been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Proposed Collection: Title: STAR METRICS (Science and Technology for America's Reinvestment: Measuring the EffecTs of Research on Innovation, Competitiveness and Science). *Type of information Collection Request:* Extension of OMB number 0925–0616, expiration date 03/31/2012. *Need and Use of Information Collection:* The aim of STAR METRICS is twofold. The goal of STAR METRICS is to continue to provide mechanisms that will allow participating universities and Federal agencies with a reliable and consistent means to account for the number of scientists and staff that are on research institution payrolls, supported by federal funds. In subsequent generations of the program, it is hoped that STAR METRICS will allow for measurement of science impact on economic outcomes (such as job creation), on knowledge generation (such as citations, and patents) as well as on social and health outcomes.

Frequency of Response: Quarterly. *Affected Public:* Universities and other research institutions. *Type of Respondents:* University administrators. The annual reporting burden is as follows:

Estimated Number of Respondent: 100. *Estimated Number of Responses per Respondent:* 4. *Average Burden Hours per Response:* 2.5. *Estimated Total Annual Burden Hours Requested:* 1,315. The annualized cost to respondents is estimated to be $65,750. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

A.12–1—ESTIMATES OF ANNUAL BURDEN HOURS

	Number of respondents	Frequency of response	Average Time per response (in hours)	Annual hour burden
Stage I: One time data input	7	1	45	315
Stage 2: Ongoing quarterly data input	100	4	2.5	1000
Total				1315

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the functioning of the National Cancer Institute, including whether the information will have practical utility; (2) 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) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to 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 Attention: NIH Desk Officer, Office of Management and Budget, at *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: George Chacko, Office of Planning, Analysis, and Evaluation, Center for Scientific Review, 6701 Rockledge Drive, Suite 3030, Bethesda, MD 20892 or call non-toll-free at 301–435–1111 or email your request, including your address to: *chackoge@mail.nih.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: February 20, 2012.

George Chacko,

Center for Scientific Review, National Institutes of Health.

[FR Doc. 2012–4536 Filed 2–24–12; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request DERT Extramural Grantee Data Collection

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute of Environmental Health Sciences (NIEHS), 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**, Vol. 76, No. 202, on Wednesday, October 19, 2011, page 64954 and allowed 60 days for public comment. No public comments 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: Title: DERT Extramural Grantee Data Collection. *Type of Information Collection Request:* New. *Need and Use of Information Collection:* In order to make informed management decisions about its research programs and to demonstrate the outputs, outcomes and impacts of its research programs NIEHS will collect, analyze and report on data from extramural grantees who are currently receiving funding or who have received funding in the past on topics such as:

- Key scientific outcomes achieved through the research and the impact on the field of environmental health science.
- Contribution of research findings to program goals and objectives.
- Satisfaction with the program support received.
- Challenges and benefits of the funding mechanism used to support the science.
- Emerging research areas and gaps in the research.

Information gained from this primary data collection will be used in conjunction with data from grantee progress reports and presentations at grantee meetings to inform internal programs and new funding initiatives.

Outcome information to be collected includes measures of agency-funded research resulting in dissemination of findings, investigator career development, grant-funded knowledge and products, commercial products and drugs, laws, regulations and standards, guidelines and recommendations, information on patents and new drug applications and community outreach and public awareness relevant to extramural research funding and emerging areas of research. Satisfaction information to be collected includes measures of satisfaction with the type of funding or program management mechanism used, challenges and benefits with the program support received, and gaps in the research. *Frequency of Response:* Once per grantee, per NIEHS research portfolio. *Affected Public:* Current or past NIEHS grantees. *Type of Respondents:* Principal Investigators with current or past NIEHS research or training grants. The annual reporting burden is as follows: *Estimated Number of Respondents:* 600; *Estimated Number of Responses per Respondent:* 1; *Average Burden Hours per Response:* .5 (30 minutes); and *Estimated Total Annual Burden Hours Requested:* 100. The annualized cost to respondents is estimated at: Approximately $17. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

(Note: The following table is acceptable for the Respondent and Burden Estimate information, if appropriate, instead of the text as shown above.)

Type of respondents	Estimated number of respondents	Estimated number of responses per respondent	Average burden hours per response (min.)	Estimated total annual burden hours requested
NIEHS Grantee	600	1	30	100

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) 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) 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) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to 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 Dr. Kristianna Pettibone, Evaluator, Program Analysis Branch, NIEHS, NIH, 530 Davis Dr., Room 3055, Morrisville, NC 20560, or call non-toll-free number 919–541–7752 or email your request, including your address to: *pettibonekg@niehs.nih.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: February 16, 2012.

Joellen M. Austin,

Associate Director for Management, NIEHS, National Institutes of Health.

[FR Doc. 2012–4543 Filed 2–24–12; 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, 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.

Model Cell Lines With and Without AKT1 Mutations Derived From Proteus Syndrome Patients

Description of Technology: The Proteus syndrome is a congenital disorder characterized by patchy overgrowth and hyperplasia (cell proliferation) of multiple tissues and organs, along with susceptibility to developing tumors. It is a rare disorder, with incidence of less than one case per million, caused by a somatic mutation. It is also a mosaic disorder, that is one in which cells of the same person have different genetic content from one another. The NHGRI inventors have generated cell lines from patients with Proteus syndrome and discovered that a somatic activating mutation in the serine-threonine kinase AKT1 is associated with Proteus syndrome. AKT1 is an oncogene and an enzyme known to mediate cell proliferation and apoptosis (programmed cell death process) and has been a target for anti-cancer therapies. A number of single-cell lines with the AKT1 mutation showing increased AKT1 phosphorylation and their matched controls without the mutation have been generated. The cell lines can be used to screen therapeutic targets for AKT1, for study design, as models of Proteus syndrome and early stages of cancerous conditions.

Potential Commercial Applications

- Cell lines generated from patients with Proteus syndrome.
- Obtained a number of single-cell lines with the AKT1 mutation and their matched controls without the mutation.