(updating metadata, performing well maintenance, and well drilling). The proposal would also require estimates of costs to complete the above tasks and a timeline for planned completion. The proposal will be reviewed by the USGS and the NGWMN Program Board who will make funding recommendations.

Title of Collection: National Ground-Water Monitoring Network Cooperative

Funding Application.

OMB Control Number: 1028–0114. *Form Number:* None.

Type of Review: Extension of a currently approved collection.

Respondents/Affected Public: Multistate, state, or local water-resources agencies who operate groundwater monitoring networks.

Total Estimated Number of Annual Respondents: 30.

Total Estimated Number of Annual Responses: 30.

Estimated Completion Time per Response: 40 hours.

Total Estimated Number of Annual Burden Hours: 1200 hours.

Respondent's Obligation: Mandatory to be considered for funding.

Frequency of Collection: Annually. Total Estimated Annual Non-hour Burden Cost: None.

An agency may not conduct or sponsor and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number.

The authorities for this action are the Paperwork Reduction Act of 1995 (44 U.S.C. 3501, et seq.).

Janice M. Fulford,

 $\label{eq:continuous} Director\ Observing\ Systems\ Division. \\ \mbox{[FR Doc. 2018-04396 Filed 3-2-18; 8:45 am]}$

BILLING CODE 4338-11-P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 731-TA-895 (Third Review)]

Pure Granular Magnesium From China; 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 the Tariff Act of 1930 ("the Act"), that revocation of the antidumping duty order on pure granular magnesium from China would be likely to lead to continuation or recurrence of material injury to an

industry in the United States within a reasonably foreseeable time.

Background

The Commission, pursuant to section 751(c) of the Act (19 U.S.C. 1675(c)), instituted this review on September 1, 2017 (82 FR 41651) and determined on December 5, 2017 that it would conduct an expedited review (83 FR 4269, Ianuary 30, 2018).

The Commission made this determination pursuant to section 751(c) of the Act (19 U.S.C. 1675(c)). It completed and filed its determination in this review on February 27, 2017. The views of the Commission are contained in USITC Publication 4761 (February 2018), entitled *Pure Granular Magnesium from China: Investigation No. 731–TA–895 (Third Review).*

By order of the Commission. Issued: February 27, 2018.

Lisa R. Barton,

Secretary to the Commission. [FR Doc. 2018–04332 Filed 3–2–18; 8:45 am]

BILLING CODE 7020-02-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration [Docket No. DEA-392]

Importer of Controlled Substances Application: Stepan Company

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 on or before April 4, 2018. Such persons may also file a written request for a hearing on the application on or before April 4, 2018.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All request for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 8701 Morrissette 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 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 Assistant Administrator of the DEA Diversion Control Division ("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 January 17, 2018, Stepan Company, Natural Products Department, 100 W Hunter Avenue, Maywood, NJ 07607 applied to be registered as an importer of coca leaves (9040), a basic class of controlled substance listed in Schedule II.

The company plans to import the listed controlled substance in bulk for the manufacture of controlled substances for distribution to its customers.

Dated: February 26, 2018.

Susan A. Gibson,

Deputy Assistant Administrator. [FR Doc. 2018–04406 Filed 3–2–18; 8:45 am] BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: PerkinElmer, 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 on or before April 4, 2018. Such persons may also file a written request for a hearing on the application on or before April 4, 2018.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement

 $^{^1}$ The record is defined in sec. 207.2(f) of the Commission's Rules of Practice and Procedure (19 CFR 207.2(f)).