

with a third response option (“yes/no/unsure”), and to include a field for any clarifying comments.

FMCSA Response

FMCSA considered the NTSB’s comments but does not believe its recommendations would enhance the quality, usefulness, and clarity of the form based on the purpose for which the form was intended to be used. The “391.41 CMV Driver Medication Form, MCSA–5895,” was developed and intended to be used as a tool to supplement the information obtained from the “Medical Examination Report Form, MCSA–5875,” from the driver during the ME’s review of the driver’s health history, and from the physical examination conducted by the ME. The “Medical Examination Report Form, MCSA–5875,” already provides the ME with a complete health history for the driver including all current medications (prescriptions, non-prescriptions, supplements) and medical conditions as reported by the driver.

The “391.41 CMV Driver Medication Form, MCSA–5895,” specifically addresses medication(s) that may impair the driver’s ability to safely operate a CMV so that the ME fully understands the reasons the medications have been prescribed and can consider the impact the medication(s) and medical conditions for which the medication(s) has been prescribed may have on the driver. This information combined with the information obtained from the “Medical Examination Report Form, MCSA–5875,” from the driver during the ME’s review of the driver’s health history, and from the physical examination conducted by the ME, is used by the ME when making a physical qualification determination.

The “391.41 CMV Driver Medication Form, MCSA–5895,” contains information regarding the driver’s role and regulation in § 391.41(b)(12) as a reference for healthcare professionals and does not indicate that the healthcare professional must interpret the regulation. The “391.41 CMV Driver Medication Form, MCSA–5895,” clearly states what is and is not expected of the healthcare professional completing the form by requesting the healthcare professional review the regulation provided, complete the form, and return it to the ME. The form explains that the final determination as to whether the individual listed on the form is physically qualified to drive a CMV will be made by the certified ME. Question 4 was specifically intended to obtain the medical opinion of the healthcare professional completing the form regarding the specific medication(s)

they have prescribed to the driver for a particular medical condition(s). It is the responsibility of the ME to use the information provided by the healthcare professional completing the “391.41 CMV Driver Medication Form, MCSA–5895,” as a tool to assist them in making a physical qualification determination.

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.

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 TRANSPORTATION

Federal Motor Carrier Safety Administration

[Docket No. FMCSA–2022–0136]

Agency Information Collection Activities; Renewal of an Information Collection Request: Transportation of Hazardous Materials; Highway Routing

AGENCY: Federal Motor Carrier Safety Administration (FMCSA), Department of Transportation (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 review and approval. FMCSA requests approval to renew an ICR titled, “Transportation of Hazardous Materials, Highway Routing.” The information reported by States and Indian Tribes is necessary to identify designated/restricted routes and restrictions or limitations affecting how motor carriers may transport certain hazardous materials on highways, including dates that such routes were established and information on subsequent changes or new hazardous materials routing designations. FMCSA did not receive any comments in response to the 60-day **Federal Register** Notice published on September 8, 2022.

DATES: Comments on this notice must be received on or before March 6, 2023.

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 information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function.

FOR FURTHER INFORMATION CONTACT: Ms. Melissa Williams, General Engineer, Office of Safety/Hazardous Materials Division, DOT, FMCSA, 6th Floor, West Building, 1200 New Jersey Avenue SE, Washington, DC 20590–0001; 202–366–4163; melissa.williams@dot.gov.

SUPPLEMENTARY INFORMATION:

Title: Transportation of Hazardous Materials, Highway Routing.

OMB Control Number: 2126–0014.

Type of Request: Renewal of a currently approved ICR.

Respondents: The reporting burden is shared by 50 States, the District of Columbia, Indian Tribes with designated routes, and U.S. Territories including Puerto Rico, American Samoa, Guam, the Commonwealth of the Northern Mariana Islands, and the U.S. Virgin Islands.

Estimated Number of Respondents: 57 [36 States + the District of Columbia, with designated hazardous materials highway routes + 19 States/U.S. Territories without designated hazardous materials highway routes + 1 Indian Tribe with a designated route = 57].

Estimated Time per Response: 15 minutes.

Expiration Date: April 30, 2023.

Frequency of Response: Once every 2 years.

Estimated Total Annual Burden: 7 hours [57 annual respondents × 1 response per 2 years × 15 minutes per response/60 minutes per response = 7.125 hours rounded to 7 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.

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 TRANSPORTATION

Federal Railroad Administration

Notice of Funding Opportunity for the Federal-State Partnership for Intercity Passenger Rail Grants Program for Projects Not Located on the Northeast Corridor

AGENCY: Federal Railroad Administration (FRA), Department of Transportation (DOT).

ACTION: Notice of Funding Opportunity (NOFO or notice); Extension and Supplemental Funding Notice.

SUMMARY: FRA is adding funding and extending the application submittal period for its Fiscal Year (FY) 2022 NOFO for the Federal-State Partnership for Intercity Passenger Rail Program (FSP Program) for projects not located on the Northeast Corridor published on December 7, 2022. The basis for the funding increase and extension is the appropriation of additional amounts for the FSP Program from the Consolidated Appropriations Act, 2023, together with the government's goal of efficiently administering such funds.

DATES: The original due date was March 7, 2023. The period for submitting applications to the NOFO published on December 7, 2022, is extended. Applications must now be submitted by 5 p.m. EDT on April 21, 2023.

FOR FURTHER INFORMATION CONTACT: For further information concerning this Notice, please contact the FRA NOFO Support program staff via email at FRA-NOFO-Support@dot.gov.

Amendment

FRA amends its FY 2022 NOFO for the Federal-State Partnership for Intercity Passenger Rail Program (FSP Program) for projects not located on the Northeast Corridor published December 7, 2022 (87 FR 75119) <https://www.govinfo.gov/content/pkg/FR-2022-12-07/pdf/2022-26610.pdf>. by: (1) adding up to an additional \$2,283,150,000 for a total of up to \$4,566,300,000 and (2) extending the period for submitting applications to April 21, 2023.

Section B.1. Federal Award Information/Available Award Amount is deleted in its entirety, and the following is substituted therefore:

B. Federal Award Information

1. Available Award Amount

The total funding available for awards under this NOFO is up to \$4,566,300,000 made available by IJA supplemental appropriations, the Appropriations Act, and the Consolidated Appropriations Act, 2023 (Pub. L. 117-328, December 29, 2022), as follows:

a. Up to \$4,464,000,000 in IJA supplemental appropriations: Title VIII of Division J of IJA provided \$36,000,000,000 in supplemental appropriations for the FSP Program, with not more than \$24,000,000,000 made available for projects for the NEC and thus at least \$12,000,000,000 of such funds made available for FSP-National (\$2,400,000,000 made available per year for fiscal years 2022 through 2026). After the funding set aside for FRA award and project management oversight and the planning and development activities authorized at 49 U.S.C. 24911(k), up to \$4,464,000,000 in funding made available for fiscal year 2022 and fiscal year 2023 is made available for FSP Program awards under this FSP-National NOFO.

b. Up to \$102,300,000 in fiscal year 2022 and fiscal year 2023 annual appropriations: The appropriations acts for fiscal year 2022 and fiscal year 2023 provided \$200,000,000 for the FSP Program. Consistent with 49 U.S.C. 24911(d)(3), a minimum of 45 percent and a maximum of 55 percent of this amount is for FSP-National, of which not less than 20 percent (a minimum of \$16,740,000) shall be for projects that benefit (in whole or in part) a long-distance route. After the funding set aside for FRA award and project management oversight and the planning and development activities authorized at 49 U.S.C. 24911(k), at least \$83,700,000 and up to \$102,300,000 in fiscal year 2022 and fiscal year 2023 annual funding is made available for FSP Program awards under this FSP-National NOFO.

Should additional FSP-National funds become available after the release of this NOFO, FRA may elect to award such additional funds to applications received under this NOFO. Any selection and award under this NOFO is subject to the availability of appropriated funds.

Issued in Washington, DC.

Amitabha Bose,
Administrator.

[FR Doc. 2023-02346 Filed 2-2-23; 8:45 am]

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DEPARTMENT OF TRANSPORTATION

Maritime Administration

[Docket No. MARAD-2023-0013]

Coastwise Endorsement Eligibility Determination for a Foreign-Built Vessel: WANDERLUST (Motor); Invitation for Public Comments

AGENCY: Maritime Administration, DOT.

ACTION: Notice.

SUMMARY: The Secretary of Transportation, as represented by the Maritime Administration (MARAD), is authorized to issue coastwise endorsement eligibility determinations for foreign-built vessels which will carry no more than twelve passengers for hire. A request for such a determination has been received by MARAD. By this notice, MARAD seeks comments from interested parties as to any effect this action may have on U.S. vessel builders or businesses in the U.S. that use U.S.-flag vessels. Information about the requestor's vessel, including a brief description of the proposed service, is listed below.

DATES: Submit comments on or before March 6, 2023.

ADDRESSES: You may submit comments identified by DOT Docket Number MARAD-2023-0013 by any one of the following methods:

- *Federal eRulemaking Portal:* Go to <http://www.regulations.gov>. Search MARAD-2023-0013 and follow the instructions for submitting comments.
- *Mail or Hand Delivery:* Docket Management Facility is in the West Building, Ground Floor of the U.S. Department of Transportation. The Docket Management Facility location address is: U.S. Department of Transportation, MARAD-2023-0013, 1200 New Jersey Avenue SE, West Building, Room W12-140, Washington, DC 20590, between 9 a.m. and 5 p.m., Monday through Friday, except on Federal holidays.

Note: If you mail or hand-deliver your comments, we recommend that you include your name and a mailing address, an email address, or a telephone number in the body of your document so that we can contact you if we have questions regarding your submission.

Instructions: All submissions received must include the agency name and specific docket number. All comments received will be posted without change to the docket at www.regulations.gov, including any personal information provided. For detailed instructions on submitting comments, or to submit comments that are confidential in