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 or request more information on the proposed project contact: Anthony Dickherber, NCI Center for Strategic Scientific Initiatives, 31 Center Drive, Rm10A33, Bethesda, MD 20892 or call non-toll-free number 301–547–9980 or Email your request, including your address to: *dickherberaj@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: Surveys and Interviews to Support an Evaluation of the Innovative Molecular Analysis Technologies (IMAT) Program (NCI), 0925–NEW, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: The purpose of the proposed evaluation is to pursue a comprehensive process and outcome assessment of the 15-year old Innovative Molecular Analysis Technologies (IMAT) program. While the program consistently offers promising indicators of success, the full program has not been evaluated since 2008, and never in as comprehensive a manner as has been formulated in the current evaluation plan. An outcome evaluation of the long-standing National Cancer Institute's (NCI) IMAT program presents a rich and unique opportunity likely to serve institutes across the National Institutes of Health (NIH), and perhaps other federal agencies, considering the costs and benefits of directing resources towards supporting technology development. An award through the NIH Evaluation Set-Aside program to support this evaluation, for which NIH-wide relevance is a principle element of determining merit for support, is testament to this. The evaluation serves as an opportunity to gauge the impact of investments in technology development and also to assess the strengths and weaknesses of phased innovation award mechanisms.

Like all institutes and centers (ICs) of the NIH, NCI seeks opportunities for improving their programs' utility for the broad continuum of researchers, clinicians and ultimately patients. NCI Director Harold Varmus and other leadership across NCI, as well as the NCI Board of Scientific Advisors, will be the primary users of the evaluation results. Findings are primarily intended for considering the long-term strategy to support innovative technology development and how to more efficiently translate emerging capabilities through such technologies into the promised benefits for cancer research and clinical care. Interviews with grantees, program officers, review officers, and other NIH awardees make up a crucial component of the evaluation plan and will largely follow set survey protocols. Specific near-term aims include the use of this information to consider the utility of continued investment through existing solicitations and in strategic planning generally for institute support for innovative technology development.

OMB approval is requested for 1 year. There are no costs to respondents other than their time. The total estimated annualized burden hours are 575.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden (in hours)
IMAT Awardee Interview	IMAT Awardees	100	1	1	100
Evaluation Web-based Survey	IMAT Applicants and Other NIH Awardees	900	1	30/60	450
Tech End Users Interview	Technology End-Users	50	1	30/60	25

Dated: February 23, 2015.

Karla Bailey,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2015–05298 Filed 3–6–15; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10555]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by May 8, 2015.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or

Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number _____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web site address at *http://www.cms.hhs.gov/PaperworkReductionActof1995*.

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov*.

3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: Reports Clearance Office at (410) 786–1326.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS–10555 Small Business Health Options Program (SHOP) Effective Date and Termination Notice Requirements

Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collection

1. *Type of Information Collection Request:* New collection (Request for a new OMB control number); *Title of Information Collection:* Small Business Health Options Program (SHOP) Effective Date and Termination Notice Requirements; *Use:* We are requiring that for plan years beginning on or after January 1, 2017, the Small Business Health Options Program (SHOP) must ensure that a qualified health plan (QHP) issuer notifies qualified employees, enrollees, and new enrollees in a QHP through the SHOP of the effective date of coverage. As required by the Patient Protection and Affordable Care Act; HHS Notice of Benefit and Payment Parameter for 2016 (CMS–9944–F), which published February 27, 2015, if any enrollee's coverage through the SHOP is terminated due to non-payment of premiums or a loss of the enrollee's or employer group's eligibility to participate in the SHOP, the SHOP must notify the enrollee or the qualified employer of the termination of such coverage. In the termination of coverage the SHOP must include the termination date and reason for termination to the enrollee or qualified employer. *Form Number:* CMS–10555 (OMB Control Number: 0938–New); *Frequency:* Annually; *Affected Public:* Private sector (Business or other for profits and Not-for-profit institutions); *Number of Respondents:* 445; *Total Annual Responses:* 1,335; *Total Annual Hours:* 46,725. (For policy questions regarding this collection contact Christelle Jang at (410) 786–8438).

Dated: March 4, 2015.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2015–05420 Filed 3–6–15; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Committee on Organ Transplantation (ACOT)

AGENCY: Health Resources and Services Administration (HRSA), HHS.

ACTION: Notice; correction.

SUMMARY: The Health Resources and Services Administration published a notice in the **Federal Register**, FR 2015–03929 (February 26, 2015), announcing the meeting for the Advisory Committee on Organ Transplantation (ACOT). This action is to add dial-in information.

Correction: In the **Federal Register**, FR 2015–03929 (February 26, 2015), please make the following addition:

Participants can also join this meeting via conference call by calling the toll-free phone number 888–324–4391 and providing the participant pass code 7744447.

Jackie Painter,

Director, Division of the Executive Secretariat.

[FR Doc. 2015–05415 Filed 3–6–15; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases; 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: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; Collaborative Interdisciplinary Team Science in NIDDK Research Areas (R24)–Hematopoiesis.

Date: March 20, 2015.

Time: 2:00 p.m. to 3:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Paul A. Rushing, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 747, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–8895, *rushingp@extra.niddk.nih.gov*.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; RFA–DK14–011 Chronic Kidney Disease Biomarkers Consortium (CKD BioCon) (U01).

Date: March 24, 2015.

Time: 11:30 a.m. to 3:30 p.m.