

**Status:** Contract  
**Location:** **Region:** Northeast **City:** East Hanover **State:** NJ  
**Zip Code:** 07936 **Area Code:** 862  
**Job Code:** bw0805sp

**Title:** Statistical Programmer

**Job Description:**

Provide statistical programming support and validation on clinical drug trials for a large pharma company. Will work in multiple therapeutic groups on various phases of clinical studies, effectively communicate with Biostatistics, statistical reporting and other clinical groups.

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**Skill Set:**

BA/BS or equivalent experience in computer science, mathematics, statistics, life sciences or related field(s). MS in Statistics a plus. Intermediate SAS programming skills with 2-5 years experience working with clinical trials data. Proficient with Macros, STAT and GRAPH. Good understanding of clinical trial practices, procedures and methodologies. Good verbal and written communication skills. Good interpersonal and organizational skills. Attention to detail and correctness. Working knowledge of office tools.

**Education:**

B.S. or M.S.

**Degree:**

Biostatistics, Computer Science, Life Sciences, Mathematics, Statistics

**Industry Experience:**

Mid Level

**Categories:**

Biotechnology and Pharmaceutical

If you are interested in this position, please send your resume to [Fred.Follett@asg-inc.com](mailto:Fred.Follett@asg-inc.com) and reference the job code.

**Status:** Contract  
**Location:** **Region:** Northern California **City:** San Francisco Peninsula  
**State:** CA **Zip Code:** 94080 **Area Code:** 650  
**Job Code:** ep0721cspa

**Title:** Clinical SAS Programmer/Analyst

**Job Description:**

Client is in need of Clinical SAS Programmer/Analyst with a minimum of 5 years of programming or related experience with at least 3 years in pharma/biotech/health care industry. Strong understanding of clinical monitoring/data query process and proficiency in writing clear, concise SAS code for the purposes of data listing review. Intermediate UNIX, Oracle Clinical or equivalent clinical DM system and relational database theory. Basic knowledge of regulatory environment and FDA/ICH guidelines e.g. 21CFR Part 11. Conversant in good programming practice and the software development life cycle. Understands the functional areas of a clinical trial and their role throughout the study. Excellent communication skills essential: Candidate should be fully capable of clear and timely written and verbal communication with peers, customers and management.

- The Clinical Programmer Analyst applies advanced level programming techniques and input to the design, development, implementation and maintenance of software in support in support of monitoring, reporting and analysis of clinical trial data quality.
- Uses creativity and ingenuity to solve complex problems (e.g. creates reusable code, develops project/department standard code, designs interconnected programs).
- Works closely with clinical research professionals to identify project/study level needs (e.g. deliverables schedule, report approval process, standard reporting styles, etc.) and to create reports needed to identify/resolve data issues.
- Analyzes data capt communicates effectively to coordinate problem resolution efforts with implementation team customers and functional peers.
- Contributes toward continuing process, business and technical enhancements.
- Mentors junior staff.

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**Skill Sets:** SAS Base, SAS Macro, SAS GRAPH

**Education:** B.S. or M.S.

**Degree:** Biostatistics, Computer Science, Life Sciences, Mathematics

**Industry Experience:** Experienced

**Categories:** Biotechnology and Pharmaceutical

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**Status:** Perm  
**Location:** **Region:** Northern California **City:** Emeryville  
**State:** CA **Zip Code:** 94612 **Area Code:** 510  
**Job Code:** ep0721spa

**Title:**  
SAS Programmer/Analyst

**Job Description:**

Our client, a successful health maintenance organization located in Northern California has an outstanding opportunity for a SAS Programmer/Analyst.

**Qualifications:**

- Experience writing and maintaining SAS macros.
- Experience with SQL, PROC REPORT and statistical procedures in SAS preferred.
- Experience with Web Development, Java, JSP and Servlets strongly preferred.
- Experience with analytical manipulation and interpretation of large databases required.
- Five or more years of related analytical consulting experience, healthcare experience preferred.
- Broad familiarity with medical practices, especially population management and process and outcomes measurement.
- Experience in leading projects strongly preferred.
- Experience with MVS/TSO/JCL, Microsoft Office.
- Exceptional analytic and critical thinking skills, writing skills, communication skills, consulting skills and ability to work within a team.
- Ability to express complex analytical and technical information to senior management or to audiences with clinical training.

Masters degree in economics, finance, health care administration, public health administration, public health administrations, statistics, mathematics, operations research, or related field required, or equivalent Bachelors degree and work experience in lieu of a Masters degree.

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**Skill Sets:** SAS Base, SAS Macro  
**Education:** B.S. or M.S.  
**Degree:** Computer Science, Life Sciences, Mathematics, Statistics  
**Industry Experience:** Experienced  
**Categories:** Healthcare

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**Status:** Contract  
**Location:** **Region:** Northern California **City:** San Francisco Peninsula  
**State:** CA **Zip Code:** 94080 **Area Code:** 650  
**Job Code:** ep0721ssp

**Title:**  
Statistician/SAS Programmer

**Job Description:**

We have a client in both Northern and Southern California that is in need of an MS level statistician for a six-month assignment.

The ideal candidate will be someone who has been with a major drug/biotech firm or a CRO (4 – 6 years of experience), with very focused experiences working through clinical protocol, analysis plan and report generation. MS level candidate will be supporting analysis plans. Must have clinical experience. Development of text, sharing tables, working on data set specifications for the analysis plan. Doing database checks. Working with the medical writers to create statistical reports. 6 plus months ready to go as soon as possible....MS level will work....SAS Programming skills necessary.

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**Skill Sets:**  
SAS Base, SAS Macro, SAS STAT

**Education:**  
B.S. or M.S.

**Degree:**  
Biostatistics, Life Sciences

**Industry Experience:**  
Mid-level

**Categories:**  
Biotechnology and Pharmaceutical

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**Status:** Perm  
**Location:** **Region:** Northern California **City:** East Bay **State:** CA  
**Zip Code:** **Area Code:** 510  
**Job Code:** mu0727scsp

**Title:** Senior Clinical SAS Programmer

**Job Description:**

Be a leader in creating meaningful biotherapeutics to improve our patient's quality of life, providing challenging and rewarding opportunities for our employees. Excellence and high ethical standards, respect for the individual and creativity with integrity. Do you share these visions and values? Then read on for an excellent fulltime career opportunity!

Our sponsor/client is seeking out an experienced clinical SAS Programmers that can provide SAS programming support and collaborate cross functionally within the study team; plan, execute and validate SAS programming in support of clinical study reports for all phases of clinical studies.

**Highlight duties:**

- Works closely with Clinical Team to understand analysis data definitions and summary table and listing specifications as they relate to SAS programming tasks.
- Works closely with Clinical Data Management to support data integrity checks and adherence to data definition standards.
- Responsible for the timely completion of SAS programming deliverables without compromising efficient use of I/O and CPU.

**Highlighted requirements:**

- Must be knowledgeable in clinical data presentations, statistical procedures and regulatory guidelines.
- Must be able to perform intermediate techniques using SAS version 8.2 modules including Base, GRAPH, STAT, the macro facility and the ODS. Working knowledge of SAS IntrNet, CONNECT and SHARE desirable.

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**Skill Sets:** SAS Base, SAS Macro, SAS STAT, SAS GRAPH

**Education:** B.S.

**Degree:** Biostatistics, Computer Science, Life Sciences and other related disciplines

**Industry Experience:** Mid-Level, Experienced

**Categories:** Biotechnology and Pharmaceutical

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**Status:** Contract  
**Location:** **Region:** Northern California **City:** South Bay **State:** CA  
**Zip Code:** **Area Code:** 408

**Title:**  
Contract Biostatistician

**Job Description:**

Interested in applying your M.S. in Statistics to the support of clinical drug trials? Have you supported a sponsor or CRO in this endeavor? Are you immediately available for contract employment? If so, read on for an excellent opportunity.

Highlighted responsibilities:

Provides strategic input into the clinical development plan. For assigned clinical development project(s), provides sound strategic, statistical and scientific input on project objectives, experimental design and data analysis to meet project needs and FDA statistical and scientific requirements. Review all project protocols, author protocol statistical analysis sections and generate study randomizations. Work in accordance with cGMP principles, ICH guidelines and Company SOP's. Work with the team to develop, review and approve, as needed, SOP's related to statistics and clinical data management.

Highlighted requirements:

M.S. in Statistics with 4-6 years of clinical trials experience or Ph.D. in Statistics with 2-4 years of clinical trials experience.

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**Skill Sets:** SAS Base, SAS Macro, SAS GRAPH, SAS STAT, Windows NT and Statistical Analysis

**Education:** M.S. or Ph.D.

**Degree:** Biostatistics

**Industry Experience:** Experienced

**Categories:** Biotechnology and Pharmaceutical

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**Status:** Perm  
**Location:** **Region:** Northern CA **City:** East Bay **State:** CA  
**Zip Code:** **Area Code:**  
**Job Code:** mu0805sspa

**Title:**  
Senior SAS Programmer/Analyst

**Job Description:**

Biotechnology is the future of medical breakthroughs. Would you like to join a company who's mission is to develop meaningful biotherapeutics to improve patient's quality of life? Would you like to apply your clinical programming skills in an exciting environment? Wouldn't you really rather be working in the East Bay? Read on for a career opportunity for a SAS professional.

**Summary:**

Principally responsible for providing strong SAS programming skills in support of clinical study reports. Also responsible for supporting colleagues across many functional areas. Ability to prioritize and monitor project timelines and tasks for statistical programming deliverables. Ability to write, document and maintain project standard macros and programs. Peer review of SAS code and validation of programs written by others.

**Requirements:**

B.A. or B.S. in Statistics, Mathematics, Computer Sciences or other related discipline. A minimum of three (3) years of SAS programming experience in clinical trials with a pharmaceutical or biotechnology company. SAS v.8.2 with modules Base, Macro, STAT and GRAPH. Knowledge of relevant FDA regulations and guidelines. Detail oriented with good organizational skills. Good written and verbal communication skills.

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**Skill Sets:** SAS Base, SAS Macro, SAS STAT, SAS GRAPH and UNIX

**Education:** B.S.

**Degree:** Biostatistics, Life Sciences, Mathematics, Statistics, and other related disciplines

**Industry Experience:** Mid-Level, Experienced

**Categories:** Biotechnology and Pharmaceutical

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