

Cherokee Indians; Kialegee Tribal Town; Poarch Band of Creeks (previously listed as the Poarch Band of Creek Indians of Alabama); The Muscogee (Creek) Nation; Thlopthlocco Tribal Town; and United Keetoowah Band of Cherokee Indians in Oklahoma (herein referred to as “The Consulted Tribes”).

History and Description of the Remains

At some time prior to February of 1837, human remains representing, at minimum, one individual were removed from the Brakebill Mound site (40KN55) in Knox County, TN, by Professor Gerard Troost (b. 1776–d.1850). Professor Troost was a founding member of the Academy of Natural Sciences in Philadelphia and state geologist for Tennessee (1831–1839). The mound is situated at the junction of the French Broad and Holston Rivers on private land. At some time prior to October of 1838, the human remains were loaned to Dr. Samuel G. Morton for his study of human crania from around the world, and accessioned into his collections between 1839 and 1840. In 1853, Dr. Morton’s collections were formally presented to the Academy of Natural Sciences of Philadelphia, loaned to the University of Pennsylvania Museum of Archaeology and Anthropology in 1966, and formally gifted to the University of Pennsylvania Museum of Archaeology and Anthropology in 1997 (UPM no. 97–606–992). The human remains consist of a cranium representing a single male, over 50 years old. No known individuals were identified.

Archival records and museum documentation do not designate a specific culture for this individual. Published anthropological information indicates that the Brakebill Mound site is a Dallas Phase archeological site dating from 1300 to 1600 CE. Based on consultation information and published ethnographic and anthropological literature, current evidence suggest that the Dallas Phase archeological culture may be associated with the Muscogee Creek and/or Cherokee cultural traditions. Today, these groups are represented by The Consulted Tribes.

Determinations Made by the University of Pennsylvania Museum of Archaeology and Anthropology

Officials of the University of Pennsylvania Museum of Archaeology and Anthropology have determined that:

- Pursuant to 25 U.S.C. 3001(9), the human remains described in this notice represent the physical remains of one

individuals of Native American ancestry.

- Pursuant to 25 U.S.C. 3001(2), there is a relationship of shared group identity that can be reasonably traced between the Native American human remains and The Consulted Tribes.

Additional Requestors and Disposition

Lineal descendants or representatives of any Indian Tribe or Native Hawaiian organization not identified in this notice that wish to request transfer of control of these human remains should submit a written request with information in support of the request to Dr. Julian Siggers, Williams Director, University of Pennsylvania Museum of Archaeology and Anthropology, 3260 South Street, Philadelphia, PA 19104, telephone (215) 898–4050, by September 7, 2017. After that date, if no additional requestors have come forward, transfer of control of the human remains to The Consulted Tribes may proceed.

The University of Pennsylvania Museum of Archaeology and Anthropology is responsible for notifying The Consulted Tribes that this notice has been published.

Dated: June 19, 2017.

Melanie O’Brien,

Manager, National NAGPRA Program.

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INTERNATIONAL TRADE COMMISSION

[Investigation No. 731–TA–1185 (Review)]

Steel Nails From the United Arab Emirates; 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 the Tariff Act of 1930 (“the Act”) to determine whether revocation of the antidumping duty order on steel nails from the United Arab Emirates would be likely to lead to continuation or recurrence of material injury within a reasonably foreseeable time.

DATES: July 7, 2017.

FOR FURTHER INFORMATION CONTACT: Calvin Chang (202–205–3062), 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 (<https://www.usitc.gov>). The public record for this review may be viewed on the Commission’s electronic docket (EDIS) at <https://edis.usitc.gov>.

SUPPLEMENTARY INFORMATION:

Background.—On July 7, 2017, the Commission determined that the domestic interested party group response to its notice of institution (82 FR 16229, April 03, 2017) 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 Tariff Act of 1930 (19 U.S.C. 1675(c)(3)).

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 and B (19 CFR part 201), and part 207, subparts A, D, E, and F (19 CFR part 207).

Staff report.—A staff report containing information concerning the subject matter of the review will be placed in the nonpublic record on August 3, 2017, 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 August 8, 2017 and may not contain new factual

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 Mid Continent Steel & Wire, Inc. to be individually adequate. Comments from other interested parties will not be accepted (*see* 19 CFR 207.62(d)(2)).

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 August 8, 2017. 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. The Commission's rules with respect to filing were revised effective July 25, 2014. See 79 FR 35920 (June 25, 2014), and the revised Commission Handbook on E-filing, available from the Commission's Web site at <https://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 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.

Determination.—The Commission has determined this review is extraordinarily complicated and therefore has determined to exercise its authority to extend the review period by up to 90 days pursuant to 19 U.S.C. 1675(c)(5)(B).

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: August 3, 2017.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2017-16677 Filed 8-7-17; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: Registrants listed below have applied for and been granted registration by the Drug Enforcement Administration (DEA) as importers of various classes of schedule I or II controlled substances.

SUPPLEMENTARY INFORMATION: The companies listed below applied to be registered as importers of various basic classes of controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted and no requests for hearing were submitted for these notices.

Company	FR docket	Published
Cambridge Isotope Laboratories	82 FR 19083	April 25, 2017.
Janssen Ortho LLC	82 FR 19083	April 25, 2017.
Galephar Pharmaceutical Research, Inc.	82 FR 23069	May 19, 2017.
Mallinckrodt LLC	82 FR 23071	May 19, 2017.
Cerilliant Corporation	82 FR 25335	June 1, 2017.

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrants to import the applicable basic classes of schedule I or II controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated each company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying each company's compliance with state and local laws, and reviewing each company's background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the DEA has granted a registration as an importer for schedule I or II controlled substances to the above listed persons.

Dated: August 2, 2017.

Demetra Ashley,

Acting Assistant Administrator.

[FR Doc. 2017-16698 Filed 8-7-17; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Leia A. Frickey, M.D.; Decision and Order

On February 28, 2017, the Assistant Administrator, Diversion Control Division, Drug Enforcement Administration (DEA), issued an Order to Show Cause to Leia A. Frickey, M.D. (Registrant), of New Orleans, Louisiana. The Show Cause Order proposed the revocation of Registrant's Certificate of Registration, the denial of any applications to renew or modify her registration, and the denial of any applications for any other DEA registration on the ground that she lacks "state authority to handle controlled substances" in Louisiana, the State in which she is registered with the DEA. Order to Show Cause, at 1 (citing 21 U.S.C. 824(a)(3)).

With respect to the Agency's jurisdiction, the Show Cause Order alleged that Registrant is registered as a practitioner in schedules II through V, pursuant to DEA Certificate of Registration BF5029574, at the address of 3312 South I-10 Service Road, Metairie, Louisiana. *Id.* The Order also

alleged that this registration does not expire until September 30, 2017. *Id.*

As substantive grounds for the proceeding, the Show Cause Order alleged that on May 6, 2016, the Louisiana State Board of Medical Examiners issued a "Notice of Summary Suspension of Medical License, summarily suspending [Registrant's] medical license." ¹ *Id.* at 1. As a result, the Order alleged that Registrant is "currently without authority to practice medicine or handle controlled substances in . . . Louisiana, the [S]tate in which [she is] registered with the DEA." *Id.* at 2. Thus, based on her "lack of authority to [dispense] controlled substances in . . . Louisiana," the Order asserted that "DEA must revoke" her

¹ The Show Cause Order also alleges that "on July 25, 2016, the Louisiana Board of Pharmacy issued a Notice of Suspension, suspending [Registrant's] Louisiana CDS license, number CDS.024813-MD, effective May 6, 2016." *Id.* at 1-2. Although those exact facts are not reflected in the record, the record does show that on November 16, 2016, the Louisiana State Board of Pharmacy issued an Order that Registrant's "LOUISIANA CONTROLLED SUBSTANCE LICENSE No. 024813 is hereby indefinitely suspended in accordance with the suspension of her medical license by the Louisiana State Board of Medical Examiners on May 6, 2016." See Government Exhibit (GX) 4, at 1.