(viii) List all modifications, if any, made to the EUT by the testing company or individual to achieve compliance with the regulations in this chapter;

(ix) Include all of the data required to show compliance with the appropriate regulations in this chapter; and

(x) Contain, on the test report, the signature of the individual responsible for testing the product along with the name and signature of an official of the responsible party, as designated in § 2.909.

(4) For equipment subject to the provisions in part 15 of this chapter, the records shall indicate if the equipment was verified pursuant to the transition provisions contained in § 15.37 of this chapter.

(b) The records listed in paragraph (a) of this section shall be retained for two years after the manufacture of said equipment item has been permanently discontinued, or until the conclusion of an investigation or a proceeding if the manufacturer or importer is officially notified that an investigation or any other administrative proceeding involving his equipment has been instituted.

The Commission needs and requires the information under FCC Rules at 47 CFR parts 15 and 18, that RF equipment manufacturers (respondents) ''self-determine'' their responsibility for adherence to these rules, as guided by the following criteria:

(a) Whether the RF equipment device that is being marketed complies with the applicable Commission Rules; and

(b) If the operation of the equipment is consistent with the initially documented test results, as reported to the Commission.

The information collection is essential to controlling potential interference to radio communications.

(a) Companies that manufacture RF equipment are the anticipated respondents to this information collection.

(b) This respondent ''public'' generally remains the same, although the types of equipment devices that they manufacture may change in response to changing technologies and to new spectrum allocations made by the Commission.

(c) In addition, the Commission may establish new technical operating standards in response to these changing technologies and in allocation spectrum, which these RF equipment manufacturers must meet to receive their equipment authorization from the FCC.

(d) However, the process that RF equipment manufacturers must follow to verify their compliance, as mandated by 47 CFR 2.955 of FCC Rules, will not change despite new technical standards established for specific equipment.

This information collection, therefore, applies to a variety of equipment, which is currently manufactured in the future, and that operates under varying technical standards.

Federal Communications Commission.

Marlene H. Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2017–21516 Filed 10–4–17; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–1242]

Information Collection Approved by the Office of Management and Budget

AGENCY: Federal Communications Commission.

ACTION: Notice.

SUMMARY: The Federal Communications Commission (Commission) has received Office of Management and Budget (OMB) approval, on an emergency basis, for a new, one-time information collection pursuant to the Paperwork Reduction Act of 1995. An agency may not conduct or sponsor a collection of information unless it displays a currently valid OMB control number, and no person is required to respond to a collection of information unless it displays a currently valid control number. Comments concerning the accuracy of the burden estimates and any suggestions for reducing the burden should be directed to the person listed in the **FOR FURTHER INFORMATION CONTACT** section below.

FOR FURTHER INFORMATION CONTACT: Cathy Williams, *Cathy.Williams@fcc.gov,* (202) 418–2918.

SUPPLEMENTARY INFORMATION: The total annual reporting burdens and costs for the respondents are as follows:

OMB Control Number: 3060–1242.

OMB Approval Date: September 27, 2017.

OMB Expiration Date: March 31, 2018.

Title: Qualified 4G LTE Coverage Data Collection for Mobility Fund Phase II.

Form Number: N/A.

Respondents: Business or other for-profit entities, not-for-profit institutions, and state, local or tribal governments.

Number of Respondents and Responses: 50 respondents; 50 responses.

Estimated Time per Response: 64 hours.

Frequency of Response: One-time reporting requirement.

Total Annual Burden: 3,200 hours.

Total Annual Cost: None.

Obligation To Respond: Required to obtain or retain benefits. Statutory authority for the currently approved information collection is contained in sections 154, 254, and 303(r) of the Communications Act, as amended, 47 U.S.C. 4, 254, 303(r).

Nature and Extent of Confidentiality: The information collected under this collection is confidential and will not be made publicly available.

Privacy Act Impact Assessment: No impact(s).

Needs and Uses: In 2011, the Commission established the Mobility Fund, which consists of two phases. Mobility Fund Phase I provided one-time universal service support payments to immediately accelerate deployment of mobile broadband services. Mobility Fund Phase II (MF–II) will use a reverse auction to provide ongoing universal service support payments to continue to advance deployment of such services. In its February 2017 *Mobility Fund II Report and Order* (*MF–II Report and Order*) (FCC 17–11), the Commission adopted the rules and framework for moving forward expeditiously with the MF–II auction and stated that, prior to the auction, it would establish a map of areas presumptively eligible for MF–II support based on the most recently available FCC Form 477 mobile wireless coverage data, and provide a limited timeframe for parties to challenge those initial determinations during the pre-auction process. In its August 2017 *Order on Reconsideration and Second Report and Order* (FCC 17–102), the Commission, among other things, reconsidered its earlier decision to use FCC Form 477 data to compile the map of areas presumptively eligible for MF–II support. The Commission decided it would instead conduct a new, one-time data collection of 4G LTE coverage data that will be used for this purpose, concluding that for purposes of implementing MF–II expeditiously, this approach will provide the Commission and interested parties with the best available starting point for the challenge process and should result in fewer and more narrowly-focused challenges regarding representations of coverage. The information collected under this collection will be used by the Commission to compile the map of areas presumptively eligible for MF–II support.

The Commission received approval from OMB for the information collection requirements contained in OMB 3060–1242 on September 27, 2017.

Federal Communications Commission.

Marlene H. Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2017–21482 Filed 10–4–17; 8:45 am]

BILLING CODE 6712–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry (BSC, NCEH/ATSDR)

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

ACTION: Notice of meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, the Centers for Disease Control and Prevention (CDC), announces the following meeting for the Board of Scientific Counselors, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry (BSC, NCEH/ATSDR). This meeting is open to the public, limited in the room by 60 people and 75 lines over the phone. The public is also welcome to listen to the meeting by 1–888–790–2009, passcode: 7865774, with 75 lines. The deadline for notification of attendance is November 10, 2017. The public comment period is scheduled on Wednesday, November 15, 2017 from 2:00 p.m. until 2:15 p.m.; and from 3:25 p.m. until 3:40 p.m. EST, and on Thursday, November 16, 2017 from 10:10 a.m. until 10:25 a.m. EST. Individuals wishing to make a comment during Public Comment period, please email your name, organization, and phone number by November 6, 2017 to William Cibulas at *wic1@cdc.gov.*

DATES: The meeting will be held on November 15, 2017, 8:30 a.m. to 4:30 p.m., EST and November 16, 2017, 8:30 a.m. to noon, EST.

ADDRESSES: CDC, 4770 Buford Hwy., Atlanta, Georgia 30341, Building 107, Room 1A or by phone: 1–888–790–2009

FOR FURTHER INFORMATION CONTACT: Shirley Little, Program Analyst, NCEH/ATSDR, CDC, 4770 Buford Hwy., Mail Stop F–45, Atlanta, Georgia 30341, telephone (770) 488–0577; *snl7@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Purpose: The Secretary, Department of Health and Human Services (HHS) and by delegation, the Director, CDC and Administrator, NCEH/ATSDR, are authorized under Section 301 (42 U.S.C. 241) and Section 311 (42 U.S.C. 243) of the Public Health Service Act, as amended, to: (1) Conduct, encourage, cooperate with, and assist other appropriate public authorities, scientific institutions, and scientists in the conduct of research, investigations, experiments, demonstrations, and studies relating to the causes, diagnosis, treatment, control, and prevention of physical and mental diseases and other impairments; (2) assist states and their political subdivisions in the prevention of infectious diseases and other preventable conditions and in the promotion of health and wellbeing; and (3) train state and local personnel in health work. The BSC, NCEH/ATSDR provides advice and guidance to the Secretary, HHS; the Director, CDC and Administrator, ATSDR; and the Director, NCEH/ATSDR, regarding program goals, objectives, strategies, and priorities in fulfillment of the agency's mission to protect and promote people's health. The board provides advice and guidance that will assist NCEH/ATSDR in ensuring scientific quality, timeliness, utility, and dissemination of results. The board also provides guidance to help NCEH/ATSDR work more efficiently and effectively with its various constituents and to fulfill its mission in protecting America's health.

Matters To Be Considered: The agenda will include discussions on NCEH/ATSDR Director Updates; Noise-Induced Hearing Loss; NCEH/ATSDR Program Responses to BSC Guidance and Action Items; CDC's Hurricane Season Response; Lead Poisoning Prevention Program Updates; Flint Registry; Revision of blood lead level reference value (status); Discussion of Legislative Requirements of new Lead Exposure Poisoning Federal Advisory Committee; Amyotrophic Lateral Sclerosis (ALS) Program Update; Environmental Health Tracking Program update; updates from the National Institute of Environmental Health Sciences, the National Institute for Occupational Safety and Health, the U.S. Department of Energy and the U.S. Environmental Protection Agency. Agenda items are subject to change as priorities dictate.

The Director, Management Analysis and Services Office, 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.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2017–21422 Filed 10–4–17; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Mine Safety and Health Research Advisory Committee (MSHRAC)

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

ACTION: Notice of meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, the Centers for Disease Control and Prevention (CDC), National Institute for Occupational Safety and Health (NIOSH), announces the following meeting for the Mine Safety and Health Research Advisory Committee (MSHRAC). This meeting is open to the public, limited only by the limited only by the space available. The meeting room accommodates approximately 50 people. If you wish to attend in person or by phone, please contact Marie Chovanec by email at *MChovanec@cdc.gov* or by phone at least 5 business days in advance of the meeting.

DATES: The meeting will be held on November 15, 2017, 8:00 a.m.–3:00 p.m., Mountain Time.

ADDRESSES: Sheraton Denver West Hotel, 360 Union Boulevard, Lakewood, CO 80228 or call 412–386–5302.

FOR FURTHER INFORMATION CONTACT: Jeffrey H. Welsh, Designated Federal Officer, MSHRAC, NIOSH, CDC, 626 Cochrans Mill Road, Pittsburgh, PA 15236, telephone 412–386–4040, fax 412–386–6614.

SUPPLEMENTARY INFORMATION:

Purpose: This committee is charged with providing advice to the Secretary, Department of Health and Human Services; the Director, CDC; and the Director, NIOSH, on priorities in mine safety and health research, including grants and contracts for such research, 30 U.S.C. 812(b)(2), Section 102(b)(2).

Matters To Be Considered: The meeting will focus on mining safety and health research projects and outcomes, including built-in-place refuge alternatives, explosion protection, domestic and international