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.

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 8366	II 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.

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 1205	II II

The company plans to bulk manufacture the listed controlled substances for distribution to their customers. No other activities for these drug codes are authorized for this registration.

Matthew Strait,

Deputy Assistant Administrator. [FR Doc. 2023–13471 Filed 6–23–23; 8:45 am] BILLING CODE P

DEPARTMENT OF LABOR

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Student Safety Assessment of Job Corps Centers

ACTION: Notice of availability; request for comments.

SUMMARY: The Department of Labor (DOL) is submitting this Employment and Training Administration (ETA)-sponsored information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (PRA). Public comments on the ICR are invited.

DATES: The OMB will consider all written comments that the agency receives on or before July 26, 2023.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting "Currently under 30-day Review—Open for Public Comments" or by using the search function.

Comments are invited on: (1) whether the collection of information is necessary for the proper performance of the functions of the Department, including whether the information will have practical utility; (2) the accuracy of the agency's estimates of the burden and cost of the collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility and clarity of the information collection; and (4) ways to minimize the burden of the collection of information on those who are to respond, including the use of automated collection techniques or other forms of information technology.

FOR FURTHER INFORMATION CONTACT: Nicole Bouchet by telephone at 202–693–0213, or by email at *DOL_PRA_PUBLIC@dol.gov*.

SUPPLEMENTARY INFORMATION: The Department of Labor's Office of Job Corps conducts a Student Safety Assessment. Thes collection of information through this assessment is necessary for program evaluation to gauge active students' sense of safety and security at centers. For additional substantive information about this ICR, see the related notice published in the Federal Register on February 7, 2023 (88 FR 7997).

This information collection is subject to the PRA. A Federal agency generally cannot conduct or sponsor a collection of information, and the public is generally not required to respond to an information collection, unless the OMB approves it and displays a currently valid OMB Control Number. In addition, notwithstanding any other provisions of law, no person shall generally be subject to penalty for failing to comply with a collection of information that does not display a valid OMB Control Number. See 5 CFR 1320.5(a) and 1320.6.

DOL seeks PRA authorization for this information collection for three (3) years. OMB authorization for an ICR cannot be for more than three (3) years without renewal. The DOL notes that information collection requirements submitted to the OMB for existing ICRs receive a month-to-month extension while they undergo review.

Agency: DOL-ETA.

Title of Collection: Student Safety Assessment of Job Corps Centers.

OMB Control Number: 1205-0542.

Affected Public: Private Sector—Individuals or Households.

Total Estimated Number of Respondents: 33,906.

Total Estimated Number of Responses: 230,072.

Total Estimated Annual Time Burden: 57,518 hours.

Total Estimated Annual Other Costs Burden: \$0.

(Authority: 44 U.S.C. 3507(a)(1)(D))

Nicole Bouchet,

 $Senior\,PRA\,Analyst.$

[FR Doc. 2023-13468 Filed 6-23-23; 8:45 am]

BILLING CODE 4510-26-P

DEPARTMENT OF LABOR

Office of Workers' Compensation Programs

Agency Information Collection Activities; Comment Request; Roentgenographic Interpretation (CM– 933), Roentgenographic Quality Rereading (CM–933b), Medical History and Examination for Coal Mine Workers' Pneumoconiosis (CM–988), Report of Arterial Blood Gas Study (CM–1159), and Report of Ventilatory Study (CM–2907)

ACTION: Notice.

SUMMARY: The Department of Labor (DOL) is soliciting comments concerning a proposed extension for the authority to conduct the information collection request (ICR) titled, "Roentgenographic Interpretation" (Form CM-933), "Roentgenographic Quality Rereading" (Form CM-933a), "Medical History and Examination for Coal Mine Workers' Pneumoconiosis' (Form CM-988 and CM-988a), "Report of Arterial Blood Gas Study" (Form CM-1159), and "Report of Ventilatory Study" (Form CM-2907). This comment request is part of continuing Departmental efforts to reduce paperwork and respondent burden in accordance with the Paperwork Reduction Act of 1995 (PRA).

DATES: Consideration will be given to all written comments received by August 25, 2023.

ADDRESSES: A copy of this ICR with applicable supporting documentation; including a description of the likely respondents, proposed frequency of response and estimated total burden may be obtained free by contacting Anjanette Suggs by telephone at 202–354–9660 or by email at suggs.anjanette@dol.gov.

Submit written comments about this ICR by mail or courier to the U.S. Department of Labor, Office of Workers' Compensation Program, Division of Coal Mine Workers' Compensation, Room S—3323, 200 Constitution Avenue NW, Washington, DC 20210; by email: suggs.anjanette@dol.gov.

FOR FURTHER INFORMATION CONTACT:

Anjanette Suggs by telephone at 202–354–9660 or by email at suggs.anjanette@dol.gov.

SUPPLEMENTARY INFORMATION: The DOL, as part of continuing efforts to reduce paperwork and respondent burden, conducts a pre-clearance consultation program to provide the general public and Federal agencies an opportunity to comment on proposed and/or continuing collections of information