

Application	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Total .....	10	1	10	6.5	60

E-mail comments to [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or mail the HRSA Reports Clearance Officer, Room 10–33, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: September 3, 2010.

**Sahira Rafiullah,**

*Director, Division of Policy and Information Coordination.*

[FR Doc. 2010–22662 Filed 9–10–10; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Health Resources and Services Administration**

**Agency Information Collection Activities: Submission for OMB Review; Comment Request**

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to

OMB for review, e-mail [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or call the HRSA Reports Clearance Office on (301) 443–1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

**Proposed Project: Organ Procurement and Transplantation Network and Scientific Registry of Transplant Recipients Data System (OMB No. 0915–0157)—Extension**

Section 372 of the Public Health Service (PHS) Act requires that the Secretary, by contract, provide for the establishment and operation of an Organ Procurement and Transplantation Network (OPTN). The OPTN, among other responsibilities, operates and maintains a national waiting list of individuals requiring organ transplants, maintains a computerized system for matching donor organs with transplant candidates on the waiting list, and operates a 24-hour system to facilitate matching organs with individuals included in the list.

Data for the OPTN data system are collected from transplant hospitals, organ procurement organizations, and tissue-typing laboratories. The information is used to indicate the

disease severity of transplant candidates, to monitor compliance of member organizations with OPTN rules and requirements, and to report periodically on the clinical and scientific status of organ donation and transplantation in this country. Data are used to develop transplant, donation and allocation policies, to determine if institutional members are complying with policy, to determine member-specific performance, to ensure patient safety when no alternative sources of data exist and to fulfill the requirements of the OPTN Final Rule. The practical utility of the data collection is further enhanced by requirements that the OPTN data must be made available, consistent with applicable laws, for use by OPTN members, the Scientific Registry of Transplant Recipients, the Department of Health and Human Services, and others for evaluation, research, patient information, and other important purposes.

No revisions of the 29 data collection forms are proposed at this time; however, the OPTN is currently undergoing a review of the forms and expects to submit proposed revisions within the next year.

The annual estimate of burden is as follows:

Form	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Deceased Donor Registration .....	58	216	12,528	0.7500	9,396.0000
Death referral data .....	58	12	696	10.0000	6,960.0000
Death Notification Referral—Eligible .....	58	161	9338	0.2000	1,867.6000
Death Notification Referral—Imminent .....	58	168	9744	0.5000	4,872.0000
Living Donor Registration .....	308	39	12,012	0.6500	7,807.8000
Living Donor Follow-up .....	308	50	15,400	0.5000	7,700.0000
Donor Histocompatibility .....	156	131	20,436	0.1000	2,043.6000
Recipient Histocompatibility .....	156	196	30,576	0.2000	6,115.2000
Heart Candidate Registration .....	127	35	4,445	0.5000	2,222.5000
Lung Candidate Registration .....	68	42	2,856	0.5000	1,428.0000
Heart/Lung Candidate Registration .....	51	2	102	0.5000	51.0000
Thoracic Registration .....	127	36	4,572	0.7500	3,429.0000
Thoracic Follow-up .....	127	320	40,640	0.6500	26,416.0000
Kidney Candidate Registration .....	241	183	44,103	0.5000	22,051.5000
Kidney Registration .....	241	83	20,003	0.7500	15,002.2500
Kidney Follow-up* .....	241	742	178,822	0.5500	98,352.1000
Liver Candidate Registration .....	129	109	14,061	0.5000	7,030.5000
Liver Registration .....	129	58	7,482	0.6500	4,863.3000
Liver Follow-up .....	129	519	66,951	0.5000	33,475.5000
Kidney/Pancreas Candidate Registration .....	143	14	2,002	0.5000	1,001.0000
Kidney/Pancreas Registration .....	143	7	1,001	0.9000	900.9000
Kidney/Pancreas Follow-up .....	143	85	12,155	0.8500	10,331.7500
Pancreas Candidate Registration .....	143	7	1,001	0.5000	500.5000
Pancreas Registration .....	143	3	429	0.7500	321.7500
Pancreas Follow-up .....	143	20	2,860	0.6500	1,859.0000
Intestine Candidate Registration .....	44	7	308	0.5000	154.0000
Intestine Registration .....	44	5	220	0.9000	198.0000
Intestine Follow-up .....	44	28	1,232	0.8500	1,047.2000

Form	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Post Transplant Malignancy .....	684	10	6,840	0.2000	1,368.0000
Total .....	463	.....	522,815	.....	278,765.9500

\* Includes an estimated 2,500 kidney transplant patients transplanted prior to the initiation of the data system.

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to the desk officer for HRSA, either by e-mail to [OIRA\\_submission@omb.eop.gov](mailto:OIRA_submission@omb.eop.gov) or by fax to 202-395-6974. Please direct all correspondence to the "attention of the desk officer for HRSA."

Dated: September 3, 2010.

**Sahira Rafiullah,**

Director, Division of Policy and Information Coordination.

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Proposed Collection; Comment Request; Generic Clearance for Surveys of Customers and Partners of the Office of Extramural Research of the National Institutes of Health

**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 Office of Extramural Research (OER), 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.

*Proposed Collection: Title:* Generic Clearance for Surveys of Customers and Partners of the Office of Extramural Research of the National Institutes of Health. *Type of Information Collection Request:* NEW. *Need and Use of Information Collection:* OER develops, coordinates the implementation of, and evaluates NIH-wide policies and procedures for the award of extramural funds. To move forward with our initiatives to ensure success in accomplishing the NIH mission, input from partners and customers is essential. Quality management principles have been integrated into OER's culture and these surveys will provide customer satisfaction input on various elements of OER's business processes. The approximately 14 (10 quantitative and 4 qualitative) customer

satisfaction surveys that will be conducted under this generic clearance will gather and measure customer and partner satisfaction with OER processes and operations. The data collected from these surveys will provide the feedback to track and gauge satisfaction with NIH's statutorily mandated operations and processes. OER/OD/NIH will present data and outcomes from these surveys to inform the NIH staff, officers, leadership, advisory committees, and other decision-making bodies as appropriate. Based on feedback from these stakeholders, OER/OD/NIH will formulate improvement plans and take action when necessary. *Frequency of Response:* 1 Response. *Affected Public:* Individuals. *Type of Respondents:* Science professionals (applicants, reviewers, Institutional Officials), adult science trainees, and the general public. The annual reporting burden is as follows:

*Quantitative surveys:*

*Estimated Number of Respondents per Survey:* 9,820; *Estimated Number of Responses per Respondent:* 1; *Average Burden Hours per Response:* 0.25; *Estimated Total Annual Burden Hours Requested per Quantitative Survey:* 2,455; *Estimated Total Annual Burden Hours Requested for 10 Quantitative Surveys:* 24,550.

*Qualitative surveys:*

*Estimated Number of Respondents per Survey:* 30; *Estimated Number of Responses per Respondent:* 1; *Average Burden Hours per Response:* 1.0; *Estimated Total Annual Burden Hours Requested per Qualitative Survey:* 30; *Estimated Total Annual Burden Hours Requested for 4 Qualitative Surveys:* 120.

*Based on an estimated 10 quantitative and 4 qualitative surveys per year:*

*Estimated Total Combined Annual Hours of Burden Requested in Each of 3 years:* 24,670.

*Estimated Total Combined Cost to Respondents:* \$728,326.

*Based on an estimated 10 quantitative and 4 qualitative surveys per year over 3 years:*

*Estimated Total Hours of Burden to Respondents for 2011, 2012, and 2013 Combined:* 74,010.

*Estimated Total Cost to Respondents for 2011, 2012, and 2013 Combined:* \$2,184,978.

There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

*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.

**FOR FURTHER INFORMATION CONTACT:** To request more information on the proposed generic clearance or to obtain a copy of example data collection instruments, contact Dr. Gwynne Jenkins, Special Assistant to the Director, Office of Extramural Programs, OER, NIH, 6705 Rockledge Drive, Suite 350, Bethesda, MD 20892, or call non-toll-free number (301) 496-9232 or E-mail your request, including your address to: [OEPMailbox@mail.nih.gov](mailto:OEPMailbox@mail.nih.gov).

*Comments 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.

Dated: August 30, 2010.

**Sherry Mills,**

Director, Office of Extramural Programs.

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