following information collection. 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 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 Office of Management and Budget (OMB) control number.

DATES: Written PRA comments should be submitted on or before December 5, 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 Nicole Ongele, FCC, via email *PRA@fcc.gov* and to *Nicole.Ongele@fcc.gov.*

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Nicole Ongele at (202) 418–2991.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060–0741.

Title: Technology Transitions, GN Docket No. 13–5, et al.

Form Number(s): N/A.

Type of Review: Revision of a currently approved collection.

Respondents: Business or other for-profit entities.

Number of Respondents and Responses: 5,357 respondents; 573,767 responses.

Estimated Time per Response: 0.5–8 hours.

Frequency of Response: On occasion and one-time reporting requirements; recordkeeping and third party disclosure requirements.

Obligation to Respond: Required to obtain or retain benefits. Statutory authority is contained in 47 U.S.C. 251.

Total Annual Burden: 575,840 hours.

Total Annual Cost: No cost.

Privacy Impact Assessment: No impact(s).

Nature and Extent of Confidentiality: The Commission is not requesting that the respondents submit confidential information to the FCC. Respondents may, however, request confidential treatment for information they believe to be confidential under 47 CFR 0.459 of the Commission's rules.

Needs and Uses: Section 251 of the Communications Act of 1934, as amended, 47 U.S.C. 251, is designed to accelerate private sector development and deployment of telecommunications technologies and services by spurring competition. These OMB collections are designed to help implement certain provisions of section 251, and to eliminate operational barriers to competition in the telecommunications services market. Specifically, these OMB collections will be used to implement (1) local exchange carriers' (''LECs'') obligations to provide their competitors with dialing parity and non-discriminatory access to certain services and functionalities; (2) incumbent local exchange carriers' (''ILECs'') duty to make network information disclosures; and (3) numbering administration. The Commission estimates that the total annual burden of the entire collection, as revised, is 575,840 hours. This revision relates to a change in one of many components of the currently approved collection—specifically, certain reporting, recordkeeping and/or third party disclosure requirements under section 251(c)(5). In August 2015, the Commission adopted new rules concerning certain information collection requirements implemented under section 251(c)(5) of the Act, pertaining to network change disclosures. The changes to those rules applied specifically to a certain subset of network change disclosures, namely notices of planned copper retirements. The changes were designed to provide interconnecting entities adequate time to prepare their networks for the planned copper retirements and to ensure that consumers are able to make informed choices. The Commission estimated that the 2015 revisions did not result in any additional burden hours or outlays of funds for hiring outside contractors or procuring equipment. In July 2016, the Commission revised section 51.329(c) of its network change disclosure rules to make available to filers new titles applicable to copper retirement notices. The Commission estimates that the revision does not result in any additional burden hours or outlays of funds for hiring outside contractors or procuring.

Federal Communications Commission.

Marlene H. Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2016–24069 Filed 10–4–16; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL DEPOSIT INSURANCE CORPORATION

FDIC Advisory Committee on Economic Inclusion (ComE–IN); Notice of Meeting

AGENCY: Federal Deposit Insurance Corporation (FDIC).

ACTION: Notice of open meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, Public Law 92–463 (Oct. 6, 1972), 5 U.S.C. App. 2, notice is hereby given of a meeting of the FDIC Advisory Committee on Economic Inclusion, which will be held in Washington, DC The Advisory Committee will provide advice and recommendations on initiatives to expand access to banking services by underserved populations.

DATES: Thursday, October 20, 2016, from 9:00 a.m. to 3:30 p.m.

ADDRESSES: The meeting will be held in the FDIC Board Room on the sixth floor of the FDIC Building located at 550 17th Street NW., Washington, DC.

FOR FURTHER INFORMATION CONTACT: Requests for further information concerning the meeting may be directed to Mr. Robert E. Feldman, Committee Management Officer of the FDIC, at (202) 898–7043.

SUPPLEMENTARY INFORMATION:

Agenda: The agenda will be focused on the FDIC's National Survey of Unbanked and Underbanked Households, the FDIC's Youth Savings Pilot, and expanding access to safe transaction accounts. The agenda may be subject to change. Any changes to the agenda will be announced at the beginning of the meeting.

Type of Meeting: The meeting will be open to the public, limited only by the space available on a first-come, first-served basis. For security reasons, members of the public will be subject to security screening procedures and must present a valid photo identification to enter the building. The FDIC will provide attendees with auxiliary aids (*e.g.,* sign language interpretation) required for this meeting. Those attendees needing such assistance should call (703) 562–6067 (Voice or TTY) at least two days before the meeting to make necessary arrangements. Written statements may be filed with the committee before or

after the meeting. This ComE–IN meeting will be Webcast live via the Internet at: *http://fdic.windrosemedia.com.* Questions or troubleshooting help can be found at the same link. For optimal viewing, a high speed internet connection is recommended. The ComE–IN meeting videos are made available on-demand approximately two weeks after the event.

Dated: September 30, 2016.

Federal Deposit Insurance Corporation.

Robert E. Feldman,

Executive Secretary, Federal Deposit Insurance Corporation.

[FR Doc. 2016–24039 Filed 10–4–16; 8:45 am]

BILLING CODE 6714–01–P

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 31, 2016.

A. Federal Reserve Bank of Dallas (Robert L. Triplett III, Senior Vice President), 2200 North Pearl Street, Dallas, Texas 75201–2272:

1. *Caldwell Holding Company,* Columbia, Louisiana; to acquire Progressive National Financial Corporation, and thereby indirectly acquire Progressive National Bank, both in Mansfield, Louisiana.

Board of Governors of the Federal Reserve System, September 30, 2016.

Margaret McCloskey Shanks,

Deputy Secretary of the Board.

[FR Doc. 2016–24055 Filed 10–4–16; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2016–N–0001]

Pharmacy Compounding Advisory 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 Pharmacy Compounding Advisory Committee (PCAC). The general function of the committee is to provide advice on scientific, technical, and medical issues concerning drug compounding, as well as any other product for which FDA has regulatory responsibility, and to make appropriate recommendations to the Agency. The meeting will be open to the public.

DATES: The meeting will be held on November 3, 2016, from 8:30 a.m. to 4:30 p.m.

ADDRESSES: FDA White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503), Silver Spring, MD 20993–0002. 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: Cindy Hong, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993–0002, 301–796–9001, FAX: 301–847–8533, *PCAC@fda.hhs.gov,* 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.

Background: Section 503A of the FD&C Act (21 U.S.C. 353a) describes the conditions that must be satisfied for human drug products compounded by a licensed pharmacist in a State licensed pharmacy or a Federal facility, or licensed physician, to be exempt from the following three sections of the Federal Food, Drug, and Cosmetic Act (FD&C Act): (1) Section 501(a)(2)(B) (21 U.S.C. 351(a)(2)(B)) concerning current good manufacturing practice (CGMP); (2) section 502(f)(1) (21 U.S.C. 352(f)(1)) concerning the labeling of drugs with adequate directions for use; and (3) section 505 (21 U.S.C. 355) concerning the approval of human drug products under new drug applications (NDAs) or abbreviated new drug applications (ANDAs).

The Drug Quality and Security Act added a new section 503B to the FD&C Act (21 U.S.C. 353b), which created a new category of compounders termed ''outsourcing facilities.'' Under section 503B of the FD&C Act, outsourcing facilities are defined, in part, as facilities that meet certain conditions described in section 503B, including registration with FDA as an outsourcing facility. If these conditions are satisfied, a drug product compounded for human use by or under the direct supervision of a licensed pharmacist in an outsourcing facility is exempt from three sections of the FD&C Act: (1) Section 502(f)(1) concerning the labeling of drugs with adequate directions for use); (2) section 505 concerning the approval of human drug products under NDAs or ANDAs; and (3) section 582 concerning the drug supply chain security requirements (21 U.S.C. 360eee–1). Outsourcing facilities are not exempt from CGMP requirements in section 501(a)(2)(B).

One of the conditions that must be satisfied to qualify for the exemptions under section 503A of the FD&C Act is that a bulk drug substance (active pharmaceutical ingredient) used in a compounded drug product must meet one of the following criteria: (1) Complies with the standards of an applicable United States Pharmacopoeia (USP) or National Formulary monograph, if a monograph exists, and the USP chapter on pharmacy compounding; (2) if an applicable