

## MedStar Research Institute-Georgetown University Oncology Institutional Review Board

### Request for Facilitated Review of Research Approved by the Central Institutional Review Board of the National Cancer Institute

#### Section One: Project and Personnel

<b>Investigator:</b>	<b>Date:</b>
<b>Title of Project:</b>	
<b>CIRB Number:</b>	<b>Cooperative Group &amp; Number:</b>

Georgetown University (GU) and MedStar Research Institute (MRI) are members of the Central Institutional Review Board (CIRB), a pilot project sponsored by the National Cancer Institute (NCI) in consultation with the DHHS Office of Human Research Protection (OHRP) to streamline the process for local IRB review of multi-center cancer treatment trials.

Consultants or Co-investigators, if any	Title	Department/ Institution	Phone

An investigator at GU or MRI who wishes to enroll subjects in a CIRB-approved protocol should download the protocol, informed consent documents, and the CIRB application from the participant side of the CIRB website (<http://www.ncicirb.org>) and submit them along with this Request for Facilitated Review to the MRI-GU Oncology IRB for local expedited review.

1. Please attach the following materials to this application:
  - CIRB approved protocol documents, including approval letter and CIRB approved consent form
  - Any survey tools or questionnaires
  - Scientific Review results
  - Informed Consent Document in MRI-GU format

**CIRB NOTE:** Local boilerplate additions to the informed consent dealing with state and local law, institutional requirements, or IRB policies may be added to the local consent form. Local IRBs may also make minor word substitutions or additions in the informed consent document, particularly to facilitate better comprehension by the local population, as long as the proposed changes do not alter the meaning of the CIRB approved contents. Revisions/changes to the local consent form other than those described above require full board review at the local level, and facilitated review may not be used. Per current OHRP/NCI guidance, any informed consent changes must be justified in the IRB minutes and sent to the Cooperative Group administering that protocol.

2. Has the Institutional Biosafety Committee approved the protocol?     Yes     No     Not Applicable
3. Has the Radiation Safety Committee approved the protocol?     Yes     No     Not Applicable

