Order

Pursuant to 28 CFR 0.100(b) and the authority vested in me by 21 U.S.C. 824(a), I hereby revoke DEA Certificate of Registration No. FH4036667 issued to William S. Husel, D.O. Further, pursuant to 28 CFR 0.100(b) and the authority vested in me by 21 U.S.C. 823(f), I hereby deny any pending application of William S. Husel to renew or modify this registration, as well as any other applications of William S. Husel for an additional registration in Ohio. This Order is effective April 9, 2020.

Dated: January 29, 2020.

Uttam Dhillon,

Acting Administrator.

[FR Doc. 2020–04837 Filed 3–9–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–582]

Bulk Manufacturer of Controlled Substances Application: S&B Pharma, Inc.; Correction

ACTION: Notice of application; correction.

SUMMARY: The Drug Enforcement Administration (DEA) published a document in the **Federal Register** on November 22, 2019, concerning a notice of application. As that document correctly indicated, the applicant, S&B Pharma, Inc., DBA Norac Pharma, 405 South Motor Avenue, Azusa, California 91702–3232 applied to be registered as a bulk manufacturer of a number of controlled substances, to include applying for authorization in order to synthetically manufacture using drug code 7360 (marihuana). However, on the notice of application published, drug code 7360 was inadvertently identified and listed as Gamma Hydroxybutyric Acid instead of Marihuana.

SUPPLEMENTARY INFORMATION:

Correction

In the **Federal Register** of November 22, 2019, in FR Doc. 2019–25402 (84 FR 64563), on page 64564, correct the listing of drug code 7360 to be identified as Marihuana, as is shown below.

Controlled substance	Drug code	Schedule
Marihuana	7360	I

Dated: February 11, 2020.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–04829 Filed 3–9–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–592]

Importer of Controlled Substances Application: Johnson Matthey Inc.

ACTION: Notice of application.

DATES: Registered importers 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 9, 2020. Such persons may also file a written request for a hearing on the application on or before April 9, 2020.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on September 11, 2019, Johnson Matthey Inc., Pharmaceutical Materials, 2003 Nolte Drive, West Deptford, New Jersey 08066–1742 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Coca Leaves	9040	II
Thebaine	9333	II
Opium, raw	9600	II
Noroxymorophone	9668	II
Poppy Straw Concentrate	9670	II
Fentanyl	9801	II

The company plans to import Coca Leaves (9040), Opium, raw (9600), and Poppy Straw Concentrate (9670) in order to bulk manufacture active pharmaceutical ingredients (API) for distribution to its customers. The company plans to also import Thebaine (9333), Noroxymorophone (9668), and Fentanyl (9801) to use as analytical reference standards, both internally and to be sold to their customers to support testing of Johnson Matthey Inc.'s active pharmaceutical ingredients (API's) only.

Dated: February 10, 2020.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–04836 Filed 3–9–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–594]

Importer of Controlled Substances Application: Arizona Department of Corrections

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic class, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before April 9, 2020. Such persons may also file a written request for a hearing on the application on or before April 9, 2020.

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