

ESTIMATED ANNUALIZED BURDEN TABLE

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
State, local, and territory registration officials (Annual Survey of Training Needs)	57	1	20/60
Training applicants (Application for Training)	100	1	15/60

Dated: March 23, 2010.

Maryam I. Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. 2010-7065 Filed 3-29-10; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; Process Evaluation of the NIH's Roadmap Interdisciplinary Research Work Group Initiatives

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the (insert name of NIH Institute or IC), the National Institutes of Health has submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. This proposed information collection was previously published in the **Federal Register** on January 5, 2010 (p. 382) and allowed 60 days for public comment. One comment was received, which included a request for additional information, and additional information was provided. No additional questions were received. The purpose of this notice is to allow an additional 30 days for public comment. The 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.

Proposed Collection: The National Institute of Dental and Craniofacial Research of the National Institutes of Health requests a two-year clearance for Title: "Process Evaluation of the NIH Roadmap Interdisciplinary Research Work Group Initiatives," *Type of information collection:* New. *Need and use of information collection:* This study will be used to determine whether the NIH's Interdisciplinary Research Work Group initiatives have been, and are being, conducted as planned, whether the expected outputs are being produced, and how the activities and

processes associated with the initiatives can be improved. Information collected during the evaluation will be used to assess whether and how these initiatives differed from existing initiatives to determine whether these unique initiatives or mechanisms are necessary, to make decisions about whether to continue and/or to modify the programs, and to make decisions about structural or procedural changes within NIH that may be necessary to support cross-cutting interdisciplinary programs.

Frequency of response: The frequency of response is once for most respondents, and twice for a limited group. **Affected public:** The affected public includes a limited number of individuals; **Type of respondents:** principal investigators, other grant investigators, and Initiative trainees. The annual reporting burden is as follows: **Estimated number of respondents:** 450; **Estimated number of responses per respondent:** PIs, 2; Other Investigators, 1; Trainees, 1; **Average burden hours per response:** 30 minutes; and **Estimated total annual burden hours requested:** 250 hours. The total annualized cost to respondents (calculated as the number of respondents * frequency of response * average time per response * approximate hourly wage rate) is estimated to be \$7,450. There are no Capital Costs, Operating Costs, and/or Maintenance Costs to report.

Request for Comments: Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological

collection techniques or other forms of information technology.

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: Desk Officer for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Sue Hamann, PhD, Science Evaluation Officer, Office of Science Policy Officer and Analysis, National Institute of Dental and Craniofacial Research (NIDCR), NIH. You may reach Dr. Hamann by telephone on 301-594-4849 (this is not a toll-free number), or you may e-mail your request to Dr. Hamann at Sue.Hamann@nih.hhs.gov.

Comments 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.

Dated: March 24, 2010.

Sue Hamann,

Science Evaluation Officer, OSPA, NIDCR, National Institutes of Health.

[FR Doc. 2010-7086 Filed 3-29-10; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: OCSE-75 Tribal Child Support Enforcement Program Annual Data Report.

OMB No.: 0970-0320.

Description: The data collected by form OCSE-75 are used to prepare the OCSE preliminary and annual data reports. In addition, Tribes administering CSE programs under Title IV-D of the Social Security Act are required to report program status and accomplishments in an annual narrative

report and submit the OCSE-75 report annually.

Respondents: Tribal Child Support Enforcement Organizations or the Department/Agency/Bureau responsible

for Child Support Enforcement in each Tribe.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
OCSE-75	37	1	60	2,220

Estimated Total Annual Burden Hours: 2,220.

Additional Information

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. E-mail address: infocollection@acf.hhs.gov.

OMB Comment

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following:

Office of Management and Budget,
Paperwork Reduction Project. Fax:
202-395-7285. E-mail:
OIRA_SUBMISSION@OMB.EOP.GOV.
Attn: Desk Officer for the
Administration for Children and
Families.

Dated: March 25, 2010.

Robert Sargis,
Reports Clearance Officer.

[FR Doc. 2010-7042 Filed 3-29-10; 8:45 am]

BILLING CODE 4184-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Government-Owned Inventions; Availability for Licensing

AGENCY: National Institutes of Health, Public Health Service, 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. 207 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.

ADDRESSES: 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.

Zscan4, a Therapeutic Target for Cancer, Regenerative Medicine and Aging

Description of Invention: This technology has broad potential for the development of therapeutics for cancer, diseases of aging, and regenerative medicine, and targets Zscan4, a gene that regulates telomere length and genomic stability in embryonic stem (ES) cells.

The ability to maintain genomic stability in ES cells and other stem cells is critical for the development of stem cell-based therapies; genomic stability and telomere length are also active areas of cancer and aging research. NIA investigators have discovered that the Zscan4 gene regulates telomere length and genomic stability in ES cells, and plays an essential role in early embryonic development; this activity is independent of telomerase activity. The investigators have shown that ablation of Zscan4 results in shortened telomere length and deterioration of the karyotype of ES cells, and that Zscan4 overexpression increases telomere length.

This technology discloses methods for increasing genome stability or increasing telomere length in an ES cell,

and methods of treating a subject in need of ES cell therapy. Also disclosed are methods of promoting blastocyst outgrowth of embryonic stem cells, as well as Zscan4 expression vectors and methods of identifying stem cells expressing Zscan4.

Applications

- Development of therapeutics for cancer treatment, aging, and regenerative medicine.
- Development of assisted reproduction technologies.
- Studies of early embryonic development.

Development Status: In vitro and in vivo studies have been performed.

Inventors: Minoru S. H. Ko *et al.* (NIA).

Publications

1. M Zalzman *et al.* Zscan4 regulates telomere elongation and genomic stability in ES cells. *Nature* 2010 Mar 24; advance online publication, doi 10.1038/nature08882.

2. G Falco *et al.* Zscan4: A novel gene expressed exclusively in late 2-cell embryos and embryonic stem cells. *Dev Biol.* 2007 Jul 15;307(2):539-550. [PubMed: 17553482.]

Patent Status

- HHS Reference No. E-088-2007/0—PCT Application No. PCT/US2008/058261 filed 26 Mar 2008.
- US Application No. 12/529,004 filed 27 Aug 2009.
- Foreign counterparts in Europe, Australia, Canada, and Japan
- HHS Reference No. E-172-2009/0—U.S. Provisional Application No. 61/275,983 filed 04 Sep 2009.

Licensing Status: Available for licensing.

Licensing Contact: Tara Kirby, PhD; 301-435-4426; tarak@mail.nih.gov.

Collaborative Research Opportunity: The National Institute on Aging, Laboratory of Genetics, is seeking statements of capability or interest from parties interested in collaborative research to further develop, evaluate, or commercialize this technology. Please contact Nicole Guyton, PhD at 301-435-