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 process 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 Dorch,
Secretary.

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

BILLING CODE 6712–01–P

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 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 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.

William Tosick,
Executive Director.

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
determined 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