Reclamation's publication of its Buy American decision is required pursuant to the Buy American Act, 2 CFR 176.80(b)(2).

Upon publication of this **Federal Register** notice, Reclamation is notifying the public of the decision to approve the Buy American waiver requested by the DRC to purchase foreign ductile iron flanges as part of the American Recovery and Reinvestment Act of 2009 (ARRA) grant for the TSID Phase III Main Canal piping project located in Sisters, Oregon.

Dated: May 20, 2011.

Grayford F. Payne,

Deputy Commissioner—Policy, Administration and Budget, Bureau of Reclamation.

[FR Doc. 2011–12997 Filed 5–26–11;8:45 am]

BILLING CODE 4310–MN–P

INTERNATIONAL TRADE COMMISSION

Investigation Nos. [731–TA–1186–1187] (Preliminary)

Certain Stilbenic Optical Brightening Agents From China and Taiwan

Determinations

On the basis of the record [1] developed in the subject investigation, the United States International Trade Commission (Commission) determines, pursuant to section 733(a) of the Tariff Act of 1930 (19 U.S.C. 1673b(a)) (the Act), that there is a reasonable indication that an industry in the United States is materially injured by reason of imports from China and Taiwan of certain stilbenic optical brightening agents, provided for in subheadings 3204.20.80, 2933.69.6050, 2921.59.40, and 2921.59.8090 of the Harmonized Tariff Schedule of the United States, that are alleged to be sold in the United States at less than fair value (LTFV).

Commencement of Final Phase Investigations

Pursuant to section 207.18 of the Commission's rules, the Commission also gives notice of the commencement of the final phase of its investigations. The Commission will issue a final phase notice of scheduling, which will be published in the **Federal Register** as provided in section 207.21 of the Commission's rules, upon notice from the Department of Commerce (Commerce) of affirmative preliminary determinations in the investigation under section 733(b) of the Act, or, if the preliminary determinations are negative, upon notice of affirmative final determinations in the investigations under section 735(a) of the Act. Parties that filed entries of appearance in the preliminary phase of the investigations need not enter a separate appearance for the final phase of the investigations. Industrial users, and, if the merchandise under investigation is sold at the retail level, representative consumer organizations have the right to appear as parties in Commission antidumping and countervailing duty investigations. The Secretary will prepare a public service list containing the names and addresses of all persons, or their representatives, who are parties to the investigations.

Background

On March 31, 2011, a petition was filed with the Commission and Commerce by Clariant Corp., Charlotte, NC, alleging that an industry in the United States is materially injured by reason of LTFV imports of certain stilbenic optical brightening agents from China and Taiwan. Accordingly, effective March 31, 2011, the Commission instituted antidumping duty investigation Nos. 731–TA–1186–1187 (Preliminary).

Notice of the institution of the Commission's investigations and of a public conference to be held in connection therewith was given by posting copies of the notice in the Office of the Secretary, U.S. International Trade Commission, Washington, DC, and by publishing the notice in the **Federal Register** of April 7, 2011 (76 FR 19383). The conference was held in Washington, DC, on April 21, 2011, and all persons who requested the opportunity were permitted to appear in person or by counsel.

The Commission transmitted its determinations in these investigations to the Secretary of Commerce on May 16, 2011. The views of the Commission are contained in USITC Publication 4236 (May 2011), entitled *Certain Stilbenic Optical Brightening Agents from China and Taiwan: Investigation Nos. 731–TA–1186–1187 (Preliminary).*

Issued: May 23, 2011.

By order of the Commission.

James R. Holbein,

Secretary to the Commission.

[FR Doc. 2011–13185 Filed 5–26–11; 8:45 am]

BILLING CODE 7020–02–P

[1] The record is defined in section 207.2(f) of the Commission's Rules of Practice and Procedure (19 CFR 207.2(f)).

INTERNATIONAL TRADE COMMISSION

[Investigation No. 337–TA–714]

In the Matter of Certain Electronic Devices With Multi-Touch Enabled Touchpads and Touchscreens; Notice of Request for Statements on the Public Interest

Section 337 of the Tariff Act of 1930 provides that if the Commission finds a violation it shall exclude the articles concerned from the United States:

> Unless, after considering the effect of such exclusion upon the public health and welfare, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, and United States consumers, it finds that such articles should not be excluded from entry.

19 U.S.C. 1337(d)(1). A similar provision applies to cease and desist orders. 19 U.S.C. 1337(f)(1).

The Commission is interested in further development of the record on the public interest in its investigations. Accordingly, the parties are invited to file submissions of no more than five (5) pages concerning the public interest in light of the administrative law judge's Recommended Determination on Remedy and Bonding issued in this investigation on April 29, 2011. Comments should address whether issuance of a limited exclusion order and/or a cease and desist order in this investigation could affect the public health and welfare in the United States, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, or United States consumers.

In particular, the Commission is interested in comments that:

(i) Explain how the articles potentially subject to the orders are used in the United States;

(ii) identify any public health, safety, or welfare concerns in the United States relating to the potential orders;

(iii) indicate the extent to which like or directly competitive articles are produced in the United States or are otherwise available in the United States, with respect to the articles potentially subject to the orders;

(iv) indicate whether Complainant, Complainant's licensees, and/or third party suppliers have the capacity to replace the volume of articles potentially subject to an exclusion order and a cease and desist order within a commercially reasonable time; and

(v) indicate whether the limited exclusion order and/or cease and desist

order could impact United States consumers.

Any submissions are due on June 6, 2011.

Issued: May 23, 2011.

By order of the Commission.

James R. Holbein,

Secretary to the Commission.

[FR Doc. 2011–13188 Filed 5–26–11; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 337–TA–760]

In the Matter of Certain Liquid Crystal Display Devices, Products Containing Same, and Methods for Using the Same; Notice of Commission Decision Not To Review an Initial Determination Terminating the Investigation

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has determined not to review the presiding administrative law judge's initial determination (''ID'') (Order No. 9) granting a joint motion to terminate the investigation.

FOR FURTHER INFORMATION CONTACT: Sidney A. Rosenzweig, Office of the General Counsel, U.S. International Trade Commission, 500 E Street, SW., Washington, DC 20436, telephone (202) 708–2532. Copies of non-confidential documents filed in connection with this investigation are or will be available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street, SW., Washington, DC 20436, telephone (202) 205–2000. General information concerning the Commission may also be obtained by accessing its Internet server at *http://www.usitc.gov*. The public record for this investigation may be viewed on the Commission's electronic docket (EDIS) at *http://edis.usitc.gov*. Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on (202) 205–1810.

SUPPLEMENTARY INFORMATION: The Commission instituted this investigation on March 2, 2011, based on a complaint filed by Sharp Corporation of Japan (''Sharp'') that named as respondents: AU Optronics Corp. of Taiwan; AU Optronics Corporation America of Houston, Texas; BenQ America of Irvine, California; BenQ Corporation of Taiwan; Haier America Trading LLC, of New York, New York; Haier Group Company of China; LG Electronics Inc. of South Korea; LG Electronics U.S.A., Inc. of Englewood Cliffs, New Jersey; SANYO Electric Co. of Japan; SANYO North America Corporation of San Diego, California; TCL Corporation of China; TTE Technology, Inc. d/b/a TCL America of Indianapolis, Indiana; and VIZIO, Inc. of Irvine, California. 76 FR 11512 (Mar. 2, 2011). The complaint alleged a violation of section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337, in the importation, sale for importation, and sale within the United States after importation of certain liquid crystal display (''LCD'') devices, products containing same, and methods for using same by reason of the infringement of certain claims of U.S. Patent Nos. 6,879,364; 7,304,626; 7,532,183; 7,283,192; 6,937,300; 7,057,689; and 7,838,881.

On April 21, 2011, Sharp and the AU Optronics respondents (''AUO'') filed a joint motion for termination of the investigation on the basis of settlement and licensing agreements. No other party opposed the motion. The agreements call for Sharp and AUO to terminate the investigation and to dismiss parallel district court proceedings. The other respondents make or sell products that contain accused AUO LCD components, and the settlement between Sharp and AUO thereby resolved all disputes in the investigation.

On May 3, 2011, the ALJ granted the motion as an ID (Order No. 9).

No petitions for review of the ID were filed. The Commission has determined not to review the ID.

The authority for the Commission's determination is contained in section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in Part 210 of the Commission's Rules of Practice and Procedure (19 CFR part 210).

By order of the Commission.

Issued: May 23, 2011.

James R. Holbein,

Secretary to the Commission.

[FR Doc. 2011–13189 Filed 5–26–11; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(i), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in schedule I or II, and prior to issuing a regulation under 21 U.S.C. 952(a)(2), authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with 21 CFR 1301.34(a), this is notice that on March 14, 2011, Almac Clinical Services Inc. (ACSI), 25 Fretz Road, Souderton, Pennsylvania 18964, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the following basic classes of controlled substances:

Drug	Schedule
Oxycodone (9143)	II
Hydromorphone (9150)	II
Tapentadol (9780)	II
Fentanyl (9801)	II

The company plans to import small quantities of the listed controlled substances in dosage form to conduct clinical trials.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43, and in such form as prescribed by 21 CFR 1316.47.

Any such comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, **Federal Register** Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than June 27, 2011.

This procedure is to be conducted simultaneously with, and independent of, the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice published in the **Federal Register** on September 23, 1975, 40 FR 43745–46, all applicants for registration to import the basic class of any controlled substance in schedule I or II are, and will continue to be, required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a); 21 U.S.C. 823(a); and 21 CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.