B. Federal Reserve Bank of Kansas City (Dennis Denney, Assistant Vice President), 1 Memorial Drive, Kansas City, Missouri 64198–0001:

1. *Kevin Page, Trustee of the Kevin Page Trust, Fort Worth, Texas, Dana Page, Co-Trustee of the Dana Page Trust, Austin, Texas, Meghan Anderson Smith, Co-Trustee of the Meghan Anderson Smith Trust, Fremont, Nebraska, Whitney Anderson, Co-Trustee of the Whitney Anderson Trust, Coppell, Texas, Eric Jones, Trustee of the Eric Jones Trust, Blue Lake, California, and Christopher Marious Jones, Co-Trustee of the Christopher Marious Jones Trust, Los Angeles, California;* to retain shares of Page Bancshares, Inc., Liberty, Missouri, and thereby retain shares of Pony Express Bank, Braymer, Missouri.

Board of Governors of the Federal Reserve System, August 6, 2018.

Yao-Chin Chao,

Assistant Secretary of the Board.

[FR Doc. 2018–17094 Filed 8–8–18; 8:45 am]

BILLING CODE P

FEDERAL RESERVE SYSTEM

Notice of Proposals To Engage in or To Acquire Companies Engaged in Permissible Nonbanking Activities

The companies listed in this notice have given notice under section 4 of the Bank Holding Company Act (12 U.S.C. 1843) (BHC Act) and Regulation Y, (12 CFR part 225) to engage *de novo,* or to acquire or control voting securities or assets of a company, including the companies listed below, that engages either directly or through a subsidiary or other company, in a nonbanking activity that is listed in § 225.28 of Regulation Y (12 CFR 225.28) or that the Board has determined by Order to be closely related to banking and permissible for bank holding companies. Unless otherwise noted, these activities will be conducted throughout the United States.

Each notice is available for inspection at the Federal Reserve Bank indicated. The notice also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the question whether the proposal complies with the standards of section 4 of the BHC Act.

Unless otherwise noted, comments regarding the applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than August 23, 2018.

A. Federal Reserve Bank of New York (Ivan Hurwitz, Vice President) 33 Liberty Street, New York, New York 10045–0001. Comments can also be sent electronically to *Comments.applications@ny.frb.org:*

1. *Standard Chartered Bank, London, England;* through its subsidiary, Standard Chartered Holdings, Inc., New York, New York, to engage through a newly formed entity, The Consortium, LLC, in data processing activities, pursuant to section 225.28(b)(14) of Regulation Y.

Board of Governors of the Federal Reserve System, August 3, 2018.

Ann Misback,

Secretary of the Board.

[FR Doc. 2018–17010 Filed 8–8–18; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket Number: NIOSH 278]

Board of Scientific Counselors, National Institute for Occupational Safety and Health (BSC, NIOSH)

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 CDC announces the following meeting of the Board of Scientific Counselors, National Institute for Occupational Safety and Health (BSC, NIOSH). This meeting is open to the public, limited only by the space available. The meeting room accommodates approximately 33 people. The meeting is also open to the public via webcast. If you wish to attend in person or by webcast, please see the NIOSH website to register (*http://www.cdc.gov/niosh/bsc/*) or call (404–498–2539) at least five business days in advance of the meeting. Teleconference is available toll-free; please dial (888) 397–9578, Participant Pass Code 63257516. Adobe Connect webcast will be available at *https://odniosh.adobeconnect.com/nioshbsc/* for participants wanting to connect remotely. This meeting is open to the public, limited only by the space available. The public is welcome to participate during the public comment period, 12:30 p.m. to 12:45 p.m., EDT, September 27, 2018. Please note that the public comment period ends at the time indicated above. Each commenter will be provided up to five minutes for comment. A limited number of time slots are available and will be assigned on a first come-first served basis. Written comments will also be accepted from those unable to attend the public session via an on-line form at the following website: *http://www.cdc.gov/niosh/bsc/contact.html.*

DATES: The meeting will be held on September 27, 2018, 8:30 a.m.–2:30 p.m., EDT.

ADDRESSES: Patriots Plaza I, 395 E Street SW, Room 9000, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Alberto Garcia, M.S., Executive Secretary, BSC, NIOSH, CDC, 1090 Tusculum Avenue, MS–R5, Cincinnati, OH 45226, telephone (513) 841–4596, fax (513) 841–4506, email: *agarcia1@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Purpose: The Secretary, the Assistant Secretary for Health, and by delegation the Director, Centers for Disease Control and Prevention, are authorized under Sections 301 and 308 of the Public Health Service Act to conduct directly or by grants or contracts, research, experiments, and demonstrations relating to occupational safety and health and to mine health. The Board of Scientific Counselors provides guidance to the Director, National Institute for Occupational Safety and Health on research and prevention programs. Specifically, the Board provides guidance on the Institute's research activities related to developing and evaluating hypotheses, systematically documenting findings and disseminating results. The Board evaluates the degree to which the activities of the National Institute for Occupational Safety and Health: (1) Conform to appropriate scientific standards, (2) address current, relevant needs, and (3) produce intended results.

Matters to be Considered: The agenda for the meeting addresses occupational safety and health issues related to: NIOSH confronts the Opioid Crisis; 21st Century Surveillance Report; and Enhancing the Transparency of NIOSH Science. An agenda is also posted on the NIOSH website (*http://www.cdc.gov/niosh/bsc/*). Members of the public who wish to address the NIOSH BSC are requested to contact the Executive Secretary for scheduling purposes (see contact information below). Alternatively, written comments to the BSC may be submitted via an on-line form at the following website: *http://www.cdc.gov/niosh/bsc/contact.html.* 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.

Sherri A. Berger,

Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2018–17042 Filed 8–8–18; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

In accordance with Section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting.

The meeting will be closed to the public in accordance with provisions set forth in Section 552b(c) (4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Name of Committee: Safety and Occupational Health Study Section (SOHSS), National Institute for Occupational Safety and Health (NIOSH).

Dates: October 16–October 18, 2018.

Time: 8:00 a.m.–5:00 p.m., EDT.

Place: Embassy Suites, 1900 Diagonal Road, Alexandria, VA 22314.

Agenda: The meeting will convene to address matters related to the conduct of Study Section business and for the study section to consider safety and occupational health-related grant applications.

For Further Information Contact: Nina Turner, Ph.D., Scientific Review Officer, NIOSH, 1095 Willowdale Road, Morgantown, WV 26506, (304) 285–5976; *nturner@cdc.gov.*

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.

Sherri Berger,

Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2018–17043 Filed 8–8–18; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–N–0558]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Disclosures in Professional and Consumer Prescription Drug Promotion

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (PRA).

DATES: Fax written comments on the collection of information by September 10, 2018.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, Fax: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–NEW and title ''Disclosures in Professional and Consumer Prescription Drug Promotion.'' Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–7726, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Disclosures in Professional and Consumer Prescription Drug Promotion

OMB Control Number 0910–NEW

I. Background

Section 1701(a)(4) of the Public Health Service Act (42 U.S.C. 300u(a)(4)) authorizes FDA to conduct research relating to health information. Section 1003(d)(2)(C) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. 393(d)(2)(C)) authorizes FDA to conduct research relating to drugs and other FDA regulated products in carrying out the provisions of the FD&C Act.

FDA regulates prescription drug advertising and promotional labeling directed to healthcare professionals (HCPs) and consumers (section 502(a) and (n), respectively, of the FD&C Act (21 U.S.C. 352(a) and (n))). In the course of promoting their products, pharmaceutical sponsors (sponsors) may present a variety of information including the indication, details about the administration of the product, efficacy information, and clinical trial data. To present often complicated information concisely, sponsors may not include relevant information in the body of the text or visual display of the claim. Additionally, sponsors may not always present limitations to the claim in the main body of the text or display. In these cases, sponsors typically include disclosures of information somewhere in the promotional piece.

There is limited published research on disclosures in prescription drug promotion, either directed to consumers or to HCPs. The use of disclosures is one method of communicating information to HCPs and consumers about scientific and clinical data, the limitations of that data, and practical utility of that information. These disclosures may influence HCP and consumer comprehension and decision making, and may affect how and what treatment HCPs prescribe for their patients. Previous research on the effectiveness of disclosures has been conducted primarily in the dietary supplement arena (Refs. 1–4). Thus, the proposed research will examine the effectiveness of clear and conspicuous disclosures in prescription drug promotion directed to both populations. The purpose of our study is to determine how useful disclosures regarding prescription drug information are when presented prominently and adjacent to claims.[1] Specifically, are HCPs and consumers able to use disclosures to effectively frame information in efficacy claims in prescription drug promotion?

To address this research question, we have designed a set of studies that cover both consumers and HCPs, as well as three presentations addressing different

[1] The Federal Trade Commission (FTC), which regulates the advertising of non-prescription drug products as well as other non-FDA regulated products (*e.g.,* package goods, cars, etc.) issued a specific position on disclosures (Ref. 5) for the advertising it regulates. Specifically, FTC explains that disclosures must be ''clear and conspicuous''; in other words, in understandable language, located near the claim to be further clarified, and not hidden or minimized by small font or other distractions.