TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Form 1: Demographic, Service Utilization, and Clinical Indicators Data	56	1	56	425	23,800
Form 2: Performance and System Outcome Benchmark Data	56	1	56	425	23,800
Total	56		56		47,600

Jackie Painter,

Director, Division of the Executive Secretariat.

[FR Doc. 2015–31936 Filed 12–18–15; 8:45 am]

BILLING CODE 4165–15–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 emailing the indicated licensing contact at the National Heart, Lung, and Blood, Office of Technology Transfer and Development Office of Technology Transfer, 31 Center Drive Room 4A29, MSC2479, Bethesda, MD 20892–2479; telephone: 301–402–5579. A signed Confidential Disclosure Agreement may be required to receive copies of the patent applications.

SUPPLEMENTARY INFORMATION: Technology description follows.

Fluorescent Nanodiamonds as Fiducial Markers for Microscopy

Description of Technology

The invention relates to fluorescent nanodiamonds (FNDs) and their uses as fiducial markers for microscopy. FNDs are bright fluorescent probes that do not blink or bleach and have broad fluorescence excitation and emission peaks. The fluorescence intensity can be readily controlled by the size of the FND, the number of fluorescent centers produced in the nanodiamonds, or *in situ* through the application of a weak magnetic field. The particular advantage of the FND compositions of this invention are that they are particularly useful for extended imaging of a single sample over time periods that can be as long as a week or more. In an exemplary embodiment, FNDs are immobilized in a substrate that are coated with an inert top coating, like silicon dioxide, or transparent polymer (*e.g.* poly-L-lysine, poly-L-arginine, or siloxanes). Generally, any suitable methods known for surface functionalization of the substrate can be used to make the composition. In another aspect of this invention, the inventors designed software for super-resolution imaging correction method is employed to precisely determine the position coordinates of each of a set of FNDs in a plurality of images by using Gaussian fitting of the point spread function comprises each of the FNDs in the plurality of images. The calculated correction is then used to displace each image to align the coordinates of the FNDs. The positions of the FNDs can be tracked with sub-nanometer precision and residual drift can be reduced to the nanometer scale over hundreds of hours of tracking.

Potential Commercial Applications

- Fluorescent Microscopy
- Super-resolution microscopy
- Correlative imaging techniques combing fluorescence microscopy with electron, x-ray, or atomic force microscopy imaging modalities

Competitive Advantages

- Non-blinking, Non-bleaching
- Chemically inert
- Chemically and physically stable
- Broad excitation
- Longevity

Development Stage

- In vitro data

Inventors

- Keir Neuman, Ambika Bumb, Han Wen, Jennifer Hong and Susanta Sarkar (all of NHLBI)
- Chang Yi, Lawrence Samelson, Asit Manna (all of NCI)

Intellectual Property: HHS Reference No. E–217–2015/0–US–01

- US Provisional Patent Application 62/262,058 filed December 2, 2015.

Licensing Contact: Michael Shmilovich, Esq, CLP; 301–435–5019; *shmilovm@mail.nih.gov.*

Collaborative Research Opportunity: The National Heart, Lung and Blood Institute seeks statements of capability or interest from parties interested in collaborative research to further develop and evaluate metallic nanoparticle vesicles for cancer phototherapy. For collaboration opportunities, please contact Vincent Kolesnitchenko, Ph.D. at *kolesniv@nhlbi.nih.gov*.

Dated: December 15, 2015.

Michael Shmilovich,

Senior Licensing and Patenting Manager, National Heart, Lung, and Blood Institute, Office of Technology Transfer and Development.

[FR Doc. 2015–31890 Filed 12–18–15; 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; Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (NIAID)

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health, 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 October 1, 2015, 80 FR 59168 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 Institute of Allergy and Infectious Diseases (NIAID), 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.

DATES: 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: Ms. Dione Washington, Health Science Policy Analyst, Office of Strategic Planning, Initiative Development and Analysis, 5601 Fishers Lane, Rockville, Maryland 20892, or call a non-toll-free number 240 669 2100 or Email your request, including your address to *washingtondi@niaid.nih.gov.* Formal requests for additional plans and instruments must be requested in writing. Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (NIAID), 0925–0668, Expiration Date 1/31/2016, EXTENSION, National Institute of Allergy and Infectious Diseases (NIAID).

Need and Use of Information Collection: There are no changes being requested for this submission. The proposed information collection activity provides a means to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study. This feedback will provide information about the NIAID's customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the NIAID and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 16,100.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of collection	Number of respondents	Annual frequency per response	Hours per response	Total hours
Estimated Annual Reporting Burden				
Customer satisfaction surveys	25,000	1	30/60	12,500
In-Depth Interviews (IDIs) or Small Discussion Groups	500	1	90/60	750
Individual Brief Interviews	200	1	15/60	50
Focus Groups	1,000	1	2	2,000
Pilot testing surveys	200	1	30/60	100
Conferences and Training Pre- and Post-surveys	1,000	1	30/60	500
Website or Software Usability Tests	100	1	2	200
Total	28,000			16,100

Dated: December 15, 2015.

Brandie Taylor Bumgardner,

Project Clearance Liaison, NIAID, NIH.

[FR Doc. 2015–31986 Filed 12–18–15; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning opportunity for public comment on proposed collections of information, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information collection plans, call the SAMHSA Reports Clearance Officer on (240) 276–1243.

Comments are invited on: (a) Whether the proposed collections of information are necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

Proposed Project: Community Support Evaluation (CSE)—New

The Substance Abuse and Mental Health Services Administration (SAMHSA), Center for Mental Health Services (CMHS), is requesting clearance for the new data collection associated with the CSE. The CSE is a multicomponent evaluation of two SAMHSA programs—Behavioral Health Treatment Court Collaborative (BHTCC) and Transforming Lives through Supported Employment (SE). SE intends to promote recovery for individuals with serious mental illness, substance