

other activities for these drug codes are authorized for this registration.

**Matthew J. Strait,**

*Deputy Assistant Administrator.*

[FR Doc. 2022-04324 Filed 3-1-22; 8:45 am]

**BILLING CODE P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-968]

**Importer of Controlled Substances  
Application: Janssen Pharmaceuticals, Inc.**

**AGENCY:** Drug Enforcement Administration, Justice.

**ACTION:** Notice of application.

**SUMMARY:** Janssen Pharmaceuticals Inc., 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 submit electronic comments on or objections to the issuance of the proposed registration on or before April 1, 2022. Such persons may also file a written request for a hearing on the application on or before April 1, 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 10, 2021, Janssen Pharmaceuticals Inc, 1440 Olympic Drive, Buildings 1-5 and 7-14, Athens, Georgia 30601-1645, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Thebaine .....	9333	II
Poppy Straw Concentrate .....	9670	II
Tapentadol .....	9780	II

The company plans to import intermediates classified under Tapentadol (9780) and Thebaine (9333) for further manufacturing to the controlled substances tapentadol and buprenorphine, respectively, prior to distribution to customers. The company plans to import Poppy Straw Concentrate (9670) to bulk manufacture other controlled substances. No other activity for this drug code 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 J. Strait,**

*Deputy Assistant Administrator.*

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-959]

**Importer of Controlled Substances  
Application: Medi-Physics Inc. dba GE Healthcare**

**AGENCY:** Drug Enforcement Administration, Justice.

**ACTION:** Notice of application.

**SUMMARY:** Medi-Physics Inc. dba GE Healthcare 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 April 1, 2022. Such persons may also file a written request for a hearing on the application on or before April 1, 2022.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also 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.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on December 16, 2021, Medi-Physics Inc. dba GE Healthcare, 3350 North Ridge Avenue, Arlington Heights, Illinois 60004-1412, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Cocaine .....	9041	II
Ecgonine .....	9180	II

The company plans to import derivatives of the listed controlled substances to be used for the manufacture of a diagnostic product and 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 J. Strait,**

*Deputy Assistant Administrator.*

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