

Vice, Rudith (CDC/NCEZID/OD)

From: Youngblood, Laura (CDC/NCEZID/OD)
Sent: Monday, June 3, 2024 12:22 PM
To: Vice, Rudith (CDC/NCEZID/OD); Nti-Berko, Sonja Mali (CDC/NCEZID/DIDRI/RRRSB)
Subject: Regulatory Status of EIP

Dear Rudith and Sonja,

The purpose of this e-mail is to confirm the regulatory status of the projects funded through the EIP Cooperative Agreement. EIP is a hybrid CoAg that allows for both research and non-research activities. I generally do not know which activities are or might be research until I receive the applications.

However, I can confirm that the Core surveillance activities for each program area are non-research and do not require IRB review.

Sincerely,

Laura

Laura Youngblood, MPH, CIP

Human Subjects Advisor | NCEZID Information Collections & Human Studies Team | lyoungblood@cdc.gov | Office: (404) 639-6394 | Mobile: (404) 510-0093