III. Requirements for Submissions

To ensure consideration, interested parties must submit comments and responses to TPSC questions electronically via REGS.GOV by the applicable deadlines in the DATES section above. The docket number is USTR–2021–0014. All submissions must be in English. USTR will not accept hand-delivered submissions.

To submit comments using REGS.GOV, enter docket number USTR–2021–0014 in the ‘search for’ field on the home page and click ‘search.’ The site will provide a search-results page listing all documents associated with this docket. Find a reference to this notice by selecting ‘notice’ under ‘document type’ in the ‘filter results by’ section on the left side of the screen and click on the link entitled ‘comment now.’ REGS.GOV offers the option of providing comments by filling in a ‘type comment’ field or by attaching a document using the ‘upload file[s]’ field. USTR prefers that you provide submissions in an attached document and, in such cases, that you write ‘see attached’ in the ‘type comment’ field on the online submission form. In addition, USTR prefers submissions in Microsoft Word (.doc) or Adobe Acrobat (.pdf). If the submission is in an application other than those two, please indicate the name of the application in the ‘type comment’ field. At the beginning of the submission, include the following text: (1) 2021 Russia WTO Implementation Report; (2) your organization’s name; and (3) whether the document is a comment or an answer to a TPSC question. Written comments should not exceed 30 single-spaced, standard letter-size pages in 12-point type, including attachments. Include any data attachments to the submission in the same file as the submission itself, and not as separate files.

When you complete the submission procedure at REGS.GOV you will receive a tracking number confirming successful transmission. For further information on using REGS.GOV, please consult the resources provided on the website by clicking on ‘How to Use Regulations.gov’ on the bottom of the home page. USTR is not able to provide technical assistance for REGS.GOV.

IV. Business Confidential Submissions

An interested party requesting that USTR treat information contained in a submission as BCI must certify that the information is business confidential and would not customarily be released to the public by the submitter. You must clearly designate BCI by marking the submission ‘BUSINESS CONFIDENTIAL’ at the top and bottom of the cover page and on each succeeding page, and indicating, via brackets, the specific information that is BCI. Additionally, you must include ‘business confidential’ in the ‘type comment’ field and add the designation BCI to the end of the file name for any attachments. For any submission containing BCI, you separately must submit a non-confidential version, i.e., not as part of the same submission with the BCI version, indicating where BCI has been redacted. USTR will post the non-confidential version in the docket for public inspection.

V. Public Viewing of Review Submissions

USTR will post comments in the docket for public inspection, except BCI. You can view comments at REGS.GOV by entering docket number USTR–2021–0014 in the search field on the home page. General information concerning USTR is available at www.ustr.gov.

Edward Gresser,
Chair of the Trade Policy Staff Committee,
Office of the United States Trade Representative.

[FR Doc. 2021–17477 Filed 8–13–21; 8:45 am]
BILLING CODE 3290–F1–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration
[Docket No. FAA–2021–0180]

Agency Information Collection Activities: Requests for Comments; Clearance of Renewed Approval of Information Collection: Report of Inspections Required by Airworthiness Directive, Part 39

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Notice and request for comments.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995, FAA invites public comments about our intention to request the Office of Management and Budget (OMB) approval to renew an information collection. The Federal Register Notice with a 60-day comment period soliciting comments on the following collection of information was published on March 31, 2021. The collection involves the member of the public that may submit an Alternative Methods of Compliance (AMOC) request to the FAA by using the ADD External website. The information to be collected will be used to support publicly disseminated information to the FAA and/or is necessary because this information supports the Department of Transportation’s strategic goal to promote the public health and safety by working toward eliminating transportation-related deaths and injuries.

DATES: Written comments should be submitted by September 15, 2021.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAmain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function.

FOR FURTHER INFORMATION CONTACT: Robert Romero by email at: Robert.A.Romero@faa.gov; phone: 817–222–5102.

SUPPLEMENTARY INFORMATION:
Public Comments Invited: You are asked to comment on any aspect of this information collection, including (a) Whether the proposed collection of information is necessary for FAA’s performance; (b) the accuracy of the estimated burden; (c) ways for FAA to enhance the quality, utility and clarity of the information collection; and (d) ways that the burden could be minimized without reducing the quality of the collected information.

OMB Control Number: 2120–0056.
Title: Report of Inspections Required by Airworthiness Directives, Part 39.
Form Numbers: There is no standard form to use for AMOC submission. However, the public may access the AMOC External website to submit an AMOC request to the FAA.

Type of Review: Renewal of an information collection.

Background: The Federal Register Notice with a 60-day comment period soliciting comments on the following collection of information was published on March 31, 2021 (FRN citation 2021–06646). Alternative Methods of Compliance (AMOC) are submitted to the FAA by the general public. While anyone may submit an AMOC there is no standard form to use. From Order 8110.103B Alternative Methods of Compliance (AMOC), Section 3–2: 3–2. AMOC Proposal. 14 CFR 39.19 states in part that “anyone may propose to FAA an alternative method of compliance or a change in the compliance time, if the proposal provides an acceptable level of safety.” a. Although a lottery is preferred, AMOC proposals may be submitted by...
other means, such as email, fax, or telephone. AMOC proposals received by telephone must be documented.

An AMOC Response Letter is written by an internal FAA user and sent to the AMOC Requester. The template may be generated from the ADD Dashboard and follows the latest Order. There is not an FAA or OMB number on this template.

A member of the public may submit an AMOC request to the FAA by using the AMOC external website. Registration is not needed to use this website. External users must consent to the “Terms of Use” statement before proceeding to the AMOC proposal webpage. An AMOC is required if an owner/operator of aircraft cannot comply with an AD or finds a different method to comply with the actions specified in an AD, as mandated by FAA Order 8110.103B.

Respondents: The respondents are a member of the public who may submit an AMOC request to FAA by using the AMOC External website. We estimate that 25 ADs yearly will require reports of information and findings. The average AD affects about 1,120 owners/operators. Therefore, 25 ADs times 1,120 owners/operators per year equal 28,000 reports.

Frequency: As needed.

Estimated Average Burden per Response: These reports, requiring an average of 1 hour each to prepare, consume 28,000 reporting hours.

Estimated Total Annual Burden: The total annualized cost to respondents is $2,380,000. We base this on the 28,000 reporting hours times an estimated hourly rate of $85/hour per respondent. The average cost to the respondents per AD per year is $85.00 ($2,380,000 divided by 28,000).

Issued in Washington, DC, on July 9, 2021.

Patrick Idlett,
ASKME Program Manager, Office of Enterprise Program Management (AEM), Project Portfolio Performance Division.

[FR Doc. 2021–17417 Filed 8–13–21; 8:45 am]
BILLING CODE 4910–13–P

DEPARTMENT OF TRANSPORTATION

Federal Motor Carrier Safety Administration


Qualification of Drivers; Exemption Applications; Epilepsy and Seizure Disorders

AGENCY: Federal Motor Carrier Safety Administration (FMCSA), Department of Transportation (DOT).

ACTION: Notice of renewal of exemptions; request for comments.

SUMMARY: FMCSA announces its decision to renew exemptions for two individuals from the requirement in the Federal Motor Carrier Safety Regulations (FMCSRs) that interstate commercial motor vehicle (CMV) drivers have “no established medical history or clinical diagnosis of epilepsy or any other condition which is likely to cause loss of consciousness or any loss of ability to control a CMV.” The exemptions enable these individuals who have had one or more seizures and are taking anti-seizure medication to continue to operate CMVs in interstate commerce.

DATES: The exemptions are applicable on August 13, 2021. The exemptions expire on August 13, 2023. Comments must be received on or before September 15, 2021.

ADDRESSES: You may submit comments identified by the Federal Docket Management System (FDMS) Docket No. FMCSA–2015–0116 or Docket No. FMCSA–2019–0027 using any of the following methods:

• Federal eRulemaking Portal: Go to www.regulations.gov, insert the docket number, FMCSA–2015–0116 or FMCSA–2019–0027 in the keyword box, and click “Search.” Next, sort the results by “Posted (Newer-Older),” choose the first notice listed, click the “Comment” button, and type your comment into the text box on the following screen. Choose whether you are submitting your comment as an individual or on behalf of a third party and then submit.

If you submit your comments by mail or hand delivery, submit them in an unbound format, no larger than 8½ by 11 inches, suitable for copying and electronic filing. If you submit comments by mail and would like to know that they reached the facility, please enclose a stamped, self-addressed postcard or envelope.

FMCSA will consider all comments and material received during the comment period.

B. Viewing Comments

To view comments go to www.regulations.gov. Insert the docket number, FMCSA–2015–0116 or FMCSA–2019–0027 in the keyword box, and click “Search.” Next, sort the results by “Posted (Newer-Older),” choose the first notice listed, and click “Browse Comments.” If you do not have access to the internet, you may view the docket online by visiting Dockets Operations in Room W12–140 on the ground floor of the DOT West Building, 1200 New Jersey Avenue SE, Washington, DC 20590–0001, between 9 a.m. and 5 p.m., ET, Monday through Friday, except Federal holidays.

To avoid duplication, please use only one of these four methods. See the “Public Participation” portion of the SUPPLEMENTARY INFORMATION section for instructions on submitting comments.

FOR FURTHER INFORMATION CONTACT: Ms. Christine A. Hydock, Chief, Medical Programs Division, (202) 366–4001, fmcsamedical@dot.gov, FMCSA, DOT, 1200 New Jersey Avenue SE, Room W64–224, Washington, DC 20590–0001.

Office hours are from 8:30 a.m. to 5 p.m., ET, Monday through Friday, except Federal holidays. If you have questions regarding viewing or submitting material to the docket, contact Dockets Operations, (202) 366–9826.

SUPPLEMENTARY INFORMATION:

I. Public Participation

A. Submitting Comments

If you submit a comment, please include the docket number for this notice (Docket No. FMCSA–2015–0116 or Docket No. FMCSA–2019–0027), indicate the specific section of this document to which each comment applies, and provide a reason for each suggestion or recommendation. You may submit your comments and material online or by fax, mail, or hand delivery, but please use only one of these means. FMCSA recommends that you include your name and a mailing address, an email address, or a phone number in the body of your document so that FMCSA can contact you if there are questions regarding your submission.

To submit your comment online, go to www.regulations.gov/, insert the docket number, FMCSA–2015–0116 or FMCSA–2019–0027 in the keyword box, and click “Search.” Next, sort the results by “Posted (Newer-Older),” choose the first notice listed, click the “Comment” button, and type your comment into the text box on the following screen. Choose whether you are submitting your comment as an individual or on behalf of a third party and then submit.

If you submit your comments by mail or hand delivery, submit them in an unbound format, no larger than 8½ by 11 inches, suitable for copying and electronic filing. If you submit comments by mail and would like to know that they reached the facility, please enclose a stamped, self-addressed postcard or envelope.

FMCSA will consider all comments and material received during the comment period.