Further, providers will use the information to ensure that roaming will work expeditiously in times of emergencies and to better understand their network capabilities related to roaming and ensure their networks roam as effectively as possible when a disaster strikes. Further, the Commission will use information as a basis for potential future improvements to the MDRI and other programs in furtherance of public safety, including by gauging providers' compliance with the MDRI's roaming provision, ensuring accountability by providers who fail to comply and for resolving disputes related to roaming agreements. Thus, the information sought in this collection is necessary and vital to ensuring that the MDRI is effective at protecting the life and property of the public.

Federal Communications Commission.

Marlene Dortch.

Secretary, Office of the Secretary. [FR Doc. 2023–07695 Filed 4–11–23; 8:45 am]

BILLING CODE 6712-01-P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060-0512; FR ID 135566]

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, the Federal Communications Commission (FCC or the Commission) invites the general public and other Federal agencies to take this opportunity to comment on the 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 June 12, 2023. 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, (202) 418–2991.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060–0512. Title: ARMIS Annual Summary Seport.

Report Number: FCC Report 43–01. Type of Review: Extension of a currently approved collection.

Respondents: Business or other forprofit entities.

Number of Respondents and Responses: 90 respondents; 90 responses.

Estimated Time per Response: 8 hours.

Frequency of Response: Annual reporting requirement.

Obligation to Respond: Mandatory. Statutory authority for this information collection is contained in 47 U.S.C. 219 and 220 of the Communications Act of 1934, as amended.

Total Annual Burden: 720 hours. Total Annual Cost: No cost.

Needs and Uses: The information contained in FCC Report 43–01 helps the Commission fulfill its regulatory responsibilities regarding pole attachment rates. The Commission has granted all carriers forbearance from ARMIS 43–01 with the exception that carriers are still required to file pole attachment cost data. These data are required to allow the Commission to fulfill its responsibilities under Section 224 of the Communications Act of 1934, as amended.

Federal Communications Commission.

Marlene Dortch,

Secretary, Office of the Secretary. [FR Doc. 2023–07693 Filed 4–11–23; 8:45 am]

BILLING CODE 6712-01-P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060-0917; FR ID 135572]

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, the Federal Communications Commission (FCC or the Commission) invites the general public and other Federal agencies to take this opportunity to comment on the 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 June 12, 2023. 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, (202) 418–2991.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060–0917. Title: CORES Registration Form, FCC Form 160. Form Number: FCC Form 160. Type of Review: Extension of a currently approved collection.

Respondents: Businesses or other forprofit entities; Individuals or households; Not-for-profit institutions; and State, Local, or Tribal Governments.

Number of Respondents and Responses: 145,726 respondents; 145,726 responses.

Estimated Time per Response: 10 minutes (0.167 hours).

Frequency of Response: One-time reporting requirement.

Obligation to Respond: Required to obtain or retain benefits. Statutory authority for this information collection is contained in the Debt Collection Act of 1996 (DCCA), Public Law 104–134, Chapter 10, Section 31001.

Total Annual Burden: 24,366 hours. Total Annual Costs: No Cost.

Needs and Uses: Respondents use FCC Form 160 to register in FCC's Commission Registration System (CORES). Entities must register in CORES to do regulatory transactions with FCC, including receiving licenses, paying fees, participating in auctions, etc. Without this collection of information, FCC would not have a database of the identity and contact information of the entities it does regulatory business with.

Federal Communications Commission.

Marlene Dortch,

Secretary, Office of the Secretary. [FR Doc. 2023–07694 Filed 4–11–23; 8:45 am]

BILLING CODE 6712-01-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 public portions of the applications listed below, as well as other related filings required by the Board, if any, are available for immediate inspection at the Federal Reserve Bank(s) indicated below and at the offices of the Board of Governors. This information may also be obtained

on an expedited basis, upon request, by contacting the appropriate Federal Reserve Bank and from the Board's Freedom of Information Office at https://www.federalreserve.gov/foia/request.htm. Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)).

Comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors, Ann E. Misback, Secretary of the Board, 20th Street and Constitution Avenue NW, Washington DC 20551–0001, not later than May 12, 2023.

A. Federal Reserve Bank of Richmond (Brent B. Hassell, Assistant Vice President) P.O. Box 27622, Richmond, Virginia 23261, or electronically to Comments.applications@rich.frb.org:

1. Churchill Bank Corporation, Clearwater, Florida; to become a bank holding company by acquiring Miners Exchange Bank, Coeburn, Virginia.

B. Federal Reserve Bank of Minneapolis (Stephanie Weber, Assistant Vice President) 90 Hennepin Avenue, Minneapolis, Minnesota 55480–0291, or electronically to MA@ mpls.frb.org:

1. First Financial Corporation, Arthur, North Dakota; to merge with HSB Financial Corporation, and thereby indirectly acquire Harwood State Bank, both of Harwood, North Dakota.

Board of Governors of the Federal Reserve System. $\,$

Ann E. Misback,

Secretary of the Board. [FR Doc. 2023–07715 Filed 4–11–23; 8:45 am]

BILLING CODE 6210-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration [Docket No. FDA-2019-D-0362]

A Risk-Based Approach To Monitoring of Clinical Investigations—Questions and Answers; Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a final guidance for industry entitled "A Risk-Based Approach to Monitoring of Clinical Investigations—Questions and Answers." This guidance provides information on risk-based approaches to

monitoring investigational studies of human drug and biological products, medical devices, and combination products. The guidance contains recommendations on planning a monitoring approach, developing the content of a monitoring plan, and addressing and communicating monitoring results. This guidance expands on the guidance for industry entitled "Oversight of Clinical Investigations—A Risk-Based Approach to Monitoring" (August 2013) by providing additional information to facilitate sponsors' implementation of risk-based monitoring. This guidance finalizes the draft guidance entitled "A Risk-Based Approach to Monitoring of Clinical Investigations: Questions and Answers," issued on March 15, 2019.

DATES: The announcement of the guidance is published in the **Federal Register** on April 12, 2023.

ADDRESSES: You may submit either electronic or written comments on Agency guidances at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

 Federal eRulemaking Portal: https://www.regulations.gov. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to https:// www.regulations.gov will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on https://www.regulations.gov.

• If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

Written/Paper Submissions

Submit written/paper submissions as follows:

• Mail/Hand Delivery/Courier (for written/paper submissions): Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.