established in the last few decades; however, FDA has become aware of an increase in the number of adverse event reports related to the misuse of these products. These reports led FDA to convene a meeting of the Ophthalmic Devices Panel of the Medical Devices Advisory Committee and the Risk Communication Advisory Committee on March 17, 2017, to discuss additional measures to mitigate the potential risk for misuse of these devices. The meeting covered a range of important issues, including appropriate labeling and packaging of these products and the importance of clearly communicating these concerns to the consumer public, which were incorporated into this draft guidance. When finalized, this guidance is intended to provide recommendations concerning the content and format of labeling for HPCPs. FDA believes that the labeling recommendations in this guidance may help manufacturers develop labeling with information about specific risks and directions for use of the HPCPs in conjunction with a user's prescribed contact lenses.

This draft guidance is being issued consistent with FDA's good guidance practices regulation (21 CFR 10.115). The draft guidance, when finalized, will represent the current thinking of FDA on ''Hydrogen Peroxide-Based Contact Lens Care Products: Consumer Labeling Recommendations—Premarket Notification (510(k)) Submissions.'' It does not establish any rights for any person and is not binding on FDA or the public. You can use an alternative approach if it satisfies the requirements of the applicable statutes and regulations.

II. Electronic Access

Persons interested in obtaining a copy of the draft guidance may do so by downloading an electronic copy from the internet. A search capability for all Center for Devices and Radiological Health guidance documents is available at *https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance/guidance-documents-medical-devices-and-radiation-emitting-products.* This guidance document is also available at *https://www.regulations.gov* or *https://www.fda.gov/regulatory-information/search-fda-guidance-documents.*

Persons unable to download an electronic copy of ''Hydrogen Peroxide-Based Contact Lens Care Products: Consumer Labeling Recommendations—Premarket Notification (510(k)) Submissions'' may send an email request to *CDRH-Guidance@fda.hhs.gov* to receive an electronic copy of the document. Please use the document number 18041 and complete title to identify the guidance you are requesting.

III. Paperwork Reduction Act of 1995

While this guidance contains no new collection of information, it does refer to previously approved FDA collections of information. Therefore, clearance by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3521) is not required for this guidance. The previously approved collections of information are subject to review by OMB under the PRA. The collections of information in the following FDA regulations, guidance, and forms have been approved by OMB as listed in the following table:

21 CFR part	Topic	OMB control No.
807, subpart E	Premarket notification	0910–0120
800, 801, and 809	Medical Device Labeling Regulations	0910–0485

Dated: August 11, 2022.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2022–17642 Filed 8–16–22; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Senior Executive Service Performance Review Board

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: HRSA, an operating division of HHS, is publishing a list of individuals who may be named to serve on the Senior Executive Service Performance Review Board that oversees the evaluation of performance appraisals for Senior Executive Service members for the Fiscal Years 2022 and 2023.

FOR FURTHER INFORMATION CONTACT: Georgia Lyons, HRSA Executive Resources, Office of Human Resources, 5600 Fishers Lane, Rm 12N06C, Rockville, Maryland 20857, or (301) 443–4618.

SUPPLEMENTARY INFORMATION: Title 5, U.S.C. Section 4314(c)(4) of the Civil Service Reform Act of 1978, Public Law 95–454, requires that the appointment of Performance Review Board Members be published in the **Federal Register**. The following individuals may be named to serve on the Senior Executive Service Performance Review Board:

Onyekachukwu Anaedozie
Anthony Archeval
Cynthia Baugh
Tonya Bowers
Adriane Burton
Tina Cheatham
Laura Cheever
Christopher Coppenbarger
Natasha Coulouris
Cheryl Dammons
Elizabeth DeVoss
Tanette Downs
Diana Espinosa
Catherine Ganey
Alexandra Garcia
Jordan Grossman
Heather Hauck
Alexandra Huttinger
Carole Johnson
Laura Kavanagh
Martin Kramer
James Macrae
Maren McBride Kahn
Susan Monarez
Thomas Morris
Suma Nair
Luis Padilla
Nisha Patel
Krista Pedley
Wendy Ponton
Sheila Pradia Williams
Michael Warren

Carole Johnson,

Administrator.

[FR Doc. 2022–17657 Filed 8–16–22; 8:45 am]

BILLING CODE 4165–15–P

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.