

or to animal or plant products (select agents and toxins). This joint effort constitutes the Federal Select Agent Program. Due to the continuing pandemic and concerns for the safety of our workshop attendees and employees, DSAT replaced in-person workshops with webinars. The purpose of the webinars is to provide guidance on completing APHIS/CDC Form 2 (Request to Transfer Select Agents and Toxins), APHIS/CDC Form 3 (Report of a Release/Loss/Theft), and APHIS/CDC Form 4 (Reporting the Identification of a Select Agent or Toxin) (APHIS/CDC Forms 2–4) for interested individuals. Two sessions covering the same agenda will be held to provide two opportunities for interested individuals to participate.

DATES: The webinars will be held October 6, 2021 from 10 a.m. to 12:30 p.m. (EDT) and November 3, 2021 from 1:30 p.m. to 4:00 p.m. (EDT). Registration instructions are found on the website, <https://www.selectagents.gov>.

ADDRESSES: The webinars will be conducted from the Centers for Disease Control and Prevention, 1600 Clifton Road NE, Atlanta, Georgia 30329.

FOR FURTHER INFORMATION CONTACT: CDC: Samuel S. Edwin, Ph.D., Director, DSAT, Center for Preparedness and Response, CDC, 1600 Clifton Road NE, MS H–21–7, Atlanta, Georgia 30329. Telephone: (404) 718–2000; email: lrsvat@cdc.gov. APHIS: Jack Taniewski, DVM, Director, DASAT, APHIS, 4700 River Road, Unit 2, Riverdale, MD 20737. Telephone: (301) 851–2070; email: DASAT@usda.gov.

SUPPLEMENTARY INFORMATION: The two public webinar sessions covering the same content, scheduled for Wednesday, October 6, 2021 and Wednesday, November 3, 2021, are opportunities for interested individuals to obtain guidance on completing the APHIS/CDC Forms 2–4 and reporting requirements related to the select agent and toxin regulations (7 CFR part 331, 9 CFR part 121 and 42 CFR part 73). For individuals not able to attend the webinars, the information will be available under the training section at <http://www.selectagents.gov>.

Representatives from the Federal Select Agent Program will be present during the webinars followed by question and answer session to address questions and concerns from the webinar participants.

Participants who want to participate in the webinar should complete their registration online by September 18, 2021. The registration instructions are

located on this website: <http://www.selectagents.gov>.

Dated: July 14, 2021.

Sandra Cashman,

Executive Secretary, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

In accordance with Section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended, and the Determination of the Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92–463.

Name of Committee: Safety and Occupational Health Study Section (SOHSS), National Institute for Occupational Safety and Health (NIOSH).

Dates: October 20–21, 2021.

Time: 11:00 a.m.–5:00 p.m., EDT.

Place: Teleconference.

Agenda: The meeting will convene to address matters related to the conduct of Study Section business and for the study section to consider safety and occupational health-related grant applications.

For Further Information Contact: Michael Goldcamp, Ph.D., Scientific Review Officer, NIOSH, CDC, 1095 Willowdale Road, Morgantown, WV 26506; Telephone: (304) 285–5951; Email: mgoldcamp@cdc.gov.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2021–15246 Filed 7–16–21; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–21–21DC]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled National Syringe Services Program (SSP) Evaluation to the Office of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on February 25, 2021 to obtain comments from the public and affected agencies. CDC received three public comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) Minimize the burden of the collection of information on those who are to respond, including, through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570. 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