

Staff (MOS), coordinates the activities of the ACF Native American Affairs Liaison Workgroup, serves as the ANA liaison to the Inter-Departmental Council on Native American Affairs, provides coordination of ANA's data driven strategic plan, and advises the Commissioner on strategic and operational activities of ANA.

The MOS provides administrative and budget support to ANA. These responsibilities include: (1) Serving as the Executive Secretariat for ANA, including managing correspondence, correspondence systems, and public requests including, but not limited to Freedom of Information Act (FOIA) requests; (2) coordinating human resources activities; (3) developing and executing the budget; (4) providing on-going administrative technical support of ANA contracts; and (5) as appropriate, developing internal policies and procedures relating to these activities.

B. The Division of Program Operations is primarily responsible for the pre-award and post-award administration of discretionary grant programs to eligible Tribes and non-profit Native American organizations. These responsibilities include: (1) Developing ANA's Funding Opportunity Announcements; (2) managing annual grant competitions, including coordination of the panel review process and internal application review; (3) on-going grantee monitoring and support; (4) administering grant award portfolio, including close-out; and (5) providing liaison to the Office of Administration, Divisions of Grants Management and Division of Grants Policy.

C. The Division of Program Evaluation and Planning is responsible for evaluations of grantee effectiveness and impact as well as ANA performance including, but not limited to Government Performance and Results Act measures. These responsibilities include: (1) Oversight of planning and implementation activities related to ANA program evaluation, including development of annual reports, which includes the annual Report to Congress on Impact and Effectiveness; (2) data analyses and special organizational studies to guide programmatic enhancements and inform training and technical assistance efforts; (3) coordination of pre-award and post-award training and technical assistance activities in Alaska, the Pacific Basin, and the lower forty-eight states; (4) coordination of ANA's data and social media tools; and (5) liaison with the Office of Planning, Research and Evaluation.

D. The Division of Policy is responsible for providing support and guidance to define, establish, and disseminate policy affecting Native American communities at large. These responsibilities include: (1) Management of a unified and effective policy formulation process, including coordination of ANA regulations and other policy issuances affecting grantees and Native American communities; (2) formulation of advice on tribal and state legislative, and other activities affecting Native Americans; (3) development of long- and short-term strategies to address issues raised in consultations and collaborations with Native Americans; (4) support of the activities of the ACF Tribal Advisory Committee and other committees, councils, and workgroups affecting Native Americans; and (5) liaison with the Office of the General Counsel and others in the Department on matters involving or affecting Native Americans.

II. Continuation of Policy. Except as inconsistent with this reorganization, all statements of policy and interpretations with respect to organizational components affected by this notice within ACF, heretofore issued and in effect on this date of this reorganization are continued in full force and effect.

III. Delegation of Authority. All delegations and redelegations of authority made to officials and employees of affected organizational components will continue in them or their successors pending further redelegations, provided they are consistent with this reorganization.

IV. Funds, Personnel, and Equipment. Transfer of organizations and functions affected by this reorganization shall be accompanied in each instance by direct and support funds, positions, personnel, records, equipment, supplies, and other resources.

This reorganization will be effective upon date of signature.

Dated: November 24, 2014.

Mark H. Greenberg,

Acting Assistant Secretary for Children and Families.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; Health Information National Trends Survey (HINTS)

Summary: In compliance with the requirement of Section 3506(c)(2)(A) of

the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI), National Institutes of Health (NIH), will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

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.

To Submit Comments and For Further Information: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project contact: Bradford W. Hesse, Ph.D., Health Communication and Informatics Research Branch, 9609 Medical Center Drive, MSC 9761, Room 3E610, Rockville, MD 20850 or call non-toll free number 240-276-6721 or Email your request, including your address, to hessb@mail.nih.gov. Formal requests for additional plans and instruments must be requested in writing.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Proposed Collection: Health Information National Trends Survey (HINTS) 0925-0538, Reinstatement with Change, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: This partnership between NCI and FDA will include assessing the public's knowledge of medical devices, communications related to product recalls, nutritional supplement labeling, and topics to inform FDA's regulatory authority over tobacco, such as risk perceptions about new tobacco products, product pack color gradations, perceptions of product harm, and tobacco product claims and labels. This HINTS survey will couple knowledge-

related questions with inquiries into the communication channels through which understanding is being obtained, and assessment of FDA-regulated material. This survey will extend the information collected and priorities from HINTS

which have been to provide a comprehensive assessment of the American public's current access to, and use of, information about cancer across the cancer care continuum from cancer

prevention, early detection, diagnosis, treatment, and survivorship.

OMB approval is requested for 1 year. There are no costs to respondents other than their time. The total estimated annualized burden hours are 2,159.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hour
Individuals	4,318	1	30/60	2,159

Dated: November 24, 2014.

Karla Bailey,

NCI Project Clearance Liaison, National Institutes of Health.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; Surveys and Interviews To Support an Evaluation of the Innovative Molecular Analysis Technologies (IMAT) Program (NCI)

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI), National Institutes of Health (NIH), will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

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.

To Submit Comments and For Further Information: To obtain a copy of the data collection plans and instruments, submit comments in writing, 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.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

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.