**Study Title:   
IRB#**

**PI:***All study team members should be trained on the protocol, protocol updates, and investigator’s brochure or device manual, including any subsequent changes. Maintain copies of agendas, attendance logs and relevant training materials with this Study-Specific Training Form. Examples are included below.*

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| **Training description** | **Trainer** | **Date completed and initials** | **Comments** |
| *Protocol version 2.1* | *Dr. Jones, Study PI* | *1/1/2022* | *Research nurse read protocol prior to training. PI discussed purpose, eligibility criteria, procedures, risks, informed consent, and investigational drug.* |
| *Investigator Brochure version 9* | *Sponsor’s monitor* | *1/10/2022* |  |
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