Federal Communications Commission.

Marlene H. Dortch,

Secretary, Office of Secretary.

[FR Doc. 2016–22874 Filed 9–21–16; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–1215]

Information Collection Being Reviewed by the Federal Communications Commission

AGENCY: Federal Communications Commission.

ACTION: Notice and request for comments.

SUMMARY: As part of its continuing effort to reduce paperwork burdens, and as required by the Paperwork Reduction Act (PRA) of 1995 (44 U.S.C. 3501–3520), the Federal Communications Commission (FCC or Commission) invites the general public and other Federal agencies to take this opportunity to comment on the following information collections. Comments are requested concerning: Whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and ways to further reduce the information collection burden on small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid OMB control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the PRA that does not display a valid OMB control number.

DATES: Written PRA comments should be submitted on or before November 21, 2016. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

ADDRESSES: Direct all PRA comments to Cathy Williams, FCC, via email *PRA@fcc.gov* and to *Cathy.Williams@fcc.gov.*

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Cathy Williams at (202) 418–2918.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060–1215.

Title: Use of Spectrum Bands Above 24 GHz for Mobile Radio Services.

Form Number: N/A.

Type of Review: Revision of an existing collection.

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

Number of Respondents: 247 respondents; 247 responses.

Estimated Time per Response: .5–10 hours.

Frequency of Response: On occasion reporting requirement; third party disclosure requirement; upon commencement of service, or within 3 years of effective date of rules; and at end of license term, or 2024 for incumbent licensees.

Obligation to Respond: Statutory authority for this collection are contained in sections 1, 2, 3, 4, 5, 7, 10, 201, 225, 227, 301, 302, 302a, 303, 304, 307, 309, 310, 316, 319, 332, and 336 of the Communications Act of 1934, 47 U.S.C. 151, 152, 153, 154, 155, 157, 160, 201, 225, 227, 301, 302, 302a, 303, 304, 307, 309, 310, 316, 319, 332, 336, Section 706 of the Telecommunications Act of 1996, as amended, 47 U.S.C. 1302.

Total Annual Burden: 363 hours.

Total Annual Cost: $196,875.

Privacy Act Impact Assessment: No impact(s).

Nature and Extent of Confidentiality: There is no need for confidentiality with this collection of information.

Needs and Uses: In this collection, the Commission adopted new licensing, service, and technical rules for bands 27.5–28.35 GHz band (28 GHz band), the 38.6–40 GHz band (39 GHz band), and the 37–38.6 GHz band (37 GHz band), to include 64–71 GHz band under Part 15. In so doing, the Commission created a consistent framework across all of the bands that can serve as a template for additional bands in the future.

The rules adopted by the Commission, in FCC 16–89, contain the following information collections:

Section 25.136—This rule contains both a third party coordination requirement and a filing requirement. Both requirements are necessary to ensure that Fixed Satellite Service earth stations can receive interference protection without having an undue impact on terrestrial deployment.

Section 30.3—This rule contains a filing requirement which is necessary to ascertain compliance with the foreign ownership restrictions contained in the Communications Act and the Commission's rules.

Section 30.8—This rule contains a requirement that each licensee file a statement describing its network security plans and related information, which shall be signed by a senior executive within the licensee's organization with personal knowledge of the security plans and practices within the licensee's organization. This statement is necessary to ensure that licensees properly take security into consideration when designing their systems.

Section 30.105—This rule contains filing requirements relating to demonstration of compliance with the Commission's buildout requirements. These filings are necessary in order to ensure that licensees are placing the spectrum in use and not warehousing spectrum.

Section 30.107—This rule contains filing requirements that apply when licensees propose to discontinue service. These filings are necessary in order to ensure that licensees are placing the spectrum in use and not warehousing spectrum.

Federal Communications Commission.

Marlene H. Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2016–22873 Filed 9–21–16; 8:45 am]

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FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Bank Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Bank Holding Company Act of 1956 (12 U.S.C. 1841 *et seq.*) (BHC Act), Regulation Y (12 CFR part 225), and all other applicable statutes and regulations to become a bank holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a bank or bank holding company and all of the banks and nonbanking companies owned by the bank holding company, including the companies listed below.

The applications listed below, as well as other related filings required by the Board, are available for immediate inspection at the Federal Reserve Bank indicated. The applications will also be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)). If the

proposal also involves the acquisition of a nonbanking company, the review also includes whether the acquisition of the nonbanking company complies with the standards in section 4 of the BHC Act (12 U.S.C. 1843). Unless otherwise noted, nonbanking activities will be conducted throughout the United States.

Unless otherwise noted, comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than October 17, 2016.

A. Federal Reserve Bank of Cleveland (Nadine Wallman, Vice President) 1455 East Sixth Street, Cleveland, Ohio 44101–2566. Comments can also be sent electronically to *Comments.applications@clev.frb.org:*

1. *F.N.B. Corporation,* Pittsburgh, Pennsylvania; to acquire Yadkin Financial Corporation, Raleigh, North Carolina, and thereby acquire Yadkin Bank, Statesville, North Carolina,

Board of Governors of the Federal Reserve System, September 19, 2016.

Michele Taylor Fennell,

Assistant Secretary of the Board.

[FR Doc. 2016–22847 Filed 9–21–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–16–0997]

Agency Forms Undergoing Paperwork Reduction Acct Review

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

ACTION: Notice with comment period; withdrawal.

SUMMARY: The Centers for Disease Control and Prevention (CDC) in the Department of Health and Human Services (HHS) announces the withdrawal of the notice published under the same title on August 25, 2016 for public comment.

DATES: Effective September 22, 2016.

FOR FURTHER INFORMATION CONTACT: 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: On August 25, 2016 CDC published a notice in the **Federal Register** titled ''Agency Forms Undergoing Paperwork Reduction Act Review'' (Vol. 81, No. 165 FR Doc. 2016–20333, Pages 58511–58512). This notice was published prematurely and inadvertently. The notice is being withdrawn immediately for public comment. A new notice will be published at a later date for public comment.

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. 2016–22866 Filed 9–21–16; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2016–N–0001]

Technical Electronic Product Radiation Safety Standards Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) announces a forthcoming public advisory committee meeting of the Technical Electronic Product Radiation Safety Standards Committee. The general function of the committee is to provide advice and recommendations to the Agency on FDA's regulatory issues. The meeting will be open to the public.

DATES: The meeting will be held on October 25, 2016, from 8:30 a.m. to 5 p.m. and October 26, 2016, from 8:30 a.m. to 5 p.m.

ADDRESSES: Gaithersburg Holiday Inn, Ballroom, Two Montgomery Village Ave., Gaithersburg, MD 20879. The hotel's telephone number is 301–948–8900. Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.*

FOR FURTHER INFORMATION CONTACT: Sara J. Anderson, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 1643, Silver Spring, MD 20993–0002, *sara.anderson@fda.hhs.gov,* 301–796–7047, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site at *http://www.fda.gov/AdvisoryCommittees/default.htm* and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

SUPPLEMENTARY INFORMATION:

Agenda: The general function of the committee is to provide advice and recommendations to the Agency on the technical feasibility, reasonableness, and practicability of performance standards for electronic products to control the emission of radiation from such products, and may recommend electronic product radiation safety standards to the Agency for consideration.

On October 25, 2016, the committee will discuss and make recommendations regarding possible FDA performance standards for the following topics: Radiofrequency (RF) radiation products, such as microwave ovens and wireless power transfer; laser products, including an update to amendments to the laser rule, light detection and ranging (LIDAR), laser data (Light Fidelity-LiFi)/energy transfer, illumination applications and infrared applications; sunlamp products including an update on the performance standards amendments; and non-coherent light sources (*e.g.,* LEDs and UVC lamps) including new initiatives.

On October 26, 2016, the committee will discuss and make recommendations regarding possible FDA performance standards for the following topics: International Electrotechnical Commission (IEC) standards versus performance standards for medical devices; computed tomography (CT); radiography and fluoroscopy; diagnostic and therapeutic ultrasound; and radiation therapy.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the