

gear that may not be protective against all hazards is being used by fire fighters.

Intuitively, the use conditions to which turnout gear would be exposed to when used by a large or medium metropolitan fire department would be very different than those of a smaller department. However, the absence of scientific data to link performance to use conditions (*e.g.*, number and type of washings, number of fire-related calls) provides a barrier to transitioning to an alternative approach to retirement.

This study will obtain a statistically meaningful sample of turnout gear from three fire departments. The use

conditions for the sampled turnout gear will be determined, and the gear will be subjected to established performance requirements. For each set of gear, its performance will be directly linked to its use condition history. This combined lab and field data will help determine if there is a relationship between turnout gear use conditions and the ability for turnout gear to effectively protect the user.

The use conditions for each set of sampled gear will be determined by:

(1) Reviewing fire department records, practices, and policies;

(2) surveying the fire fighters assigned to each set of sampled gear to obtain one-month of retrospective information about the use conditions to which it was likely exposed; and

(3) a 6-month prospective data collection where the fire fighters assigned to each set of sampled gear provide information about their shift-specific exposures.

The estimated annualized Burden Hours for this information collection is 1,050. There is no cost to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Individual Fire Fighter	Turnout Gear Safety Survey—Retrospective Exposures for past month.	100	1	30/60
	Turnout Gear Safety Survey—Prospective Exposures for six months.	100	60	10/60

Jeffrey M. Zirger,

Acting Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2018–N–3305]

Allergenic Products Advisory Committee; Cancellation

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The meeting of the Allergenic Products Advisory Committee scheduled for November 7, 2018, is cancelled. This meeting was announced in the **Federal Register** of September 26, 2018.

FOR FURTHER INFORMATION CONTACT:

Serina Hunter-Thomas, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 6338, Silver Spring, MD 20993–0002, 240–402–5771, serina.hunter-thomas@fda.hhs.gov, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), and follow the

prompts to the desired center or product area. Please call the Information Line for up-to-date information on this meeting.

Dated: October 16, 2018.

Leslie Kux,

Associate Commissioner for Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2018–N–3728]

Agency Information Collection Activities; Proposed Collection; Comment Request; Collection of Conflict of Interest Information for Participation in Food and Drug Administration Non-Employee Fellowship and Traineeship Programs

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and

to allow 60 days for public comment in response to the notice. This notice solicits comments on the “Collection of Conflict of Interest Information for Participation in FDA Non-Employee Fellowship and Traineeship Programs.”

DATES: Submit either electronic or written comments on the collection of information by December 21, 2018.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before December 21, 2018]. The <https://www.regulations.gov> electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of December 21, 2018. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- **Federal eRulemaking Portal:** <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a