conducted annually over a three-year period, beginning in FY15.

Likely Respondents: Survey respondents will consist of Medicare beneficiaries, providers, suppliers, or their representatives who participated in a hearing before an OMHA ALJ. OMHA will draw a representative, non-redundant sample of appellants whose cases have been closed in the last six months.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Type of respondent	Form name	Number of respondents	Number responses per respondent	Average burden per response (in hours)	Total burden hours
Charged at rate of Healthcare Providers and Suppliers	Form A	240	1	11/60	44
Charged at rate of Beneficiaries	Form A	160	1	11/60	29
Total		400	1	11/60	73

OS specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Darius Taylor,

Information Collection Clearance Officer.

[FR Doc. 2014–20769 Filed 8–29–14; 8:45 am]

BILLING CODE 4150–46–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Notice of Interest Rate on Overdue Debts

Section 30.18 of the Department of Health and Human Services' claims collection regulations (45 CFR Part 30) provides that the Secretary shall charge an annual rate of interest, which is determined and fixed by the Secretary of the Treasury after considering private consumer rates of interest on the date that the Department of Health and Human Services becomes entitled to recovery. The rate cannot be lower than the Department of Treasury's current value of funds rate or the applicable rate determined from the ''Schedule of Certified Interest Rates with Range of Maturities'' unless the Secretary waives interest in whole or part, or a different rate is prescribed by statute, contract, or repayment agreement. The Secretary of the Treasury may revise this rate quarterly. The Department of Health and Human Services publishes this rate in the **Federal Register**.

The current rate of 10⅜%, as fixed by the Secretary of the Treasury, is certified for the quarter ended June 30, 2014. This rate is based on the Interest Rates for Specific Legislation, ''National Health Services Corps Scholarship Program (42 U.S.C. 250(B)(1)(A))'' and ''National Research Service Award Program (42 U.S.C. 288(c)(4)(B)).'' This interest rate will be applied to overdue debt until the Department of Health and Human Services publishes a revision.

Dated: August 11, 2014.

Teresa Miranda,

Director, Financial Management Policy Division.

[FR Doc. 2014–20773 Filed 8–29–14; 8:45 am]

BILLING CODE 4150–04–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Decision To Evaluate a Petition To Designate a Class of Employees From the Dow Chemical Co. Facility in Walnut Creek, California (Also Known as Pittsburg, CA), To Be Included in the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: NIOSH gives notice as required by 42 CFR 83.12(e) of a decision to evaluate a petition to designate a class of employees from the Dow Chemical Co. facility in Walnut Creek, California (also known as Pittsburg, CA), to be included in the Special Exposure Cohort under the Energy Employees Occupational Illness Compensation Program Act of 2000. The initial proposed definition for the class being evaluated, subject to revision as warranted by the evaluation, is as follows:

Facility: Dow Chemical Co. facility.

Location: Walnut Creek, California (also known as Pittsburg, CA).

Job Titles and/or Job Duties: All employees who worked in any area.

Period of Employment: January 1, 1947 through December 31, 1957.

FOR FURTHER INFORMATION CONTACT: Stuart L. Hinnefeld, Director, Division of Compensation Analysis and Support, National Institute for Occupational Safety and Health, 1090 Tusculum Avenue, MS C–46, Cincinnati, OH 45226–1938, Telephone 877–222–7570. Information requests can also be submitted by email to *DCAS@CDC.GOV.*

John Howard,

Director, National Institute for Occupational Safety and Health.

[FR Doc. 2014–20740 Filed 8–29–14; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–14–0800]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (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 or send an email to *omb@cdc.gov.* Written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Focus Group Testing to Effectively Plan and Tailor Cancer Prevention and Control Communication Campaigns (OMB No. 0920–0800, exp. 11/30/2014)—Extension—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The mission of the CDC's Division of Cancer Prevention and Control (DCPC) is to reduce the burden of cancer in the United States through cancer prevention, reduction of risk, early detection, better treatment, and improved quality of life for cancer survivors. Toward this end, the DCPC supports the scientific development, implementation, and evaluation of various health communication campaigns with an emphasis on specific cancer burdens. This process requires testing of messages, concepts, and materials prior to their final development and dissemination. Communication campaigns vary according to the type of cancer, the qualitative dimensions of the message described above, and the type of respondents.

CDC is currently approved to collect information needed to plan and tailor cancer communication campaigns (OMB No. 0920–0800, exp. 11/30/2014), and seeks OMB approval to extend the existing generic clearance. No changes to the scope of the clearance or data collection methodology are proposed. There are small decreases in the annualized estimates for the number of respondents and burden hours.

Information will be collected primarily through focus groups, and will be used to assess numerous qualitative dimensions of cancer prevention and control messages, including, but not limited to, knowledge, attitudes, beliefs, behavioral intentions, information needs and sources, and compliance to recommended screening intervals. Insights gained from the focus groups will assist in the development and/or refinement of future campaign messages and materials.

DCPC plans to conduct or sponsor up to 80 focus groups per year over a three-year period. An average of 10 respondents will participate in each focus group discussion. Screening will be conducted to recruit respondents for specific target audiences, e.g., the general public or health care providers. The estimated burden per response for screening is three minutes. Each focus group discussion will be facilitated by a written discussion guide, and will last approximately two hours. CDC will submit an information collection request to OMB for approval of each focus group activity.

OMB approval is requested for three years. There are no changes to information collection purpose or methodology. There are minor reductions in the annualized estimates for the number of respondents and corresponding burden hours.

Participation is voluntary and there are no costs to respondents except their time. The total estimated annualized burden hours are 1,680.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
General Public	Screening Form	960	1	3/60
	Focus Group Guide	480	1	2
Health Care Professionals	Screening Form	640	1	3/60
	Focus Group Guide	320	1	2

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. 2014–20718 Filed 8–29–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–D–0329]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on Fees for Human Drug Compounding Outsourcing Facilities Under the Federal Food, Drug, and Cosmetic Act

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (the PRA).

DATES: Fax written comments on the collection of information by October 2, 2014.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs,