

cannot guarantee that we will be able to do so.

Authority: 40 CFR 1506.6, 40 CFR 1506.10, 43 CFR 1610.2, 43 CFR 1610.5.

Ruth Welch,

BLM Colorado State Director.

[FR Doc. 2015-07013 Filed 3-26-15; 8:45 am]

BILLING CODE 4130-JB-P

INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 704-TA-1 and 734-TA-1 (Review)]

Sugar from Mexico; Determinations

On the basis of the record¹ developed in the subject reviews, the United States International Trade Commission ("Commission") determines, pursuant to sections 704(h) and 734(h) of the Tariff Act of 1930 (19 U.S.C. 1671c(h) and 1673c(h)) ("the Act"), that agreements the U.S. Department of Commerce ("Commerce") has entered into with Mexican exporters of sugar and the government of Mexico suspending antidumping and countervailing duty investigations concerning sugar from Mexico eliminate completely the injurious effect of subject imports.²

Background

The Commission instituted these investigations effective January 8, 2015, following receipt of a petition filed with the Commission by Imperial Sugar Company ("Imperial"), Sugar Land, Texas and AmCane Sugar LLC ("AmCane"), Taylor, Michigan. The Commission determined that Imperial and AmCane are interested parties who were parties to the underlying investigations at the time the petitions were filed, and consequently are appropriate petitioning parties. Notice of the scheduling of these reviews and of a public oral presentation 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 January 26, 2015 (80 FR 3977). The oral presentation was held in Washington, DC, on February 19, 2015, and all persons who requested the opportunity were permitted to appear in person or by counsel.

The Commission completed and filed its determination in these reviews on

March 24, 2015. The views of the Commission are contained in USITC Publication 4523 (April 2015), entitled *Sugar From Mexico: Investigation Nos. 704-TA-1 and 734-TA-1 (Review)*.

By order of the Commission.

Issued: March 24, 2015.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2015-07071 Filed 3-26-15; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: HOSPIRA

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic class, 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 April 27, 2015. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before April 27, 2015.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODW, 8701 Morrisette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrisette 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, 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 pt. 0, subpt. R, App.

In accordance with 21 CFR 1301.34(a), this is notice that on October 31, 2014, Hospira, 1776 North Centennial Drive, McPherson, Kansas

67460-1247, applied to be registered as an importer of remifentanyl (9739), a basic class of controlled substance listed in schedule II.

The company plans to import remifentanyl for use in dosage form manufacturing.

Dated: March 20, 2015.

Joseph T. Rannazzisi,

Deputy Assistant Administrator.

[FR Doc. 2015-06969 Filed 3-26-15; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: Meda Pharmaceuticals, Inc.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic class, 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 April 27, 2015. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before April 27, 2015.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODXL, 8701 Morrisette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrisette 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, 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 of subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on December 5, 2014, Meda

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 All six Commissioners voted in the affirmative.