destruction of APO materials or conversion to judicial protective order is hereby requested. Failure to comply with the regulations and terms of an APO is a violation which is subject to sanction.

This determination is published pursuant to sections 705(d) and 777(i) of the Act.

Dated: June 12, 2009.

Ronald K. Lorentzen,

Acting Assistant Secretary for Import Administration.

APPENDIX

List of Comments and Issues in the Decision Memorandum

Comment 1: Application of CVD Law to a Country that the Department Treats as an NME in a Parallel AD Investigation
Comment 2: Double Counting/ Overlapping Remedies
Comment 3: Cut–off Date for Countervailing Subsidies
Comment 4: Discount Rate Used for Benefit Calculations
Comment 5: Public Authority Status of Hot–Rolled Steel Producer
Comment 6: Preferential Tax Policies for Enterprises with Foreign Investment (Two Free, Three Half Program)
Comment 7: Refund of Enterprise Income Taxes on FIE Profits Reinvested in an Export Oriented Enterprise
Comment 8: Import Tariff and VAT Exemptions for Encouraged Industries Importing Equipment for Domestic Operations
Comment 9: Export Incentive Payments Characterized as ''VAT Rebates''
Comment 10: Amortization of Startup Costs in the PRC Tax Law
Comment 11: Calculation of the All Others Rate
Comment 12: Whether to Clarify the Scope Language for Hitches

[FR Doc. E9–14471 Filed 6–18–09; 8:45 am]

BILLING CODE 3510–DS–S

DEPARTMENT OF COMMERCE

International Trade Administration

The Manufacturing Council: Meeting

AGENCY: International Trade Administration, U.S. Department of Commerce.

ACTION: Notice of an open meeting.

SUMMARY: The Manufacturing Council will hold an introductory meeting with the Secretary of Commerce to discuss topics related to the U.S. manufacturing sector.

DATES: June 23, 2009.

Time: TBD.
Location: TBD.

FOR FURTHER INFORMATION CONTACT: J. Marc Chittum, The Manufacturing Council, Room 4043, 1401 Constitution Avenue, NW., Washington, DC 20230; telephone: 202–482–4501; and e-mail: *Marc.Chittum@mail.doc.gov*.

Dated: June 15, 2009.

J. Marc Chittum,

Executive Secretary, The Manufacturing Council.

[FR Doc. E9–14392 Filed 6–18–09; 8:45 am]

BILLING CODE 3510–DR–P

DEPARTMENT OF COMMERCE

National Institute of Standards and Technology

Announcing a Workshop on the Protocol for Lightweight Authentication of Identity (PLAID)

AGENCY: National Institute of Standards and Technology, Commerce.

ACTION: Notice of public workshop.

SUMMARY: The National Institute of Standards and Technology (NIST), in cooperation with the Australian Federal Government agency, Centrelink, will hold a public workshop on July 13–15, 2009, at the NIST Gaithersburg campus. The workshop is open to the public but requires registration. The goal of the 3-day workshop is to explore potential commercial implementations of the Protocol for Lightweight Authentication of Identity (PLAID) and the potential usefulness of this protocol to U.S. Federal agencies.

PLAID public resources are available from the following site: *http://www.govdex.gov.au.*

The principle of the workshop is that each attending vendor has two days, with the assistance of Centrelink, to ''port'' existing source code, or develop new code for their device or card. All the information vendors need will be available to them ahead of the workshop on the *http://www.govdex.gov.au site.* Vendors may obtain as much or as little assistance as they please from other vendors in the workshop. (Card vendors might, for instance, assist device vendors.)

On the third day, end-users are invited to view the efforts of all of the vendors. Each vendor will use their space to demonstrate their PLAID implementation to attendees. Attendees are invited to interact with the vendors and discuss their PLAID implementation.

DATES: The workshop will be held on July 13–15, 2009, 9 a.m. till 5 p.m.

ADDRESSES: The workshop will be held in the Employees' Lounge and the Poster Hallway in the Administration Building on the NIST Gaithersburg campus, 100 Bureau Drive, Gaithersburg, Maryland 20899. Please note registration and admittance instructions under the **SUPPLEMENTARY INFORMATION** section of this notice.

FOR FURTHER INFORMATION CONTACT: Tanya Brewer, T: (301) 975–4534, E: *tbrewer@nist.gov.*

SUPPLEMENTARY INFORMATION:

The National Institute of Standards and Technology (NIST), in cooperation with the Australian Federal Government agency, Centrelink, will hold a public workshop on July 13–15, 2009, at the NIST Gaithersburg campus. The workshop is open to the public but requires registration. The goal of the 3-day workshop is to explore potential commercial implementations of the Protocol for Lightweight Authentication of Identity (PLAID) and the potential usefulness of this protocol to U.S. Federal agencies.

PLAID public resources are available from the following site: *http://www.govdex.gov.au.* The principle of the workshop is that each attending vendor has two days, with the assistance of Centrelink, to ''port'' existing source code, or develop new code for their device or card. All the information vendors need will be available to them ahead of the workshop on the *www.govdex.gov.au* site. Vendors may obtain as much or as little assistance as they please from other vendors in the workshop. (Card vendors might, for instance, assist device vendors.)

On the third day, end-users are invited to view the efforts of all of the vendors. Each vendor will use their space to demonstrate their PLAID implementation to attendees. Attendees are invited to interact with the vendors and discuss their PLAID implementation.

This workshop is open to the public, but all attendees, vendors and others, must pre-register in advance. All visitors to the NIST campus are required to register in advance. No late or same-day registrations will be accepted for this reason. All attendees must present a government-issued ID when gaining access to the campus. There is no registration fee. Please submit your name, time of arrival, e-mail address and phone number to Tanya Brewer or Annie Sokol, and one of them will provide you with further logistics information. Non-U.S. citizens must also submit their country of citizenship, title, employer/sponsor, and address. The registration deadline is July 7, 2009. Tanya Brewer's e-mail address is

tbrewer@nist.gov, and Annie Sokol's is *annie.sokol@nist.gov.*

Dated: June 15, 2009.

Patrick Gallagher,

Deputy Director.

[FR Doc. E9–14459 Filed 6–18–09; 8:45 am]

BILLING CODE 3510–13–P

DEPARTMENT OF COMMERCE

National Oceanic and Atmospheric Administration

RIN 0648–XP45

2009 European Union Export Certification for Fishery Products

AGENCY: Seafood Inspection Program (SIP), National Marine Fisheries Service (NMFS), National Oceanic and Atmospheric Administration (NOAA), Commerce.

ACTION: Notice of procedural change.

SUMMARY: The National Oceanic and Atmospheric Administration Seafood Inspection Program (NOAA SIP) will become the sole certifying agency for all fish and fishery products for export to European Union (EU) or European Free Trade Association (EFTA) member countries. Due to the large volume of demand for these certificates and the need for expedient service, SIP, through this notice, is announcing a change from current practices, including fee structure, for providing Health Certificates for the EU and EFTA.

DATES: Effective June 16, 2009.

FOR FURTHER INFORMATION CONTACT: Timothy Hansen, *Timothy.hansen@noaa.gov*, Program Director SIP NMFS/NOAA (301) 713–2355 EXT. 214

SUPPLEMENTARY INFORMATION:

Background

On January 15, 2009 (74 FR 2600), the U.S. Food and Drug Administration (FDA) published a **Federal Register** Notice announcing that after February 17, 2009, FDA will no longer issue health certificates required by the EU for export of fish or fishery products to the EU or the EFTA. By subsequent notice in the **Federal Register** on February 11, 2009 (74 FR 6902), FDA announced a 120-day delay in the effective date of the January 15, 2009 notice. FDA now intends to cease issuing EU Health Certificates on June 17, 2009. The U. S. Department of Commerce Seafood Inspection Program will continue to issue these certificates upon request on a fee-for-service basis.

The Seafood Inspection Program of the National Marine Fisheries Service (NMFS), National Oceanic and Atmospheric Administration (NOAA), Department of Commerce, operating under authority of the Agricultural Marketing Act (7 U.S.C. 1621 *et seq.*) and the Fish and Wildlife Act (16 U.S.C. 742a *et seq.*), is responsible for the development and advancement of commercial grade standards for fishery products and better health and sanitation standards in the industry and for furnishing inspection, evaluation, analytical, grading, and certification services to interested parties. Its major purpose is to encourage and assist the industry in improving the quality, wholesomeness, safety, proper labeling, and marketability of its products.

In 1993, the EU began requiring health certificates for fish and fishery products that entered the EU. Both the FDA and SIP were recognized by the EU as competent U.S. Government authorities and acceptable sources for EU health certificates. The EU also required that shippers to the EU be on a list of firms that demonstrated compliance with the U.S. food safety laws and regulations. Since 1993, FDA has issued health certificates for seafood processing firms appearing on the EU Export Certificate List free of charge. By contrast, SIP examined the product and labeling, confirmed all the shipping information and issued health certificates on a fee for service basis. FDA initially issued approximately 3000 certificates per year, but as European demand for U.S. fishery products increased over the years, the number of certificates issued annually by FDA has grown ten-fold to over 30,000. FDA currently issues about 80 percent of all EU health certificates. The increased volume of certificates issued and concomitant decrease in agency resources has made FDA reassess its involvement in the issuance of EU health certificates.

New Procedures for Receiving EU Certificates From SIP

Effective immediately, SIP policy is as follows: SIP, upon request, will issue EU Health Certificates to SIP program participants and rely on inspection results or an approved control system, e.g. the Hazard Analysis and Critical Control Points Quality Management Program (HACCP QMP) or the Integrated Quality Assurance (IQA) Program, to issue the certificate. Seafood processors and other entities that are not SIP program participants may receive EU Health Certificates from SIP based on a periodic verification of the information provided, compliance of the product labeling to EU requirements and the condition of the product. Instructions for requesting an EU Health Certificate can be found on the SIP Web site at:*http://www.seafood.nmfs.noaa.gov*.

All applicants for EU Health Certificates must be in regulatory good standing with the FDA and must be on the FDA's EU Export Certificate List. In addition, prior to the issuance of EU Health Certificates, all applicants will be required to sign an agreement including, but not limited to, the following provisions:

- The applicant agrees to allow SIP auditors or EC Food and Veterinary auditors entrance to the processing facility at reasonable times when periodic audits occur.
- The applicant agrees to keep information about the origin of foreign raw material to ensure that it was produced in a firm and country that are approved by the EC, make this information available to SIP auditors upon request and provide this information for each certificate request when foreign product is to be certified by SIP.
- The applicant acknowledges that s/he has read the terms and conditions of the agreement and understands that making false statements in connection with issuance of an EU Health Certificate would be a violation of 7 U.S.C.1622(h), punishable by a fine of not more than $1,000 or imprisonment for not more than one year, or both.

Fee Structure

Program Participants

For participants in SIP's continuous on-site inspection service program, certificates will be provided at no extra cost assuming that the work demands can be adequately addressed in the agreed upon contract hours. If additional time is needed for EU Health Certificate completion, it will be charged at the appropriate hourly rate, published on the SIP Web site. EU Health Certificates for facilities operating under the HACCP QMP or the IQA Program will be charged $50 for each EU Health Certificate request. Participants may choose to contract specifically for EU Health Certificate services if there is a significant volume.

Non-Program Participants

Seafood processors and other entities that are not SIP program participants will be charged $69 for each EU Health Certificate request. Fees and charges may be adjusted as necessary to recover costs. Changes in this fee structure will be announced via notice in the **Federal Register**.