

people of the United States, without discrimination on the basis of race, color, religion, national origin, or sex.” The term “equity” is used here consistent with Executive Order 13985 as the consistent and systematic fair, just, and impartial treatment of all individuals, including individuals who belong to underserved communities that have been denied such treatment, such as Black, Latino, and Indigenous and Native American persons, Asian Americans and Pacific Islanders and other persons of color; members of religious minorities; lesbian, gay, bisexual, transgender, and queer (LGBTQ+) persons; persons with disabilities; persons who live in rural areas; and persons otherwise adversely affected by persistent poverty or inequality. Specifically, the Commission seeks comment on how its proposals may promote or inhibit advances in diversity, equity, inclusion, and accessibility, as well the scope of the Commission’s relevant legal authority.

20. Authority for this Notice of Inquiry may be found in sections 1, 4(i)–(j), 4(n), 7, and 403 of the Communications Act of 1934, as amended, 47 U.S.C. 151, 154(i)–(j), 154(n), 157 and Section 1.430 of the Commission’s rules, 47 CFR 1.430.

Federal Communications Commission.

**Marlene H. Dortch,**

*Secretary, Office of the Secretary.*

[FR Doc. 2022–05121 Filed 3–10–22; 8:45 am]

**BILLING CODE 6712–01–P**

## FEDERAL COMMUNICATIONS COMMISSION

[FR ID 76259]

### SES Performance Review Board

**AGENCY:** Federal Communications Commission.

**ACTION:** Notice.

**SUMMARY:** As required by the Civil Service Reform Act of 1978, Chairwoman Jessica Rosenworcel has appointed the following executives to the Senior Executive Service (SES) Performance Review Board (PRB):

Trent Harkrader

Debra Jordan

Holly Saurer

Federal Communications Commission.

**Katura Jackson,**

*Federal Register Liaison Officer.*

[FR Doc. 2022–05229 Filed 3–10–22; 8:45 am]

**BILLING CODE 6712–01–P**

## FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–1298; FR ID 75434]

### 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 May 10, 2022. 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](mailto:PRA@fcc.gov) and to [nicole.ongele@fcc.gov](mailto: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 No.:* 3060–1298.

*Title:* Volunteer Service Agreement Form, FCC Form A–384.

*Form No.:* FCC Form A–384.

*Type of Review:* Extension of a currently approved collection.

*Respondents:* Individuals or households.

*Number of Respondents and Responses:* 140 respondents and 140 responses.

*Estimated Time Per Response:* 0.25 hours.

*Frequency of Response:* One-time reporting requirement.

*Obligation to Respond:* Mandatory. The statutory authority to collect this information derives from 5 U.S.C. 3111, Acceptance of volunteer service. Certification of compliance with COVID–19 vaccine requirements for Federal workers derives from several sources, including most recently Executive Order 13991, Protecting the Federal Workforce and Requiring Mask-Wearing; Executive Order 14043, Requiring Coronavirus Disease 2019 Vaccination for Federal Employees; and OMB Memorandum M 21–15, COVID–19 Safe Federal Workplace: Agency Model Safety Principles (Jan. 24, 2021), as amended.

*Total Annual Burden:* 35 hours.

*Total Annual Cost:* No Cost.

*Privacy Act Impact Assessment:* Yes. Records of current and former Federal employees as defined in 5 U.S.C. 2105, including volunteers, grantees, and contract employees on whom the agency maintains records, are covered by OPM’s governmentwide System of Records Notice (SORN) OPM/GOVT–1 General Personnel Records, posted at <https://www.opm.gov/information-management/privacy-policy/sorn/opm-sorn-govt-1-general-personnel-records.pdf>. The Privacy Impact Assessment (PIA) for the Electronic Official Personnel Folder is posted at <https://www.opm.gov/information-management/privacy-policy/privacy-policy/eopf-pia.pdf>.

*Nature and Extent of Confidentiality:* As Privacy Act-protected records, these records are kept confidential and will not be disclosed except under applicable Privacy Act exceptions, including the routine uses identified in the OPM/GOVT–1 SORN.

*Needs and Uses:* The Civil Service Reform Act of 1978 authorized Federal agencies to establish programs designed to provide educationally related work assignments for students in a non-pay status. The Act provides that heads of agencies may accept, subject to regulations issued by the Office of Personnel Management, volunteer service for the United States if the service (1) is performed by a student,

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](http://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](mailto: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 Bennis, Telephone: 301-427-1496 or Email: [epc@ahrq.hhs.gov](mailto: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