Dated: June 27, 2016.

Jennifer Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–15643 Filed 6–30–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Alcohol Abuse and Alcoholism; Notice of Closed Meetings

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 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: National Institute on Alcohol Abuse and Alcoholism Special Emphasis Panel; NIAAA Member Conflict Reviews—Clinical Sciences and Epidemiology.

Date: July 11, 2016.

Time: 11:00 a.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health–NIAAA, 5635 Fishers Lane, Rockville, MD 20892 (Telephone Conference Call).

Contact Person: Richard A. Rippe, Ph.D., Scientific Review Officer, National Institute on Alcohol Abuse and Alcoholism, National Institutes of Health, 5635 Fishers Lane, Room 2109, Rockville, MD 20852, 301–443–8599, *rippera@mail.nih.gov.*

Name of Committee: National Institute on Alcohol Abuse and Alcoholism Special Emphasis Panel; NIAAA Fellowship Review.

Date: July 26, 2016.

Time: 8:00 a.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, National Institute on Alcohol Abuse and Alcoholism, Terrace Level Conference Room 508, 5635 Fishers LN, Rockville, MD 20892.

Contact Person: Richard A. Rippe, Ph.D., Scientific Review Officer, National Institute on Alcohol Abuse and Alcoholism, National Institutes of Health, 5635 Fishers Lane, Room 2109, Rockville, MD 20852, 301–443–8599, *rippera@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.271, Alcohol Research Career Development Awards for Scientists and Clinicians; 93.272, Alcohol National Research Service Awards for Research Training; 93.273, Alcohol Research Programs; 93.891, Alcohol Research Center Grants; 93.701, ARRA Related Biomedical Research and Research Support Awards, National Institutes of Health, HHS)

Dated: June 27, 2016.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–15592 Filed 6–30–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; Cancellation of Meeting

Notice is hereby given of the cancellation of the National Cancer Institute Special Emphasis Panel, June 27, 2016, 11:00 a.m. to June 27, 2016, 12:00 p.m., National Cancer Institute Shady Grove, 9609 Medical Center Drive, 7W624, Rockville, MD, 20850 which was published in the **Federal Register** on May 2, 2016, 81FR26240.

This meeting is canceled because the applications were reassigned.

Dated: June 27, 2016.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–15591 Filed 6–30–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Alcohol Abuse And Alcoholism; Notice of Closed Meetings

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 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: National Institute on Alcohol Abuse and Alcoholism Special Emphasis Panel; U01 Application Review.

Date: July 13, 2016.

Time: 11:00 a.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, National Institute on Alcohol Abuse and Alcoholism, 5635 Fishers Lane, Rockville, MD 20892, (Telephone Conference Call).

Contact Person: Richard A. Rippe, Ph.D., Scientific Review Officer, National Institute on Alcohol Abuse and Alcoholism, 5635 Fishers Lane, Room 2109, Rockville, MD 20852, 301–443–8599, *rippera@mail.nih.gov.*

Name of Committee: National Institute on Alcohol Abuse and Alcoholism Special Emphasis Panel; Targets of Low Dose Alcohol in the Brain.

Date: July 21, 2016.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, National Institute on Alcohol Abuse and Alcoholism, Conference Room 3002/3004, 5635 Fishers Lane, Rockville, MD 20892.

Contact Person: Philippe Marmillot, Ph.D., National Institutes of Health, National Institute on Alcohol Abuse and Alcoholism, 5635 Fishers Lane, Rm 2017, Bethesda, MD 20892, 301–443–2861, *marmillotp@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.271, Alcohol Research Career Development Awards for Scientists and Clinicians; 93.272, Alcohol National Research Service Awards for Research Training; 93.273, Alcohol Research Programs; 93.891, Alcohol Research Center Grants; 93.701, ARRA Related Biomedical Research and Research Support Awards, National Institutes of Health, HHS)

Dated: June 27, 2016.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–15593 Filed 6–30–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; Population Assessment of Tobacco and Health (PATH) Study—Fourth Wave of Data Collection

AGENCY: National Institute of Health, HHS.

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information

collection was previously published in the **Federal Register** on April 15, 2016, pages 22290—22291 and allowed 60-days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Direct Comments To OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, *OIRA_Submission@omb.eop.gov* or by fax to (202) 395–6974, Attention: NIH Desk Officer.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, submit comments in writing or request more information on the proposed project, contact: Kevin P. Conway, Ph.D., Deputy Director, Division of Epidemiology, Services, and Prevention Research, National Institute on Drug Abuse, 6001 Executive Boulevard, Room 5185; or call non-toll-free number (301) 443–8755; or Email your request, including your address to: *PATHprojectofficer@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: Population Assessment of Tobacco and Health (PATH) Study—Fourth Wave of Data Collection (NIDA), 0925–0664, expiration date 8/31/2018—REVISION—NIDA, NIH, in partnership with the Food and Drug Administration (FDA).

Need and Use of Information Collection: This is a revision request (OMB number 0925–0664, expiration date 8/31/2018) for the Population Assessment of Tobacco and Health (PATH) Study to conduct the fourth wave of data collection. The PATH Study is a large national longitudinal cohort study on tobacco use behavior and health among the U.S. household population of adults age 18 and older and youth ages 12 to 17. On an annual basis, the PATH Study conducts interviews with and collects biospecimens from adults and youth to help inform the development, implementation, and evaluation of tobacco-product regulations by FDA in meeting its mission under the Family Smoking Prevention and Tobacco Control Act (TCA) to regulate tobacco products, including tobacco-product advertising, labeling, marketing, constituents, ingredients, and additives. The longitudinal design of the PATH Study provides it with the capacity to measure and report within-person changes and between-person differences in tobacco product use behaviors and health effects within the cohort over time. These data will help to inform regulatory decisions and actions by FDA and FDA's evaluations of associations between its regulations and tobacco use behaviors and health indicators in the population.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 94,798.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
1. Household Screener	Households	48,018	1	14/60	11,204
2. Individual Screener	Adults—New adults	9,152	1	6/60	915
3. Extended Interview	Adults—New adults and Wave 1 youth respondents who age up to adult cohort at Wave 4.	10,737	1	68/60	12,169
4. Extended Interview	Adults—Adult respondents at previous wave	23,414	1	1	23,414
5. Parent Interview	Adults—Parents of new youth and parents of shadow youth who age up to youth cohort at Wave 4.	6,561	1	19/60	2,078
6. Parent Interview	Adults—Parents of youth respondents at previous wave.	8,800	1	16/60	2,347
7. Extended Interview	Youth—New youth and shadow youth who age up to youth cohort at Wave 4.	6,432	1	45/60	4,824
8. Extended Interview	Youth—Youth respondents at previous wave	8,627	1	35/60	5,032
9. Tobacco Use Form	Adults	23,133	1	5/60	1,928
10. Tobacco Use Form	Youth	10,239	1	5/60	853
11. Shadow Youth Only Screener	Households	41,207	1	5/60	3,434
12. Verification Interview	Adults	33,889	1	2/60	1,130
13. Validation Interview	Adults	301	1	4/60	20
14. Biospecimen Collection: Blood	Adults—New adults and Wave 1 youth respondents who age up to adult cohort at Wave 4.	4,832	1	18/60	1,450
15. Biospecimen Collection: Urine	Adults	18,301	1	10/60	3,050
16. Biospecimen Collection: Urine	Youth	10,239	1	10/60	1,707
17. Follow-up/Tracking Participant Information Form.	Adults	34,151	2	8/60	9,107
18. Follow-up/Tracking Participant Information Form for Youth (completed by parents).	Adults—Parents of youth respondents	15,059	2	8/60	4,016

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Form name	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
19. Follow-up/Tracking Participant Information Form for sample shadow youth (completed by parents).	Adults—Parents of shadow youth	4,684	2	8/60	1,249
20. Consent for Extended Interview	Adults—New adults and Wave 1 youth respondents who age up to adult cohort at Wave 4.	13,984	1	4/60	932
21. Parent Permission and Consent for Parent Interview.	Adults—Parents of new youth and parents of Shadow youth who age up to youth cohort at Wave 4.	7,657	1	5/60	638
22. Assent for Extended Interview	Youth—New youth and shadow youth who age up to youth cohort at Wave 4.	7,657	1	3/60	383
23. Consent for Biological Samples	Adults—New adults and Wave 1 youth respondents who age up to adult cohort at Wave 4.	10,737	1	5/60	895
24. Parent permission for urine collection.	Adults—Parents of youth respondents at previous wave.	15,360	1	3/60	768
25. Assent for urine collection	Youth	15,059	1	5/60	1,255
Total		388,229	442,123		94,798

Dated: June 27, 2016.

Genevieve deAlmeida,

Project Clearance Liaison, National Institute on Drug Abuse, NIH.

[FR Doc. 2016–15644 Filed 6–30–16; 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 Certain Network Cables and Transceivers

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 certain transceivers imported separately and certain imported network cables containing transceivers. Based upon the facts presented, CBP has concluded in both instances that the country of origin of the merchandise is China for purposes of U.S. Government procurement.

DATES: The final determination was issued on June 14, 2016. 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 1, 2016.

FOR FURTHER INFORMATION CONTACT: Grace A. Kim, Valuation and Special Programs Branch, Regulations and Rulings, Office of International Trade (202) 325–7941.

SUPPLEMENTARY INFORMATION: Notice is hereby given that on June 14, 2016, 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 certain network cables and transceivers, which may be offered to the U.S. Government under an undesignated government procurement contract. This final determination, HQ H273091, 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). In the final determination, CBP concluded that the processing of the imported merchandise in the U.S. does not result in a substantial transformation. Therefore, the country of origin of the transceivers and of the network cables containing transceivers is China for purposes of U.S. Government procurement.

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 14, 2016.

Joanne R. Stump,

Acting Executive Director, Regulations and Rulings, Office of International Trade.

HQ H273091

June 14, 2016

OT:RR:CTF:VS H273091 GaK

CATEGORY: Origin

Janet C. Wallett
FCI USA LLC.
825 Old Trail Road
Etters, PA 17319

RE: U.S. Government Procurement; Country of origin of copper cables containing transceivers and of the fiber optic transceiver; Substantial Transformation

Dear Ms. Wallett:

This is in response to your letter dated January 6, 2016, requesting a final determination on behalf of FCI USA LLC (''FCI''), pursuant to subpart B of part 177 of the U.S. Customs & Border Protection (''CBP'') Regulations (19 CFR part 177). Under these regulations, which implement Title III of the Trade Agreements Act of 1979 (''TAA''), as amended (19 U.S.C. § 2511 *et seq.*), CBP issues country of origin advisory rulings and final determinations as to whether an article is or would be a product of a designated country or instrumentality for the purposes of granting waivers of certain ''Buy American'' restrictions in U.S. law or practice for products offered for sale to the U.S. Government. This final determination concerns the country of origin of FCI's Copper Direct Attach Copper (''DAC'') cable—HPL500 (''Cable'') and Fiber Optic Transceivers—HPL512 (''Transceivers''). We note that as a U.S. importer, FCI is a party-at-interest within the meaning of 19 CFR § 177.22(d)(1) and is entitled to request this final determination.