Commission's notice of investigation named as respondents Aker BioMarine AS of Oslo, Norway; Aker BioMarine Antarctic USA, Inc. of Issaquah, Washington; Aker BioMarine Antarctic AS of Stamsund, Norway (collectively, ''the Aker Respondents''); Olympic Seafood AS of Fosnavag, Norway; Olympic Biotec Ltd. of New Zealand; Avoca, Inc. of Merry Hill, North Carolina; Rimfrost USA, LLC of Merry Hill, North Carolina; Bioriginal Food & Science Corp. of Saskatoon, Saskatchewan, Canada (collectively, ''the Olympic Respondents''); Enzymotec Ltd. of Industrial Zone K'far Baruch, Israel; and Enzymotec USA, Inc. of Morristown, New Jersey (collectively, ''the Enzymotec Respondents'').

The Olympic Respondents were terminated from the investigation on the basis of a settlement agreement on November 5, 2013 (Order No. 31, affirmed by the Commission on December 17, 2013). The Aker Respondents were terminated from the investigation on the basis of a settlement agreement on December 17, 2013 (Order No. 40, not reviewed by the Commission on January 15, 2014).

On May 2, 2014, Complainants and the Enzymotec Respondents filed a joint motion to terminate the investigation based on a settlement and license agreement. On May 13, 2014, the ALJ issued the subject ID (Order No. 48) granting the joint motion to terminate the investigation. No petitions for review were filed.

After considering the subject ID and the relevant portions of the record, the Commission has determined not to review the subject ID. The Commission agrees with the ALJ that the joint motion to terminate the investigation complies with the Commission's rules for termination and that the settlement does not adversely affect the public health and welfare, competitive conditions in the U.S. economy, the production of like or directly competitive articles in the United States, and U.S. consumers.

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: June 12, 2014.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2014–14147 Filed 6–16–14; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 731–TA–1021 (Second Review)]

Malleable Iron Pipe Fittings From China Scheduling of an Expedited Five-Year Review

AGENCY: United States International Trade Commission.

ACTION: Notice.

SUMMARY: The Commission hereby gives notice of the scheduling of an expedited review pursuant to section 751(c)(3) of the Tariff Act of 1930 (19 U.S.C. 1675(c)(3)) (the Act) to determine whether revocation of the antidumping duty order on malleable iron pipe fittings from China would be likely to lead to continuation or recurrence of material injury within a reasonably foreseeable time. For further information concerning the conduct of this review and rules of general application, consult the Commission's Rules of Practice and Procedure, part 201, subparts A through E (19 CFR part 201), and part 207, subparts A, D, E, and F (19 CFR part 207).

DATES: *Effective Date:* June 6, 2014.

FOR FURTHER INFORMATION CONTACT: Christopher J. Cassise (202–708–5408), Office of Investigations, U.S. International Trade Commission, 500 E Street SW., Washington, DC 20436. Hearing-impaired persons can obtain information on this matter by contacting the Commission's TDD terminal on 202–205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its internet server (*http://www.usitc.gov*). The public record for this review may be viewed on the Commission's electronic docket (EDIS) at *http://edis.usitc.gov.*

SUPPLEMENTARY INFORMATION:

Background

On June 6, 2014, the Commission determined that the domestic interested party group response to its notice of institution (79 FR 11819, March 3, 2014) of the subject five-year review was adequate and that the respondent interested party group response was inadequate. The Commission did not find any other circumstances that would warrant conducting a full review.[1] Accordingly, the Commission determined that it would conduct an expedited review pursuant to section 751(c)(3) of the Act.

Staff Report

A staff report containing information concerning the subject matter of the review will be placed in the nonpublic record on July 2, 2014, and made available to persons on the Administrative Protective Order service list for this review. A public version will be issued thereafter, pursuant to section 207.62(d)(4) of the Commission's rules.

Written Submissions

As provided in section 207.62(d) of the Commission's rules, interested parties that are parties to the review and that have provided individually adequate responses to the notice of institution,[2] and any party other than an interested party to the review may file written comments with the Secretary on what determination the Commission should reach in the review. Comments are due on or before July 8, 2014 and may not contain new factual information. Any person that is neither a party to the five-year review nor an interested party may submit a brief written statement (which shall not contain any new factual information) pertinent to the review by July 8, 2014. However, should the Department of Commerce extend the time limit for its completion of the final results of its review, the deadline for comments (which may not contain new factual information) on Commerce's final results is three business days after the issuance of Commerce's results. If comments contain business proprietary information (BPI), they must conform with the requirements of sections 201.6, 207.3, and 207.7 of the Commission's rules. Please be aware that the Commission's rules with respect to electronic filing have been amended. The amendments took effect on November 7, 2011. See 76 Fed. Reg. 61937 (Oct. 6, 2011) and the revised Commission's Handbook on E-Filing, available on the Commission's Web site at *http://edis.usitc.gov.*

In accordance with sections 201.16(c) and 207.3 of the rules, each document filed by a party to the review must be served on all other parties to the review (as identified by either the public or BPI

[1] A record of the Commissioners' votes, the Commission's statement on adequacy, and any individual Commissioner's statements will be available from the Office of the Secretary and at the Commission's Web site.

[2] The Commission has found the responses submitted by Anvil International LLC and Ward Manufacturing to be individually adequate. Comments from other interested parties will not be accepted (*see* 19 CFR 207.62(d)(2)).

service list), and a certificate of service must be timely filed. The Secretary will not accept a document for filing without a certificate of service.

Authority: This review is being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to section 207.62 of the Commission's rules.

By order of the Commission.

Issued: June 12, 2014.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2014–14148 Filed 6–16–14; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 731–TA–991 (Second Review)]

Silicon Metal From Russia

Determination

On the basis of the record [1] developed in the subject five-year review, the United States International Trade Commission (Commission) determines, pursuant to section 751(c) of the Tariff Act of 1930 (19 U.S.C. § 1675(c)), that revocation of the antidumping duty order would be likely to lead to continuation or recurrence of material injury to an industry in the United States within a reasonably foreseeable time.[2]

Background

The Commission instituted this review on June 3, 2013 (78 FR 33064) and determined on September 6, 2013 that it would conduct a full review (78 FR 61384, October 3, 2013). Notice of the scheduling of the Commission's review and of a public hearing 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** on December 19, 2013 (78 FR 76856). The hearing was cancelled, on April 7, 2014 (79 FR 19921, April 10, 2014).

The Commission completed and filed its determination in this review on June 11, 2014. The views of the Commission are contained in USITC Publication 4471 (June 2014), entitled *Silicon Metal from Russia: Investigation No. 731–TA–991 (Second Review).*

By order of the Commission.

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

[2] Commissioner Rhonda K. Schmidtlein did not participate in the vote.

Issued: June 12, 2014.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2014–14146 Filed 6–16–14; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: CATALENT CTS, LLC

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.34(a) on or before July 17, 2014. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before July 17, 2014.

ADDRESS: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, and dispensers of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Deputy Assistant Administrator of the DEA Office of Diversion Control (''Deputy Assistant Administrator'') pursuant to sec. 7(g) of 28 CFR pt. 0, subpt. R, App.

In accordance with 21 CFR 1301.34(a), this is notice that on May 7, 2014, Catalent CTS, LLC, 10245 Hickman Mills Drive, Kansas City, Missouri 64137, applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
Marihuana (7360)	I
Poppy Straw Concentrate (9670)	II

The company plans to import a finished pharmaceutical product containing cannabis extracts in dosage form for a clinical trial study.

In reference to drug code 7360, the company plans to import a synthetic cannabidiol. This compound is listed under drug code 7360. No other activity for this drug code is authorized for this registration.

In addition, the company plans to import an ointment for the treatment of wounds which contain trace amounts of the controlled substances normally found in poppy straw concentrate for packaging and labeling to be used in clinical trials.

Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007).

Dated: June 10, 2014.

Joseph T. Rannazzisi,

Deputy Assistant Administrator.

[FR Doc. 2014–14123 Filed 6–16–14; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: ALKERMES GAINESVILLE LLC

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.34(a) on or before July 17, 2014. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before July 17, 2014.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, and