**Foreign Site Information**

*If the Pilot Award will have multiple foreign sites, complete this form for EACH research site*

|  |  |
| --- | --- |
| **Principal Investigator Name:** |  |
| **Project Title:** |  |
| **Institution:** | This is the institution that will be administering the award  |
| **Foreign Research Site:** | Location of foreign research activities |
| **Point of Contact for Research Site:** | Name, organization, address, email, and phone number |

**Introduction (3-4 sentences):**

**Provide a simple description of the overall goals of the project including the work that will be done at the foreign site. Indicate if any of the NIAID grant funds will be sent to the foreign site.**

The goals of this project are…

To achieve these goals, the investigator will collaborate with…

$X of grant funds will be sent to the site for these studies.

**Foreign component (3-4 sentences):**

**Describe the specific role of the foreign site (i.e. conduct critical experiments?, conduct data analyses?, provide consultations?, provide samples?, study human populations?, collect data to be brought to the US?).**

The site will…

**Human Subjects (1 word or 1 sentence per bullet):**

* **Parent Study IRB approval**
	+ **IRB approval number for parent study:** #
	+ **IRB approval date:** DD-MMM-YYYY
	+ **Human Subject Assurance Number:** #
* **Does this study require a modification to the IRB approval of a parent study (Yes or No)?**
	+ **?**
* **Will existing samples from human subjects will be used: (Yes or No)?**
	+ **?**
	+ **How many subjects provided the existing samples to be used?** #
* **Population parameters:**
	+ **Gender:** # males, # females
	+ **Age Group:** ages # - #
	+ **Race/Ethnicity:** # Caucasian subjects, # subjects of African ancestry, # subjects of Asian ancestry, etc.
* **Will local IRB approval be obtained prior to engaging in any research involving human subjects (Yes or No)? If No, describe alternate approvals being sought or state why local IRB approval is not required.**
	+ ?
* **Will informed consent be obtained prior to engaging in any research involving human subjects (Yes or No)? If No, describe alternate approvals/consents being sought or state why informed consent is not required.**
	+ ?
* **Will data be brought back to the US (Yes or No)?**
	+ ?