

On June 10, 2016, Complainants moved to terminate the investigation based on their withdrawal of the complaint. No party responded to the motion.

On June 14, 2016, the ALJ issued the subject ID, granting Complainants' motion and terminating the investigation. No petitions for review were filed.

The Commission has determined not to review the subject 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: July 7, 2016.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2016-16523 Filed 7-12-16; 8:45 am]

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INTERNATIONAL TRADE COMMISSION

[Investigation No. 731-TA-287 (Second Review)]

Raw In-Shell Pistachios From Iran; Notice of Commission Determination To Conduct a Full Five-Year Review

AGENCY: United States International Trade Commission.

ACTION: Notice.

SUMMARY: The Commission hereby gives notice that it will proceed with a full review pursuant to the Tariff Act of 1930 to determine whether revocation of the antidumping duty order on raw in-shell pistachios from Iran would be likely to lead to continuation or recurrence of material injury within a reasonably foreseeable time. A schedule for the review will be established and announced at a later date.

DATES: *Effective Date:* July 5, 2016.

FOR FURTHER INFORMATION CONTACT: Chris 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>.

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).

SUPPLEMENTARY INFORMATION: On July 5, 2016, the Commission determined that it should proceed to a full review in the subject five-year review pursuant to section 751(c) of the Tariff Act of 1930 (19 U.S.C. 1675(c)). The Commission found that the domestic interested party group response to its notice of institution (81 FR 18882, April 1, 2016) was adequate. Although the Commission received a response to its notice of institution from respondent interested parties, the Commission found that the respondent interested party group response with respect to the reviews on subject imports from Iran was inadequate.¹ The Commission also found that other circumstances warranted conducting a full review. 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.

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: July 7, 2016.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2016-16525 Filed 7-12-16; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: Janssen Pharmaceutical, Inc.

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

¹ Commissioner Broadbent determined that the respondent group response was adequate.

accordance with 21 CFR 1301.34(a) on or before August 12, 2016. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before August 12, 2016.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/ODW, 8701 Morrisette Drive, Springfield, Virginia 22152. Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007).

SUPPLEMENTARY INFORMATION: The Attorney General has delegated her 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, dispensers importers, and exporters 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 section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on April 11, 2016, Janssen Pharmaceutical, Inc., 1440 Olympic Drive, Athens, Georgia 30601 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
Phenylacetone (8501)	II
Thebaine (9333)	II
Poppy Straw Concentrate (9670)	II
Tapentadol (9780)	II

The company plans to import thebaine derivatives (9333) as reference standards. The company plans to import an intermediate form of tapentadol (9780) to bulk manufacture tapentadol for distribution to its customers. The