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[FR Doc. 2021-15792 Filed 7-23-21; 8:45 am]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[60Day-21-0800; Docket No. CDC-2021-  
0072]

#### 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 effort to reduce public  
burden and maximize the utility of  
government information, invites the  
general public and other Federal  
agencies the opportunity to comment on  
a proposed and/or continuing  
information collection, as required by  
the Paperwork Reduction Act of 1995.  
This notice invites comment on a  
proposed information collection project  
titled Focus Group Testing to Effectively  
Plan and Tailor Cancer Prevention and  
Control Communication Campaigns.  
CDC is requesting a Revision to this  
Generic Clearance to include an  
additional cancer-related  
communications campaign, expand the  
modes of data collection to include  
online focus groups and in-depth  
interviews (in-person, phone, and  
online), and to focus on respondents  
from the general public.

**DATES:** CDC must receive written  
comments on or before September 24,  
2021.

**ADDRESSES:** You may submit comments,  
identified by Docket No. CDC-2021-  
0072 by any of the following methods:

- *Federal eRulemaking Portal:*  
*Regulations.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;
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; and
5. Assess information collection costs.

#### Proposed Project

Focus Group Testing to Effectively  
Plan and Tailor Cancer Prevention and  
Control Communications Campaigns—  
(OMB Control No. 0920-0800, Exp. 10/  
31/2021)—Revision—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, and improved quality of life  
for cancer survivors. Toward this end,  
the DCPC supports the scientific  
development and implementation 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, as described in the  
second step of the health  
communication process. The health  
communication process is a scientific  
model developed by the U.S.  
Department of Health and Human  
Services' National Cancer Institute to  
guide sound campaign development.  
The communication literature supports  
various data collection methods to  
conduct credible formative, concept,  
message, and materials testing. This  
process ensures that the public clearly  
understands cancer-specific information  
and concepts, are motivated to take the  
desired action, and do not react  
negatively to the messages. CDC is  
currently approved to collect  
information needed to plan and tailor  
cancer communication campaigns (OMB  
Control No. 0920-0800, Exp. 10/31/  
2021), and seeks OMB approval to  
revise the existing generic clearance to  
include another cancer-related  
communications campaign, expand the  
modes of data collection to include  
online focus groups and in-depth  
interviews (in-person, phone, and  
online), and to focus on respondents  
from the general public.

Information collection will involve  
discussions to assess numerous  
qualitative dimensions of cancer  
prevention and control messages  
including, but not limited to, cancer  
knowledge, attitudes, beliefs, behavioral  
intentions, information needs and  
sources, and compliance with cancer  
screening as recommended by the  
United States Preventive Services Task  
Force. Insights gained from these  
discussions will assist in the  
development and/or refinement of  
future campaign messages and  
materials. Communication campaigns  
and messages will vary according to the  
type of cancer and the qualitative  
dimensions of the message described  
above. A separate information collection

request will be submitted to OMB for approval of each discussion activity. The request will describe the purpose of the activity and include the customized information collection instruments.

OMB approval is requested for three years. There is no change in burden hours or respondents. Participation is voluntary and there are no costs to respondents except their time. CDC

requests approval for an estimated 1,680 annual burden hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
General Public .....	Screening Form .....	1600	1	3/60	80
General Public .....	Discussion Guide .....	800	1	2	1,600
Total .....	.....	.....	.....	.....	1,680

**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**

**Mine Safety and Health Research Advisory Committee (MSHRAC); Cancellation of Meeting**

Notice is hereby given of a change in the meeting of the Mine Safety and Health Research Advisory Committee (MSHRAC); June 21, 2021, 10:00 a.m.–2:30 p.m., EDT, in the original FRN.

The meeting was published in the **Federal Register** on April 23, 2021, Volume 86, Number 77, page 21739.

This meeting is being canceled in its entirety.

**FOR FURTHER INFORMATION CONTACT:**

George W. Luxbacher, Designated Federal Officer, MSHRAC, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, 2400 Century Parkway NE, Atlanta, GA 30345; Telephone: (404) 498-2808; email: [gluxbacher@cdc.gov](mailto:gluxbacher@cdc.gov).

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and

Prevention and the Agency for Toxic Substances and Disease Registry.

**Kalwant Smagh,**

*Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[30Day-21-0556]

**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 “Assisted Reproductive Technology (ART) Program Reporting System” 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 12, 2021 to obtain comments from the public and affected agencies. CDC did not receive comments 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](http://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**

Assisted Reproductive Technology (ART) Program Reporting System (OMB Control No. 0920-0556, Exp. 8/31/2021)—Revision—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).