

Pursuant to 49 U.S.C. 33106 and 49 CFR 543.7(b), the agency grants a petition for exemption from the parts-marking requirements of Part 541 either in whole or in part, if it determines that, based upon substantial evidence, the standard equipment antitheft device is likely to be as effective in reducing and deterring motor vehicle theft as compliance with the parts-marking requirements of Part 541. The agency finds that Nissan has provided adequate reasons for its belief that the antitheft device for the Infiniti QX50 vehicle line is likely to be as effective in reducing and deterring motor vehicle theft as compliance with the parts-marking requirements of the Theft Prevention Standard (49 CFR part 541). This conclusion is based on the information Nissan provided about its device.

For the foregoing reasons, the agency hereby grants in full Nissan's petition for exemption for the Nissan Infiniti QX50 vehicle line from the parts-marking requirements of 49 CFR part 541. The agency notes that 49 CFR part 541, Appendix A-1, identifies those lines that are exempted from the Theft Prevention Standard for a given model year. 49 CFR part 543.7(f) contains publication requirements incident to the disposition of all Part 543 petitions. Advanced listing, including the release of future product nameplates, the beginning model year for which the petition is granted and a general description of the antitheft device is necessary in order to notify law enforcement agencies of new vehicle lines exempted from the parts-marking requirements of the Theft Prevention Standard.

If Nissan decides not to use the exemption for this line, it must formally notify the agency. If such a decision is made, the line must be fully marked according to the requirements under 49 CFR parts 541.5 and 541.6 (marking of major component parts and replacement parts).

NHTSA notes that if Nissan wishes in the future to modify the device on which this exemption is based, the company may have to submit a petition to modify the exemption. Part 543.7(d) states that a Part 543 exemption applies only to vehicles that belong to a line exempted under this part and equipped with the antitheft device on which the line's exemption is based. Further, Part 543.9(c)(2) provides for the submission of petitions "to modify an exemption to permit the use of an antitheft device similar to but differing from the one specified in that exemption."

The agency wishes to minimize the administrative burden that Part 543.9(c)(2) could place on exempted

vehicle manufacturers and itself. The agency did not intend in drafting Part 543 to require the submission of a modification petition for every change to the components or design of an antitheft device. The significance of many such changes could be *de minimis*. Therefore, NHTSA suggests that if the manufacturer contemplates making any changes, the effects of which might be characterized as *de minimis*, it should consult the agency before preparing and submitting a petition to modify.

Issued in Washington, DC, under authority delegated in 49 CFR part 1.95.

**Raymond R. Posten,**

*Associate Administrator for Rulemaking.*

[FR Doc. 2017-22658 Filed 10-18-17; 8:45 am]

**BILLING CODE 4910-59-P**

## DEPARTMENT OF THE TREASURY

### Internal Revenue Service

#### Proposed Collection; Comment Request for Form 1024-A; Extension of Comment Period

**AGENCY:** Internal Revenue Service (IRS), Treasury.

**ACTION:** Notice and request for comments; extension of comment period.

**SUMMARY:** This document extends the comment period for a notice and request for comments that was published in the **Federal Register** on Monday, August 28, 2017. The notice and request for comments relates to the Application for Recognition of Exemption Under Section 501(c)(4) of the Internal Revenue Code.

**DATES:** The comment period for the notice and request for comments published on August 28, 2017 (82 FR 40228), is extended to November 28, 2017.

**ADDRESSES:** Direct all written comments to L. Brimmer, Internal Revenue Service, Room 6526, 1111 Constitution Avenue NW., Washington, DC 20224.

**FOR FURTHER INFORMATION CONTACT:** Requests for additional information or copies of the form and instructions should be directed to LaNita Van Dyke at (202) 317-6009, Internal Revenue Service, Room 6526, 1111 Constitution Avenue NW., Washington, DC 20224, or through the Internet at [lanita.vandyke@irs.gov](mailto:lanita.vandyke@irs.gov).

**SUPPLEMENTARY INFORMATION:** A notice and request for comments that appeared in the **Federal Register** on Monday, August 28, 2017 (82 FR 40228) announced that written comments are to

be received by October 23, 2017. In order to provide the public with a sufficient opportunity to submit comments, the due date to receive written comments has been extended to Tuesday, November 28, 2017.

Approved: October 12, 2017.

**L. Brimmer,**

*Senior Tax Analyst.*

[FR Doc. 2017-22596 Filed 10-16-17; 4:15 pm]

**BILLING CODE 4830-01-P**

## DEPARTMENT OF VETERANS AFFAIRS

### Cooperative Studies Scientific Evaluation Committee; Notice of Meeting

The Department of Veterans Affairs gives notice under the Federal Advisory Committee Act that the Cooperative Studies Scientific Evaluation Committee will hold a meeting on December 13, 2017, at the American Association of Airport Executives, 601 Madison Street, Alexandria, VA. The meeting will begin at 8:30 a.m. and end at 3:30 p.m.

The Committee advises the Chief Research and Development Officer on the relevance and feasibility of proposed projects and the scientific validity and propriety of technical details, including protection of human subjects.

The session will be open to the public for approximately 30 minutes at the start of the meeting for the discussion of administrative matters and the general status of the program. The remaining portion of the meeting will be closed to the public for the Committee's review, discussion, and evaluation of research and development applications.

During the closed portion of the meeting, discussions and recommendations will deal with qualifications of personnel conducting the studies, staff and consultant critiques of research proposals and similar documents, and the medical records of patients who are study subjects, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy. As provided by section 10(d) of Public Law 92-463, as amended, closing portions of this meeting is in accordance with 5 U.S.C. 552b(c)(6) and (c)(9)(B).

The Committee will not accept oral comments from the public for the open portion of the meeting. Those who plan to attend or wish additional information should contact Dr. Grant Huang, Acting Director, Cooperative Studies Program (10P9CS), Department of Veterans Affairs, 810 Vermont Avenue NW., Washington, DC 20420, at (202) 443-

5700 or by email at [grant.huang@va.gov](mailto:grant.huang@va.gov). Those wishing to submit written comments may send them to Dr. Huang at the same address and email.

Dated: October 16, 2017.

**LaTonya L. Small,**  
Federal Advisory Committee Management  
Officer.

[FR Doc. 2017-22679 Filed 10-18-17; 8:45 am]

BILLING CODE 8320-01-P

## DEPARTMENT OF VETERANS AFFAIRS

[OMB Control No. 2900-0556]

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

**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:** Comments must be submitted on or before November 20, 2017.

**ADDRESSES:** Submit written comments on the collection of information through [www.Regulations.gov](http://www.Regulations.gov), or to Office of Information and Regulatory Affairs, Office of Management and Budget, Attn: VA Desk Officer; 725 17th St. NW., Washington, DC 20503 or sent through electronic mail to [oir\\_submission@omb.eop.gov](mailto:oir_submission@omb.eop.gov). Please refer to "OMB Control No. 2900-0556" in any correspondence.

**FOR FURTHER INFORMATION CONTACT:** Cynthia Harvey-Pryor, Enterprise Records Service (005R1B), Department of Veterans Affairs, 810 Vermont Avenue NW., Washington, DC 20420, (202) 461-5870 or email [cynthia.harvey-pryor@va.gov](mailto:cynthia.harvey-pryor@va.gov). Please refer to "OMB Control No. 2900-0556" in any correspondence.

#### SUPPLEMENTARY INFORMATION:

**Authority:** 38 U.S.C. 7331.

**Title:** Living Will and Durable Power of Attorney for Health Care; VA Form 10-0137

**OMB Control Number:** 2900-0556.

**Type of Review:** Extension of a currently approved collection.

**Abstract:** VA Form 10-0137, VA Advance Directive: Durable Power of Attorney for Health Care and Living Will, is the Department of Veterans Affairs (VA) recognized legal document that permits VA patients to designate a health care agent and/or specify preferences for future health care. The VA Advance Directive is invoked if a patient becomes unable to make health care decisions for him or herself. Use of the VA Advance Directive is specified in VHA Handbook 1004.02, Advance Care Planning and Management of Advance Directives. Veterans' rights to designate a health care agent and specify health care preferences in advance are codified in 38 CFR 17.32. This regulation also obligates VA to recognize advance directives and to use the information contained therein when health care decisions must be made for a patient that has lost decision making capacity. Use of advance directives is a well-established standard within clinical practice in the U.S. Offering the opportunity to complete an advance directive and the requirement to honor such documents is supported by Joint Commission standards and the Patient Self Determination Act of 1990 (applicable to Medicare providers.) Use of advance directives is also consistent with the health care ethics standard that patients have autonomy in health care decision making and have a right to control what is done to them in a medical setting.

An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. The **Federal Register** Notice with a 60-day comment period soliciting comments on this collection of information was published at 82 FR 151 on August 8, 2017, page 37167.

**Affected Public:** Individuals and households.

**Estimated Annual Burden:** 171,811 hours.

**Estimated Average Burden per Respondent:** 30 minutes.

**Frequency of Response:** Annually.

**Estimated Number of Respondents:** 343,622.

By direction of the Secretary.

**Cynthia Harvey-Pryor,**

Department Clearance Officer, Office of  
Quality, Privacy and Risk, Department of  
Veterans Affairs.

[FR Doc. 2017-22689 Filed 10-18-17; 8:45 am]

BILLING CODE 8320-01-P

## DEPARTMENT OF VETERANS AFFAIRS

[OMB Control No. 2900-NEW]

### Agency Information Collection Activity: VA Disaster Resilience Survey of Community Dwelling Veterans

**AGENCY:** Veterans Health  
Administration, Department of Veterans  
Affairs.

**ACTION:** Notice.

**SUMMARY:** Veterans Health Administration, Department of Veterans Affairs (VA), is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act (PRA) of 1995, Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of a currently approved collection, and allow 60 days for public comment in response to the notice.

**DATES:** Written comments and recommendations on the proposed collection of information should be received on or before December 18, 2017.

**ADDRESSES:** Submit written comments on the collection of information through Federal Docket Management System (FDMS) at [www.Regulations.gov](http://www.Regulations.gov) or to Brian McCarthy, Office of Regulatory and Administrative Affairs (10B4), Department of Veterans Affairs, 810 Vermont Avenue NW., Washington, DC 20420 or email to [Brian.McCarthy4@va.gov](mailto:Brian.McCarthy4@va.gov). Please refer to "OMB Control No. 2900-NEW" in any correspondence. During the comment period, comments may be viewed online through FDMS.

**FOR FURTHER INFORMATION CONTACT:** Brian McCarthy at (202) 461-6345.

**SUPPLEMENTARY INFORMATION:** Under the PRA of 1995, Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor.

With respect to the following collection of information, VHA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of VHA's functions, including whether the information will have practical utility; (2) the accuracy of VHA's estimate of the burden of the proposed collection of information; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use