connection therewith was given by posting copies of the notice in the Office of the Secretary, U.S. International Trade Commission, Washington, DC, and by publishing the notice in the **Federal Register** on November 18, 2022 (87 FR 69338). The Commission conducted its hearing on April 11, 2023. All persons who requested the opportunity were permitted to participate.

The Commission made these determinations pursuant to section 751(c) of the Act (19 U.S.C. 1675(c)). It completed and filed its determinations in these reviews on June 20, 2023. The views of the Commission are contained in USITC Publication 5432 (June 2023), entitled *Frozen Warmwater Shrimp from China, India, Thailand, and Vietnam: Investigation Nos. 731–TA–1064 and 1066–1068 (Third Review).*

By order of the Commission.

Issued: June 20, 2023.

Katherine Hiner,

Acting Secretary to the Commission.

[FR Doc. 2023–13444 Filed 6–23–23; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1221]

Bulk Manufacturer of Controlled Substances Application: American Radiolabeled Chem

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: American Radiolabeled Chem 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 August 25, 2023. Such persons may also file a written request for a hearing on the application on or before August 25, 2023.

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 May 19, 2023, American Radiolabeled Chem, 100 Arc Drive, St. Louis, Missouri 63146–3502, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid	2010	I
Ibogaine	7260	I
Lysergic acid diethylamide	7315	I
Tetrahydrocannabinols	7370	I
Dimethyltryptamine	7435	I
1-[1-(2-Thienyl)cyclohexyl]piperidine	7470	I
Noroxymorphone	9145	I
Heroin	9200	I
Normorphine	9313	I
Amphetamine	1100	II
Methamphetamine	1105	II
Amobarbital	2125	II
Phencyclidine	7471	II
Phenylacetone	8501	II
Cocaine	9041	II
Codeine	90.50	II
Dihydrocodeine	9120	II
Oxycodone	9143	II
Hydromorphone	9150	II
Ecgonine	9180	II
Hydrocodone	9193	II
Meperidine	9230	II
Metazocine	9240	II
Methadone	9250	II
Dextropropoxyphene, bulk (non-dosage forms)	9273	II
Morphine	9300	II
Oripavine	9330	II
Thebaine	9333	II
Oxymorphone	9652	II
Phenazocine	9715	II
Carfentanil	9743	II
Fentanyl	9801	II

The company plans to bulk manufacture the listed controlled substances for the internal use intermediates or for sale to its customers. The company plans to manufacture small quantities of the above-listed controlled substances as radiolabeled compounds for biochemical research. No other activities for these drug codes are authorized for this registration.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2023–13473 Filed 6–23–23; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1203]

Bulk Manufacturer of Controlled Substances Application: Arista Biologicals

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Arista Biologicals 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 August 25, 2023. Such persons may also file a written request for a hearing on the application on or before August 25, 2023.

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 April 6, 2023, Arista Biologicals, 1101 Hamilton Street, Allentown, Pennsylvania 18101–1043, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
4-Anilino-N-Phenethyl-4-Piperidine (ANPP)	8333	II
Norfentanyl (N-phenyl-N-(piperidin-4-yl)propionamide)	8366	II

The company plans to bulk manufacture the listed controlled substances for internal use as intermediates for formulation and analytical development purposes. No other activities for these drug codes are authorized for this registration.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2023–13466 Filed 6–23–23; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1220]

Bulk Manufacturer of Controlled Substances Application: Olon Ricerca Bioscience LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Olon Ricerca Bioscience 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 August 25, 2023. Such persons may also file a written request for a hearing on the application on or before August 25, 2023.

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 May 17, 2023, Olon Ricerca Bioscience LLC, 7528 Auburn Road, Concord Township, Ohio 44077–9176 applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Amphetamine	1100	II
Lisdexamfetamine	1205	II