requirements; and Third party disclosure requirement.

Obligation to Respond: Required to obtain or retain benefits. Statutory authority for this information collection is contained in 47 U.S.C. Sections 4, 302, 303, 307 and 336 of the Communications Act of 1934, as amended.

Total Annual Burden: 3,474 hours.

Total Annual Cost: $52,150.

Privacy Act Impact Assessment: This information collection affects individuals or households. The Commission has a System of Records, FCC/OET–1 ''Experimental Radio Station License Files'' which covers the personally identifiable information (PII) that individual applicants may include in their submissions for experimental radio authorizations. The system of records notice (SORN) was published in the **Federal Register** on April 5, 2006, *see* 71 FR 17234, 17241. The SORN may be viewed at *https://www.fcc.gov/general/privacy-act-information.*

Nature and Extent of Confidentiality: Applicants may request that any information supplied be withheld from public inspection, *e.g.,* granted confidentiality, pursuant to 47 CFR Section 0.459 of the Commission's rules.

Needs and Uses: The Commission will submit this revised information collection to the Office of Management and Budget (OMB) after this 60-day comment period to obtain the three-year clearance.

On June 29, 2016, the Commission adopted a Second Report and Order, in ET Docket No. 10–236 and 06–155; FCC 16–86, which updates Part 5 of the CFR—''Experimental Radio Service'' (ERS).[1] The Commission's recent Report and Order revises and streamlines the rule part under for the ERS. This rule change allows licensees operation under frequency bands mentioned in Section 5.303 and as state, within rule part 15.205(a). These rule changes update procedures used to obtain and use an experimental license.

§ 5.303 Frequencies.

(a) Licensees may operate in any frequency band, including those above 38.6 GHz, except for frequency bands exclusively allocated to the passive services (including the radio astronomy service). In addition, licensees may not use any frequency or frequency band below 38.6 GHz that is listed in § 15.205(a) of this chapter.

(b) *Exception:* Licensees may use frequencies listed in § 15.205(a) of this chapter for testing medical devices (as defined in § 5.402(b) of this chapter), if the device is designed to comply with all applicable service rules in Part 18, Industrial, Scientific, and Medical Equipment; Part 95, Personal Radio Services Subpart H—Wireless Medical Telemetry Service; or Part 95, Subpart I—Medical Device Radiocommunication Service.

Federal Communications Commission.

Marlene Dortch,

Secretary.

[FR Doc. 2018–23853 Filed 10–31–18; 8:45 am]

BILLING CODE 6712–01–P

[1] *See* In the Matter of Promoting Expanded Opportunities for Radio Experimentation and Market Trials Under Part 5 of the Commission's Rules and Streamlining Other Related Rules, ET Docket No. 10–236; 2006 Biennial Review of Telecommunications Regulations—Part 2, Administered by the Office of Engineering and Technology (OET), ET Docket No. 06–155; 31 FCC Rcd 7529 (2016), FCC 16–86.

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–0669]

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 Commission) invites the general public and other Federal agencies to take this opportunity to comment on the following information collections. 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 OMB 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 OMB control number.

DATES: Written PRA comments should be submitted on or before December 31, 2018. 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 Cathy Williams, FCC, via email *PRA@fcc.gov* and to *Cathy.Williams@fcc.gov.*

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Cathy Williams at (202) 418–2918.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060–0669.

Title: Section 76.946, Advertising of Rates.

Form Number: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Business and other for-profit entities.

Number of Respondents and Responses: 8,250 respondents; 8,250 responses.

Estimated Time per Response: 30 minutes (0.5 hours).

Frequency of Response: On occasion reporting requirement; Third party disclosure requirement.

Total Annual Burden to Respondents: 4,125 hours.

Total Annual Costs: None.

Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this collection is contained in Section 4(i) of the Communications Act of 1934, as amended.

Nature and Extent of Confidentiality: There is no need for confidentiality with this collection of information.

Privacy Impact Assessment: No impact(s).

Needs and Uses: The information collection requirements contained in 47 CFR 76.946 states that cable operators that advertise rates for basic service and cable programming service tiers shall be required to advertise rates that include all costs and fees. Cable systems that cover multiple franchise areas having differing franchise fees or other franchise costs, different channel line-ups, or different rate structures may advertise a complete range of fees without specific identification of the rate for each individual area. In such circumstances, the operator may advertise a ''fee plus'' rate that indicates the core rate plus the range of possible additions, depending on the particular location of the subscriber.

Federal Communications Commission.

Marlene Dortch,

Secretary.

[FR Doc. 2018–23845 Filed 10–31–18; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL LABOR RELATIONS AUTHORITY

Senior Executive Service Performance Review Board

AGENCY: Federal Labor Relations Authority.

ACTION: Notice.

SUMMARY: The Federal Labor Relations Authority (FLRA) publishes the names of the persons selected to serve on its SES Performance Review Board (PRB). This notice supersedes all previous notices of the PRB membership.

DATES: Upon publication.

ADDRESSES: Written comments about this final rule can be mailed to the Case Intake and Publication Office, Federal Labor Relations Authority, 1400 K Street NW, Washington, DC 20424. All written comments will be available for public inspection during normal business hours at the Case Intake and Publication Office.

FOR FURTHER INFORMATION CONTACT: William Tosick, Executive Director, Federal Labor Relations Authority, 1400 K St. NW, Washington, DC 20424, (202) 218–7791, *wtosick@flra.gov.*

SUPPLEMENTARY INFORMATION: Section 4314(c) of Title 5, U.S.C. requires each agency to establish, in accordance with regulations prescribed by the Office of Personnel Management, one or more PRBs. The PRB shall review and evaluate the initial appraisal of a senior executive's performance by the supervisor, along with any response by the senior executive, and make recommendations to the final rating authority relative to the performance of the senior executive.

The persons named below have been selected to serve on the FLRA's PRB.

PRB Chairman:

William Tosick, Executive Director

PRB Members:

Kimberly D. Moseley, Executive Director, Federal Service Impasses Panel; Douglas Fitzgerald, Director, Division of Longshore and Harbor Workers' Compensation at U.S. Department of Labor; Richard Jones, Atlanta Regional Director; and Paula Chandler, Director, Human Resources Division, FLRA, as an ex officio member.

Dated: October 29, 2018.

William Tosick,

Executive Director.

[FR Doc. 2018–23898 Filed 10–31–18; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–19–0969]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled ''Monitoring Changes in Attitudes and Practices among Family Planning Providers and Clinics'' to the Office of Management and Budget (OMB) for review and approval. CDC previously published a ''Proposed Data Collection Submitted for Public Comment and Recommendations'' notice on June 8, 2018 to obtain comments from the public and affected agencies. CDC received one substantive and five non-substantive comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) Minimize the burden of the collection of information on those who are to respond, including, through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to *omb@cdc.gov.* Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395–5806. Provide written comments within 30 days of notice publication.

Proposed Project

Monitoring Changes in Attitudes and Practices among Family Planning Providers and Clinics (OMB Number 0920–0969, Expiration Date: 05/31/2014)—Reinstatement with Change—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Division of Reproductive Health (DRH) at the Centers for Disease Control and Prevention (CDC) and the HHS Office of Population Affairs (OPA) develop and disseminate guidance to improve the use of contraception and the delivery of quality family planning services. The *U.S. Medical Eligibility Criteria for Contraceptive Use* (US MEC), the first national guidance on family planning containing evidence-based recommendations for the safe use of contraceptive methods for women and men with specific characteristics and medical conditions, was first published by the CDC in June 2010. The *US Selected Practice Recommendations for Contraceptive Use* (US SPR), which provides guidance on how to use contraceptive methods safely and effectively once they are deemed to be medically appropriate, was first published by the CDC in June 2013. The US MEC and US SPR were updated after review of the scientific evidence and consultation with national experts in family planning; the revised US MEC and US SPR were published in August 2016.

Providing Quality Family Planning Services (QFP), which provides evidence-informed recommendations to improve client care and service delivery infrastructure to support the provision of quality family planning services to women and men of reproductive age in the United States, was published by CDC and OPA in April 2014. The US MEC, US SPR, and QFP have been widely disseminated to health care providers and other constituents via professional organizations, federal program grantees, scientific and programmatic meetings, scientific