TABLE 2—NEW ENTRIES TO THE LIST OF RECOGNIZED STANDARDS—Continued

Recognition No.	Title of standard ¹	Reference No. and date		
11–393	Standard Test Method for Evaluating Mobile Bearing Knee Dislocation	ASTM F2724–21.		
	O. Physical Medicine			
16–232	Medical electrical equipment—Part 2–78: Particular requirements for basic safety and essential performance of medical robots for rehabilitation, assessment, compensation or alleviation.	IEC 80601-2-78 Edition 1.0 2019-07		
	P. Radiology			
	No new entries at this time.			
	Q. Software/Informatics			
13–120	Medical Devices and Medical Systems—Essential safety requirements for equipment comprising the patient-centric integrated clinical environment (ICE)—Part 1: General requirements and conceptual model.	ANSI/AAMI 2700-1:2019.		
	R. Sterility			
14–571	Sterilization of health care products—Biological indicators—Part 8: Method for validation of a reduced incubation time for a biological indicator.	ISO 11138–8 First edition 2021–07.		
	S. Tissue Engineering			
	No new entries at this time.			

¹ All standard titles in this table conform to the style requirements of the respective organizations.

IV. List of Recognized Standards

FDA maintains the current list of FDA Recognized Consensus Standards in a searchable database that may be accessed at https://www.accessdata. fda.gov/scripts/cdrh/cfdocs/ cfStandards/search.cfm. Such standards are those that FDA has recognized by notice published in the Federal Register or that FDA has decided to recognize but for which recognition is pending (because a periodic notice has not yet appeared in the Federal Register). FDA will announce additional modifications and revisions to the list of recognized consensus standards, as needed, in the Federal Register once a year, or more often if necessary.

V. Recommendation of Standards for Recognition by FDA

Any person may recommend consensus standards as candidates for recognition under section 514 of the FD&C Act by submitting such recommendations, with reasons for the recommendation, to CDRHStandardsStaff@fda.hhs.gov. To be considered, such recommendations should contain, at a minimum, the information available at https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance/standards-and-conformity-assessment-program#process.

Dated: April 18, 2022.

Lauren K. Roth,

Associate Commissioner for Policy. [FR Doc. 2022–08571 Filed 4–21–22; 8:45 am] BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Indian Health Service

[OMB No. 0917-0040]

Request for Public Comment: 30-Day Information Collection: Request for Reinstatement of Indian Health Service Purchased/Referred Care Proof of Residency

AGENCY: Indian Health Service, HHS. **ACTION:** Notice and request for comments; request for reinstatement.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, the Indian Health Service (IHS) invites the general public to take this opportunity to comment on the reinstatement of the information collection, Office of Management and Budget (OMB) Control Number 0917–0040, titled, Purchased/Referred Care Proof of Residency. The IHS is requesting OMB to approve a reinstatement of this collection. Notice regarding the information collection was last published in the Federal Register on January 24, 2022, and allowed 60 days for public comment. The purpose

of this notice is to announce the IHS's intent to reinstate this collection to OMB and to allow 30 days for public comment to be submitted directly to OMB. A copy of the supporting statement is available at www.regulations.gov (see Docket ID: IHS_FRDOC_0001).

comments received by May 23, 2022. ADDRESSES: Direct Your Comments to OMB: Send your comments and suggestions regarding the proposed information collection contained in this notice, especially regarding the estimated public burden and associated response time to: Office of Management and Budget, Office of Regulatory Affairs, New Executive Office Building, Room 10235, Washington, DC 20503, Attention: Desk Officer for IHS.

DATES: Consideration will be given to all

FOR FURTHER INFORMATION CONTACT: To request additional information, please contact Evonne Bennett, Information Collection Clearance Officer at: Evonne.Bennett@ihs.gov or 301–443–4750.

SUPPLEMENTARY INFORMATION: This previously approved information collection project was last published in the **Federal Register** on January 24, 2022, and allowed 60 days for public comment (87 FR 3562). No public comment was received in response to the notice. This notice announces our intent to reinstate this collection, which expired March 31, 2022; to submit this

collection to OMB for approval of reinstatement; and to solicit comments on specific aspects for the proposed information collection.

Title: Purchased/Referred Care Proof of Residency.

OMB Control Number: 0917–0040. Need and Use of Information Collection: The IHS Purchased/Referred Care Program needs the information requested on the PRC Proof of Residency form to verify that individuals seeking medical services through a PRC program meet the residency requirements specific to PRC under 42 CFR 136.23.

Agency Form Number: IHS 976.

Members of Affected Public: Individuals/Households.

Status of the Proposed Information Collection: Renewal request.

Type of Respondents: Individuals. The table below provides: Types of data collection instruments, Estimated number of respondents, Number of responses per respondent, Annual number of responses, Average burden hour per response, and Total annual burden hours.

Data collection instrument(s)	Estimated number of respondents	Responses per respondent	Annual number of responses	Average burden hour per response*	Total annual burden hours
Individual Patient Count	77,185	1	77,185	3/60	3,859.25
Total	77,185	1	77,185	3/60	3,859.25

^{*} For ease of understanding, the average burden per response is 3 minutes.

There are no direct costs to respondents to report.

Requests for Comments: Your written comments and/or suggestions are invited on one or more of the following points:

- (a) Whether the information collection activity is necessary to carry out an agency function;
- (b) whether the agency processes the information collected in a useful and timely fashion;
- (c) the accuracy of the public burden estimate (the estimated amount of time needed for individual respondents to provide the requested information);
- (d) whether the methodology and assumptions used to determine the estimates are logical;
- (e) ways to enhance the quality, utility, and clarity of the information being collected; and
- (f) ways to minimize the public burden through the use of automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Elizabeth A. Fowler,

Acting Director, Indian Health Service. [FR Doc. 2022–08619 Filed 4–21–22; 8:45 am]

BILLING CODE 4165-16-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Nursing Research; Notice of Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the National Advisory Council for Nursing Research. The meeting will be open to the public as indicated below, with attendance limited to space available. The URL link to this meeting is https://videocast.nih.gov/. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the 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 Advisory Council for Nursing Research.

Date: May 24, 2022.

Open: 11:00 a.m. to 3:00 p.m. Agenda: Discussion of Program Policies and Issues.

Place: National Institute of Nursing Research, National Institutes of Health, 6701 Democracy Boulevard, One Democracy Plaza, Bethesda, MD 20892, https:// videocast.nih.gov/watch=45025 (Virtual Meeting).

Closed: 3:00 p.m. to 4:00 p.m.

Agenda: To review and evaluate review of applications.

*Place: National Institute of Nursing Research, National Institutes of Health, 6701 Democracy Boulevard, One Democracy Plaza, Bethesda, MD 20892, https:// videocast.nih.gov/watch=45025 (Virtual Meeting).

Contact Person: Dr. Elizabeth Tarlov, Ph.D., R.N., Director, Division of Extramural Science Programs, National Institute of Nursing Research/NIH, 6701 Democracy Blvd., Bethesda, MD 20892, (301) 594–1580, Elizabeth.tarlov@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: https://www.ninr.nih.gov/aboutninr/nacnr, where an agenda and any additional information for the meeting will be posted when available. (Catalogue of Federal Domestic Assistance Program Nos. 93.361, Nursing Research, National Institutes of Health, HHS)

Dated: April 18, 2022.

Victoria E. Townsend,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2022–08554 Filed 4–21–22; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Eye Institute; 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