

Principal Research Associate - Biological Technologies

Job ID: 00414698

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

The Principal Research Associate within the Biological Technologies group in Analytical Development and Quality Control will be responsible for all aspects of automation for all phases of clinical trials utilizing precision robotics. The Sr QC Associate will support the execution of vendor IQ, OQ/PQ documents, write and execute instrument, assay validation protocols and IQ, OQ/PQ documents. The associate will also support the development of R&D assays onto high throughput automation platforms utilizing liquid handlers, plate readers and high content analysis instruments. The candidate will identify and evaluate new technologies and work collaboratively with different research and development departments. Expectations will also include writing test procedures, SOPs, protocols, technical reports, and summarizing/presenting data. Additional responsibilities may include presentation of data at internal and external meetings, as well as reports in support of IND/BLA regulatory filings, and supporting collaborations with other groups/sites.

Who You Are

BSc or MSc with 10 years experience in a GLP, GxP, Clinical and R&D development lab. Experience in automation, liquid handling, and process throughput a must. Experience in Assay development, Assay validation, Equipment, Reagent, Computer qualification & validation preferred.

Genentech is an Equal Opportunity Employer.