61. Kediala Magassouba, New York City, New York, Court of Federal Claims No: 20–0762V
62. James Sokol, Seattle, Washington, Court of Federal Claims No: 20–0763V
63. Joanie Arseneault, Melbourne, Florida, Court of Federal Claims No: 20–0765V
64. Gregory Carter, Raleigh, North Carolina, Court of Federal Claims No: 20–0767V
65. Destiny Maynard and Elijah Bunch on behalf of C. B., Deceased, East Moline, Illinois, Court of Federal Claims No: 20–0768V
66. Candice E. Hutchison, Belfast, Maine, Court of Federal Claims No: 20–0773V
67. Kelly Steele, Baltimore, Maryland, Court of Federal Claims No: 20–0775V
68. Lorraine Ferrucci on behalf of J. B., Orchard Park, New York, Court of Federal Claims No: 20–0776V
69. Lisa Black, St. Paul, Minnesota, Court of Federal Claims No: 20–0777V
70. Sharon Danberry, Mankato, Minnesota, Court of Federal Claims No: 20–0778V
71. Brianna Meyers, Albany, New York, Court of Federal Claims No: 20–0779V
72. Sandra Francis, Leesburg, Florida, Court of Federal Claims No: 20–0780V
73. Renee Byndloss, Chicago, Illinois, Court of Federal Claims No: 20–0781V
74. Juliet Wolf, Dresher, Pennsylvania, Court of Federal Claims No: 20–0782V
75. Larry Alex Klickstein, Pasadena, California, Court of Federal Claims No: 20–0785V
76. Antonios Tsamasiros, Rockaway Beach, New York, Court of Federal Claims No: 20–0786V
77. Patricia Parks, White Plains, New York, Court of Federal Claims No: 20–0788V
78. Cheree Roach and Jason Roach on behalf of I. R., Eagle River, Alaska, Court of Federal Claims No: 20–0789V
79. Jacqueline Stokes, Chicago, Illinois, Court of Federal Claims No: 20–0790V
80. Niberley Walton, Fort Belvoir, Virginia, Court of Federal Claims No: 20–0791V
81. Adam Smith, San Antonio, Texas, Court of Federal Claims No: 20–0794V
82. Nydia Ellentuch, Saint Cloud, Florida, Court of Federal Claims No: 20–0795V
83. Julie Leibold, Blue Springs, Missouri, Court of Federal Claims No: 20–0796V
84. Keri H. Daigle, Richmond, Virginia, Court of Federal Claims No: 20–0797V

[FR Doc. 2020–15830 Filed 7–21–20; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–0324]

Agency Information Collection Request: 30-Day Public Comment Request

AGENCY: Office of the Secretary, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.

DATES: Comments on the ICR must be received on or before August 21, 2020.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain* . Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: Sherrette Funn, *Sherrette.Funn@hhs.gov* or (202) 795–7714. When submitting comments or requesting information, please include the document identifier 0990–New–30D and project title for reference.

SUPPLEMENTARY INFORMATION: Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Title of the Collection: Resport of Dental Examination of Applicants to the Commissioned Corps of the U.S. Public Health Service.

Type of Collection: Reinstatement.

OMB No.: 0990–0324.

Abstract: The Commissioned Corps of the U.S. Public Health Service has a need for the information in order to assess the qualifications of each applicant and make a determination whether the applicant meets the requirements to receive a commission. The information is used to make determinations on candidates/applicants seeking appointment to the Corps to assess their medical suitability. The purpose is to evaluate the medical suitability of applicants on the basis of the Corps' medical accession standards and policy. The protected information is accessed by appropriate personnel and clinical reviewers. The form is not disclosed to external entities, other than for uses authorized by law.

Type of respondent; frequency (annual); Applicants/Candidates to the Commissioned Corps of the U.S. Public Health Service.

ESTIMATED ANNUALIZED BURDEN TABLE

Type of respondent	Number of respondents	Number responses per respondent	Average burden per response (in hours)	Total burden hours
Annual	1000	1	1	1000
Total	1000	1	1	1000

Sherrette A. Funn,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.

[FR Doc. 2020–15786 Filed 7–21–20; 8:45 am]

BILLING CODE 4150–49–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Guidance on Elimination of Institutional Review Board (IRB) Review of Research Applications and Proposals: 2018 Requirements

AGENCY: The Office for Human Research Protections, Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services (HHS).

ACTION: Notice

SUMMARY: The Office for Human Research Protections (OHRP), Office of the Assistant Secretary for Health, is announcing the availability of a guidance document titled, ''Guidance on Elimination of Institutional Review Board (IRB) Review of Research Applications and Proposals: 2018 Requirements.'' The guidance document provides OHRP's first formal guidance on this topic. The document, which is available on OHRP's website at *https://www.hhs.gov/ohrp/regulations-and-policy/guidance/index.html,* is intended primarily for institutions, IRBs, investigators, HHS funding agencies, and others that may be responsible for the review, conduct, or oversight of nonexempt research involving human subjects conducted or supported by HHS. The guidance document announced in this notice finalizes the draft guidance that was made available for public comment through a notice in the **Federal Register** on July 25, 2018 (83 FR 35278). OHRP received 2 comments from individuals or organizations on the draft document and those comments were considered as the guidance was finalized.

DATES: Comments on OHRP guidance documents are welcome at any time.

ADDRESSES: Submit written requests for a single copy of the guidance document titled ''Guidance on Elimination of Institutional Review Board (IRB) Review of Research Applications and Proposals: 2018 Requirements'' to the Division of Policy and Assurances, Office for Human Research Protections, 1101 Wootton Parkway, Suite 200, Rockville, MD 20852. Send one self-addressed adhesive label to assist that office in processing your request, or fax your request to 240–453–8420. See the **SUPPLEMENTARY INFORMATION** section for information on electronic access to the guidance document.

Submit written comments to: Comments on Elimination of Institutional Review Board (IRB) Review of Research Applications and Proposals: 2018 Requirements Guidance, Office for Human Research Protections, 1101 Wootton Parkway, Suite 200, Rockville, MD 20852. Comments also may be sent via email to *ohrp@hhs.gov* or via facsimile at 240–453–8420.

FOR FURTHER INFORMATION CONTACT: Irene Stith-Coleman, Ph.D., Office for Human Research Protections, 1101 Wootton Parkway, Suite 200, Rockville, MD 20852, 240–453–6700; email *Irene.Stith-Coleman@hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

OHRP is announcing the availability of a guidance document titled ''Guidance on Elimination of Institutional Review Board (IRB) Review of Research Applications and Proposals: 2018 Requirements.'' The guidance document provides OHRP's first formal guidance on this topic. The document is intended primarily for institutions, IRBs, investigators, HHS funding agencies, and others that may be responsible for the review, conduct, or oversight of nonexempt research involving human subjects conducted or supported by HHS.

The guidance document applies to nonexempt research involving human subjects that is conducted or supported by HHS. It provides guidance on the elimination of the requirement in section 45 CFR 46.103(f) of the pre-2018 Requirements that each application or proposal for research undergo IRB review and approval as part of the certification process. This guidance also addresses the requirement in the 2018 Requirements for certification of each proposed research study prior to initiation. In particular, the guidance addresses the following two topics: (1) Pre-2018 Requirements; and, (2) 2018 Requirements.

The guidance document announced in this notice finalizes the draft guidance that was made available for public comment through a notice in the **Federal Register** on July 25, 2018 (83 FR 35278).

II. Electronic Access

Persons with access may obtain the draft guidance documents on OHRP's website at OHRP's website at *https://www.hhs.gov/ohrp/regulations-and-policy/guidance/index.html.*

Dated: July 16, 2020.

Jerry Menikoff,

Director, Office for Human Research Protections.

[FR Doc. 2020–15808 Filed 7–21–20; 8:45 am]

BILLING CODE 4150–36–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Eunice Kennedy Shriver National Institute of Child Health & Human Development; 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/or contract proposals 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 of Child Health and Human Development Initial Review Group Biobehavioral and Behavioral Sciences Subcommittee.

Date: October 23, 2020.

Time: 10:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: NIH/NICHD, 6710B Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Clayton W. Mash, Ph.D., Scientific Review Branch, Eunice Kennedy Shriver National Institute of Child Health and Human Development, NIH 6710B, Rockledge Drive, Rm. 2131A, Bethesda, MD 20892, (301) 496–6866, *mashc@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.864, Population Research; 93.865, Research for Mothers and Children; 93.929, Center for Medical Rehabilitation Research; 93.209, Contraception and Infertility Loan Repayment Program, National Institutes of Health, HHS)

Dated: July 16, 2020.

Ronald J. Livingston, Jr.,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2020–15783 Filed 7–21–20; 8:45 am]

BILLING CODE 4140–01–P