evidence that environmental chemicals, physical substances, or mixtures (collectively referred to as ''substances'') cause adverse health effects and provides opinions on whether these substances may be of concern given what is known about current human exposure levels. OHAT also organizes workshops or state-of-the-science evaluations to address issues of importance in environmental health sciences. Information about OHAT is found at *http://ntp.niehs.nih.gov/go/ohat.*

Background Information on NTP Peer Review Panels: NTP panels are technical, scientific advisory bodies established on an ''as needed'' basis to provide independent scientific peer review and advise the NTP on agents of public health concern, new/revised toxicological test methods, or other issues. These panels help ensure transparent, unbiased, and scientifically rigorous input to the program for its use in making credible decisions about human hazard, setting research and testing priorities, and providing information to regulatory agencies about alternative methods for toxicity screening. The NTP welcomes nominations of scientific experts for upcoming panels. Scientists interested in serving on an NTP panel should provide current *curriculum vitae* to the **FOR FURTHER INFORMATION CONTACT**. The authority for NTP panels is provided by 42 U.S.C. 217a; section 222 of the Public Health Service (PHS) Act, as amended. The panel is governed by the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), which sets forth standards for the formation and use of advisory committees.

Dated: February 22, 2016.

John R. Bucher,

Associate Director, NTP.

[FR Doc. 2016–04102 Filed 2–25–16; 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: The National Physician Survey of Precision Medicine in Cancer Treatment (NCI)

Summary: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Cancer Institute, the National Institutes of Health, 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 November 18, 2015 (80 FR 72077), 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 Cancer Institute (NCI), 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, or request more information on the proposed project, contact: Janet S. de Moor, Ph.D., MPH, Project Officer, Division of Cancer Control and Population Sciences, 9609 Medical Center Drive, 3E438, MSC 9764, Rockville, MD, 20850 or call non-toll-free number 240–276–6806 or Email your request, including your address to: *janet.demoor@nih.gov*. Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: The National Physician Survey of Precision Medicine in Cancer Treatment 0925–NEW, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: The purpose of this study is to investigate the current practice of precision medicine in cancer treatment among medical oncologists in the U.S. This is a nationally representative survey designed to assess oncologists' current and potential use of genomic testing, to inform the development of interventions to facilitate optimal use of genomic testing and to improve patient-physician discussions of the risks, possible benefits, and uncertainties surrounding the use of these tests. Current knowledge of this topic is limited as there are no nationally-representative studies on this topic to date. There are only two non-federal studies two that have examined physicians' knowledge and attitudes regarding somatic genetic and genomic testing. The survey will be administered by mail and web to medical oncology physicians across the U.S. Non-respondents will be invited to complete a follow-back survey to share their reasons for not participating. The study findings will inform NCI of relevant issues and concerns relating to the application of precision medicine to current and future cancer treatment patterns and practice. This information will also inform the development of new funding initiatives to optimize the use of precision medicine in cancer treatment. Additionally, information collected as part of this survey will be used to develop physician educational materials to address barriers to precision medicine in cancer care delivery.

OMB approval is requested for 2 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 350.

TOTAL 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 hour
Telephone Screener	Receptionists	775	1	3/60	39
Precision Medicine Survey—Pilot Study	Oncology Physicians	175	1	20/60	58
Precision Medicine Survey—Full Study	Oncology Physicians	600	1	20/60	200
Non-response Follow-back Survey	Oncology Physicians	40	1	5/60	3

TOTAL 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 hour
Telephone Reminder Script	Receptionists	600	1	5/60	50
Total		1,375	2,190		350

Dated: February 11, 2016.

Karla Bailey,

Project Clearance Liaison, National Cancer Institute, NIH.

[FR Doc. 2016–04105 Filed 2–25–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

[1651–0008]

Agency Information Collection Activities: Application for Identification Card

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

ACTION: 30-Day notice and request for comments; Extension of an existing collection of information.

SUMMARY: U.S. Customs and Border Protection (CBP) of the Department of Homeland Security will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act: Application for Identification Card (CBP Form 3078). This is a proposed extension of an information collection that was previously approved. CBP is proposing that this information collection be extended with no change to the burden hours or to the information collected. This document is published to obtain comments from the public and affected agencies.

DATES: Written comments should be received on or before March 28, 2016 to be assured of consideration.

ADDRESSES: Interested persons are invited to submit written comments on this proposed information collection to the Office of Information and Regulatory Affairs, Office of Management and Budget. Comments should be addressed to the OMB Desk Officer for Customs and Border Protection, Department of Homeland Security, and sent via electronic mail to *oira_submission@omb.eop.gov* or faxed to (202) 395–5806.

FOR FURTHER INFORMATION CONTACT: Requests for additional information should be directed to Tracey Denning, U.S. Customs and Border Protection, Regulations and Rulings, Office of International Trade, 90 K Street NE., 10th Floor, Washington, DC 20229–1177, at 202–325–0265.

SUPPLEMENTARY INFORMATION: This proposed information collection was previously published in the **Federal Register** (80 FR 66915) on October 30, 2015, allowing for a 60-day comment period. This notice allows for an additional 30 days for public comments. This process is conducted in accordance with 5 CFR 1320.10. CBP invites the general public and other Federal agencies to comment on proposed and/or continuing information collections pursuant to the Paperwork Reduction Act of 1995 (Pub. L. 104–13; 44 U.S.C. 3507). The comments should address: (a) Whether the collection of information is 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 estimates of the burden of the collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden, including the use of automated collection techniques or the use of other forms of information technology; and (e) the annual costs to respondents or record keepers from the collection of information (total capital/startup costs and operations and maintenance costs). The comments that are submitted will be summarized and included in the CBP request for OMB approval. All comments will become a matter of public record. In this document, CBP is soliciting comments concerning the following information collection:

Title: Application for Identification Card.

OMB Number: 1651–0008.

Form Number: CBP Form 3078.

Abstract: CBP Form 3078, *Application for Identification Card,* is filled out in order to obtain an Identification Card which is used to gain access to CBP security areas. This form collects biographical information and is usually completed by licensed Cartmen or Lightermen whose duties require receiving, transporting, or otherwise handling imported merchandise which has not been released from CBP custody. This form is submitted to the local CBP office at the port of entry that the respondent will be requesting access to the Federal Inspection Section. Form 3078 is authorized by 19 U.S.C. 66, 1551, 1555, 1565, 1624, 1641; and 19 CFR 112.42, 118, 122.182, and 146.6. This form is accessible at: *http://www.cbp.gov/sites/default/files/documents/CBP%20Form%203078.pdf.*

Action: CBP proposes to extend the expiration date of this information collection with no change to the estimated burden hours or to CBP Form 3078.

Type of Review: Extension (without change).

Affected Public: Businesses.

Estimated Number of Respondents: 150,000.

Estimated Number of Total Annual Responses: 150,000.

Estimated Time per Response: 17 minutes.

Estimated Total Annual Burden Hours: 42,450.

Dated: February 22, 2016.

Tracey Denning,

Agency Clearance Officer, U.S. Customs and Border Protection.

[FR Doc. 2016–04121 Filed 2–25–16; 8:45 am]

BILLING CODE 9111–14–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

[1651–0014]

Agency Information Collection Activities: Declaration for Free Entry of Unaccompanied Articles

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

ACTION: 60-Day notice and request for comments; extension of an existing collection of information.