

## DEPARTMENT OF JUSTICE

## Drug Enforcement Administration

[Docket No. DEA-614]

Importer of Controlled Substances  
Application: Shertech Laboratories,  
LLC**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturer 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 May 1, 2020. Such persons may also file a written request for a hearing on the application on or before May 1, 2020.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on February 21, 2020, Shertech Laboratories, LLC, 1185 Woods Chapel Road, Duncan, South Carolina 29334 applied to be registered as an importer of the following basic class(es) of a controlled substance:

Controlled substance	Drug code	Schedule
Cocaine .....	9041	II

The company plans to import synthetic derivatives of the listed controlled substance in bulk form to conduct clinical trials. Approval of permit applications will occur only when the registrant's activity is consistent with what is authorized under to 21 U.S.C. 952(a)(2). Authorization will not extend to the import of FDA approved or non-approved finished dosage forms for commercial sale.

**William T. McDermott,**  
Assistant Administrator.

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## DEPARTMENT OF LABOR

## Office of the Secretary

Agency Information Collection  
Activities; Submission for OMB  
Review; Comment Request; Cadmium  
in General Industry Standard**ACTION:** Notice of availability; request for comments.

**SUMMARY:** The Department of Labor (DOL) is submitting this Occupational Safety and Health Administration (OSHA)-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 agency receives on or before May 1, 2020.

**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](http://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) if the information will be processed and used in a timely manner; (3) 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; (4) ways to enhance the quality, utility and clarity of the information collection; and (5) 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:** Frederick Licari by telephone at 202-693-8073, TTY 202-693-8064, (these are not toll-free numbers) or by email at [DOL\\_PRA\\_PUBLIC@dol.gov](mailto:DOL_PRA_PUBLIC@dol.gov).

**SUPPLEMENTARY INFORMATION:** The Occupational Safety and Health Act of 1970 (the OSH Act) (29 U.S.C. 651 *et seq.*) authorizes information collection by employers as necessary or appropriate for enforcement of the OSH Act, or for developing information regarding the causes and prevention of occupational injuries, illnesses, and accidents (see 29 U.S.C. 657). The OSH

Act also requires OSHA to obtain such information with a minimum burden upon employers, especially those operating small businesses, and to reduce to the maximum extent feasible unnecessary duplication of effort in obtaining said information (see 29 U.S.C. 657). The collection of information specified in the Cadmium in General Industry Standard (29 CFR 1910.1027) protects workers from the adverse health effects that may result from their exposure to cadmium. The major collection of information of the standard include: Conducting worker exposure monitoring; notifying workers of their cadmium exposures; implementing a written compliance program; implementing medical surveillance of workers; providing examining physicians with specific information; ensuring that workers receive a copy of their medical surveillance results; maintaining workers' exposure monitoring and medical surveillance records for specific periods; and providing access to these records to the workers who are the subject of the records, the worker's representative, and other designated parties. For additional substantive information about this ICR, see the related notice published in the **Federal Register** on November 1, 2019 (84 FR 58747).

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-OSHA.

*Title of Collection:* Cadmium in General Industry Standard.

*OMB Control Number:* 1218-0185.

*Affected Public:* Private Sector: Businesses or other for-profits.

*Total Estimated Number of Respondents:* 50,679.

*Total Estimated Number of Responses:* 234,036.