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| Human Research Compliance |  |
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| New York, NY 10022 |  |
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**CLINICALTRIALS.GOV DISSEMINATION PLAN**

Investigators submitting NIH FORM E must utilize the following template dissemination plan language in their application.

*The WCM Clinicaltrials.gov administrator will facilitate the Principal Investigator’s dissemination of study results through ClinicalTrials.gov registration and reporting.*

* *X (insert your name or PI designee) will be responsible for handling ClinicalTrials.gov requirements for this project according to WCM clinicaltrials.gov SOP. (insert your name or PI designee) will register the trial prior to enrolling the first subject. Once a record is established, (insert your name or PI designee) will confirm accuracy of record content; resolve problems; and maintain records including content update and modifications.  (insert your name or PI designee) will also be responsible for results reporting and Adverse Events reporting at the conclusion of the project.*
* *Informed consent documents for the clinical trial(s) will include a specific statement relating to posting of clinical trial information at ClinicalTrials.gov; and*
* *WCM has an internal policy in place to ensure that clinical trials registration and results reporting occur in compliance with requirements contained in NIH Policy on the Dissemination of Clinical Trial Information (NOT-OD-16-149).*
* *Add any additional specifics related to this trial.*