

quality of service offered to the public. If this information is not collected, vital feedback from customers and stakeholders on the Agency's services will be unavailable.

The agency will only submit a collection for approval under this generic clearance if it meets the following conditions:

- The collections are voluntary;
- The collections are low-burden for respondents (based on considerations of total burden hours, total number of respondents, or burden-hours per respondent) and are low-cost for both the respondents and the Federal Government;
- The collections are non-controversial and do not raise issues of concern to other Federal agencies;
- Any collection is targeted to the solicitation of opinions from respondents who have experience with the program or may have experience with the program in the near future;
- Personally identifiable information (PII) is collected only to the extent necessary and is not retained;
- Information gathered will be used only internally for general service improvement and program management purposes and is not intended for release outside of the agency;
- Information gathered will not be used for the purpose of substantially informing influential policy decisions; and
- Information gathered will yield qualitative information; the collections will not be designed or expected to yield statistically reliable results or used as though the results are generalizable to the population of study.

Feedback collected under this generic clearance provides useful information, but it does not yield data that can be generalized to the overall population. This type of generic clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance. Such data uses require more rigorous designs that address: The target population to which generalizations will be made, the sampling frame, the sample design (including stratification and clustering), the precision requirements or power calculations that justify the proposed sample size, the expected response rate, methods for assessing potential non-response bias, the protocols for data collection, and any testing procedures that were or will be undertaken prior to fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be

eligible for submission for other generic mechanisms that are designed to yield quantitative results.

As a general matter, information collections will not result in any new system of records containing privacy information and will not ask questions of a sensitive nature, such as sexual behavior and attitudes, religious beliefs, and other matters that are commonly considered private.

Title: Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery.

OMB Control Number: 2126–0049.

Type of Request: Renewal of a currently-approved information collection.

Respondents: State and local agencies, the general public and stakeholders, original equipment manufacturers and suppliers to the commercial motor vehicle (CMV) industry, CMV fleet owners, CMV owner-operators, state CMV safety agencies, research organizations and contractors, news organizations, safety advocacy groups, and other Federal agencies.

Estimated Number of Respondents: 9,270.

Estimated Time per Response: Range from 5 to 30 minutes.

Expiration Date: August 31, 2021.

Frequency of Response: Generally, on an annual basis.

Estimated Total Annual Burden: 2,233 hours.

Public Comments Invited: You are asked to comment on any aspect of this information collection, including: (1) Whether the proposed collection is necessary for the performance of FMCSA's functions; (2) the accuracy of the estimated burden; (3) ways for FMCSA to enhance the quality, usefulness, and clarity of the collected information; and (4) ways that the burden could be minimized without reducing the quality of the collected information. The agency will summarize or include your comments in the request for OMB's clearance of this information collection.

Issued under the authority of 49 CFR 1.87.

Thomas P. Keane,

Associate Administrator, Office of Research and Registration.

[FR Doc. 2021–02850 Filed 2–11–21; 8:45 am]

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DEPARTMENT OF TRANSPORTATION

Federal Motor Carrier Safety Administration

[Docket No. FMCSA–2020–0226]

Agency Information Collection Activities; Renewal of a Currently-Approved Information Collection Request: Application for Certificate of Registration for Foreign Motor Carriers and Foreign Motor Private Carriers

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

ACTION: Notice and request for comments.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995, FMCSA announces its plan to submit the Information Collection Request (ICR) described below to the Office of Management and Budget (OMB) for its review and approval and invites public comment. The purpose of this ICR titled, “Application for Certificate of Registration for Foreign Motor Carriers and Foreign Motor Private Carriers,” requires foreign (Mexico-based) for-hire and private motor carriers to file an application Form OP–2 if they wish to register to transport property only within municipalities in the United States on the U.S.-Mexico international borders or within the commercial zones of such municipalities.

DATES: We must receive your comments on or before April 13, 2021.

ADDRESSES: You may submit comments identified by Federal Docket Management System (FDMS) Docket Number FMCSA–2020–0226 using any of the following methods:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the online instructions for submitting comments.

- *Fax:* 1–202–493–2251.

- *Mail:* Docket Operations; U.S. Department of Transportation, 1200 New Jersey Avenue SE, West Building, Ground Floor, Room W12–140, Washington, DC 20590–0001.

- *Hand Delivery or Courier:* U.S. Department of Transportation, 1200 New Jersey Avenue SE, West Building, Ground Floor, Room W12–140, Washington, DC 20590–0001 between 9 a.m. and 5 p.m. e.t., Monday through Friday, except Federal holidays.

Instructions: All submissions must include the Agency name and docket number. For detailed instructions on submitting comments, see the Public Participation heading below. Note that all comments received will be posted without change to <http://www.regulations.gov>, including any

personal information provided. Please see the Privacy Act heading below.

Docket: For access to the docket to read background documents or comments received, go to <http://www.regulations.gov>, and follow the online instructions for accessing the dockets, or go to the street address listed above.

Privacy Act: In accordance with 5 U.S.C. 553(c), DOT solicits comments from the public to better inform its rulemaking process. DOT posts these comments, without edit, including any personal information the commenter provides, to www.regulations.gov, as described in the system of records notice (DOT/ALL-14 FDMS), which can be reviewed at www.dot.gov/privacy.

Public Participation: The Federal eRulemaking Portal is available 24 hours each day, 365 days each year. You can obtain electronic submission and retrieval help and guidelines under the "help" section of the Federal eRulemaking Portal website. If you want us to notify you that we received your comments, please include a self-addressed, stamped envelope or postcard, or print the acknowledgement page that appears after submitting comments online. Comments received after the comment closing date will be included in the docket and will be considered to the extent practicable.

FOR FURTHER INFORMATION CONTACT: Ms. Dora Tambo-Gonzales, Office of Registration, Licensing and Insurance Division, Department of Transportation, Federal Motor Carrier Safety Administration, West Building 6th Floor, 1200 New Jersey Avenue SE, Washington, DC 20590. Telephone: 202-366-2577; email: dora.tambo.gonzales@dot.gov.

SUPPLEMENTARY INFORMATION:

Background: Title 49 U.S.C. 13902(c) contains basic licensing procedures for registering foreign (Mexico-based) motor carriers to operate across the U.S.-Mexico international border into the United States. 49 CFR pt. 368 contains the regulations that require foreign (Mexico-based) motor carriers to apply to the FMCSA for a Certificate of Registration to provide interstate transportation in municipalities in the United States on the U.S.-Mexico international border or within the commercial zones of such municipalities as defined in 49 U.S.C. 13902(c)(4)(A). The FMCSA carries out this registration program under authority delegated by the Secretary of Transportation.

Foreign (Mexico-based) motor carriers use Form OP-2 to apply for Certificate of Registration authority with the

FMCSA. The form requests information on the foreign motor carrier's name, address, U.S. DOT Number, form of business (e.g., corporation, sole proprietorship, partnership), locations where the applicant plans to operate, types of registration requested (e.g., for-hire motor carrier, household goods carrier, motor private carrier), insurance, safety certifications, household goods arbitration certifications, and compliance certifications.

Title: Application for Certificate of Registration for Foreign Motor Carriers and Foreign Motor Private Carriers.

OMB Control Number: 2126-0019.

Type of Request: Renewal of a currently-approved information collection.

Respondents: Foreign motor carriers.

Estimated Number of Respondents: 31.

Estimated Time per Response: 1.5 hours to complete or update Form OP-2.

Expiration Date: October 31, 2021.

Frequency of Response: Occasionally.

Estimated Total Annual Burden: 47 hours [31 responses × 1½ hours to complete Form OP-2 = 47 hours].

Public Comments Invited: You are asked to comment on any aspect of this information collection, including: (1) Whether the proposed collection is necessary for the performance of FMCSA's functions; (2) the accuracy of the estimated burden; (3) ways for FMCSA to enhance the quality, usefulness, and clarity of the collected information; and (4) ways that the burden could be minimized without reducing the quality of the collected information. The agency will summarize or include your comments in the request for OMB's clearance of this information collection.

Issued under the authority of 49 CFR 1.87.

Thomas P. Keane,

Associate Administrator, Office of Research and Registration.

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DEPARTMENT OF VETERANS AFFAIRS

[OMB Control No. 2900-0556]

Agency Information Collection Activity Under OMB Review: VA Advance Directive: Durable Power of Attorney for Health Care and Living Will

AGENCY: Veterans Health Administration, Department of Veterans Affairs.

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act (PRA) of 1995, this notice announces that the Veterans Health Administration, Department of Veterans Affairs, will submit the collection of information abstracted below to the Office of Management and Budget (OMB) for review and comment. The PRA submission describes the nature of the information collection and its expected cost and burden, and it includes the actual data collection instrument.

DATES: 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. Refer to "OMB Control No. 2900-0556."

FOR FURTHER INFORMATION CONTACT: Maribel Aponte, Office of Enterprise and Integration, Data Governance Analytics (008), 1717 H Street NW, Washington, DC 20006, (202) 266-4688 or email maribel.aponte@va.gov. Please refer to "OMB Control No. 2900-0556" in any correspondence.

SUPPLEMENTARY INFORMATION:

Authority: 44 U.S.C. 3501-21.

Title: VA Advance Directive: Durable Power of Attorney for Health Care and Living Will, VA Form 10-0137.

OMB Control Number: 2900-0556.

Type of Review: Extension of a currently approved collection.

Abstract: Section 7331 of title 38, United States Code (U.S.C.), requires, in relevant part, that the Secretary of Veterans Affairs, upon the recommendation of the Under Secretary for Health, prescribe regulations to ensure, to the maximum extent practicable, that all Department of Veterans Affairs (VA) patient care be carried out only with the full and informed consent of the patient, or in appropriate cases, a representative thereof. Based on VA's interpretation of this statute and our mandate in 38 U.S.C. 7301(b) to provide a complete medical and hospital service, we recognize that patients with decision-making capacity have the right to state their treatment preferences in a VA or other valid advance directive.

VA Form 10-0137, *VA Advance Directive: Durable Power of Attorney for Health Care and Living Will*, is the VA recognized legal document that permits VA patients to designate a health care agent and/or specify preferences for