1, 2009, through June 30, 2010.[1] The final results of administrative review are currently due August 27, 2011.

Extension of Time Limit of Preliminary Results

Section 751(a)(3)(A) of the Tariff Act of 1930, as amended (the Act), requires that the Department issue final results within 120 days after the date on which the preliminary results are published. However, if it is not practicable to complete the review within this time period, section 751(a)(3)(A) of the Act allows the Department to extend the time period to a maximum of 180 days. Completion of the final results of the administrative review within the 120-day period is not practicable because the Department needs additional time to analyze complex issues regarding affiliation and knowledge of U.S. destination. Given the complexity of these issues, and in accordance with section 751(a)(3)(A) of the Act, we are extending the time period for issuing the final results of this review to 180 days. Therefore, the final results are now due no later than October 26, 2011.

We are publishing this notice pursuant to sections 751(a)(3)(A) and 777(i)(1) of the Act.

Dated: August 19, 2011.

Christian Marsh,

Deputy Assistant Secretary for Antidumping and Countervailing Duty Operations.

[FR Doc. 2011–21833 Filed 8–24–11; 8:45 am]

BILLING CODE 3510–DS–P

DEPARTMENT OF COMMERCE

International Trade Administration

Virginia Polytechnic Institute, et al.; Notice of Decision on Applications for Duty-Free Entry of Scientific Instruments

This is a decision pursuant to Section 6(c) of the Educational, Scientific, and Cultural Materials Importation Act of 1966 (Pub. L. 89–651, as amended by Pub. L. 106–36; 80 Stat. 897; 15 CFR part 301). Related records can be viewed between 8:30 a.m. and 5 p.m. in Room 3720, U.S. Department of Commerce, 14th and Constitution Ave., NW., Washington, DC.

Docket Number: 11–039. *Applicant:* Virginia Polytechnic Institute, Department of Engineering Science and Mechanics, Blacksburg, VA 24061. *Instrument:* Nano test platform. *Manufacturer:* Micro Materials Ltd., United Kingdom. *Intended Use:* See notice at 76 FR 43263, July 20, 2011. *Comments:* None received. *Decision:* Approved. We know of no instruments of equivalent scientific value to the foreign instruments described below, for such purposes as this is intended to be used, that was being manufactured in the United States at the time of its order. *Reasons:* This instrument is unique in that it can support the technical requirements for high temperature nanoindentations, nanoimpact, nanofatigue and wet stage nanoindentation.

Docket Number: 11–040. *Applicant:* University of Colorado at Boulder, Procurement Service Center, Denver, CO 80202. *Instrument:* Low-temperature atomic force microscope. *Manufacturer:* Attocube Systems AG, Germany. *Intended Use:* See notice at 76 FR 43263, July 20, 2011. *Comments:* None received. *Decision:* Approved. We know of no instruments of equivalent scientific value to the foreign instruments described below, for such purposes as this is intended to be used, that was being manufactured in the United States at the time of its order. *Reasons:* This instrument must be compatible with high magnetic fields, which requires a special selection of non-magnetic materials the instrument has to be built from. The low-temperature capability requires special piezoelectric scanners and sample mounting and cooling techniques, unique to this instrument.

Dated: August 22, 2011.

Gregory W. Campbell,

Director, Subsidies Enforcement Office, Import Administration.

[FR Doc. 2011–21757 Filed 8–24–11; 8:45 am]

BILLING CODE 3510–DS–P

CORPORATION FOR NATIONAL AND COMMUNITY SERVICE

Information Collection; Submission for OMB Review, Comment Request

AGENCY: Corporation for National and Community Service.

ACTION: Notice.

SUMMARY: The Corporation for National and Community Service (hereinafter the ''Corporation''), has submitted a public information collection request (ICR) entitled Current Population Survey Civic Engagement Supplement for review and approval in accordance with the Paperwork Reduction Act of 1995, Public Law 104–13, (44 U.S.C. Chapter 35). Copies of this ICR, with applicable supporting documentation, may be obtained by calling the Corporation for National and Community Service, Nathan Dietz, at (202) 606–6633 or e-mail to *ndietz@cns.gov.* Individuals who use a telecommunications device for the deaf (TTY–TDD) may call (202) 606–3472 between 8:30 a.m. and 5 p.m. Eastern Time, Monday through Friday.

ADDRESSES: Comments may be submitted, identified by the title of the information collection activity, to the Office of Information and Regulatory Affairs, Attn: Ms. Sharon Mar, OMB Desk Officer for the Corporation for National and Community Service, by any of the following two methods within 30 days from the date of publication in the **Federal Register**:

(1) *By fax to:* (202) 395–6974, Attention: Ms. Sharon Mar, OMB Desk Officer for the Corporation for National and Community Service; and

(2) *Electronically by e-mail to: smar@omb.eop.gov.*

SUPPLEMENTARY INFORMATION: The OMB is particularly interested in comments which:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the Corporation, including whether the information will have practical utility;
- Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Propose ways to enhance the quality, utility, and clarity of the information to be collected; and
- Propose ways to minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submissions of responses.

Comments

A 60-day public comment Notice was published in the **Federal Register** on June 17, 2011. This comment period ended August 16, 2011. No public comments were received from this Notice.

Description: The Corporation is seeking approval for the Civic Engagement Supplement, which is conducted by the U.S. Census Bureau in conjunction with the annual November Current Population Survey (CPS). The Civic Engagement Supplement provides information on the extent to which American communities are places where individuals are civically active. The Corporation uses the Civic

[1] *See Certain Pasta From Turkey: Notice of Preliminary Results of Antidumping Duty Administrative Review,* 76 FR 23974 (April 29, 2011) (*Preliminary Results*).

Engagement Supplement to collect data for the Civic Health Assessment, an annual report that is mandated by the Serve America Act.

Type of Review: Renewal.

Agency: Corporation for National and Community Service.

Title: Current Population Survey Civic Engagement Supplement.

OMB Number: # 0607–0466 [existing Census clearance number].

Agency Number: None.

Affected Public: Individuals or households.

Total Respondents: 54,000.

Frequency: Annual.

Average Time Per Response: Ten minutes per household.

Estimated Total Burden Hours: 9,000 hours.

Total Burden Cost (capital/startup): None.

Total Burden Cost (operating/maintenance): None.

Dated: August 22, 2011.

John Kim,

Director of Strategic Initiatives, Strategy Office.

[FR Doc. 2011–21734 Filed 8–24–11; 8:45 am]

BILLING CODE 6050–$$–P

DEPARTMENT OF DEFENSE

Office of the Secretary

[Docket ID: DoD–2011–HA–0096]

Proposed Collection; Comment Request

AGENCY: Office of the Assistant Secretary of Defense for Health Affairs, DoD.

ACTION: Notice.

SUMMARY: In compliance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Assistant Secretary of Defense for Health Affairs announces the proposed extension of a public information collection and seeks public comment on the provisions thereof. Comments are invited on: (a) Whether the proposed 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 estimate of the burden of the proposed information collection; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the information collection on respondents, including through the use of automated collection techniques or other forms of information technology.

DATES: Consideration will be given to all comments received by October 24, 2011.

ADDRESSES: You may submit comments, identified by docket number and title, by any of the following methods:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Federal Docket Management System Office, 4800 Mark Center Drive, Suite 02G09, Alexandria, VA 22350–3100.

Instructions: All submissions received must include the agency name, docket number and title for this **Federal Register** document. The general policy for comments and other submissions from members of the public is to make these submissions available for public viewing on the Internet at *http://www.regulations.gov* as they are received without change, including any personal identifiers or contact information.

FOR FURTHER INFORMATION CONTACT: To request more information on this proposed information collection or to obtain a copy of the proposal and associated collection instruments, please write to the Office of the Chief Medical Officer (OCMO), TRICARE Management Activity, ATTN: Ms. Judy George, Skyline 5, Suite 810, 5111 Leesburg Pike, Falls Church, VA 22041–3206, or call OCMO, Patient Safety Division, at (703) 681–0064.

Title; Associated Form; and OMB Number: DoD Patient Safety Survey; OMB Number 0720–0034.

Needs and Uses: The 2001 National Defense Authorization Act contains specific sections addressing patient safety in military and veterans health care systems. This legislation states that the Secretary of Defense shall establish a patient care error reporting and management system to study occurrences of errors in patient care and that one of the purposes of the system should be ''To identify systemic factors that are associated with such occurrences'' and ''To provide for action to be taken to correct the identified systemic factors'' (Sec. 754, items b2 and b3). In addition, the legislation states that the Secretary shall ''Continue research and development investments to improve communication, coordination, and team work in the provision of health care'' (Sec. 754, item d4).

In its ongoing response to this legislation and in support of its mission to ''promote a culture of safety to eliminate preventable patient harm by engaging, educating and equipping patient-care teams to institutionalize evidence-based safe practices,'' the DoD Patient Safety Program plans to field the Tri-Service Patient Safety Culture Survey. The Culture Survey is based on the Department of Health and Human Services' Agency for Healthcare Research and Quality's validated survey instrument. Previously administered in 2005/6 and 2008, the survey obtains MHS staff opinions on patient safety issues such as teamwork, communications, medical error occurrence and response, error reporting, and overall perceptions of patient safety. The purpose of the survey is to assess the current status of patient safety in MHS facilities and to assess patient safety improvement over time. Two versions of the survey will be available for administration. The inpatient survey tool is the same, OMB-approved tool that was administered in previous years. There will also be a corresponding outpatient survey tool, with congruous questions tailored to the ambulatory or clinic setting. Respondents will select the survey corresponding to their care survey.

Affected Public: Federal government; individuals or households.

Annual Burden Hours: 2,337 hours.

Number of Respondents: 14,022.

Responses per Respondent: 1.

Average Burden per Response: 10 minutes.

Frequency: On occasion.

SUPPLEMENTARY INFORMATION: Respondent's obligation—voluntary.

Summary of Information Collection

The Web-based survey will be administered on a voluntary-basis to all staff working in Army, Navy, and Air Force Military Health System (MHS) direct care facilities in the U.S. and internationally, including Military Treatment Facility (MTF) hospitals as well as ambulatory and dental services. Responses and respondents will remain anonymous. There are two versions of the survey that may be administered, corresponding to the setting in which care is delivered, either Hospital (inpatient) or Ambulatory (outpatient/clinic setting).

Dated: August 22, 2011.

Aaron Siegel,

Alternate OSD Federal Register Liaison Officer, Department of Defense.

[FR Doc. 2011–21744 Filed 8–24–11; 8:45 am]

BILLING CODE 5001–06–P