|  |  |  |
| --- | --- | --- |
| Name | Study Role | Study Tasks (select from list) |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |

## RCR-Registration Types:

**Investigator** (**IVR)** *MD, DO, or international equivalent*

**Non-Physician Investigator** (**NPIVR)** *advanced practice providers (e.g., NP or PA) or graduate level researchers (e.g., PhD)*

**Associate Plus (AP)** *clinical site staff*

## Study Tasks List for Personnel on DTL:

1. DTL and RCR Administrator (IVR, NPIVR, AP)

2. Primary Study/Site Contact (IVR, NPIVR, AP)

3. Person Authorized to Obtain Consent (IVR, NPIVR, AP)

4. Data Management/EDC (IVR, NPIVR, AP)

5. Eligibility Assessments (IVR, NPIVR, AP) **\*\***

6. Endpoint Assessments (IVR, NPIVR)

7. AE/SAE and Toxicity Assessments (IVR)

8. H&P Assessments (IVR)

9. IND Agent Prescribing (IVR)

10. Non-IND Agent Prescribing (IVR)

11. Investigational Product Accountability (IVR, NPIVR, AP)   
12. Study-Related Interventions (IVR, NPIVR, AP)

13. Patient Screening/Recruiting (IVR, NPIVR, AP)

14. Source Documentation Completion (IVR, NPIVR, AP)

15. Regulatory Contact (IVR, NPIVR, AP)

16. Specimen Tracking System (IVR, NPIVR, AP)

17. Specimen Processing/Shipping (IVR, IVR, AP)

18. Pathology Lab Support (IVR, NPIVR, AP)

19. Other*(d)*:

20. Other*(d)*:

*\*\*Eligibility Assessments:*

Screening (IVR, NPIVR, AP); Randomization (IVR)