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## **IRB COOPERATIVE AUTHORIZATION AGREEMENT**

### **Institution or Organization Providing IRB Review**

Name (Institution/Organization A): \_\_\_\_\_

IRB Registration/Protocol #: \_\_\_\_\_

Federalwide Assurance(FWA)#, if any: \_\_\_\_\_

### **Institution Relying on the Designated IRB (Institution B)**

Name: \_\_\_\_\_

FWA#: \_\_\_\_\_

The Officials signing below agree that \_\_\_\_\_(name of Institution B) may rely on the designated IRB for review and continuing oversight of its human subjects research described below: (check one):

☐ This agreement applies to all human subjects research covered by Institution B's FWA.

☐ This agreement is limited to the following specific protocol(s):

Name of Research Project: \_\_\_\_\_

Name of Principal Investigator: \_\_\_\_\_

Sponsor or Funding Agency: \_\_\_\_\_

Award Number, if any: \_\_\_\_\_

☐ Other (describe): \_\_\_\_\_

The review performed by the designated IRB will meet the human subject protection requirements of Institution B's OHRP-approved FWA. The IRB at Institution/Organization A will follow written procedures for reporting its findings and actions to appropriate officials at Institution B. Relevant minutes of IRB meetings will be made available to Institution B upon request. Institution B remains responsible for ensuring compliance with the IRB's determinations and with the Terms of its OHRP-approved FWA. This document must be kept on file by both parties and provided to OHRP upon request.

Signature of Signatory Official (Institution/Organization A):

\_\_\_\_\_

Date: \_\_\_\_\_

Print Full Name: \_\_\_\_\_

Institutional Title: \_\_\_\_\_

Signature of Signatory Official (Institution B):

\_\_\_\_\_

Date: \_\_\_\_\_

Print Full Name: \_\_\_\_\_

Institutional Title: \_\_\_\_\_