DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-674]

Bulk Manufacturer of Controlled Substances Application: Purisys, LLC

ACTION: Notice of application.

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 **[INSERT 60 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER]**.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on May 14, 2020, Purisys, LLC, 1550 Olympic Drive Athens, Georgia 30601-1602, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled Substance	Drug Code	Schedule
Gamma-hydroxybutyic acid	2010	I
Marihuana Extract	7350	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Codeine-N-Oxide	9053	I
Dihydromorphine	9145	I
Hydromorphinol	9301	I
Nabilone	7379	II
Codeine	9050	II
Dihydrocodeine	9120	II
Oxycodone	9143	II

Hydromorphone	9150	II
Hydrocodone	9193	II
Levorphanol	9220	II
Morphine	9300	II

The company plans to manufacture 7360, 7370, and 7379 as bulk active pharmaceutical ingredients and manufacture the remaining above-listed controlled substances as analytical reference standards for distribution to customers. The company also plans to use these substances for lab scale research and development activities. In reference to drug codes 7360 and 7370, the company plans to bulk manufacture these as synthetic. No other activities for these drug codes are authorized for this registration.

William T. McDermott,

Assistant Administrator.

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