response bias, the protocols for data collection, and any testing procedures that were or will be undertaken prior fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other generic mechanisms that are designed to yield quantitative results.

The Agency received no comments in response to the 60-day notice published

in the **Federal Register** on April 30, 2014 (75 FR 24432).

Respondents will be screened and selected from Individuals and Households, Businesses, Organizations, and/or State, Local or Tribal Government. There is no cost to respondents other than their time. ATSDR is requesting an increase in the annual burden hours from 2,425 to 7,075 and an increase in the annual

number of respondents from 2,800 to 8,300. These estimates of burden hours and respondents are based on an anticipated increase in the number of the Agency's generic information collections (GenICs) each year over the next three years. The estimated annualized burden hours for this data collection activity are 7,075.

Type of respondents	Form name	No. of re- spondents	No. of re- sponses per respondent	Avg. burden per response (in hrs.)
Individuals and Households; Businesses and Organizations; State, Local or Tribal Government.	Small discussion groups	300	1	90/60
	Request for customer comment cards/complaint forms/post-conference or training surveys.	1,500	1	15/60
	Focus groups of customers, potential customers, delivery partners, or other stakeholders.	2,000	1	2
	Qualitative customer satisfaction surveys or interviews.	3,000	1	30/60
	Usability testing/in-person observation testing	1,500	1	30/60

Leroy A. Richardson,

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.

[FR Doc. 2015–25920 Filed 10–9–15; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

Statement of Organization, Functions, and Delegations of Authority

Part J (Agency for Toxic Substances and Disease Registry) of the Statement of Organization, Functions, and Delegations of Authority of the Department of Health and Human Services (50 FR 25129–25130, dated June 17, 1985, as amended most recently at 77 FR 68125–68127, dated November 12, 2012) is amended to reflect the Order of Succession for the Agency for Toxic Substances and Disease Registry.

Section J–C, Order of Succession: Delete in its entirety the Section C–C, Order of Succession, and insert the following:

During the absence or disability of the Administrator, Agency for Toxic Substances and Disease Registry (ATSDR), or in the event of a vacancy in that office, the first official listed below who is available shall act as

Administrator, except during a planned period of absence, the Administrator may specify a different order of succession:

- 1. Administrator, ATSDR
- 2. Principal Deputy Administrator, ATSDR
- 3. Assistant Administrator, ATSDR
- 4. Deputy Director for Noncommunicable Diseases, Injury and Environmental Health

Iames Seligman.

Acting Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2015–25775 Filed 10–9–15; 8:45 am]

BILLING CODE 4160-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60 Day-16-15BHH; Docket No. CDC-2016-0087]

Proposed Data Collection Submitted for Public Comment and Recommendations

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

ACTION: Cancellation of notice with comment period

SUMMARY: The notice "Proposed Data Collection Submitted for Public Comment and Recommendations" on Personal Protective Equipment

Information (PPE-Info) Database (80 FR 60906, October 8, 2015) is cancelled. This noticed invited comment on the Personal Protective Equipment Information (PPE-Info) Database which is a compendium of personal protective equipment (PPE) Federal regulations and consensus standards. This proposed data collection will be resubmitted at a later date for public comment once the review to include one additional standard is completed on the data collection instrument.

FOR FURTHER INFORMATION CONTACT:

(404) 639–7570 or send comments to CDC, Leroy Richardson, 1600 Clifton Road, MS D–74, Atlanta, GA 30333 or send an email to *omb@cdc.gov*.

Dated: October 8, 2015.

Leroy A. Richardson,

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.

[FR Doc. 2015–26067 Filed 10–8–15; 4:15 pm]

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