Large Healthcare organization hiring for a SAS Programmer. This position supports RTG Medical department clinical trials and regulatory submissions for pharmaceutical products and medical devices.

EXPERIENCE AND REQUIRED SKILLS:
* 2 - 5 years' related experience; or an advanced degree without experience; or equivalent directly related work experience.
* Good SAS programming skills in data manipulation, statistical analysis, graph and macro
* Knowledge of ICH/FDA regulations regarding data structure for submission, statistical analysis, clinical practices and other pertinent guidance

PRINCIPAL DUTIES AND RESPONSIBILITIES:
* Develops professional expertise in writing and validating programs, primarily SAS,for analysis datasets, tables, figures, and listings for clinical studies, or for regulatory submissions; applies company policies and procedures to resolve a variety of issues.
* Works with general instructions on routine jobs, detailed instructions on new projects or assignments. Work is reviewed for soundness.
* Solves problems of moderate scope in statistical programming for clinical trials conducted by RTG Medical Department. Exercises judgment within defined procedures and practices to determine appropriate action.
* Provides assistance to junior level staff with general tasks that require a better understanding of statistical programming, as directed by immediate supervisor.

Contact:

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