

Sr. SAS programmer  
Lexington, MA

**Responsibilities:**

- Provide general SAS programming support to Clinical Research and Development
- Work closely with statisticians to generate analysis data sets and create tables, listings and graphs for interim analyses, preliminary reports and Clinical Study Reports for in-house use, publications, and/or regulatory submissions
- Perform quality control checks on program code and outputs produced by other team members
- Conduct data edit checks to ensure the quality of source data
- Enhance existing utility programs and develop new utility macros to standardize data processing and output procedure
- Review and provide input on programming related documentation including plans and specifications
- Manage programming timelines, schedules, and activities on specific programming tasks
- Report to Associate Director of Statistical Programming

**Requirements:**

- Bachelor's degree in life science, statistics, mathematics, computer science, or related field required; Master's degree preferred
- 5+ years of SAS programming experience in biotech or pharmaceutical industry
- Experience in SAS programming language, macros, and procedures
- Working knowledge of ISS/ISE, CDISC standards (SDTM, ADaM)
- Detail oriented with strong organizational and communication skills
- Ability to work in a team environment

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