Regulation under delegated authority, May 27, 2020.

Ann Misback,

Secretary of the Board.

[FR Doc. 2020–11771 Filed 6–1–20; 8:45 am]

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FEDERAL TRADE COMMISSION

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Extension

AGENCY: Federal Trade Commission.

ACTION: Notice.

SUMMARY: The Federal Trade Commission (''FTC'' or ''Commission'') requests that the Office of Management and Budget (''OMB'') extend for an additional three years the current Paperwork Reduction Act (''PRA'') clearance for information collection requirements associated with its Funeral Industry Practice Rule (''Funeral Rule'' or ''Rule''). That clearance expires on June 30, 2020.

DATES: Comments must be filed by July 2, 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.* Find this particular information collection by selecting ''Currently under Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: Patricia H. Poss, Division of Marketing Practices, Bureau of Consumer Protection, Federal Trade Commission, 600 Pennsylvania Ave. NW, Washington, DC 20580, *pposs@ftc.gov,* (202) 326–2413.

SUPPLEMENTARY INFORMATION: Pursuant to the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*), the FTC has submitted to the Office of Management and Budget (''OMB'') this request for extension of the previously approved collection of information discussed below.

Title of Collection: Funeral Industry Practice Rule, 16 CFR 453.

OMB Control Number: 3084–0025.

Type of Review: Extension without change of currently approved collection.

Affected Public: Private Sector: Businesses and other for-profit entities.

Estimated Number of Annual Respondents: 19,136.

Estimated Annual Burden Hours: 164,006.

Estimated Annual Labor Costs: $5,429,859.

Abstract

The Funeral Rule ensures that consumers who are purchasing funeral goods and services have access to accurate itemized price information so they can purchase only the funeral goods and services they want or need. Among other things, the Rule requires a funeral provider to: (1) Provide consumers a copy of the funeral provider's General Price List that itemizes the goods and services it offers; (2) show consumers a Casket Price List and an Outer Burial Container Price List at the outset of any discussion of those items or their prices, and in any event before showing consumers caskets or vaults; (3) provide price information from its price lists over the telephone; and (4) give consumers a Statement of Funeral Goods and Services Selected after determining the funeral arrangements with consumers. The Rule requires that funeral providers disclose this information to consumers and maintain records documenting their compliance with the Rule.

Request for Comment

On February 4, 2020, the FTC sought public comment on the information collection requirements in the Funeral Rule. 85 FR 6185 (Feb. 4, 2020). No relevant comments were received. Pursuant to the OMB regulations, 5 CFR part 1320, the FTC is providing this second opportunity for public comment while seeking OMB approval to renew clearance for the Rule's information collection requirements.

Your comment—including your name and your state—will be placed on the public record of this proceeding. Because your comment will be made public, you are solely responsible for making sure that your comment does not include any sensitive personal information, like anyone's Social Security number, date of birth, driver's license number or other state identification number or foreign country equivalent, passport number, financial account number, or credit or debit card number. You are also solely responsible for making sure that your comment does not include any sensitive health information, like medical records or other individually identifiable health information. In addition, do not include any ''[t]rade secret or any commercial or financial information which is . . . privileged or confidential'' as provided in Section 6(f) of the FTC Act 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16CFR 4.10(a)(2). In particular, do not include competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns devices, manufacturing processes, or customer names.

Josephine Liu,

Assistant General Counsel for Legal Counsel.

[FR Doc. 2020–11877 Filed 6–1–20; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–20–1204; Docket No. CDC–2020–0053]

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: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on the ''Behavioral Risk Factor Surveillance System (BRFSS) Asthma Call-back Survey (ACBS)'' (OMB Control No. 0920–1204, expiration date 11/30/2020). The ACBS is an in-depth asthma survey conducted on a subset of BRFSS respondents with an asthma diagnosis. The goal of this survey is to strengthen the existing body of asthma data and to address critical questions surrounding the health and experiences of persons with asthma.

DATES: Written comments must be received on or before August 3, 2020.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2020–0053 by any of the following methods:

- *Federal eRulemaking Portal: Regulation.gov.* Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *Regulations.gov*.

Please note: Submit all comments through the Federal eRulemaking portal

(*regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329; phone: 404–639–7570; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. 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;

2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

3. Enhance the quality, utility, and clarity of the information to be collected; and

4. 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 submissions of responses.

5. Assess information collection costs.

Proposed Project

Behavioral Risk Factor Surveillance System (BRFSS) Asthma Call-back Survey (ACBS)—Revision—National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The CDC's National Center for Environmental Health (NCEH) is requesting a three-year Paperwork Reduction Act (PRA) clearance to revise and continue to collect information under the ''Behavioral Risk Factor Surveillance System (BRFSS) Asthma Call-back Survey (ACBS)'' (OMB Control No. 0920–1204, expiration date 11/30/2020). The ACBS is funded by the NCEH National Asthma Control Program (NACP) in the Asthma and Community Health Branch (ACHB). The NACP provides its 40 participating states with technical and methodological assistance.

The ACBS is a follow-up survey on asthma and is administered on behalf of NCEH by the CDC's National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP) BRFSS Program. The BRFSS (OMB Control No. 0920–1061, expiration date 3/31/2021) is a nationwide system of customized, cross-sectional telephone health surveys. The BRFSS information collection is conducted in a continuous, three-part telephone interview process: screening, participation in a common BRFSS core survey, and participation in optional question modules that states use to customize survey content. BRFSS coordinators in the health departments in U.S. states, territories, and the District of Columbia (collectively referred to as ''states'' and ''jurisdictions'') are responsible for both the BRFSS and the ACBS administration. The ACBS is conducted within two days after the BRFSS survey.

The purpose of ACBS is to gather state-level asthma data and to make them available to track the burden of the disease, to monitor adherence to asthma guidelines, and to direct and evaluate interventions undertaken by asthma control programs located in state health departments. Beyond asthma prevalence estimates, for most states, the ACBS provides the only sources of adult and child asthma data on the state and local level.

Data collection for ACBS involves screening, obtaining permission, consenting, and telephone interviewing on a subset of the BRFSS respondents from participating states. The ACBS eligible respondents are BRFSS adults, 18 years and older, who report ever being diagnosed with asthma. In addition, some states include children, below 18 years of age, who are randomly selected subjects in the BRFSS household. Parents or guardians serve as ACBS proxy respondents for their children ever diagnosed with asthma. If both the BRFSS adult respondent and the selected child in the household have asthma, then only one or the other is eligible for the ACBS.

State BRFSS Coordinators submit de-identified data files to CDC on a monthly or quarterly basis for cleaning and weighting. The CDC BRFSS ACBS operation team returns clean, weighted data files to the state of origin for its use. The ACBS adds considerable state-level depth to the existing body of asthma data. It addresses critical questions surrounding the health and experiences of persons with asthma. Health data include symptoms, environmental factors, and medication use among persons with asthma. Data on their experiences include activity limitation, health system use, and self-management education. These asthma data are needed to direct and evaluate interventions undertaken by asthma control programs located in state health departments. Federal agencies and other entities also rely on this critical information for planning and evaluating efforts and to reduce the burden from this disease. The CDC makes annual ACBS datasets available for public use and provides guidance on statistically appropriate uses of the data.

The time burden estimates are based on the 2016 ACBS data collection, which is the most recent data released. The burden table reflects the landline and cell phone data collection methods used in 2016 and later years. Additionally, the time burden accounts for reporting burden incurred by the states for the monthly or quarterly adult and child ACBS data submissions to CDC. The total estimated annualized time burden for all respondents is 6,615 hours. Participation in the ACBS is voluntary and there are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN TO RESPONDENTS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)	Total burden hours
BRFSS Adults	ACBS Landline Screener—Adult	17,800	1	1/60	297
	ACBS Cell Phone Screener—Adult	16,733	1	1/60	279
BRFSS Parents or Guardians of Children.	ACBS Landline Screener—Child	2,576	1	1/60	43
	ACBS Cell Phone Screener—Child	3,824	1	1/60	64
ACBS Adults	ACBS Adult Consent and Questionnaire.	23,166	1	10/60	3,861
ACBS Parents or Guardians of Children.	ACBS Child Consent and Questionnaire.	3,787	1	10/60	631
State BRFSS Coordinators	ACBS Adult Data Submission Layout.	40	12	155/60	1,240
	ACBS Child Data Submission Layout.	40	12	25/60	200
Total					6,615

Jeffrey M. Zirger,

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

[FR Doc. 2020–11803 Filed 6–1–20; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–20–1027]

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 ''Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery'' 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 March 9, 2020 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

Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (OMB Control No. 0920–1027, Exp. 7/31/2020)—Extension—National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC is requesting a three-year extension of this generic information collection request. During the past three-year approval period, the generic clearance facilitated the approval of seven projects (''GenICs'') involving 13,574 respondents. The projects included web-based surveys, focus groups, and assessments. The information collection activities conducted under this extension will continue to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery. By qualitative feedback, we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training, or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative, and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management. Feedback collected under this generic clearance will provide useful information, but it will not yield data