with permission of the institution at which the student is enrolled; (2) is to be uncompensated; and (3) will not displace any employee. Form A–384 establishes the responsibility of students, their institutions, and the FCC as a precondition to accepting individuals as unpaid volunteers.

One such precondition now included on Form A–384, for which the FCC previously received Emergency approval, is the requirement that students comply with regulations and policies pertaining to COVID–19 vaccination requirements for Federal workers. On September 9, 2021, the President issued Executive Order (E.O.) 14043, ''Requiring Coronavirus Disease 2019 Vaccination for Federal Employees,'' requiring all Federal employees, as defined by 5 U.S.C. 2105, to be vaccinated against COVID–19, with exceptions only as required by law. Although the vaccination requirement is currently the subject of a nationwide injunction, the FCC will continue to develop and implement health and safety protocols to ensure and maintain the safety of all occupants during standard operations and public health emergencies or similar health and safety incidents, such as a pandemic. As relevant here, this includes requiring unpaid employees to report their vaccination status.

Federal Communications Commission.

Marlene Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2022–05155 Filed 3–10–22; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL MARITIME COMMISSION

Sunshine Act Meetings

TIME AND DATE: March 16, 2022; 10:00 a.m.

PLACE: This meeting will be held by video-conference only.

STATUS: Part of the meeting will be open to the public and available to view, streamed live, accessible from *www.fmc.gov.* The rest of the meeting will be closed to the public.

MATTERS TO BE CONSIDERED:

Portions Open to the Public:

1. Commissioner Sola, Update on Fact Finding 30: COVID–19 Impact on Cruise Industry
2. Staff Briefing on Ongoing Enforcement Activities

Portions Closed to the Public:

1. Staff Briefing on Ongoing Enforcement Activities
2. Area Representative Regional Activity Updates

CONTACT PERSON FOR MORE INFORMATION: William Cody, Secretary, (202) 523–5725.

William Cody,

Secretary.

[FR Doc. 2022–05261 Filed 3–9–22; 11:15 am]

BILLING CODE 6730–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Supplemental Evidence and Data Request on Postpartum Care for Women Up to One Year After Pregnancy

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

ACTION: Request for supplemental evidence and data submissions.

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 *Postpartum Care for Women Up to One Year After Pregnancy,* 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 April 11, 2022.

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: Jenae Benns, Telephone: 301–427–1496 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 *Postpartum Care for Women Up to One Year After Pregnancy.* AHRQ is conducting this technical brief 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 *Postpartum Care for Women Up to One Year After Pregnancy,* including those that describe adverse events. The entire research protocol is available online at: *https://effectivehealthcare.ahrq.gov/products/postpartum-care-one-year/protocol.*

This is to notify the public that the EPC Program would find the following information on *Postpartum Care for Women Up to One Year After Pregnancy* helpful:

- A list of completed studies that your organization has sponsored for this indication. 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: Study number, study period, design, methodology, indication and diagnosis, proper use instructions, inclusion and exclusion criteria, primary and secondary outcomes, baseline characteristics, number of patients screened/eligible/enrolled/lost to follow-up/withdrawn/analyzed, effectiveness/efficacy, and safety results.
- *A list of ongoing studies that your organization has sponsored for this indication.* In the list, please provide the *ClinicalTrials.gov* trial number or, if the trial is not registered, the protocol for the study including a study number, the study period, design, methodology, indication and diagnosis, proper use instructions, inclusion and exclusion criteria, and primary and secondary outcomes.
- Description of whether the above studies constitute *ALL Phase II and above clinical trials* sponsored by your organization for this indication and an index outlining the relevant information in each submitted file.

Your contribution is very beneficial to the Program. Materials submitted must be publicly available or able to be made public. Materials that are considered confidential; marketing materials; study types not included in the review; or information on indications not included in the review cannot be used by the EPC Program. This is a voluntary request for information, and all costs for complying