

Biostatistician II

Job ID: 00410115

Job Function

Development

Schedule

Full-time

Location

United States-California
South San Francisco

Job type

Regular Employee

Company/Division

Pharmaceutical

Job Level

Experienced

Who We Are

At the Roche Group, about 80,000 people across 150 countries are pushing back the frontiers of healthcare. Working together, we've become one of the world's leading research-focused healthcare groups. A member of the Roche Group, Genentech has been at the forefront of the biotechnology industry for more than 30 years, using human genetic information to develop novel medicines for serious and life-threatening diseases. The headquarters for Roche pharmaceutical operations in the United States, Genentech has multiple therapies on the market for cancer and other serious illnesses. Please take this opportunity to learn about Genentech, where we believe that our employees are our most important asset and are dedicated to remaining a great place to work.

The Position

Responsibilities:

Works with senior biostatistics staff and clinical monitors on clinical development plans, the design and conduct of clinical studies and in the evaluation, interpretation and preparation of study results. As part of a clinical assessment team, collaborates in the preparation and review of clinical assessments. For assigned clinical development project(s), provides statistically sound experimental design and data analysis input to meet project objectives and Health Authority's statistical requirements. Reviews all project protocols, author protocol statistical analysis sections and generate study randomizations. Develops study analysis plans as a team member; lead this effort for selected studies. Reviews case report forms to ensure that protocol objectives are met and project standards are maintained. Develops statistical programs as necessary to perform analyses, and verify data accuracy and validity. Co-author efficacy analysis results in the clinical study report. Supplies statistical input for filing submissions and in response to FDA questions. Provides support for publications. Keeps abreast of new developments in statistics, drug development, and regulatory

guidance through literature review, workshop attendance, etc.

Who You Are

Qualifications:

Ph.D. in statistics/biostatistics. Experience with statistical software packages such as SAS and S-Plus. Sound knowledge of theoretical and applied statistics. Effective communication skills. Effective team contributor. Interested in learning scientific and medical knowledge related to the studies.

Genentech is an Equal Opportunity Employer.