their online accounts (e.g., address change). Respondents are State and local government agencies, and some private sector business entities.

<table>
<thead>
<tr>
<th>Modality of completion</th>
<th>Number of respondents</th>
<th>Frequency of response</th>
<th>Average burden per response (minutes)</th>
<th>Estimated total annual burden (hours)</th>
</tr>
</thead>
</table>
| SSA–159
                        | 1,151                 | 1                     | 15                                   | 288                                  |
| SSA–160
                        | 410                   | 1                     | 15                                   | 103                                  |
| Totals                  | 1,561                 |                       |                                      | 391                                  |

9. Evidence From Excluded Medical Sources of Evidence—20 CFR 404.1503b and 416.903b—0960–0803. Pursuant to its broad authority to regulate under sections 205(a), 702(a)(5), and 1631(d)(1) of the Act, SSA implemented section 223(d)(5)(C), as amended, through regulations at 20 CFR 404.1503b and 416.903b. These regulations require excluded medical sources to self-report their excluded status in writing each time they submit evidence related to a claim for benefits under Titles II or XVI of the Act. Excluded medical sources’ duty to self-report their excluded status apply to evidence they submit to SSA directly or through a representative, claimant, or other individual or entity. The respondents for this collection are medical sources that: (1) Meet one of the exclusionary categories set forth in section 223(d)(5)(C)(i) of the Act, as amended; and (2) furnish evidence related to a claim for benefits under Titles II or XVI of the Act.

<table>
<thead>
<tr>
<th>Regulation section(s)</th>
<th>Number of respondents</th>
<th>Frequency of response</th>
<th>Number of responses</th>
<th>Average burden per response (minutes)</th>
<th>Estimated total annual burden (hours)</th>
</tr>
</thead>
<tbody>
<tr>
<td>404.1503b(c), 416.903b(c)</td>
<td>50</td>
<td>60</td>
<td>3,000</td>
<td>20</td>
<td>1,000</td>
</tr>
</tbody>
</table>

DEPARTMENT OF STATE

[Public Notice 10843]


SUMMARY: Notice is hereby given of the following determinations: I hereby determine that certain objects to be included in the exhibition “Lari Pittman: Declaration of Independence,” imported from abroad for temporary exhibition within the United States, are of cultural significance. The objects are imported pursuant to a loan agreement with the foreign owner or custodian. I also determine that the exhibition or display of the exhibit objects at the Hammer Museum, Los Angeles, California, from on or about September 29, 2019, until on or about January 5, 2020, and at possible additional exhibitions or venues yet to be determined, is in the national interest. I have ordered that Public Notice of these determinations be published in the Federal Register.


Marie Therese Porter Royce,
Assistant Secretary, Bureau of Educational and Cultural Affairs, Department of State.

[FR Doc. 2019–17335 Filed 8–12–19; 8:45 am]
BILLING CODE 4710–05–P

DEPARTMENT OF STATE

[Delegation of Authority No. 474]

Delegation of Authority To Concur With Decisions by the Secretary of Energy To Enter Into Agreements Relating to Contributions for Certain Nonproliferation Programs

By virtue of the authority vested in the Secretary of State, including section 1 of the State Department Basic Authorities Act, as amended (22 U.S.C. 2651a), and to the extent authorized by law, I hereby delegate to the Under Secretary for Arms Control and International Security the authority to concur with decisions by the Secretary of Energy to enter into agreements relating to contributions for certain Department of Energy nonproliferation programs, as described in 50 U.S.C. 2569(f).

The Secretary or the Deputy Secretary may at any time exercise any authority or function delegated by this delegation of authority. Any act, authority, or procedure subject to, or affected by, this delegation shall be deemed to be such act, authority, or procedure as amended from time to time.

This delegation of authority shall be published in the Federal Register.
DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

[Docket No. 2019–0599]

Agency Information Collection Activities: Requests for Comments; Clearance of Renewed Approval of Information Collection: Medical Standards and Certification

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 collection involves information applicants must provide on an application for a FAA medical certificate. The information to be collected will be used to evaluate an applicant’s medical fitness.

DATES: Written comments should be submitted by October 15, 2019.

ADDRESSES: Please send written comments:

By Electronic Docket: www.regulations.gov (Enter docket number into search field)

By mail: Nicole Harrison, Federal Aviation Administration, AAM–120, 800 Independence Ave. SW, Washington, DC 20591

FOR FURTHER INFORMATION CONTACT: Judi Citrenbaum by email at: judi.m.citrenbaum@faa.gov phone: 202–267–9689.

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. The agency will summarize and/or include your comments in the request for OMB’s clearance of this information collection.

OMB Control Number: 2120–0034.

Title: Medical Standards and Certification


Type of Review: Renewal of an information collection.

Background: The Secretary of Transportation collects this information under the authority of 49 U.S.C. 40113; 44701; 44510; 44702; 44703; 44709; 45303; and 80111. The FAA medical certification program is implemented by Title 14 Code of Federal Regulations (CFR) parts 61 and 67 (14 CFR parts 61 and 67). The Federal Aviation Administration (FAA) determines if applicants are medically qualified to perform the duties associated with the class of medical certificate sought by evaluating the information applicants provide on FAA Form 8500–8. Also, the agency uses two vision forms, as indicated, for individuals who may need further eye evaluation.

Respondents: 405,345 (all three forms).

Frequency: On occasion.

Estimated Average Burden per Response: 1.5 Hours.

Estimated Total Annual Burden: 585,517 Hours.

Issued in Washington, DC, on August 7, 2019.

Nicole Harrison,

Management and Program Analyst, Office of Aerospace Medicine, Management and Personnel Systems Branch, AAM–120.

[FR Doc. 2019–17235 Filed 8–12–19; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF TRANSPORTATION

Federal Motor Carrier Safety Administration

[Docket No. FMCSA–2019–0140]

Agency Information Collection Activities: Extension of an Approved Information Collection Request: Transportation of Hazardous Materials, Highway Routing

AGENCY: Federal Motor Carrier Safety Administration (FMCSA), U.S. 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 its review and approval and invites public comment. FMCSA requests approval to extend an existing 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 their highways, including dates that such routes were established and information on subsequent changes or new hazardous materials routing designations.

DATES: We must receive your comments on or before October 11, 2019.

ADDRESSES: You may submit comments identified by Federal Docket Management System (FDMS) Docket Number FMCSA–2019–0140 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 Management Facility; 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 docket, 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