rules are satisfied to receive various levels of pricing flexibility. These showings, which focus on unaffiliated collocations in wire centers, are intended to measure the presence of competition in a given Metropolitan Statistical Area (MSA) or non-MSA so that regulatory relief is granted where warranted. The Commission's rules provide that petitions for pricing flexibility for special access services that are not denied within 90 days after the close of the pleading cycle are deemed granted.

In an August 2012 *Report and Order,* FCC 12–92, 57 FR 57504 (Sept. 12, 2012), the Commission suspended, on an interim basis, the 90-day deadline for the granting of pricing flexibility pending adoption of a new regulatory framework. Notwithstanding the temporary suspension, the Commission is seeking Office of Management and Budget (OMB) approval for an extension to obtain the full three-year clearance for this expiring collection in the event the suspension is lifted. The Commission will separately seek OMB's approval for any subsequent modification of this collection, as a result of changes to the pricing flexibility rules, in accordance with the Paperwork Reduction Act of 1995, Public Law 104–13.

Federal Communications Commission.

Gloria J. Miles,

Federal Register Liaison, Office of the Secretary, Office of Managing Director.

[FR Doc. 2014–13975 Filed 6–13–14; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL COMMUNICATIONS COMMISSION

Information Collection Being Reviewed by the Federal Communications Commission Under Delegated Authority

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 August 15, 2014. 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 <mailto:PRA@fcc.gov>* and to *Cathy.Williams@fcc.gov <mailto: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 No.: 3060–1008.

Title: Section 27.50, Power and Antenna Height Limits; Section 27.602, Guard Band Manger Agreements.

Form No.: Not applicable.

Type of Review: Extension of a currently approved collection.

Respondents: Business or other for-profit, and State, Local or Tribal Government.

Number of Respondents and Responses: 395 respondents and 444 responses.

Estimated Time per Response: 30 minutes up to 6 hours.

Frequency of Response: Recordkeeping requirement, On occasion reporting requirement and Third party disclosure requirement.

Obligation To Respond: Required to obtain or retain benefits. The statutory authority for this collection is contained in 47 U.S.C. 151, 154(i), 157 and 309(j), as amended.

Total Annual Burden: 519 hours.

Annual Cost Burden: None.

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: The information gathered in this collection will be used to support the development of new services in the Lower 700 MHz Band. Further, Guard Band Managers are required to enter into written agreements with other licensees who plan on using their licensed spectrum by others, subject to certain conditions outlined in the rules. They must retain these records for at least two years after the date such agreement expire. Such records need to be kept current and be made available upon request for inspection by the Commission or its representatives.

Federal Communications Commission.

Gloria J. Miles,

Federal Register Liaison, Office of the Secretary, Office of Managing Director.

[FR Doc. 2014–13972 Filed 6–13–14; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisitions of Shares of a Bank or Bank Holding Company

The notificants listed below have applied under the Change in Bank Control Act (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire shares of a bank or bank holding company. The factors that are considered in acting on the notices are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The notices are available for immediate inspection at the Federal Reserve Bank indicated. The notices also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for that notice or to the offices of the Board of Governors. Comments must be received not later than July 1, 2014.

A. Federal Reserve Bank of Cleveland (Nadine Wallman, Vice President) 1455 East Sixth Street, Cleveland, Ohio 44101–2566:

1. *George and Georgia Gia Bavelis, both of Upper Arlington, Ohio; Tina Bavelis Manokas and Tasos Manokas; and Niki Bavelis Callahan,* all of Columbus, Ohio; to retain voting shares of Heartland Bancorp, and thereby indirectly retain voting shares of Heartland Bank, both in Gahanna, Ohio.

Board of Governors of the Federal Reserve System, June 11, 2014.

Michael J. Lewandowski,

Associate Secretary of the Board.

[FR Doc. 2014–13999 Filed 6–13–14; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Initial Review

The meeting announced below concerns the NIOSH Childhood Agriculture, RFA OH–14–005, initial review.

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 aforementioned meeting:

Time and Date: 1:00 p.m.–4:00 p.m., July 8, 2014 (Closed)

Place: Teleconference

Status: 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.

Matters To Be Discussed: The meeting will include the initial review, discussion, and evaluation of applications received in response to ''NIOSH Childhood Agriculture, RFA OH–14–005.''

Contact Person for More Information: Nina Turner, Ph.D., Scientific Review Officer, 1095 Willowdale Road, Morgantown, WV 26506, Telephone: (304) 285–5976.

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. 2014–13998 Filed 6–13–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0420]

Agency Information Collection Activities; Proposed Collection; Comment Request; Testing Communications on Food and Drug Administration-Regulated Products Used in Animals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information and to allow 60 days for public comment in response to the notice. This notice solicits comments on communication studies involving FDA-regulated products intended for use in animals. This information will be used to explore concepts of interest and assist in the development and modification of communication messages and campaigns to fulfill the Agency's mission to protect the public health.

DATES: Submit written or electronic comments on the collection of information by August 15, 2014.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Testing Communications on FDA/Center for Veterinary Medicine (CVM)—Regulated Products Used in Animals—21 U.S.C. 393(d)(2)(D) (OMB Control Number 0910–0689)—Extension

FDA is authorized by section 393(d)(2)(D) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 393(d)(2)(D)) to conduct educational and public information programs relating to the safety of CVM-regulated products. FDA must conduct needed research to ensure that such programs have the highest likelihood of being effective. FDA expects that improving communications about the safety of regulated animal drugs, feed, food additives, and devices will involve many research methods, including individual indepth interviews, mall-intercept interviews, focus groups, self-administered surveys, gatekeeper reviews, and omnibus telephone surveys.

The information collected will serve three major purposes. First, as formative research it will provide critical knowledge needed about target audiences to develop messages and campaigns about the use of FDA-regulated products for use in animals. Knowledge of consumer and veterinary professional decision-making processes will provide the better understanding of target audiences that FDA needs to design effective communication