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]

BILLING CODE 6210–01–P

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 The notice is being withdrawn immediately for public comment.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