

IX. LIASON OFFICERS

- A. Liaison Office for FDA
The Office of the Commissioner
- B. Liaison Office for Johns Hopkins
Office of the Vice Provost for Research

X. DURATION OF MOU

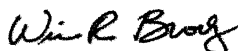
This MOU shall become effective upon the signature of all the Parties and will continue in effect for five (5) years. It may be extended by mutual written agreement of the Parties in writing. It may be modified by mutual consent or terminated by either Party upon a 30-day advance notice to the other Party.

XI. REGULATIONS

This MOU and all associated agreements will be subject to the applicable federal and state laws and regulations.

JOHNS HOPKINS UNIVERSITY

BY

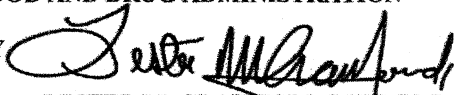


WILLIAM R. BRODY Ph.D.
PRESIDENT OF THE JOHNS HOPKINS UNIVERSITY

DATE

FOOD AND DRUG ADMINISTRATION

BY



LESTER M. CRAWFORD DVM, Ph.D.
DEPUTY COMMISSIONER FOOD AND DRUG ADMINISTRATION

DATE April 30, 2002

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BILLING CODE 4160-01-C

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Health Resources and Services Administration****Agency Information Collection Activities: Proposed Collection: Comment Request**

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, as amended, 44 U.S.C. Chapter 35), the Health Resources and Services Administration (HRSA) publishes periodic summaries of proposed projects being developed for submission to OMB under the Paperwork Reduction

Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, call the HRSA Reports Clearance Officer on (301) 443-1129.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

Proposed Project: Data System for Organ Procurement and Transplantation Network (42 CFR Part 121, OMB No. 0915-0184): Extension

The operation of the Organ Procurement and Transplantation Network (OPTN) necessitates certain record keeping and reporting requirements in order to perform the functions related to organ transplantation under contract to HHS. This is a request for an extension of the current record keeping and reporting requirements associated with the OPTN. These data will be used by HRSA in monitoring the contracts for the OPTN and the Scientific Registry of Transplant Recipients (SRTR) and in carrying out other statutory responsibilities. Information is needed to match donor organs with recipients, to monitor compliance of member organizations with OPTN rules and requirements, and to ensure that all qualified entities are accepted for membership in the OPTN.