Name of Committee: National Institute on Deafness and Other Communication Disorders Special Emphasis Panel; NIDCD Clinical Research Center Grant (P50) Review.

Date: August 2, 2017.

Time: 1:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852 (Telephone Conference Call).

Contact Person: Katherine Shim, Ph.D., Scientific Review Officer, Division of Extramural Activities, NIH/NIDCD, 6001 Executive Blvd., Room 8351, Bethesda, MD 20892, 301–496–8683, *katherine.shim@nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.173, Biological Research Related to Deafness and Communicative Disorders, National Institutes of Health, HHS)

Dated: July 3, 2017.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–14296 Filed 7–6–17; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), 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 of Allergy and Infectious Diseases Special Emphasis Panel; Development of Multipurpose Prevention Technologies (R61/R33).

Date: August 3, 2017.

Time: 11:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 5601 Fishers Lane, Rockville, MD 20892 (Telephone Conference Call).

Contact Person: Brenda Lange-Gustafson, Ph.D., Scientific Review Officer NIAID/NIH/DHHS, Scientific Review Program, 5601 Fishers Lane, Room 3G13, Rockville, MD 20852, 240–669–5047, *bgustafson@niaid.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.855, Allergy, Immunology, and Transplantation Research; 93.856, Microbiology and Infectious Diseases Research, National Institutes of Health, HHS)

Dated: June 29, 2017.

Natasha M. Copeland,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–14229 Filed 7–6–17; 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 a Digital Radiography System

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 a digital radiography system, (also commonly referred to as an x-ray system), known as the Carestream DRX-Ascend Digital Radiography system. Based upon the facts presented for purposes of U.S. Government procurement, CBP has concluded that the United States is the country of origin of the fully assembled and installed DRX-Ascend Digital Radiography system.

DATES: The final determination was issued on June 30, 2017. 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 within August 7, 2017.

FOR FURTHER INFORMATION CONTACT: Robert Dinerstein, Valuation and Special Programs Branch, Regulations and Rulings, Office of Trade, at (202) 325–0132.

SUPPLEMENTARY INFORMATION: Notice is hereby given that on June 30, 2017 pursuant to subpart B of Part 177, U.S. Customs and Border Protection Regulations (19 CFR part 177, subpart B), CBP issued a final determination concerning the country of origin of a digital radiography system known as the Carestream DRX–Ascend Digital Radiography system, which may be offered to the U.S. Government under an undesignated government procurement contract. This final determination, HQ H283088, was issued 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). The major components of the DRX-Ascend Digital Radiography system include a Chinese-origin high-voltage generator, a U.S.-origin wireless DRX detector, a Chinese-origin elevating float-top table, a Chinese-origin tubestand, a Chinese-origin wall stand, and either a U.S. or a Japanese-origin x-ray tube. These components are combined with software that is largely developed in the United States. In the final determination, CBP concluded that the components are substantially transformed in the United States when the fully functioning digital radiography system is completely assembled and installed at an on-site location. Thus, the fully assembled digital radiography system becomes a product of the United States. Therefore, for purposes of U.S. Government procurement, the United States is the country of origin of the installed and assembled Carestream DRX-Ascend Digital Radiography system.

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**.

Dated: June 30, 2017.

Alice A. Kipel,

Executive Director, Regulations and Rulings, Office of Trade.

HQ H283088

OT:RR:CTF:VS H283088 RSD

CATEGORY: Origin

Gunjan R. Talati, Esq. Kilpatrick Townsend & Stockton 607 14th Street NW. Suite 900 Washington, DC 20005–2018

RE: U.S. Government Procurement; Title III, Trade Agreements Act of 1979 (19 U.S.C. 2511); Subpart B, Part 177, CBP Regulations; Digital Radiography System

Dear Mr. Talati:

This is in response to your letter of January 11, 2017, forwarded to the National Commodity Specialist Division on behalf of Carestream Health, Inc. (Carestream), requesting a final determination concerning the country of origin of a Digital Radiography System, pursuant to subpart B of Part 177, U.S. Customs and Border Protection (CBP) Regulations (19 CFR 177.21, *et seq.*). The National Commodity Specialist Division transmitted your request to the Office of