



DEPARTMENT OF HEALTH & HUMAN SERVICES
Administration

Public Health Service
Food and Drug

Memorandum

Date April 25,2006

From Director, Division of Management Systems, Paperwork reduction Act and Records Management Branch, FDA

Subject Request for OMB Approval for “Guidance on Informed Consent for In Vitro Diagnostic Studies Using Leftover Human Specimens That Are Not Individually Identifiable.”

To Chief, Human Resources and Housing Branch
Office of Information and Regulatory Affairs, OMB
Through: Reports Clearance Officer, HHS_____

The purpose of this memorandum is to request approval of the collection of information requirements for "Guidance on Informed Consent for IN Vitro Diagnostic Studies Using Leftover Human Specimens That Are Not Individually Identifiable." The estimate of the burden for this collection of information will be 2,800 hours.

Paul Jones

Attachment

