CFAR CLINICAL CLEARANCE

I. Studies that cannot be funded through the CFAR

- Any clinical trial (NIH definition of a Clinical Trial (NOT-OD-15-015)
- Studies involving new drugs, treatments, or devices

II. Studies that can be funded via CFAR but require additional NIH review

- Studies involving new ways of using known drugs, treatments, or devices (allowed on a case-by-case basis)
- Studies that are deemed above minimal risk by the Institutional IRB
- Studies involving vulnerable populations (children, pregnant women, transgender, sex workers, prisoners, refugees, individuals who are unable to provide informed consent, etc.)
- Studies involving behavioral interventions (above minimal risk)

No human subject work may be initiated until clinical approval is received.

III. Studies that do not require additional NIH review

Research activities that do not include vulnerable populations (see Category II above) and present no more than minimal risk to human subjects as described in the OHRP Expedited Review Categories. Examples include but are not limited to the following:

- routine blood draws
- non-invasive procedures routinely employed in clinical practice (e.g. ultrasound, MRI)
- surveys, focus groups