Local Exchange Carriers, CC Docket No. 01–338 and WC Docket No. 04–313, Order on Remand.

Form Number: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Business or other for-profit entities, Not-for-profit institutions and State, Local or Tribal government.

Number of Respondents and Reponses: 645 respondents; 645 responses.

Estimated Time per Response: 8 hours.

Frequency of Response: Recordkeeping requirement, third party disclosure requirement and on occasion reporting requirement.

Obligation to Respond: Required to obtain or retain benefits. Statutory authority for this information collection is contained in *47 U.S.C. Section 251* of the Communications Act of 1934, as amended.

Total Annual Burden: 5,160 hours.

Total Annual Cost: No Cost.

Privacy Act Impact Assessment: No impact(s).

Nature and Extent of Confidentiality: The Commission is not requesting respondents to submit or disclose confidential information. However, in certain circumstances, respondents may voluntarily choose to submit confidential information pursuant to applicable confidentiality rules.

Needs and Uses: In the Order on Remand, the Commission imposed unbundling obligations in a more targeted manner where requesting carriers have undertaken their own facilities-based investments and will be using UNEs (unbundled network elements) in conjunction with self-provisioned facilities. The Commission also eliminated the subdelegation of authority to state commissions adopted in the previous order. Prior to the issuance of the Order, the Commission sought comment on issues relating to combinations of UNEs, called ''enhanced extended links'' (EELs), in order to effectively tailor access to EELs to those carriers seeking to provide significant local usage to end users. In the Order, the Commission adopted three specific service eligibility criteria for access to EELs in accordance with Commission rules.

Federal Communications Commission.

Katura Jackson,

Federal Register Liaison Officer.

[FR Doc. 2021–23145 Filed 10–22–21; 8:45 am]

BILLING CODE 6712–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–22–22AD; Docket No. CDC–2021–0113]

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 to take this opportunity to comment on proposed information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on the Research Data Center (RDC) Proposal for Access to Confidential Data for the National Center for Health Statistics (NCHS). The proposed collection will be used to assess researcher's requests for access to confidential NCHS data for their research projects.

DATES: Written comments must be received on or before December 27, 2021.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2021–0113 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 H21–8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to *Regulations.gov*, including any personal information provided. For access to the docket to read background documents or comments received, go to *Regulations.gov*.

Please note: All public comment should be submitted 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–H21–8, 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 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

Research Data Center (RDC) Proposal for Access to Confidential Data for the National Center for Health Statistics—Existing Collection in use without an OMB Control Number—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Section 306(b)(4) of the Public Health Service (PHS) Act (42 U.S.C. 242k(b)(4)), as amended, authorizes that the Secretary of Health and Human Services (DHHS), acting through NCHS, receive requests for providing data and statistics to the public. NCHS receives requests

for confidential data from the public through the Research Data Center (RDC) Proposal for Access to Confidential Data. This is a request for approval from OMB to collect information via the RDC proposal over the next three years.

As part of a comprehensive data dissemination program, the Research Data Center (RDC), National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC), requires prospective researchers who need access to confidential data to complete a research proposal. Researchers self-select whether they need access to confidential data to answer their research questions. The RDC requires the researcher to complete a research proposal so NCHS understands the research proposed, whether confidential data are available to address the research questions, how the confidential data will be used, and what data outputs the researcher needs to satisfy their project. The completed proposal is sent to NCHS for adjudication on whether the proposed research is possible.

To capture the information needed to adjudicate researchers' need for access to confidential NCHS data, CDC requests OMB approval for a total estimated annual burden total of 330 hours (990 hours for a three-year clearance period). The resulting information will be for NCHS internal use. There is no cost to respondents other than their time to complete the proposal.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Avg. burden per response (in hrs.)	Total burden (in hrs.)
Researcher	Research Data Center proposal	110	1	3	330
Total					330

Jeffrey M. Zirger,

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

[FR Doc. 2021–23186 Filed 10–22–21; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Board on Radiation and Worker Health (ABRWH), National Institute for Occupational Safety and Health (NIOSH)

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

ACTION: Notice of meeting and request for comment.

SUMMARY: In accordance with the Federal Advisory Committee Act, the CDC announces the following meeting of the Advisory Board on Radiation and Worker Health (ABRWH or the Advisory Board). This meeting is open to the public, limited only by the number of audio conference lines and internet conference accesses available, which is 200 combined. The public is welcome to submit written comments in advance of the meeting, to the contact person below. Written comments received in advance of the meeting will be included in the official record of the meeting. The public is also welcomed to listen to the meeting by joining a teleconference line and/or computer connection (information below).

DATES: The meeting will be held on December 8, 2021, from 1:00 p.m. to 6:00 p.m., EST, and December 9, 2021, from 1:00 p.m. to 4:00 p.m., EST. A public comment session will be held on December 8, 2021 at 5:00 p.m., EST, and will conclude at 6:00 p.m., EST, or following the final call for public comment, whichever comes first.

Written comments must be received on or before December 1, 2021.

ADDRESSES: You may submit comments by mail to: Sherri Diana, National Institute for Occupational Safety and Health, 1090 Tusculum Avenue, MS C–34, Cincinnati, Ohio 45226.

Meeting Information: The USA toll-free dial-in numbers are: +1 669 254 5252 US (San Jose); +1 646 828 7666 US (New York); +1 551 285 1373 US; +1 669 216 1590 US (San Jose); The Meeting ID is: 161 731 2093 and the Passcode is: 45481965; Web conference by Zoom meeting connection: *https://cdc.zoomgov.com/j/1617312093?pwd=eDREUG5JaGl6Y1Z2YUVyNnJmYllHUT09.*

FOR FURTHER INFORMATION CONTACT: Rashaun Roberts, Ph.D., Designated Federal Officer, NIOSH, CDC, 1090 Tusculum Avenue, Mailstop C–24, Cincinnati, Ohio 45226, Telephone: (513) 533–6800, Toll Free: 1(800) CDC–INFO, Email: *ocas@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Background: The Advisory Board was established under the Energy Employees Occupational Illness Compensation Program Act of 2000 to advise the President on a variety of policy and technical functions required to implement and effectively manage the new compensation program. Key functions of the Advisory Board include providing advice on the development of probability of causation guidelines which have been promulgated by the Department of Health and Human Services (HHS) as a final rule, advice on methods of dose reconstruction which have also been promulgated by HHS as a final rule, advice on the scientific validity and quality of dose estimation and reconstruction efforts being performed for purposes of the compensation program, and advice on petitions to add classes of workers to the Special Exposure Cohort (SEC). In December 2000, the President delegated responsibility for funding, staffing, and operating the Advisory Board to HHS, which subsequently delegated this authority to the CDC. NIOSH implements this responsibility for CDC.

The Advisory Board's charter was issued on August 3, 2001, renewed at appropriate intervals, rechartered on March 22, 2020, and will terminate on March 22, 2022.

Purpose: The Advisory Board is charged with (a) providing advice to the Secretary, HHS, on the development of guidelines under Executive Order 13179; (b) providing advice to the Secretary, HHS, on the scientific validity and quality of dose reconstruction efforts performed for this program; and (c) upon request by the Secretary, HHS, advising the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class.