

include the EPA's PSD preconstruction permit program, Part 71 Title V operating permit program, NSPS and NESHAP promulgated under section 112 of the CAA. State and local air pollution control agencies are usually requested to provide information concerning regulation of offshore sources and are provided opportunities to comment on the proposed determinations. The public is also provided an opportunity to comment on the proposed determinations.

Form numbers: None.

Respondents/affected entities: Entities potentially affected by this action are those that must apply for and obtain an OCS permit pursuant to the OCS permit program. In addition, State and local agencies that have been delegated authority to implement and enforce the OCS permit program, which must review permit applications and issue permits, are affected entities.

Respondent's obligation to respond: Mandatory (CFR part 55).

Estimated number of respondents: 74 industrial facilities and 7 State and local permitting agencies.

Frequency of response: On occasion, as necessary.

Total estimated burden: 36,001 hours (per year). Burden is defined at 5 CFR 1320.03(b).

Total estimated cost: \$3,755,783.00 (per year), includes \$380,372 annualized capital or operation & maintenance costs.

Changes in estimates: There is a projected increase of 15,778 hours in the total estimated respondent burden compared with the ICR currently approved by OMB. This increase is primarily due to the projected number of OCS sources subject to the program mainly related to alternative energy sources including new wind power farms along the eastern seaboard of the United States.

Courtney Kerwin,

Director, Information Engagement Division.

[FR Doc. 2024-11806 Filed 5-29-24; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OMS-2023-0605; FRL-12011-01-OMS]

Agency Information Collection Activities; Submission to the Office of Management and Budget for Review and Approval; Comment Request; Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (Renewal)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency (EPA) has submitted an information collection request (ICR), Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (EPA ICR Number 2434.204, OMB Control Number 2030-0051) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act. This is a proposed extension of the ICR, which is currently approved through May 31, 2024. Public comments were previously requested via the **Federal Register** on January 26, 2024 during a 60-day comment period. This notice allows for an additional 30 days for public comments.

DATES: Comments may be submitted on or before July 1, 2024.

ADDRESSES: Submit your comments, referencing Docket ID Number: EPA-HQ-OMS-2023-0605, to EPA online using www.regulations.gov (our preferred method) or by mail to: EPA Docket Center, Environmental Protection Agency, Mail Code 28221T, 1200 Pennsylvania Ave. NW, Washington, DC 20460. EPA's policy is that all comments received will be included in the public docket without change including any personal information provided, unless the comment includes profanity, threats, information claimed to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute.

Submit written comments and recommendations to OMB for the proposed information collection within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting "Currently under 30-day Review—Open for Public Comments" or by using the search function.

FOR FURTHER INFORMATION CONTACT:

Aaron Jackson, Information Engagement Division (IED), Office of Information

Management (OIM), 282T, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460; telephone number: (202) 566-0348; email address: jackson.aaron@epa.gov.

SUPPLEMENTARY INFORMATION: This is a proposed extension of the ICR, which is currently approved through May 31, 2024. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number.

Public comments were previously requested via the **Federal Register** on January 26, 2024 during a 60-day comment period (89 FR 5228). This notice allows for an additional 30 days for public comments. Supporting documents, which explain in detail the information that the EPA will be collecting, are available in the public docket for this ICR. The docket can be viewed online at www.regulations.gov or in person at the EPA Docket Center, WJC West, Room 3334, 1301 Constitution Ave. NW, Washington, DC. The telephone number for the Docket Center is 202-566-1744. For additional information about EPA's public docket, visit <http://www.epa.gov/dockets>.

Abstract: The information collection activity provides EPA with an opportunity to efficiently engage its customers and stakeholders by gathering qualitative information about their current or potential future interaction with Agency. Getting such feedback in a timely manner is critical if the Agency is to know how and where it should focus while seeking to improve, or expand upon, its products and services. Following the pathway established by OMB for fast-track generic ICRs, the Agency will submit individual requests for specific information collections on an as-needed basis. Those requests will describe the collection and the public burden created. The Agency will submit a collection request for approval under this generic clearance only if the collections are: voluntary; low burden and low-cost for both the respondents and the federal government; noncontroversial; targeted to respondents who have experience with the program or may have experience with the program in the near future; and abstain from collecting personally identifiable information (PII) to the greatest extent possible. Information gathered will be used internally for general service improvement and program management purposes and released publicly only in an anonymized or aggregated fashion. Information gathered will not be used in

statistical analysis intended to yield results that can be generalized to the population of study, nor will it be used to substantially inform influential policy decisions.

Form Numbers: To be provided in individual collection requests.

Respondents/affected entities: Individuals, businesses, organizations, and state, local, and Tribal representatives that are stakeholders in, consumers of, or partners in providing, EPA or EPA-supported current or potential services and programs.

Respondent's obligation to respond: Voluntary.

Estimated number of respondents: 180,000 (total).

Frequency of response: Varies.

Total estimated burden: 45,000 hours (per year). Burden is defined at 5 CFR 1320.03(b).

Total estimated cost: There are no expected capital or operation & maintenance costs.

Changes in the Estimates: There are no changes or adjustments reported in the burden or capital/O&M cost estimates.

Courtney Kerwin,

Director, Information Engagement Division.

[FR Doc. 2024-11805 Filed 5-29-24; 8:45 am]

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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 received are subject to public disclosure. In general, comments received will be made available without change and will not be modified to remove personal or business information including confidential, contact, or other identifying information. Comments should not include any information such as confidential information that would not be appropriate for public disclosure.

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 July 1, 2024.

A. Federal Reserve Bank of Kansas City (Jeffrey Imgarten, Assistant Vice President) 1 Memorial Drive, Kansas City, Missouri, 64198-0001. Comments can also be sent electronically to KCApplicationComments@kc.frb.org:

1. *West 4 Bancshares, Inc., Healy, Kansas*; to become a bank holding company by acquiring First State Bank, Healy, Kansas.

Board of Governors of the Federal Reserve System.

Erin Cayce,

Assistant Secretary of the Board.

[FR Doc. 2024-11903 Filed 5-29-24; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Supplemental Evidence and Data Request on Peripheral Nerve Blocks for Postoperative Pain Management in Cardiothoracic Surgery

AGENCY: Agency for Healthcare Research and Quality (AHRQ), HHS.

ACTION: Request for supplemental evidence and data submission.

SUMMARY: The Agency for Healthcare Research and Quality (AHRQ) is seeking scientific information submissions from the public. Scientific information is being solicited to inform our review on *Peripheral Nerve Blocks for Postoperative Pain Management in Cardiothoracic Surgery*, which is currently being conducted by the AHRQ's Evidence-based Practice Centers (EPC) Program. Access to published and unpublished pertinent

scientific information will improve the quality of this review.

DATES: *Submission Deadline* on or before July 1, 2024.

ADDRESSES:

Email submissions: epc@ahrq.hhs.gov.

Print submissions:

Mailing Address: Center for Evidence and Practice Improvement, Agency for Healthcare Research and Quality, ATTN: EPC SEADs Coordinator, 5600 Fishers Lane, Mail Stop 06E53A, Rockville, MD 20857.

Shipping Address (FedEx, UPS, etc.): Center for Evidence and Practice Improvement, Agency for Healthcare Research and Quality, ATTN: EPC SEADs Coordinator, 5600 Fishers Lane, Mail Stop 06E77D, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Kelly Carper, Telephone: 301-427-1656 or Email: epc@ahrq.hhs.gov.

SUPPLEMENTARY INFORMATION: The Agency for Healthcare Research and Quality has commissioned the Evidence-based Practice Centers (EPC) Program to complete a review of the evidence for *Peripheral Nerve Blocks for Postoperative Pain Management in Cardiothoracic Surgery*. AHRQ is conducting this review pursuant to Section 902 of the Public Health Service Act, 42 U.S.C. 299a.

The EPC Program is dedicated to identifying as many studies as possible that are relevant to the questions for each of its reviews. In order to do so, we are supplementing the usual manual and electronic database searches of the literature by requesting information from the public (*e.g.*, details of studies conducted). We are looking for studies that report on *Peripheral Nerve Blocks for Postoperative Pain Management in Cardiothoracic Surgery*.

The entire research protocol is available online at: <https://effectivehealthcare.ahrq.gov/products/peripheral-nerve-blocks/protocol>.

This is to notify the public that the EPC Program would find the following information on *Peripheral Nerve Blocks for Postoperative Pain Management in Cardiothoracic Surgery* helpful:

- A list of completed studies that your organization has sponsored for this topic. In the list, please *indicate whether results are available on ClinicalTrials.gov along with the ClinicalTrials.gov trial number.*

- *For completed studies that do not have results on ClinicalTrials.gov, a summary, including the following elements, if relevant: study number, study period, design, methodology,*