

VA; Vana Solutions, LLC., Beaver Creek, OH; VG IT Services, Inc., Ashburn, VA; and W5 Technologies, Inc., Scottsdale, AZ have been added as parties to this venture.

Also, 1901 Group LLC, Reston, VA; Alteryx, Inc., Irvine, CA; Aptima, Inc., Woburn, MA; Assured Wireless Corp., San Diego, CA; AT&T Government Solutions, Inc., Vienna, VA; Broadband Antenna Tracking Systems, Inc., Indianapolis, IN; Clemson University, Clemson, SC; Decisive Analytics Corp., Arlington, VA; Dover Microsystems, Inc., Waltham, MA; EPS Corp., Tinton Falls, NJ; Global Planning Initiatives LLC, Virginia Beach, VA; HigherEchelon, Inc., Huntsville, AL; Inonde, McLean, VA; IT Partners, Inc., Herndon, VA; KNC Strategic Services, Oceanside, CA; Kopis Mobile LLC, Flowood, MS; Kudu Dynamics LLC, Chantilly, VA; L3Harris Technologies, Palm Bay, FL; Lexington Solutions Group LLC, Lexington, VA; Motorola Solutions, Inc. US Federal Markets Division, Linthicum, MD; NewSat North America LLC, Indian Harbour Beach, FL; Nobletech Solutions, Huntsville, AL; Poplicus, Inc. dba Govini, Arlington, VA; ProSync Technology Group, Ellicott City, MD; Rincon Research Corp., Tucson, AZ; SafeFlights, Inc. dba 14bis Supply Tracking, Burlington, MA; Shield AI, Inc., San Diego, CA; Shift5, Inc., Rosslyn, VA; Si2 Technologies, Inc., North Billerica, MA; Southern Methodist University, Dallas, TX; Streif Enterprise, Inc. dba ibeeto, El Cajon, CA; Taurean General Services, Boerne, TX; XATOR Corp., Reston, VA; and XR 2 LEAD LLC, Dumfries VA have withdrawn from this venture.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and IWRP intends to file additional written notifications disclosing all changes in membership.

On October 15, 2018, IWRP filed its original notification pursuant to section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to section 6(b) of the Act on October 23, 2018 (83 FR 53499).

The last notification was filed with the Department on April 6, 2022. A notice was published in the **Federal Register** pursuant to section 6(b) of the Act on May 13, 2022 (87 FR 29385).

Suzanne Morris,

Chief, Premerger and Division Statistics Antitrust Division.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1061]

Bulk Manufacturer of Controlled Substances Application: Purisys, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Purisys, LLC, has applied to be registered as a bulk manufacturer 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 October 31, 2022. Such persons may also file a written request for a hearing on the application on or before October 31, 2022.

ADDRESSES: The Drug Enforcement Administration 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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on June 6, 2022, 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 substance(s):

Controlled substance	Drug code	Schedule
lbogaine	7260	I
5-Methoxy-N,N-diisopropyltryptamine.	7439	I
Cocaine	9041	II
Ecgonine	9180	II
Meperidine	9230	II
Meperidine intermediate-A.	9232	II
Meperidine intermediate-B.	9233	II

Controlled substance	Drug code	Schedule
Meperidine intermediate-C.	9234	II

The company plans to bulk manufacture the listed controlled substances for the production of active pharmaceutical ingredients (API) and analytical reference standards for sale to its customers. The company plans to manufacture the above listed controlled substances as clinical trial and starting materials to make compounds for distribution to its customers. No other activities for these drug codes are authorized for this registration.

Kristi O'Malley,

Assistant Administrator.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1069]

Importer of Controlled Substances Application: Caligor Coghlan Pharma Services

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Caligor Coghlan Pharma Services 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 September 29, 2022. Such persons may also file a written request for a hearing on the application on or before September 29, 2022.

ADDRESSES: The Drug Enforcement Administration 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