Dissemination Plan

INSTRUCTIONS FOR COMPLETING THIS TEMPLATE:

1. Insert study-specific text where included below in *blue italics*.

2. Once this template has been completed:
    a. Delete the purple instructional text
    b. Change all text to black

3. Save the template as a PDF and upload it in item *4.6 Dissemination Plan* of your PHS Human Subjects and Clinical Trials Information form.

In compliance with the *NIH Policy on Dissemination of NIH-Funded Clinical Trial Information* (https://grants.nih.gov/policy-and-compliance/policy-topics/clinical-trials/reporting/nih-policy), the project PI for the clinical trial ***"[Insert study title]"*** will:

1) Register the trial in ClinicalTrials.gov no later than 21 calendar days after enrollment of the first participant.

2) Submit the trial results in ClinicalTrials.gov no later than one year after the trial's primary completion date.

The project PI's home institution, ***[Insert PI's institution],*** has an internal policy in place to ensure compliance with the *NIH Policy on Dissemination of NIH-Funded Clinical Trial Information* and its requirements for registration and submission of results information to ClinicalTrials.gov.

[If the proposed project includes informed consent, include this sentence. If the project does not include informed consent, delete this sentence]: The informed consent form(s) for this clinical trial will describe that the project and its results will be posted in ClinicalTrials.gov.