

open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by April 21, 2014.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Luis G. Bravo at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at <http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm> for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: February 24, 2014.

Jill Hartzler Warner,

Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2014-04523 Filed 2-28-14; 8:45 am]

BILLING CODE 4160-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Health Resources and Services Administration

(HRSA) has submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public during the review and approval period.

DATES: Comments on this ICR should be received within 30 days of this notice.

ADDRESSES: Submit your comments, including the Information Collection Request Title, to the desk officer for HRSA, either by email to OIRA_submission@omb.eop.gov or by fax to 202-395-5806.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, email the HRSA Information Collection Clearance Officer at paperwork@hrsa.gov or call (301) 443-1984.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: Organ Procurement and Transplantation Network (OPTN) Application Form

OMB No.: 0915-0184 – Revision

Abstract: This is a request for OMB approval for revisions of the application documents used to collect information for determining if the interested party is compliant with membership and transplant program requirements contained in the Final Rule Governing the Operation of the Organ Procurement and Transplantation Network (OPTN), “the OPTN final rule”.

Need and Proposed Use of the Information: Membership in the OPTN is determined by submission of application materials to the OPTN (not to HRSA) demonstrating that the applicant meets all required criteria for membership and transplant program requirements and will agree to comply with all applicable provisions of the National Organ Transplant Act, as amended, 42 U.S.C. 273, et seq. Section 1138 of the Social Security Act, as amended, 42 U.S.C. 1320b-8 (section 1138) requires that hospitals in which transplants are performed be members of, and abide by, the rules and requirements (as approved by the

Secretary of the HHS) of the OPTN as a condition of participation in Medicare and Medicaid for the hospital. Section 1138 contains a similar provision for the organ procurement organizations (OPOs) and makes membership in the OPTN and compliance with its operating rules and requirements (that have been approved by the Secretary), including those relating to data collection, mandatory for *all* transplant hospitals and OPOs. These applications are developed to prompt submission of all the information required to make such membership approval decisions. In addition, hospitals wishing to obtain designation for particular (e.g., organ specific) transplant programs must submit applications to the OPTN.

Likely Respondents: Parties seeking initial OPTN membership approval and then maintenance of the existing OPTN approval. Applicants will include: every hospital seeking to perform organ transplants; every non-profit organization seeking to become an organ procurement organization; and every medical laboratory seeking to become a histocompatibility laboratory. In addition, there are other OPTN membership categories for organizations and individuals who want to participate in the organ transplant system and they too are required to fill out an appropriate application.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

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
A New Transplant Member/Program Application—General	8	1	8	8	64
B Kidney (KI) Designated Program Application	94	2	188	4	752
B Liver (LI) Designated Program Application	73	2	146	4	584
B Pancreas (PA) Designated Program Application	56	2	112	4	448
B Heart (HR) Designated Program Application	43	2	86	4	344
B Lung (LU) Designated Program Application	50	2	100	4	400

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS—Continued

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
B Islet (PI) Designated Program Application	4	2	8	3	24
B Living Donor (LD) Recovery Program Application	46	2	92	3	276
C OPO New Program Application	0	1	0	4	0
D Histocompatibility Lab Application	2	2	4	4	16
E Change in Transplant Program Key Personnel	377	2	754	4	3016
F Change in Histocompatibility Lab Director	8	1	8	2	16
G Change in OPO Key Personnel	10	1	10	1	10
H Medical Scientific Org Application	16	1	16	2	72
I Public Org Application	6	1	6	2	12
J Business Member Application	3	1	3	2	6
K Individual Member Application	6	1	6	1	6
Total =17 forms	802	26	1547	56	6046

Dated: February 21, 2014.

Jackie Painter,

Deputy Director, Division of Policy and Information Coordination.

[FR Doc. 2014-04576 Filed 2-28-14; 8:45 am]

BILLING CODE 4165-15-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Health Resources and Services Administration (HRSA) has submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public during the review and approval period.

DATES: Comments on this ICR should be received within 30 days of this notice.

ADDRESSES: Submit your comments, including the Information Collection Request Title, to the desk officer for HRSA, either by email to OIRA_submission@omb.eop.gov or by fax to 202-395-5806.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, email the HRSA Information Collection Clearance Officer at paperwork@hrsa.gov or call (301) 443-1984.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: NURSE Corps Loan Repayment Program OMB No.: 0915-0140—Revision

Abstract: The NURSE Corps Loan Repayment Program (NURSE Corps LRP), formerly known as the Nursing Education Loan Repayment Program (NELRP), assists in the recruitment and retention of professional Registered Nurses (RNs), including advanced practice RNs (i.e., nurse practitioners, certified registered nurse anesthetists, certified nurse-midwives, clinical nurse specialists), dedicated to working at eligible health care facilities with a critical shortage of nurses (i.e., a Critical Shortage Facility) or working as nurse faculty in eligible, accredited schools of nursing, by decreasing the financial barriers associated with pursuing a nursing profession. The NURSE Corps LRP provides loan repayment assistance to these nurses to repay a portion of their qualifying educational loans in exchange for full-time service at a public or private nonprofit Critical Shortage Facility or in an eligible, accredited school of nursing.

Need and Proposed Use of the Information: The need and purpose of this information collection is to obtain information for NURSE Corps LRP applicants and participants. The information is used to consider an applicant for a NURSE Corps LRP contract award and to monitor a participant's compliance with the service requirements. Individuals must

submit an application in order to participate in the program. The application asks for personal, professional, educational, and financial information required to determine the applicant's eligibility to participate in the NURSE Corps LRP. The semi-annual employment verification form asks for personal and employment information to determine if a participant is in compliance with the service requirements.

Likely Respondents: Professional RNs or advanced practice RNs (i.e., nurse practitioners, certified registered nurse anesthetists, certified nurse-midwives, clinical nurse specialists) who are interested in participating in the NURSE Corps LRP, and official representatives at their service sites.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

Total Estimated Annualized burden hours:

The estimates of reporting burden for Applicants are as follows: