

*Legal authorization and confidentiality:* The filing requirements under the FR 1400 are authorized by sections 10 and 11 of the Federal Reserve Act (FRA)<sup>5</sup> and section 342(c) of the Dodd-Frank Act.<sup>6</sup> Registering in the Supplier Registration System (FR 1400A) is voluntary. The remaining portions of the FR 1400 (FR 1400B, FR 1400C, and FR 1400D) are required to obtain a benefit for prospective suppliers to the Board.

A prospective supplier may request confidential treatment of information submitted as part of its Procurement Solicitation Package under exemption 4 of the Freedom of Information Act (FOIA), which protects commercial or financial information that is both customarily and actually treated as private.<sup>7</sup> In addition, a prospective supplier may request confidential treatment of information pursuant to exemption 6 of the FOIA, which protects personal information, the disclosure of which would “constitute a clearly unwarranted invasion of privacy.”<sup>8</sup> Determinations of confidentiality based on exemption 4 or exemption 6 of the FOIA would be made on a case-by-case basis.

*Current actions:* On November 23, 2021, the Board published a notice in the **Federal Register** (86 FR 66557) requesting public comment for 60 days on the extension, without revision, of the Procurement Solicitation Package. The comment period for this notice expired on January 24, 2022. The Board did not receive any comments.

Board of Governors of the Federal Reserve System, April 1, 2022.

**Margaret Shanks,**

*Deputy Secretary of the Board.*

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conclusions, and recommendations for legislative or agency action.

<sup>5</sup> Section 10(3) and section 11 of the FRA authorize the Board to manage its buildings and staff. 12 U.S.C. 243 and 248(1). Section 10(4) of the FRA authorizes the Board to determine and prescribe the manner in which its obligations shall be incurred and its disbursements and expenses allowed and paid. 12 U.S.C. 244.

<sup>6</sup> 12 U.S.C. 5452(c) (requiring the Board to develop and implement standards and procedures for the review and evaluation of contract proposals and for hiring service providers that include a component that gives consideration to the diversity of a prospective supplier and the fair inclusion of women and minorities in the workforce of such supplier and any subcontractor).

<sup>7</sup> 5 U.S.C. 552(b)(4).

<sup>8</sup> 5 U.S.C. 552(b)(6).

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10416]

#### Agency Information Collection Activities: Proposed Collection; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

**ACTION:** Notice.

**SUMMARY:** The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS’ intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency’s functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**DATES:** Comments must be received by June 6, 2022.

**ADDRESSES:** When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to <http://www.regulations.gov>. Follow the instructions for “Comment or Submission” or “More Search Options” to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number: \_\_\_\_\_, Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS’ website address at website address at <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing>.

**FOR FURTHER INFORMATION CONTACT:** William N. Parham at (410) 786-4669.

**SUPPLEMENTARY INFORMATION:**

#### Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection’s supporting statement and associated materials (see **ADDRESSES**).

#### *CMS-10416 Blueprint for Approval of State-Based Exchange*

Under the PRA (44 U.S.C. 3501-3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term “collection of information” is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

#### Information Collection

1. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Blueprint for Approval of State-based Exchange; *Use:* The Patient Protection and Affordable Care Act (ACA) and its implementing regulations provide states with flexibility in the design and operation of Exchanges to ensure states are implementing Exchanges that best meet the needs of their consumers. States can choose to establish and operate a State-based Exchange (SBE) or a State-based Exchange on the Federal Platform (SBE-FP). To ensure a state can operate a successful and compliant SBE or SBE-FP, it is critical that states provide CMS with a complete and thorough Exchange

Blueprint Application, Declaration of Intent Letter, and attest to demonstrate operational readiness. The information collected from states will be used by CMS, IRS, SSA and reviewed by other Federal agencies to determine if a state can implement a complete and fully operational Exchange. *Form Number:* CMS-10416 (OMB control number: 0938-1172); *Frequency:* Annually; *Affected Public:* State, Local, or Tribal governments; *Number of Respondents:* 4; *Total Annual Responses:* 21; *Total Annual Hours:* 126. (For policy questions regarding this collection contact Shilpa Gogna at 301-492-4257.)

Dated: March 30, 2022.

**William N. Parham, III,**

*Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.*

[FR Doc. 2022-07100 Filed 4-4-22; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Center for Scientific Review; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* Center for Scientific Review Special Emphasis Panel; Therapeutic Development and Preclinical Studies.

*Date:* April 14, 2022.

*Time:* 10:00 a.m. to 2:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

*Contact Person:* Richard D Schneiderman, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4138, Bethesda, MD 20817, 301-402-3995, [richard.schneiderman@nih.gov](mailto:richard.schneiderman@nih.gov).

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393-93.396, 93.837-93.844, 93.846-93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: March 30, 2021.

**David W. Freeman,**

*Program Analyst, Office of Federal Advisory Committee Policy.*

[FR Doc. 2022-07115 Filed 4-4-22; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Center for Scientific Review; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* Center for Scientific Review Special Emphasis Panel; RFA Panel: Tobacco Regulatory Science A.

*Date:* May 5, 2022.

*Time:* 10:00 a.m. to 7:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

*Contact Person:* Sepandarmaz Aschrafi, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4040D, Bethesda, MD 20892, (301) 451.4251, [Armaz.aschrafi@nih.gov](mailto:Armaz.aschrafi@nih.gov).

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393-93.396, 93.837-93.844, 93.846-93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: March 31, 2022.

**Melanie J. Pantoja,**

*Program Analyst, Office of Federal Advisory Committee Policy.*

[FR Doc. 2022-07149 Filed 4-4-22; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### National Institute on Aging; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* National Institute on Aging Special Emphasis Panel; Stem Cell III.

*Date:* April 29, 2022.

*Time:* 1:30 p.m. to 5:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institute on Aging, Gateway Building, 7201 Wisconsin Avenue, Bethesda, MD 20892 (Virtual Meeting).

*Contact Person:* Nijaguna Prasad, Ph.D., Scientific Review Officer, Scientific Review Branch, National Institute on Aging, 7201 Wisconsin Avenue, Gateway Bldg., Suite 2W200, Bethesda, MD 20892, (301) 496-9667 [prasadnb@nia.nih.gov](mailto:prasadnb@nia.nih.gov).

(Catalogue of Federal Domestic Assistance Program Nos. 93.866, Aging Research, National Institutes of Health, HHS)

Dated: March 30, 2022.

**Miguelina Perez,**

*Program Analyst, Office of Federal Advisory Committee Policy.*

[FR Doc. 2022-07110 Filed 4-4-22; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Center for Scientific Review; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material,