Protection of Human Subjects Assurance Identification/IRB Certification/Declaration of Exemption (Common Rule)

Policy. Research activities involving human subjects may not be conducted or supported by the Institutions must have an assurance of compliance that applies to the research to be conducted and Departments and Agencies adopting the Common Rule (56FR28003, June 18, 1991) unless the should submit certification of IRB review and approval with each application or proposal unless activities are exempt from or approved in accordance with the Common Rule. See section 101(b) otherwise advised by the Department or Agency.

of the Common Rulefor exemptions. Institutions submitting applications or proposals for support must submit certification of appropriate Institutional Review Board (IRB) review and approval to the Department or Agency in accordance with the Common Rule.	
1. Request Type 2. Type of Mechanism [] ORIGINAL [] GRANT [] CONTRACT [] FELLOWSHIF [] CONTINUATION [] COOPERATIVE AGREEMENT [] OTHER:	Name of Federal Department or Agency and, if known, Application or Proposal Identification No.
4. Title of Application or Activity	5. Name of Principal Investigator, Program Director, Fellow, or Other
6. Assurance Status of this Project (Respond to one of the following)	
[] This Assurance, on file with Department of Health and Human Services, of Assurance Identification No, the expiration	covers this activity: n date IRB Registration No
[] This Assurance, on file with <i>(agency/dept)</i> Assurance No	, covers this activity.
Assurance NoB&@A\¢] aaaaaaaaaaa	IRB Registration/Identification No(if applicable)
[] No assurance has been filed for this institution. This institution declares that it will provide an Assurance and Certification of IRB review and approval upon request.	
Exemption Status: Human subjects are involved, but this activity qualifies for exemption under Section 101(b), paragraph	
7. Certification of IRB Review (Respond to one of the following IF you have an Assurance on file)	
[] This activity has been reviewed and approved by the IRB in accordance with the Common Rule and any other governing regulations. by: [] Full IRB Review on (date of IRB meeting) or [] Expedited Review on (date) [] If less than one year approval, provide expiration date [] This activity contains multiple projects, some of which have not been reviewed. The IRB has granted approval on condition that all projects covered by the Common Rule will be reviewed and approved before they are initiated and that appropriate further certification will be submitted.	
8. Comments	
9. The official signing below certifies that the information provided above is correct and that, as required, future reviews will be performed until study closure and certification will be provided.	10. Name and Address of Institution
11. Phone No. (with area code)	
12. Fax No. (with area code)	
13. Email:	
14. Name of Official	15. Title
16. Signature	17. Date
Authorized for local Poproduction	Spannared by HUS

According to the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number. The valid OMB control number for this information collection is 0990-0263. The time required to complete this information collection is estimated to average 30 minutes per response. If you have comments concerning the accuracy of the time estimate(s) or suggestions for improving this form, please write to: U.S. Department of Health & Human Services, OS/OCIO/PRA, 200 Independence Ave., S.W., Suite 336-E, Washington D.C. 20201, Attention: PRA Reports Clearance Officer.