

practices regulation (21 CFR 10.115). The guidance represents the current thinking of FDA on “Clarification of Radiation Control Regulations for Manufacturers of Diagnostic X-Ray Equipment.” 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 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 “Clarification of Radiation Control Regulations for Manufacturers of Diagnostic X-Ray Equipment” may send an email request to CDRH-Guidance@fda.hhs.gov to receive an electronic copy of the

document. Please use the document number GUI01500029 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. The previously approved collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3521). The collections of information in the following FDA table have been approved by OMB:

21 CFR Part or FDA Form	Topic	OMB Control No.
1002, 1005, 1010, 1020, 1030, 1040, and 1050; form FDA 2579 and form FDA 2877.	Reporting and Recordkeeping for Electronic Products—General Requirements.	0910–0025
800, 801, and 809	Labeling	0910–0485
812	Investigational Device Exemption	0910–0078
“Allegations of Regulatory Misconduct” form	Voluntary Allegations of Regulatory Misconduct	0910–0769

Dated: September 25, 2024.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2024–22332 Filed 9–27–24; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA–2024–N–0758, FDA–2024–N–2032, FDA–2023–N–3847, and FDA–2024–N–1201]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approvals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is publishing a list of information collections that have been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT:

Amber Sanford, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–8867, PRAStaff@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: The following is a list of FDA information collections recently approved by OMB under section 3507 of the Paperwork Reduction Act of 1995 (44 U.S.C. 3507).

The OMB control number and expiration date of OMB approval for each information collection are shown in table 1. Copies of the supporting statements for the information collections are available on the internet at <https://www.reginfo.gov/public/do/PRAMain>. 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.

TABLE 1—LIST OF INFORMATION COLLECTIONS APPROVED BY OMB

Title of collection	OMB control No.	Date approval expires
New Plant Varieties Intended for Food Use	0910–0583	9/30/2027
Food and Cosmetic Export Certificates	0910–0793	9/30/2027
Biological Products; General Records and Postmarket Adverse Experience Reporting	0910–0308	9/30/2027
Voluntary Total Product Life Cycle Advisory Program Pilot	0910–0930	9/30/2027

Dated: September 24, 2024.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2024–22291 Filed 9–27–24; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Recharter for the Council on Graduate Medical Education

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

ACTION: Notice.

SUMMARY: In accordance with the Federal Advisory Committee Act, the Department of Health and Human Services is hereby giving notice that the Council on Graduate Medical Education (COGME or Council) is rechartered.

DATES: The effective date of the renewed charter is September 30, 2024.

FOR FURTHER INFORMATION CONTACT:

Shane Rogers, Designated Federal Official (DFO), Division of Medicine and Dentistry, Bureau of Health Workforce, HRSA, 5600 Fishers Lane, Room 15N142, Rockville, Maryland 20857; 301–443–5260; or strogers@hrsa.gov.

SUPPLEMENTARY INFORMATION: COGME makes recommendations to the Secretary of Health and Human Services and Congress on matters specified by section 762 of the Public Health Service Act. Issues addressed by COGME include: (1) the supply and distribution of physicians in the United States; (2) current and future shortages or excesses of physicians in medical and surgical specialties and subspecialties; (3) issues relating to international medical school graduates; (4) appropriate federal policies with respect to the matters specified in (1), (2), and (3) above, including policies concerning changes in the financing of undergraduate and graduate medical education programs and changes in the types of medical education training in graduate medical education programs; (5) appropriate efforts to be carried out by hospitals, schools of medicine, schools of osteopathic medicine, and accrediting bodies with respect to the matters specified in (1), (2), and (3) above, including efforts for changes in undergraduate and graduate medical education programs; and (6) deficiencies in, and needs for improvements in, existing databases concerning the supply and distribution of, and

postgraduate training programs for, physicians in the United States and steps that should be taken to eliminate those deficiencies. Not later than September 30, 2023, and not less than every 5 years thereafter, COGME shall submit a report on the recommendations made by the committee to the Secretary of Health and Human Services, and to the Committee on Health, Education, Labor, and Pensions of the Senate and the Committee on Energy and Commerce of the House of Representatives. Additionally, COGME encourages entities providing graduate medical education to conduct activities to voluntarily achieve the recommendations of the Council; develops, publishes, and implements performance measures; develops and publishes guidelines for longitudinal evaluations; and recommends appropriation levels for certain programs under title VII of the Public Health Service Act.

The renewed charter for COGME was approved on September 18, 2024. The filing date is September 30, 2024. The recharter of COGME gives authorization for the Council to operate until September 30, 2026.

A copy of the COGME charter is available on the COGME website at <https://www.hrsa.gov/sites/default/files/hrsa/advisory-committees/graduate-medical-edu/cogme-charter.pdf>. A copy of the charter also can be obtained by accessing the FACA database that is maintained by the Committee Management Secretariat under the General Services Administration. The website address for the FACA database is <http://www.facadatabase.gov/>.

Maria G. Button,

Director, Executive Secretariat.

[FR Doc. 2024–22256 Filed 9–27–24; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director, National Institutes of Health; Notice of Meeting

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the Office of AIDS Research Advisory Council.

The meeting will be held as a virtual meeting and will be open to the public as indicated below. Individuals who plan to view the virtual meeting and need special assistance or other reasonable accommodations to view the meeting, should notify the Contact

Person listed below in advance of the meeting. The meeting can be accessed from the NIH Videocast at the following link: <https://videocast.nih.gov/>.

Name of Committee: Office of AIDS Research Advisory Council.

Date: October 24, 2024.

Time: 12:00 p.m. to 5:00 p.m.

Agenda: Report from the OAR Director; Update on the Development of the NIH Strategic Plan and Research Priorities for HIV Research.

Place: Office of AIDS Research, Office of the Director, NIH, 5601 Fishers Lane, Rockville, MD 20852, (Virtual Meeting) <https://videocast.nih.gov/>.

Contact Person: CAPT Mary Glenshaw, Ph.D., M.P.H., OTR/L, Office of AIDS Research, Office of the Director, NIH, 5601 Fishers Lane, Room 2E61, Rockville, MD 20852, (301) 496–0357, OARACinfo@nih.gov.

Any interested person may file written comments with the committee by forwarding the statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person.

Information is also available on the Institute's/Center's home page: www.oar.nih.gov, where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.14, Intramural Research Training Award; 93.22, Clinical Research Loan Repayment Program for Individuals from Disadvantaged Backgrounds; 93.232, Loan Repayment Program for Research Generally; 93.39, Academic Research Enhancement Award; 93.936, NIH Acquired Immunodeficiency Syndrome Research Loan Repayment Program; 93.187, Undergraduate Scholarship Program for Individuals from Disadvantaged Backgrounds, National Institutes of Health, HHS)

Dated: September 24, 2024.

Lauren A. Fleck,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2024–22274 Filed 9–27–24; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings 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