

Period of Review

Pursuant to 19 CFR 351.214(c), an exporter or producer may request a new shipper review within one year of the date on which its subject merchandise was first entered. Moreover, 19 CFR 351.214(d)(1) states that if the request for the review is made during the six-month period ending with the end of the semiannual anniversary month, the Secretary will initiate a new shipper review in the calendar month immediately following the semiannual anniversary month. Further, 19 CFR 351.214(g)(1)(i)(B) states that if the new shipper review was initiated in the month immediately following the semiannual anniversary month, the POR will be the six-month period immediately preceding the semiannual anniversary month. Within one year of the dates on which their multilayered wood flooring was first entered Zhangshi and Muyun made the requests for new shipper reviews in June, which is the semiannual anniversary month of the *Order*. Therefore, the Secretary must initiate these reviews in July and the POR is December 1, 2014, through May 31, 2015.

Initiation of New Shipper Reviews

Pursuant to section 751(a)(2)(B) of the Act and 19 CFR 351.214(b), and the information on the record, the Department finds that the requests submitted by Zhangshi and Muyun meet the threshold requirements for initiation of new shipper reviews for the shipments of multilayered wood flooring from the PRC produced and exported by these companies.⁹ However, if the information supplied by Zhangshi and Muyun is later found to be incorrect or insufficient during the course of this proceeding, the Department may rescind the review or apply adverse facts available pursuant to section 776 of the Act, depending upon the facts on record. The Department intends to issue the preliminary results of these new shipper reviews no later than 180 days from the date of initiation, and the final results no later than 90 days from the issuance of the preliminary results.¹⁰

It is the Department's usual practice, in cases involving non-market economies, to require that a company seeking to establish eligibility for an

antidumping duty rate separate from the country-wide rate provide evidence of *de jure* and *de facto* absence of government control over the company's export activities. Accordingly, the Department will issue questionnaires to Zhangshi and Muyun which will include a section requesting information with regard to these companies' export activities for separate rates purposes. The review of each exporter will proceed if the response provides sufficient indication that it is not subject to either *de jure* or *de facto* government control with respect to its export of subject merchandise.

The Department will instruct CBP to allow, until the completion of the review, at the option of the importer, the posting of a bond or security in lieu of a cash deposit for each entry of the subject merchandise from Zhangshi and Muyun, in accordance with section 751(a)(2)(B)(iii) of the Act and 19 CFR 351.214(e). Because Zhangshi and Muyun certified that they produced and exported the subject merchandise, the Department will apply the bonding privilege only for subject merchandise that the respondent both produced and exported. To assist in its analysis of the *bona fides* of Zhangshi and Muyun's sales, upon initiation of this NSR, the Department will require Zhangshi and Muyun to submit on an ongoing basis complete transaction information concerning any sales of subject merchandise to the United States that were made subsequent to the POR.

Interested parties requiring access to proprietary information in these new shipper reviews should submit applications for disclosure under administrative protective order in accordance with 19 CFR 351.305 and 19 CFR 351.306.

This initiation and notice are in accordance with section 751(a)(2)(B) of the Act and 19 CFR 351.214 and 19 CFR 351.221(c)(1)(i).

Dated: July 21, 2015.

Christian Marsh,

Deputy Assistant Secretary for Antidumping and Countervailing Duty Operations.

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DEPARTMENT OF COMMERCE

National Institute of Standards and Technology

Judges Panel of the Malcolm Baldrige National Quality Award

AGENCY: National Institute of Standards and Technology, Department of Commerce.

ACTION: Notice of closed meeting.

SUMMARY: The Judges Panel of the Malcolm Baldrige National Quality Award (Judges Panel) will meet in closed session on Wednesday, August 26, 2015, from 9:00 a.m. until 3:30 p.m. Eastern Time. The purpose of this meeting is to review the results of examiners' scoring of written applications. Panel members will vote on which applicants merit site visits by examiners to verify the accuracy of quality improvements claimed by applicants. The meeting is closed to the public in order to protect the proprietary data to be examined and discussed at the meeting.

DATES: The meeting will be held on Wednesday, August 26, 2015, from 9:00 a.m. until 3:30 p.m. Eastern Time. The entire meeting will be closed to the public.

ADDRESSES: The meeting will be held at the National Institute of Standards and Technology, 100 Bureau Drive, Gaithersburg, MD 20899.

FOR FURTHER INFORMATION CONTACT: Robert Fangmeyer, Director, Baldrige Performance Excellence Program, National Institute of Standards and Technology, 100 Bureau Drive, Mail Stop 1020, Gaithersburg, Maryland 20899-1020, telephone number (301) 975-2360, email robert.fangmeyer@nist.gov.

SUPPLEMENTARY INFORMATION:

Authority: 15 U.S.C. 3711a(d)(1) and the Federal Advisory Committee Act, as amended, 5 U.S.C. App.

Pursuant to the Federal Advisory Committee Act, as amended, 5 U.S.C. App., notice is hereby given that the Judges Panel of the Malcolm Baldrige National Quality Award will meet on Wednesday, August 26, 2015, from 9:00 a.m. until 3:30 p.m. Eastern Time. The Judges Panel is composed of twelve members, appointed by the Secretary of Commerce, chosen for their familiarity with quality improvement operations and competitiveness issues of manufacturing companies, services companies, small businesses, health care providers, and educational institutions. Members are also chosen who have broad experience in for-profit and nonprofit areas. The purpose of this meeting is to review the results of examiners' scoring of written applications. Panel members will vote on which applicants merit site visits by examiners to verify the accuracy of quality improvements claimed by applicants. The meeting is closed to the public in order to protect the

concurrently with this notice; Memorandum to the File entitled, "Initiation of Antidumping New Shipper Review of Multilayered Wood Flooring from the People's Republic of China: Huzhou Muyun Wood Co., Ltd." ("Muyun Initiation Checklist") dated concurrently with this notice.

⁹ See Zhangshi Initiation Checklist; see also Muyun Initiation Checklist.

¹⁰ See section 751(a)(2)(B)(iv) of the Act.

proprietary data to be examined and discussed at the meeting.

The Chief Financial Officer and Assistant Secretary for Administration, with the concurrence of the Acting, Assistant General Counsel for Administration, formally determined on May, 19 2015, pursuant to Section 10(d) of the Federal Advisory Committee Act, as amended by Section 5(c) of the Government in Sunshine Act, Public Law 94–409, that the meeting of the Judges Panel may be closed to the public in accordance with 5 U.S.C. 552b(c)(4) because the meeting is likely to disclose trade secrets and commercial or financial information obtained from a person which is privileged or confidential and 5 U.S.C. 552b(c)(9)(B) because for a government agency the meeting is likely to disclose information that could significantly frustrate implementation of a proposed agency action. The meeting, which involves examination of current Award applicant data from U.S. organizations and a discussion of these data as compared to the Award criteria in order to recommend Award recipients, will be closed to the public.

Richard R. Cavanagh,

Acting Associate Director for Laboratory Programs.

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DEPARTMENT OF COMMERCE

National Institute of Standards and Technology

Genome in a Bottle Consortium—Progress and Planning Workshop

AGENCY: National Institute of Standards & Technology (NIST), Commerce.

ACTION: Notice of public workshop.

SUMMARY: NIST announces the Genome in a Bottle Consortium meeting to be held on Thursday and Friday, August 27 and 28, 2015. The Genome in a Bottle Consortium is developing the reference materials, reference methods, and reference data needed to assess confidence in human whole genome variant calls. A principal motivation for this consortium is to enable performance assessment of sequencing and science-based regulatory oversight of clinical sequencing. The purpose of this meeting is to update participants about progress of the consortium work, continue to get broad input from individual stakeholders to update or refine the consortium work plan, continue to broadly solicit consortium membership from interested

stakeholders, and invite members to participate in work plan implementation. Topics of discussion at this meeting will include progress and planning of the Analysis Group, which is analyzing and integrating the large variety of sequencing data for four candidate NIST Reference Materials, as well as potential future Reference Materials.

DATES: The Genome in a Bottle Consortium meeting will be held on Thursday, August 27, 2015 from 9:00 a.m. to 5:30 p.m. Eastern Time and Friday, August 28, 2015 from 9:00 a.m. to 12:45 p.m. Eastern Time. Attendees must register by 5:00 p.m. Eastern Time on Thursday, August 20, 2015.

ADDRESSES: The meeting will be held in the Green Auditorium, Building 101, National Institute of Standards and Technology, 100 Bureau Drive, Gaithersburg, MD 20899. Please note admittance instructions under the **SUPPLEMENTARY INFORMATION** section of this notice.

FOR FURTHER INFORMATION CONTACT: For further information contact Justin Zook by email at jzook@nist.gov or by phone at (301) 975–4133 or Marc Salit by email at salit@nist.gov or by phone at (650) 350–2338. To register, go to: https://www-s.nist.gov/CRS/conf_disclosure.cfm?&conf_id=8473.

SUPPLEMENTARY INFORMATION: Clinical application of ultra high throughput sequencing (UHTS) for hereditary genetic diseases and oncology is rapidly growing. At present, there are no widely accepted genomic standards or quantitative performance metrics for confidence in variant calling. These standards and quantitative performance metrics are needed to achieve the confidence in measurement results expected for sound, reproducible research and regulated applications in the clinic. On April 13, 2012, NIST convened the workshop “Genome in a Bottle” to initiate a consortium to develop the reference materials, reference methods, and reference data needed to assess confidence in human whole genome variant calls (www.genomeinabottle.org). On August 16–17, 2012, NIST hosted the first large public meeting of the Genome in a Bottle Consortium, with about 100 participants from government, academic, and industry. This meeting was announced in the **Federal Register** (77 FR 43237) on July 24, 2012. A principal motivation for this consortium is to enable science-based regulatory oversight of clinical sequencing.

At the August 2012 meeting, the consortium established work plans for

four technical working groups with the following responsibilities:

(1) Reference Material (RM) Selection and Design: Select appropriate sources for whole genome RMs and identify or design synthetic DNA constructs that could be spiked-in to samples for measurement assurance.

(2) Measurements for Reference Material Characterization: Design and carry out experiments to characterize the RMs using multiple sequencing methods, other methods, and validation of selected variants using orthogonal technologies.

(3) Bioinformatics, Data Integration, and Data Representation: Develop methods to analyze and integrate the data for each RM, as well as select appropriate formats to represent the data.

(4) Performance Metrics and Figures of Merit: Develop useful performance metrics and figures of merit that can be obtained through measurement of the RMs.

The products of these technical working groups will be a set of well-characterized whole genome and synthetic DNA RMs along with the methods (documentary standards) and reference data necessary for use of the RMs. These products will be designed to help enable translation of whole genome sequencing to regulated clinical applications. The pilot NIST whole genome RM was released in May 2015 and is available at <http://tinyurl.com/giabpilot>. The consortium is currently analyzing and integrating data from two trios that are candidate NIST RMs. The consortium meets in workshops two times per year, in January at Stanford University in Palo Alto, CA, and in August at the National Institute of Standards and Technology in Gaithersburg, MD. At these workshops, including the last meetings at Stanford in January 2015 and at NIST in August 2014, participants in the consortium have discussed progress developing well-characterized genomes for NIST Reference Materials and planned future experiments and analysis of these genomes (see <https://federalregister.gov/a/2012-18064>, <https://federalregister.gov/a/2013-18934>, <https://federalregister.gov/a/2014-18841> and <https://federalregister.gov/a/2015-01158> for past workshops at NIST and Stanford). The January 2015 meeting was announced in the **Federal Register** (80 FR 3220) on January 22, 2015, and the meeting is summarized at <https://docs.google.com/document/d/19j6YDg1MH1iD-8Q8mmV9L7wHOfuyUC3aogctZ2Nh87U/edit?usp=sharing>.

There is no cost for participating in the consortium. No proprietary