

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
Morrisette Drive, Springfield, Virginia
22152; and (2) Drug Enforcement
Administration, Attn: DEA Federal
Register Representative/DPW, 8701
Morrisette Drive, Springfield, Virginia
22152. All requests for a hearing should
also be sent to: Drug Enforcement
Administration, Attn: Administrator,
8701 Morrisette 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 animporter 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]

BILLING CODE P**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, 90K 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]

BILLING CODE 4410-31-P**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