ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Peer Recruiters	Recruiter Debriefing	3,333	1	2/60	112
Total					6,600

Jeffrey M. Zirger,
Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket Number CDC–2022–0063, NIOSH 063–D]

National Institute for Occupational Safety and Health (NIOSH) Fire Fighter Fatality Investigation and Prevention Program (FFFIPP) Fire Service Community Meeting

AGENCY: The Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of meeting and request for comment.

SUMMARY: The National Institute for Occupational Safety and Health (NIOSH) in the Centers for Disease Control and Prevention (CDC), an operating division of the Department of Health and Human Services (HHS), announces the following web-based meeting and request for comment on the NIOSH Fire Fighter Fatality Investigation and Prevention Program (FFFIPP).

DATES: Written comments must be received by July 27, 2022. The public meeting will be held on Monday, June 27, 2022, 10 a.m. to 3:30 p.m. EDT, or after the last public commenter in attendance has spoken, whichever occurs first. The public meeting will be held as a web-based teleconference available by remote access.

ADDRESSES: This is a virtual meeting. You may submit comments, identified by Docket No. CDC–2022–0063; NIOSH 063–D, by either of the following methods:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* National Institute for Occupational Safety and Health, NIOSH Docket Office, 1090 Tusculum Avenue, MS C–34, Cincinnati, Ohio 45226–1998.

Instructions: On June 27, 2022, NIOSH will hold a virtual (web-based) meeting to seek input. The meeting will be open to the fire service community and interested parties, limited only by web conference lines (500 web conference lines are available) using Audio/LiveMeeting Conferencing. Web-based meeting requirements include: A computer with audio capabilities and an internet connection or a telephone, preferably with mute capability. Each participant is required to register for the meeting at the NIOSH website *https://www.cdc.gov/niosh/fire/fsc.html* by June 15, 2022, 5:00 p.m. EDT. NIOSH will reply by email confirming registration and the details needed to participate in the web-based meeting.

All information received in response to this notice must include the agency name and docket number (CDC–2022–0063; NIOSH 063–D). All relevant comments and submissions provided will be reviewed and posted at *http://www.regulations.gov.* Do not submit comments by email. CDC does not accept comments by email.

FOR FURTHER INFORMATION CONTACT: Jeff Funke, Team Lead, Surveillance and Field Investigations Branch, Division of Safety Research; Telephone: 304–285–5894; Email: *NIOSHFireTrauma@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Purpose: The purpose of this web-based meeting and docket is to request public comment from the fire service community and interested parties on the NIOSH Fire Fighter Fatality Investigation and Prevention Program (FFFIPP).

Matters To Be Considered: NIOSH will provide a brief presentation and will facilitate discussion on the following two topics: (1) The primary audiences for NIOSH line-of-duty death investigation reports and the strengths and weaknesses of those reports, including report content, format, and length; and (2) specific feedback on how the NIOSH FFFIPP prioritization guideline for planning investigations can be enhanced to meet the needs of the fire service community.

Additional time will be given for invited and registered participants to bring other topics to the attention to the NIOSH FFFIPP.

Background: Since its inception in 1998, the NIOSH FFFIPP has held periodic meetings to seek input about the program with the fire service community and interested parties. These meetings have been an important component of the program and are vital to ensure the program is meeting the needs and expectations of those it serves. The FFFIPP has posted the results of these periodic meetings on its website at: *https://www.cdc.gov/niosh/fire/abouttheprogram/ourworkreviewed/ourworkreviewed.html.*

Written Fire Service Community and Interested Parties Participation

Interested fire service persons and organizations are invited to participate by submitting written views, recommendations, and data. Please note that comments received, including attachments and other supporting materials, are part of the public record and are subject to public disclosure. Comments will be posted on *https://www.regulations.gov.* Therefore, do not include any information in your comment or supporting materials that you consider confidential or inappropriate for public disclosure. CDC will review all submissions and may choose to redact, or withhold, submissions containing private or proprietary information such as Social Security numbers, medical information, inappropriate language, or duplicate/near duplicate examples of a mass-mail campaign. CDC will carefully consider all comments submitted into the docket and may modify the FFFIPP and operations.

Written fire service community and interested parties comments: The docket will be opened to receive written comments on May 13, 2022 through July 27, 2022, 5:00 p.m. EDT.

Oral Fire Service Community and Interested Parties Comments

This meeting will include time for members of the fire service community and interested parties to provide comments about the NIOSH FFFIPP,

including investigation reports and program products to improve firefighter safety and health, and suggestions for enhancing the impact of the program. A discussion period will be provided to enable the audience to contribute to any of the topics discussed. The time allotted for speakers during the discussion period will be at the discretion of the NIOSH moderator based upon overall time constraints. A chat box will also be available during the meeting for participants to submit questions or comments to the speakers or NIOSH. This chat will be part of the official record. Questions will be read by the moderator and answered by the speaker and/or NIOSH as time allows.

John J. Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–22–22BY]

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 ''Importation Regulations (42 CFR 71 Subpart F)'' to the Office of Management and Budget (OMB) for review and approval. CDC previously published a ''New Information Collection Submitted for Public Comment and Recommendations'' notice on January 24, 2022 to obtain comments from the public and affected agencies. CDC received one comment 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 for Public Comments'' or by using the search function. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395–5806. Provide written comments within 30 days of notice publication.

Proposed Project

Importation Regulations (42 CFR 71 Subpart F)—New—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

This is a request for a new information collection to consolidate forms and information collections related to the importation of animals, animal products, and human remains into one information collection. This information collection was previously part of three separate, approved information collections (0920–1034, expires March 31, 2022, 0920–0263 expires September 30, 2023, and 0920–0199 expires August 31, 2024). CDC is requesting a three-year OMB clearance for this new, combined information collection.

Section 361 of the Public Health Service Act (PHSA) (42 U.S.C. 264) authorizes the Secretary of Health and Human Services to make and enforce regulations necessary to prevent the introduction, transmission or spread of communicable diseases from foreign countries into the United States. Statute and the existing regulations governing foreign quarantine activities (42 CFR 71) authorizes quarantine officers and other personnel to inspect and undertake necessary control measures with respect to conveyances, persons, and shipments of animals and etiologic agents in order to protect the public's health.

CDC regulations govern the importation of animals and animal products capable of causing human disease. Animals that are regulated by CDC include dogs, cats, turtles, snakes, lizards, non-human primates (NHP), civets, African rodents, and bats. CDC controls the importation of these animals to ensure that these animals, or animal products, being imported into the United States meet CDC regulations. CDC does this through a permitting process for certain animals.

On June 16, 2021 CDC published a **Federal Register** Notice informing the public about a temporary suspension of dogs entering the United States from high-risk rabies countries. The canine rabies virus variant (CRVV) was declared eliminated in the United States in 2007. The importation of just one dog infected with CRVV risks re-introduction of the virus into the United States resulting in a potential public health risk with consequent monetary cost and potential loss of human and animal life. Since 2015 there have been four known rabid dogs imported into the United States.

During the suspension period, CDC will issue permits for importers with dogs who have been in a high-risk CRVV country within the last six months and do not have a current, valid U.S.-issued rabies vaccination certificate. Only importers who are permanently relocating to the United States, are a U.S. government employee traveling on official orders, are an owner of a service dog that is trained to assist them with a disability, are an individual importing dogs for science, education, exhibition, or law enforcement purposes, or people who traveled with their dog before July 31, 2021 are eligible to apply for a permit. Dogs from CRVV-free or low risk countries and dogs with valid U.S.-issued rabies vaccination certificates that are microchipped, healthy, and at least six months of age do not require a permit. The current permit application to import a dog is under collection 0920–1034. When a dog or cat arrives at an airport and is sick or dead, importers are required to notify CDC. There is no form for this notification.

Other animals that require a permit and are included in this information collection are NHPs, which can carry of number of diseases that can cause