

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 or request more information on the proposed project contact: Jackie Lavigne, Office of Education, Division of Cancer Epidemiology and Genetics, 9609 Medical Center Drive, MSC 9776, Bethesda, MD 20892-9776 or call non-toll-free number 240-376-7237 or Email your request, including your address to: [lavignej@mail.nih.gov](mailto:lavignej@mail.nih.gov). Formal requests for additional plans and instruments must be requested in writing.

*Proposed Collection:* Division of Cancer Epidemiology and Genetics (DCEG) Fellowship Program and Summer Student Applications (NCI), Existing Collection in Use without OMB Control Number, National Cancer Institute (NCI), National Institutes of Health (NIH).

*Need and Use of Information Collection:* The Division of Cancer Epidemiology and Genetics (DCEG) Office of Education (OE) administers a variety of programs and initiatives to recruit pre-college through post-doctoral educational level individuals into the Intramural Research Program to facilitate their development into future biomedical scientists. DCEG trains post-doctoral, doctoral candidates, graduate and baccalaureate students, through full time fellowships, summer fellowships, and internships in preparation for research careers in cancer epidemiology and genetics. The proposed information collection involves brief online applications completed by applicants to the full time and the summer fellowship programs. Full-time fellowships include: Full-time Equivalents (FTE) and non-FTE fellowships for US citizens, permanent residents and international fellows. These

applications are essential to the administration of these training programs as they enable OE to determine the eligibility and quality of potential awardees; to assess their potential as future scientists; to determine where mutual research interests exist; and to make decisions regarding which applicants will be proposed and approved for traineeship awards. In each case, completing the application is voluntary, but in order to receive due consideration, the prospective trainee is encouraged to complete all relevant fields. The information is for internal use to make decisions about prospective fellows and students that could benefit from the DCEG program.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 175.

**ESTIMATED ANNUALIZED BURDEN HOURS**

Form name	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
Fellowship Program Application .....	Full-time Fellows .....	150	1	30/60	75
Summer Program Application .....	Summer Students .....	300	1	20/60	100

Dated: February 19, 2015.

**Karla Bailey,**

*NCI Project Clearance Liaison, National Institutes of Health.*

[FR Doc. 2015-03789 Filed 2-24-15; 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; Assessment of Oncology Nursing Education and Training in Low and Middle Income Countries (NCI)**

**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 July 8, 2014, Vol. 79, page 38542 and allowed 60-days for public comment. One public comment was received on July 9, 2014. The purpose of this notice is to allow an additional 30 days for public comment.

The National Cancer Institute (NCI), 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](mailto: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 or request more information on the proposed project, contact: Annette Galassi, Center for Global Health, National Cancer Institute, 9609 Medical Center Dr., Rm. 3W250, Rockville, MD 20850 or call non-toll-free number 240-276-6632 or Email your request, including your address to:

[agalassi@mail.nih.gov](mailto:agalassi@mail.nih.gov). Formal requests for additional plans and instruments must be requested in writing.

*Proposed Collection:* Assessment of Oncology Nursing Education and Training in Low and Middle Income Countries, 0925-NEW, National Cancer Institute (NCI), National Institutes of Health (NIH).

*Need and Use of Information Collection:* This submission is a request for OMB to approve the Assessment of Oncology Nursing Education and Training in Low and Middle Income Countries (LMICs). NCI-Designated Cancer Centers have a range of international activities, some of which are funded by NCI, but many of which are not. These international activities may include oncology nursing education and training in LMICs, but the extent of these activities across cancer centers is unknown. The proposed assessment requests information about oncology nursing education and training projects including: descriptions of projects, partner organizations, types of activities, cost, and impact. The information will be collected annually. NCI's Center for Global Health (CGH) is in the process of developing its strategic plan for oncology nursing education in LMICs.

This information will help inform this strategic planning process and provide evidence to inform decisions on potential investments in grants for oncology nursing education in LMICs. Additionally, this information will be used in an online, interactive map that is being developed by CGH which will

allow external organizations, such as cancer centers, to explore what projects are being done in which countries, which will facilitate collaborations and minimize duplication. The frequency of the data collection will be once per year although respondents may have more

than one response if they have up to three projects.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 51.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Number of respondents/year	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
Directors of Nursing .....	68	3	15/60	51

Dated: February 19, 2015.

**Karla Bailey,**

*NCI Project Clearance Liaison, National Institutes of Health.*

[FR Doc. 2015-03788 Filed 2-24-15; 8:45 am]

**BILLING CODE 4140-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Government-Owned Inventions; Availability for Licensing**

**AGENCY:** National Institutes of Health, HHS.

**ACTION:** Notice.

**SUMMARY:** The inventions listed below are owned by an agency of the U.S. Government and are available for licensing in the U.S. in accordance with 35 U.S.C. 209 and 37 CFR part 404 to achieve expeditious commercialization of results of federally-funded research and development. Foreign patent applications are filed on selected inventions to extend market coverage for companies and may also be available for licensing.

**FOR FURTHER INFORMATION CONTACT:** Licensing information and copies of the U.S. patent applications listed below may be obtained by writing to the indicated licensing contact at the Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, Maryland 20852-3804; telephone: 301-496-7057; fax: 301-402-0220. A signed Confidential Disclosure Agreement will be required to receive copies of the patent applications.

**SUPPLEMENTARY INFORMATION:** Technology descriptions follow.

**HbF Induction Therapy for Sickle Cell Disease and Thalassemias**

Description of Technology: Sickle cell disease and thalassemia are hereditary

disorders marked by the disruption in the pathways responsible for carrying oxygen to red blood cells. Symptoms associated with these disorders include anemia, jaundice, and severe pain. It has been shown that mutations during the development of fetal to adult hemoglobin can contribute to a delay in red blood cell maturity underlying sickle cell disease. As a result, there has been an increased focus on treatments that promote the induction of fetal hemoglobin (HbF) to improve clinical symptoms and ameliorate the severity of the diseases. Researchers at the National Institute of Diabetes and Digestive and Kidney Diseases have identified methods of increasing fetal hemoglobin by increasing the expression of Lin28 or decreased expression of let-7 micro-RNAs. The lead inventor and colleagues have developed novel lentiviral expression vectors containing hemoglobin regulators under the control of erythroid-specific promoters that can be used to increase HbF expression without affecting the maturity of red blood cells. In addition, they have found, through the use of tough decoy inhibition of Let-7 micro-RNAs, a selection of Let-7 genes with greater involvement in HbF expression. This technology could lead to development of novel HbF induction therapies that reactivate and reduce the aberrant pathologies associated with human sickle-cell anemia and beta thalassemia.

**Potential Commercial Applications:**

- Ex vivo and in vivo therapeutics for treatment of sickle-cell anemia and beta thalassemias.
  - Potential use in combination with other transduction methods for unique therapeutic strategies.
- Competitive Advantages:**
- Reduced production of symptom-associated adult hemoglobin.
  - Lin28 overexpression at defined stage of hematopoietic cell development.

• Therapeutic increases in patient HbF expression at lower viral titers than current direct transduction methods.

• Improved safety and reduced toxicity as a result of erythroid-specific expression.

**Development Stage:**

- Early-stage
- In vitro data available
- In vivo data available (animal)

**Inventors:** Jeffery L. Miller, Yuanwei T. Lee, Jaira F. de Vasconcellos, Colleen K. Byrnes (all of NIDDK)

**Intellectual Property:** HHS Reference No. E-249-2014/0—US Provisional Application No. 62/046,247 filed September 5, 2014

**Related Technology:** HHS Reference No. E-456-2013/2—PCT Application No. PCT/US2013/067811 filed October 31, 2013, which published as WO 2014/200557 on December 18, 2014

**Licensing Contact:** Vince Contreras, Ph.D.; 301-435-4711; [contrerasv@mail.nih.gov](mailto:contrerasv@mail.nih.gov)

**Collaborative Research Opportunity:** The National Institute of Diabetes and Digestive and Kidney Diseases is seeking statements of capability or interest from parties interested in collaborative research to further develop, evaluate or commercialize this technology. For collaboration opportunities, please contact Marguerite J. Miller at [miller marg@nid dk.nih.gov](mailto:miller marg@nid dk.nih.gov) or 301-496-9003.

**T Cell-Based Adoptive Transfer Immunotherapy for Polyomavirus-Associated Pathologies**

Description of Technology: Available for licensing are methods to generate T cells responsive to multiple polyomaviruses. The resulting T cell populations could be useful in treating immunosuppressed individuals with polyomavirus infections or polyomavirus-associated pathologies such as Merkel cell carcinoma (MCC), polyomavirus-associated nephropathy (PVAN), hemorrhagic cystitis,