

Title: Sr. Lead Study Biostatistical Programmer
Fulltime Postion
Location: Northern NJ – Major Pharmaceutical Firm

Function of the position:

To function as study lead programmer in multiple clinical phases
Ensure all programming activities within study adhere to required guidelines and standards
Create and/or review programming specifications and plans at the study level
Execute and fully understand department, product and study level Macros (SAS Macro usage a must)
Lead technical and process improvement initiatives within and across multiple functions.

Required:

Must have CDISC experience
5+ years clinical R&D experience
Multi-therapeutic knowledge and experience.
Drug Development –early, late and/or observational in related industry studies
Biostatistical Programming –recent version – preferably Version 9.1
In depth understanding of computing operating systems, predominately UNIX. (strong plus)

If interested, please contact:

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