# [INSERT SHORT NAME]Protocol Summary

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| **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](http://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](http://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] |