intervene in this proceeding shall file a petition for leave to intervene in accordance with Rule 68 of the Commission's Rules of Practice and Procedure, 46 CFR 502.68. Such petition shall be accompanied by the petitioner's memorandum of law, affidavit of fact, and documentary evidence, if any, and shall be filed no later than the date fixed below;

It is further ordered That Great Northern & Southern Navigation Co., LLC DBA French America Line be named as Respondent in this proceeding. Affidavits of fact, memoranda of law, and documentary evidence shall be filed by Respondent and any intervenors in support of Respondent no later than November 26, 2019;

It is further ordered That the Commission's Bureau of Enforcement (BOE) be made a party to this proceeding;

It is further ordered That reply affidavits, memoranda of law, and documentary evidence shall be filed by BOE and intervenors in opposition to Respondent no later than December 11, 2019;

It is further ordered That:

(a) Should any party believe that the submission of testimony or additional evidence is required, that party must submit a request together with a statement setting forth in detail the facts to be proved, the relevance of those facts to the issues in this proceeding, a description of the evidence which would be adduced, and why such testimony or other evidence cannot be submitted by affidavit; and

(b) Any request for submission of testimony or additional evidence shall be filed no later than December 11, 2019;

It is further ordered That notice of this Order to Show Cause be published in the **Federal Register**, and that a copy thereof be served upon Respondent at its last known address;

It is further ordered That all documents submitted by any party of record in this proceeding shall be filed in accordance with Rule 2 of the Commission's Rules of Practice and Procedure, 46 CFR 502.2, as well as mailed directly to all parties of record;

Finally, it is ordered That pursuant to the terms of Rule 61 of the Commission's Rules of Practice and Procedure, 46 CFR502.61, the final decision of the Commission in this proceeding shall be issued no later than February 27, 2020.

By the Commission.

Rachel Dickon,

Secretary.

[FR Doc. 2019–24177 Filed 11–5–19; 8:45 am]

BILLING CODE 6731–AA–P

FEDERAL MINE SAFETY AND HEALTH REVIEW COMMISSION

Sunshine Act Notice

TIME AND DATE: 10:00 a.m., Thursday, December 12, 2019.

PLACE: The Richard V. Backley Hearing Room, Room 511N, 1331 Pennsylvania Avenue NW, Washington, DC 20004 (enter from F Street entrance).

STATUS: Open.

MATTERS TO BE CONSIDERED: The Commission will consider and act upon the following in open session: *Secretary of Labor* v. *The Monongalia County Coal Company,* Docket Nos. WEVA 2015–509 et al. (Issues include whether the Judge erred in denying the Secretary of Labor the opportunity to present evidence of prior violations to support his allegation that repeated flagrant violations had occurred.).

Any person attending this meeting who requires special accessibility features and/or auxiliary aids, such as sign language interpreters, must inform the Commission in advance of those needs. Subject to 29 CFR 2706.150(a)(3) and 2706.160(d).

CONTACT PERSON FOR MORE INFO: Emogene Johnson (202) 434–9935/(202) 708–9300 for TDD Relay/1–800–877–8339 for toll free.

PHONE NUMBER FOR LISTENING TO MEETING: 1–(866) 236–7472
Passcode: 678–100.

Authority: 5 U.S.C. 552b.

Dated: November 4, 2019.

Sarah L. Stewart,

Deputy General Counsel.

[FR Doc. 2019–24344 Filed 11–4–19; 4:15 pm]

BILLING CODE 6735–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) approve the proposed information collection project ''Outcome Measure Harmonization and Data Infrastructure for Patient Centered Outcomes Research in Depression.''

This proposed information collection was previously published in the **Federal Register** on August 22, 2019 and allowed 60 days for public comment. No substantive comments were received by AHRQ during these 60 days. The purpose of this notice is to allow an additional 30 days for public comment.

DATES: Comments on this notice must be received by 30 days after date of publication.

ADDRESSES: Written comments should be submitted to: AHRQ's OMB Desk Officer by fax at (202) 395–6974 (attention: AHRQ's desk officer) or by email at *OIRA_submission@omb.eop.gov* (attention: AHRQ's desk officer).

FOR FURTHER INFORMATION CONTACT: Doris Lefkowitz, AHRQ Reports Clearance Officer, (301) 427–1477, or by email at *doris.lefkowitz@AHRQ.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Proposed Project

Outcome Measure Harmonization and Data Infrastructure for Patient Centered Outcomes Research in Depression

The Agency for Healthcare Research and Quality's (AHRQ) mission is to produce evidence to make health care safer, higher quality, more accessible, equitable, and affordable, and to work within the U.S. Department of Health and Human Services and with other partners to make sure that the evidence is understood and used.

In support of this mission, AHRQ funded a prior project to harmonize the outcome measures collected across patient registries and routine clinical practice, with the goals of supporting the development of a robust data infrastructure that can consistently and efficiently collect high-quality data on outcome measures that are relevant to patients and clinicians and supporting patient-centered outcomes research and quality improvement. Harmonized outcome measures would also form the foundation for learning healthcare systems. Of note, AHRQ has supported the development of the Outcome Measures Framework (OMF). The OMF is a conceptual model for classifying outcomes that are relevant to patients and providers across most conditions. AHRQ, in collaboration with the U.S. Food and Drug Administration and the National Library of Medicine, recently