DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request: Financial Sustainability of Human Tissue Biobanking (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 or request more information on the proposed project contact: Chana Rabiner, Ph.D., Biorepositories and Biospecimen Research Branch, Cancer Diagnosis Program, 9609 Medical Center Drive, Rockville, MD 20892 or call non-toll-free number 240–276–5715 or Email your request, including your address to: *chana.rabiner@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: Financial Sustainability of Human Tissue Biobanking, 0925–NEW, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: The purpose of this web-based survey is to collect information regarding the challenges that human tissue biobanks encounter in achieving financially sustainable operations. The information will be used to assist the National Cancer Institute (NCI) in strategizing program plans to provide increased and tailored support for national and international biobanks. The survey will collect a combination of structured, quantitative, and free-text descriptive data that characterize the type and maturity of respondent biobanks, their sources of funding, and their usage of funding in conducting operations. The survey will also collect information describing the difficulties in maintaining funding sources and establishing new ones. Finally, the survey will elicit descriptions of techniques used to overcome the difficulties.

It is expected that the information generated by this survey will be used to inform published guidance to biobanks regarding the financial hazards to sustained operations and the means by which these hazards can be avoided or overcome.

OMB approval is requested for 1 year. There are no costs to respondents other than their time. The total estimated annualized burden hours are 822.

ESTIMATED ANNUALIZED BURDEN HOURS

Category of respondents	Number of respondents	Frequency of response	Average time per response (in hours)	Total burden hours
Estimates of Annualized Burden Hours				
Private Sector	548	1	90/60	822
Totals	548			822

Dated: May 1, 2013.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, NCI, NIH.

[FR Doc. 2013–10772 Filed 5–6–13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request: The Framingham Heart Study (FHS)

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 Heart, Lung, and Blood Institute (NHLBI), the 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 request more information on the proposed project or to obtain a copy of data collection plans and instruments, contact Dr. Gina Wei, Division of Cardiovascular Sciences, NHLBI, NIH, Two Rockledge Center, 6701 Rockledge Drive, MSC 7936, Bethesda, MD 20892–7936, or call non-toll-free number (301) 435–0456, or email your request, including your address to: *weig@nhlbi.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: The Framingham Heart Study, 0925–0216, Extension, National Heart, Lung, and Blood Institute (NHLBI), National Institutes of Health (NIH).

Need and Use of Information Collection: The Framingham Heart Study will continue to conduct morbidity and mortality follow-up, as well as examinations, for the purpose of studying the determinants of cardiovascular disease. Morbidity and mortality follow-up will continue to occur in all of the cohorts (Original, Offspring, Third Generation, Omni Group 1, and Omni Group 2). Examinations will continue to be conducted on the Original, Offspring, and Omni Group 1 Cohorts.

OMB approval is requested for 3 years. There is no cost to the respondents other than their time. The total estimated annualized burden hours are 4264.

ESTIMATED ANNUALIZED BURDEN HOURS, ORIGINAL COHORT

Type of respondent	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
I. Participant Components				
A. Pre-Exam:				
a. Telephone contact to set up appointment	60	1	10/60	10
b. Exam Appointment, Scheduling, Reminder, and Instructions	55	1	35/60	32
B. Exam—Cycle 32:				
a. Clinic exam	25	1	45/60	19
b. Home or nursing home visit	25	1	65/60	27
C. Annual Follow-up:				
a. Records Request	60	1	15/60	15
b. Health Status Update	45	1	15/60	11
Sub-Total: Participant Components	*60			114
II. Non-Participant Components				
A. Informant Contact (Pre-exam and Annual Follow-up)	25	1	10/60	4
B. Records Request (Annual follow-up)	50	1	15/60	13
Sub-Total: Non-Participant Components	75			17

*Number of participants as reflected in Rows I.A.a and I.C.a. above.

ESTIMATED ANNUALIZED BURDEN HOURS, OFFSPRING COHORT AND OMNI GROUP 1 COHORT

Type of respondent	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
I. Participant Components:				
A. Pre-Exam:				
a. Telephone contact to set up apt or Health status update	300	1	10/60	50
b. Appt. or update Confirmation	250	1	10/60	42
c. Food Frequency Form	250	1	10/60	42
B. Exam:				
a. Clinic Exam	100	1	175/60	292
b. Home or nursing home visit	100	1	60/60	100
c. Consent Forms	200	1	20/60	67
C. Annual Follow-Up:				
a. Records Request	2292	1	15/60	573
b. Health Status Update	1833	1	15/60	458
Sub-Total: Participant Components	*2292			1624
II. Non-Participant Components:				
A. Informant contact (Pre-exam and Annual Follow-up)	229	1	10/60	38
B. Records Request (Annual follow-up)	2292	1	15/60	573
Sub-Total: Non-Participant Components:	2521			611

*Number of participants as reflected in Rows I.C.a. above.

ESTIMATED ANNUALIZED BURDEN HOURS, GENERATION 3 COHORT AND OMNI GROUP 2 COHORT

Type of respondent	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
I. Participant Components				
Annual Follow-up:		1		
A. Records Request	3212	1	15/60	803

ESTIMATED ANNUALIZED BURDEN HOURS, GENERATION 3 COHORT AND OMNI GROUP 2 COHORT—Continued

Type of respondent	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
B. Health Status Update	3212	1	15/60	803
Sub-Total: Participant Components	*3212			1606
II. Non-Participant Components				
Annual Follow-up:				
A. Informant contacts	160	1	10/60	27
B. Records Request	1060	1	15/60	265
Sub-Total: Non-Participant Components	1220			292

*Number of participants as reflected in Rows I.A. and I.B. above.

SUMMARY OF 3 TABLES COMBINED—TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
Participants	5564	1	36/60	3344
Non-Participants	3816	1	14.5/60	920
Totals	9380			4264

(Note: reported and calculated numbers differ slightly due to rounding.)

Dated: April 25, 2013.

Lynn Susulske,

NHLBI Project Clearance Liaison, National Institutes of Health.

Michael Lauer,

Director, DCVS, National Institutes of Health.

[FR Doc. 2013–10771 Filed 5–6–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases, Diabetes Mellitus Interagency Coordinating Committee Notice of Workshop

SUMMARY: The Diabetes Mellitus Interagency Coordinating Committee (DMICC) will hold a 2-day workshop on June 6–7, 2013. The workshop will be open to the public, with attendance limited to space available.

DATES: The workshop will be held on June 6, 2013 from 8:15 a.m. to 6:00 p.m., and June 7, 2013 from 8:15 a.m. to 4:00 p.m.

ADDRESSES: The workshop will be held at the National Institutes of Health Neuroscience Center, Conference Room B1/B2, 6001 Executive Boulevard, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: For further information concerning this workshop, contact Dr. B. Tibor Roberts, Executive Secretary of the Diabetes Mellitus Interagency Coordinating Committee, National Institute of Diabetes and Digestive and Kidney Diseases, 31 Center Drive, Building 31A, Room 9A19, MSC 2560, Bethesda, MD 20892–2560, telephone: 301–496–6623; FAX: 301–480–6741; email: *dmicc@mail.nih.gov.*

SUPPLEMENTARY INFORMATION: The DMICC facilitates cooperation, communication, and collaboration on diabetes among government entities. The June 6–7, 2013, DMICC workshop will discuss new and emerging opportunities for type 1 diabetes research supported by the Special Statutory Funding Program for Type 1 Diabetes Research. An agenda for the DMICC workshop will be available by contacting Mary Allen, The Scientific Consulting Group, Inc. (*mallen@scgcorp.com;* please put ''Agenda Request for DMICC T1D Meeting'' in the subject line).

Any interested person may file written comments with the Committee by forwarding their statement to the contact person listed on this notice. The statement should include the name, address, telephone number and, when applicable, the business or professional affiliation of the interested person. Because of time constraints for the workshop, there will not be time on the agenda for oral comments from members of the public.

Members of the public who would like to receive email notification about future DMICC meetings may register on a listserv on the DMICC Web site: *www.diabetescommittee.gov.*

Please note that seating is limited and attendance will be first-come, first-served. Non-federal individuals planning to attend the workshop should register by email to Mary Allen, The Scientific Consulting Group, Inc. (*mallen@scgcorp.com;* please put ''Registration DMICC T1D Meeting'' in the subject line) at least 7 days prior to the workshop. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should inform the Contact Person listed below at least 10 days in advance of the workshop.

Dated: April 26, 2013.

B. Tibor Roberts,

Executive Secretary, DMICC, Office of Scientific Program and Policy Analysis, NIDDK, National Institutes of Health.

[FR Doc. 2013–10770 Filed 5–6–13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases; Notice of Closed Meeting

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 meeting.