This template is based on the NIH instructions here https://grants.nih.gov/grants/how-to-apply-application-guide/forms-g/general/g.500-phs-human-subjects-and-clinical-trials-information.htm#4.7

This attachment is required only if you answered "Yes" to all the questions in the "Clinical Trial Questionnaire." (i.e. your study is an NIH-defined clinical trial).

When document is complete, delete this box and save as PI Last Name_4.7 Dissemination Plan DDMOYEAR (example-01JAN2022)

NIH-recommended fonts: Arial, Georgia, Helvetica, Palatino Linotype, 11 pt or larger, margins .05 or larger https://grants.nih.gov/grants/how-to-apply-application-guide/format-and-write/format-attachments.htm

For more information:

- See the NIH Policy on the Dissemination of NIH-Funded Clinical Trial Information
- See the NIH Guide Notice on the <u>Delayed Enforcement and Short-Term Flexibilities for Some</u>
 Requirements Affecting Prospective Basic Science Studies Involving Human Participants
- See the NIH Grants Policy Statement, Section 4.1.3.1 NIH Policy on Dissemination of NIH-Funded Clinical Trial Information.

4.7 DISSEMINATION PLAN

Explain briefly your plan for the dissemination of NIH-funded clinical trial information and address how the expectations of the policy will be met. The plan must contain sufficient information to assure the following:

- the applicant will ensure that clinical trial(s) under the award are registered and results information is submitted to ClinicalTrials.gov as outlined in the <u>policy</u> and according to the specific timelines stated in the policy;
- informed consent documents for the clinical trial(s) will include a specific statement relating to posting of clinical trial information at ClinicalTrials.gov; and
- the recipient institution has an internal policy in place to ensure that clinical trials registration and results reporting occur in compliance with policy requirements.

Note: Do not include informed consent documents in the Dissemination Plan attachment.