### **Senior Clinical Program Manager**

Location: Blacksburg, VA

Salary: Will commensurate with experience

Employment Status: Full-Time
Travel Required: Occasional

Please submit resumes to <a href="https://nrpharma.com">hr@nrpharma.com</a>
Subject Heading: Senior Clinical Program Manager

Fax: (540) 633-7939

### **Job Summary:**

New River Pharmaceuticals Inc. (NASDAQ: NRPH) is seeking a Senior Clinical Program Manager to oversee all aspects of clinical trials from study start-up through closure.

## ESSENTIAL DUTIES AND RESPONSIBILITIES include the following:

- Direct and manage all clinical study activities including tasks performed by the CRO, vendors, and other staff members assigned to the program.
- Lead multidisciplinary project teams, ensuring thorough study team communication.
- Review and approve all study related documents, and verify that study plans, procedures, forms, instructions, and reports are consistent with the protocols, company SOPs, and applicable regulatory guidelines.
- Responsible for meeting project timelines and budgets, managing issue resolution, and ensuring that appropriate QC and QA measures are incorporated into the program.
- Additional duties and responsibilities that may be assigned from time to time.

**QUALIFICATIONS** To perform this job successfully, an individual must be able to perform each essential duty satisfactorily. The qualifications listed below are representative of the knowledge, skill, and/or ability required or desired.

- A working knowledge of GCP and ICH Guidelines
- Expertise that includes exposure to all aspects of clinical programs
- Ability to successfully mentor, lead, and manage interdisciplinary project teams

## **EXPERIENCE**

- 10+ years of relevant clinical trial experience
- Minimum of 5 years of full scale project management of phase II and III clinical trials, including:
  - o Vendor selection
  - Clinical supplies
  - o Site recruitment, qualification, and initiation
  - Vendor and site budget and contract negotiations
  - Development of study related documents
  - Site monitoring and management
  - Central laboratories
  - Data cleaning and finalization
  - o Content of trial master files

# **EDUCATION, CERTIFICATION, LICENSES and/or REGISTRATIONS**

Bachelor's degree required – B.S. or B.A. degree in a scientific or health related field

New River Pharmaceuticals Inc. is an equal opportunity employer.