National Institutes of Health National Cancer Institute Bethesda, Maryland 20892

## **NCI, DCP Delegation of Tasks Log**

**Study Information I** 

♦ A separate Delegation of Tasks Log must be completed by each Study Staff member.         Study Information II         Protocol Lead Investigator Name       Site       Protocol Lead Investigator Name         ♦ I have delegated to the staff member below the authority to perform the task(s) indicated, under my supervision. As qualified to perform the delegated task(s) on the basis of education, training or experience.         Delegations         Staff Member Name       Staff Member       Staff Member Signature       Staff Member Signature	CP Protocol No:  Investigator Signat  of the start date, the		Date ber was	
Study Information II  Protocol Lead Investigator Name  Site  Protocol Lead      I have delegated to the staff member below the authority to perform the task(s) indicated, under my supervision. As qualified to perform the delegated task(s) on the basis of education, training or experience.   Delegations  Staff Member Name  Staff Member Staff Member Staff Member Signature  Staff Member Signature				
Protocol Lead Investigator Name       Site       Protocol Lead         ◆ I have delegated to the staff member below the authority to perform the task(s) indicated, under my supervision. As qualified to perform the delegated task(s) on the basis of education, training or experience.         Delegations         Staff Member Name       Staff Member       Staff Member Signature       Staff Member Signature				
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The Protocol Lead Investigator will sign above at the beginning of study and sign below when study is complete. If the staff n study lifecycle, use additional lines to record new position/tasks. (Reference: FDA Guidance for Industry Investigator Respon Welfare of Study Subjects, 2009)  Study Information III				

## Task Codes

1. Obtain & Administer Informed Consent

**Protocol Lead Investigator Signature** 

2. Perform Study Drug Accountability

at conclusion of the study:

- 3. Determine Patient Eligibility
- 4. Recruit Patients

- 5. Obtain/Prepare Lab Samples
- 6. Complete Source Documents
- 7. Review & Correct Source Documents
- 8. Perform Physical Examinations
- 9. Maintain Regulatory Documents
- 10. Obtain Medical History
- 11. Dispense Study Medication
- 12. Report SAEs

- 13. Instruct Patients on Study Procedures
- 14. Complete Case Report Forms
- 15. Review & Correct Case Report Forms
- 16. Sign/Approve Data Correction Forms
- 17. Data Analysis

Date:

- 18. Research Analysis
- 19. Other (specify):