CBP has concluded that the last substantial transformation of the Lumens VC-TR60A Camera occurs in Taiwan.

DATES: The final determination was issued on December 19, 2025. A copy of the final determination is attached. Any party-at-interest, as defined in 19 CFR 177.22(d), may seek judicial review of this final determination no later than February 12, 2026.

FOR FURTHER INFORMATION CONTACT:

Anna Hedstrom, Valuation and Special Programs Branch, Regulations and Rulings, Office of Trade, at (202) 325-0227.

SUPPLEMENTARY INFORMATION: Notice is hereby given that on December 19, 2025, CBP issued a final determination concerning the country of origin of the Lumens VC-TR60A Camera for purposes of Title III of the Trade Agreements Act of 1979. This final determination, Headquarters Ruling Letter (HQ) H350894, was issued at the request of Lumens Digital Optics, Inc. under procedures set forth at 19 CFR part 177, subpart B, which implements Title III of the Trade Agreements Act of 1979, as amended (19 U.S.C. 2511-18). In the final determination, CBP has concluded that the last substantial transformation of the Lumens VC-TR60A Camera occurs in Taiwan.

Section 177.29, CBP Regulations (19 CFR 177.29), provides that a notice of final determination shall be published in the **Federal Register** within 60 days of the date the final determination is issued. Section 177.30, CBP Regulations (19 CFR 177.30), provides that any party-at-interest, as defined in 19 CFR 177.22(d), may seek judicial review of a final determination within 30 days of publication of such determination in the **Federal Register**.

Alice A. Kipel,

Executive Director, Regulations and Rulings, Office of Trade.

90 K Street NE – 10th Floor
Washington, DC 20229-1177



U.S. Customs and Border Protection

HQ H350894

December 19, 2025

OT:RR:CTF:VS H350894 ACH

CATEGORY: Origin

Gary Zheng, Lumens Digital Optics, Inc., 5F No. 20 Taiyung Street, Jhubei, 301 Taiwan

RE: U.S. Government Procurement; Title III, Trade Agreements Act of 1979 (19 U.S.C. 2511); Subpart B, Part 177, CBP Regulations; Country of Origin of Lumens Camera

Dear Mr. Zheng,

This is in response to your July 8, 2025 request for a final determination concerning the country of origin of the Pan Tilt Zoom (“PTZ”) Lumens VC-TR60A Camera (“Camera”) pursuant to Title III of the Trade Agreements Act of 1979 (“TAA”), as amended (19 U.S.C. 2511 *et seq.*), and subpart B of Part 177, U.S. Customs and Border Protection

(“CBP”) Regulations (19 CFR 177.21, *et seq.*). Lumens is a party-at-interest within the meaning of 19 CFR 177.22(d)(1) and 177.23(a) and is therefore entitled to request this final determination.

Facts

The merchandise under consideration is the Lumens PTZ Camera, Model Number TR60A. This camera is designed for video conferencing in office environments. It features speaker tracking that allows the camera to move toward the person speaking during a discussion. The camera also can combine two video images in a picture-in-picture display, providing both a wide-angle view and a close-up shot of the meeting. Additionally, it has a dual lens 12 times optical zoom, wide-angle fix lens, Artificial Intelligence (“AI”) motion detection, and simultaneous

output for High-Definition Multimedia Interface (“HDMI”), Ethernet, and Universal Serial Bus (“USB”).

The camera production consists of nine main steps: (1) zoom lens module assembly and calibration; (2) PTZ camera body assembly; (3) panoramic lens module assembly; (4) zoom lens and panoramic lens optical axis alignment; (5) run in test; (6) firmware programming to camera; (7) system function test; (8) quality control inspection; and (9) final packaging and shipping. One camera takes around three hours to produce.

The camera’s initial source code, which includes requirements for the graphical user interface that provides product status information and the user operation interface, is developed in China and sent to Taiwan for burning. This software has been developed over a year and a half and has undergone