DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA 950]

Importer of Controlled Substances Application: Meridian Medical Technologies

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Meridian Medical Technologies has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before May 6, 2022. Such persons may also file a written request for a hearing on the application on or before May 6, 2022.

ADDRESSES: The DEA requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to *https://www.regulations.gov* and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on December 6, 2021, Meridian Medical Technologies, 2555 Hermelin Drive, Saint Louis, Missouri 63144, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Morphine	9300	II

The company manufactures a product containing morphine in the United States. The company exports this product to customers around the world. The company has been asked to ensure that its product, which is sold to European customers, meets the standards established by the European Pharmacopeia, administered by the Directorate for the quality of Medicines (EDQM). In order to ensure that its product will meet European specifications, the company seeks to import morphine supplied by EDQM for use as reference standards. No other activity for these drug codes is authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2022–07207 Filed 4–5–22; 8:45 am]

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DEPARTMENT OF JUSTICE

Parole Commission

Sunshine Act Meeting

DATE AND TIME: Thursday April 14, 2022, at 2 p.m.

PLACE: U.S. Parole Commission, 90 K Street NE, 3rd Floor, Washington, DC.

STATUS: Open.

MATTERS TO BE CONSIDERED:

1. Approval of October 14, 2021 Quarterly Meeting minutes.
2. Verbal Pandemic Updates since October Quarterly Meeting from the Acting Chairman, Commissioner, Acting Chief of Staff/Case Operations Administrator, Case Services Administrator, Executive Officer, and General Counsel.
3. Verbal update from Jordana Cunningham regarding RSAT and other treatment programs being utilized.
4. Update on the PAVER program.

CONTACT PERSON FOR MORE INFORMATION: Jacquelyn Graham, Staff Assistant to the Chairman, U.S. Parole Commission, 90 K Street NE, 3rd Floor, Washington, DC 20530, (202) 346–7010.

Dated: April 4, 2022.

Patricia K. Cushwa,

Chairman (Acting), U.S. Parole Commission.

[FR Doc. 2022–07441 Filed 4–4–22; 4:15 pm]

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DEPARTMENT OF LABOR

Employment and Training Administration

Workforce Innovation and Opportunity Act (WIOA) 2021 Lower Living Standard Income Level (LLSIL)

AGENCY: Employment and Training Administration (ETA), Labor.

ACTION: Notice.

SUMMARY: Title I of WIOA requires the U.S. Secretary of Labor (Secretary) to update and publish the LLSIL tables annually, for uses described in the law (including determining eligibility for youth). WIOA defines the term ''low-income individual'' as (*inter alia*) one whose total family annual income does not exceed the higher level of the poverty line or 70 percent of the LLSIL. This issuance provides the Secretary's annual LLSIL for 2022 and references the current 2022 Health and Human Services ''Poverty Guidelines.''

DATES: This notice is effective *April 6, 2022.*

FOR FURTHER INFORMATION CONTACT: Contact Samuel Wright, Department of Labor, Employment and Training Administration, 200 Constitution Avenue NW, Room C–4526, Washington, DC 20210; Telephone: 202–693–2870; Fax: 202–693–3015 (these are not toll-free numbers); Email address: *wright.samuel.e@dol.gov.* Individuals with hearing or speech impairments may access the telephone number above via Text Telephone (TTY/TDD) by calling the toll-free Federal Information Relay Service at 1–877–889–5627 (TTY/TDD).

Federal Youth Employment Program Information: Sara Hastings, Department of Labor, Employment and Training Administration, 200 Constitution Avenue NW, Room N–4464, Washington, DC 20210; Telephone: 202–693–3599; Email: *hastings.sara@dol.gov.* Individuals with hearing or speech impairments may access the telephone number above via TTY by calling the toll-free Federal Information Relay Service at 1–877–889–5627 (TTY/TDD).

SUPPLEMENTARY INFORMATION: The purpose of WIOA is to provide