

the expanded tool, NCI-AD, has been piloted in 3 states. Fourteen states are currently slated to begin using the tool in the summer of 2015. NASUAD is the lead partner for the NCI-AD project. This one-year of grant funding, through a continuation grant, will support NASUAD in their efforts to develop and perform comprehensive and rigorous validity/reliability testing; provide support for NCI-AD regional meetings (in conjunction with ACL regional meetings); engage in additional technical assistance including bi-monthly TA calls, continuous quality improvement activities, stakeholder engagement, survey customization, in-person interviewer training, in-person meeting with state staff, refresher webinars, and monthly update calls; recruit states to participate in the project; enhance person centered planning measures in the tool; sampling methodologies, outcome reports by state, quality improvement activities by state, and others.

This program is authorized under the Older Americans Act of 1965, as amended in 2006, Public Law 109-365 as well as Title II of the Rehabilitation Act of 1973, as amended (Pub. L. 113-128), and Title I, Subtitle E of the Developmental Disabilities Assistance and Bill of Rights Act of 2000, Public Law 106-402 (Developmental Disabilities Projects of National Significance).

Dated: August 12, 2015.

**Aaron Bishop,**

*Commissioner Administration for Intellectual and Developmental Disabilities,  
Administration Community Living.*

[FR Doc. 2015-20392 Filed 8-20-15; 8:45 am]

**BILLING CODE 4154-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Center for Scientific Review; Notice of Closed Meetings**

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), 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 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:* Cardiovascular and Respiratory Sciences Integrated Review Group; Respiratory Integrative Biology and Translational Research Study Section.

*Date:* October 1, 2015.

*Time:* 8:00 a.m. to 6:30 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Holiday Inn Riverwalk, 217 N. St. Mary's Street, San Antonio, TX 78205.

*Contact Person:* Bradley Nuss, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4142, MSC7814, Bethesda, MD 20892, 301-451-8754, [nussb@csr.nih.gov](mailto:nussb@csr.nih.gov).

*Name of Committee:* Musculoskeletal, Oral and Skin Sciences Integrated Review Group; Musculoskeletal Tissue Engineering Study Section.

*Date:* October 5, 2015.

*Time:* 8:00 a.m. to 5:30 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Pier 2620 Hotel, 2620 Jones Street, San Francisco, CA 94133.

*Contact Person:* Baljit S Moonga, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4214, MSC 7806, Bethesda, MD 20892, 301-435-1777, [moongabs@mail.nih.gov](mailto:moongabs@mail.nih.gov).

*Name of Committee:* Risk, Prevention and Health Behavior Integrated Review Group; Psychosocial Risk and Disease Prevention Study Section.

*Date:* October 5-6, 2015.

*Time:* 8:00 a.m. to 5:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Embassy Suites at the Chevy Chase Pavilion, 4300 Military Road NW., Washington, DC 20015.

*Contact Person:* Stacey FitzSimmons, Ph.D., MPH, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3114, MSC 7808, Bethesda, MD 20892, (301) 451-9956, [fitsimmonss@csr.nih.gov](mailto:fitsimmonss@csr.nih.gov).

*Name of Committee:* Digestive, Kidney and Urological Systems Integrated Review Group; Pathobiology of Kidney Disease Study Section.

*Date:* October 5-6, 2015.

*Time:* 8:00 a.m. to 6:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Gaithersburg Marriott Washingtonian Center, 9751 Washingtonian Boulevard, Gaithersburg, MD 20878.

*Contact Person:* Atul Sahai, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2188, MSC 7818, Bethesda, MD 20892, 301-435-1198, [sahaia@csr.nih.gov](mailto:sahaia@csr.nih.gov).

*Name of Committee:* Vascular and Hematology Integrated Review Group; Vascular Cell and Molecular Biology Study Section.

*Date:* October 5-6, 2015.

*Time:* 8:00 a.m. to 5:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Embassy Row Hotel, 2015 Massachusetts Avenue NW., Washington, DC 20036.

*Contact Person:* Larry Pinkus, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4132, MSC 7802, Bethesda, MD 20892, (301) 435-1214, [pinkusl@csr.nih.gov](mailto:pinkusl@csr.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: August 17, 2015.

**Melanie J. Gray,**

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

[FR Doc. 2015-20645 Filed 8-20-15; 8:45 am]

**BILLING CODE 4140-01-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 (5 U.S.C. App.), 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 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 contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* National Institute on Aging Special Emphasis Panel; Age-Induced Bone Loss.

*Date:* September 22, 2015.

*Time:* 12:00 p.m. to 4:00 p.m.

*Agenda:* To review and evaluate contract proposals.

*Place:* National Institution on Aging, Gateway Building, 2C212, 7201 Wisconsin Avenue, Bethesda, MD 20892, (Telephone Conference Call).

*Contact Person:* Isis S. Mikhail, MD, MPH, DRPH National Institute On Aging Gateway Building, 7201 Wisconsin Avenue Suite 2C212, Bethesda, MD 20892, 301-402-7704, [mikhaili@mail.nih.gov](mailto:mikhaili@mail.nih.gov).

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

Dated: August 17, 2015.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2015-20643 Filed 8-20-15; 8:45 am]

BILLING CODE 4140-01-P

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Submission for OMB Review; 30-Day Comment Request; Web-Based Resource for Youth About Clinical Research**

**SUMMARY:** Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on 3/12/2015 pages 13013-13014, and allowed 60-days for public comment. One public comment was received. The purpose of this notice is to allow an additional 30 days for public comment. The National Heart, Lung and Blood Institute (NHLBI), National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

*Direct Comments to OMB:* Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office

of Management and Budget, Office of Regulatory Affairs, *OIRA\_submission@omb.eop.gov* or by fax to 202-395-6974, Attention: NIH Desk Officer.

*Comment Due Date:* Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

**FOR FURTHER INFORMATION CONTACT:** To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Ms. Victoria Pemberton, National Heart, Lung, and Blood Institute, National Institutes of Health, 6701 Rockledge Dr., Room 8102, MSC 7940, Bethesda, MD 20892-7940, or call non-toll-free number 301-435-0510, or Email your request, including your address to *pembertonv@nhlbi.nih.gov*. Formal requests for additional plans and instruments must be requested in writing.

*Proposed Collection:* Web-based Resource for Youth about Clinical Research (NHLBI), 0925-New, National Heart, Lung, and Blood Institute (NHLBI), the National Institutes of Health (NIH).

*Need and Use of Information Collection:* The purpose and use of the information collection for this project is to develop a comprehensive web-based resource for youth with chronic illnesses or diseases that will attempt to increase knowledge, self-efficacy, and positive attitudes towards participation in various clinical trials and research. As a result of the proposed web-based resource, the knowledge gained from developing and testing this web-based resource will ultimately help equip youth to make informed decisions about clinical research and increase motivation to participate in that

research. In addition, the knowledge gained will be invaluable to the field of clinical research given the need for more clinical trials with youth. Specifically, the proposed web-based resource will be an interactive, multimedia, developmentally appropriate resource for youth to be educated about pediatric clinical trials. The resource will be developed for youth aged 8 to 14 years. The theme of “investigative cyber-reporting” will be used throughout and will include youth making a series of decisions about different aspects of participating in clinical research studies. Youth will be tasked with the responsibility of learning all they can about clinical research trials in order to facilitate their knowledge and decision-making processes. Language typically used in journalism and design elements reminiscent of journalism will be incorporated into the content, design, and layout of the resource. There are three main components that will comprise the web-based resource. These include an interactive leaning module, full length video testimonials, and an electronic comic book. The benefits and necessities for this particular research on pediatric clinical trials are congruent with NHLBI’s research goals and mission statement: Attempting to assist in the enhancement of the health of individuals so that they can live longer and more fulfilling lives. The current lack of knowledge surrounding pediatric clinical trials can be dangerous and unhealthy towards the lives of youth, becoming a large public health need.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 172.

**ESTIMATES OF HOUR BURDEN**

Form name	Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hour
Individual Interview Parent Permission Form .....	parents .....	9	1	5/60	1
One-to-One Evaluation Study Parent Permission Form .....	parents .....	5	1	5/60	0.42
Pre-Post Feedback Study Parent Permission .....	parents .....	34	1	5/60	3
Individual Interview Child Assent Form .....	youth .....	9	1	5/60	1
One-to-One Evaluation Study Child Assent Form .....	youth .....	5	1	5/60	0.42
Pre-Post Feedback Study Child Assent Form .....	youth .....	34	1	5/60	3
Individual Interview Questions (Feature Stories) .....	youth .....	3	1	2	6
Individual Interview Questions (Family Spotlights) .....	youth .....	3	1	2	6
Individual Interview Questions (Comic Book) .....	youth .....	3	1	2	6
One-to-One Evaluation Study Questionnaire .....	youth .....	5	1	2	10
Pre-Post Feedback Study Questionnaire .....	youth .....	34	1	4	136