[INSERT SHORT NAME]Protocol Summary

Official Protocol Title	[This can be copied and pasted from OnCore]
Short Title (if applicable)	[This can be copied and pasted from OnCore]
Sponsor	[This can be copied and pasted from OnCore or Clinical Trials.gov]
Sponsor Protocol No.	[This is the sponsor's protocol number, and can be found usually on first page of protocol, in OnCore in the sponsor tab or on www.clinicaltrials.gov]
IRB Number	[This can be copied and pasted from OnCore]
NCT Number	[This can be copied and pasted from OnCore or Clinical Trials.gov] (For more information about this clinical trial, go to **www.clinicaltrials.gov** and search by the above number)
Purpose	[This should be copied from the purpose section on the clinicaltrials.gov website. Any more information than this could be proprietary information.]
Length of Study	[This can be found in the protocol or clinicaltrials.gov site]
Investigational Test Article	☐ N/A ☐ Investigational New Drug Drug Name: Mechanism of Action/Class: ☐ Investigational Device Device Name: Description of Device: [This area should include a brief description of the test article. This information may be found on clinical trials.gov or a protocol synopsis. This should just provide basic information for the care provider caring for a patient. If more information is needed, they may contact the research team]
Contraindications:	[In this area, document if there are any contraindication during participation in a trial. For example, if a study involves a novel anticoagulant, the study may prohibit the use of Coumadin during participation. Remember to not include any proprietary or confidential information]