

**Scientist, Small Molecule, Development Tox**

Job ID: 00390096

**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 Department of Safety Assessment at Genentech provides scientific leadership and plays an active role in the process of drug development from the discovery period through marketed products.

We are seeking a Scientist in the Department of Safety Assessment to support drug development. This role will involve working in a collaborative team environment as part of the comprehensive safety assessment of Genentech therapeutics by providing toxicology representation to small molecule programs.

**Responsibilities:**

The successful candidate for Scientist will provide scientific leadership and play an active role in the process of drug development from late-stage discovery through marketed products. Working in a collaborative team environment, the Scientist/toxicologist leads the safety assessment of Genentech therapeutics by creating a best-evidence synthesis of existing knowledge and comprehensive investigations of toxicologic activity. The successful

candidate will design and supervise toxicology studies to support the goals of small molecule project teams, work closely with Study Monitors, Pathologists, Pharmacokineticists and Pharmaceutics to deliver high quality GLP and non-GLP in vivo study reports and, as needed, represent Genentech in meetings with regulatory authorities. Responsibilities include interdisciplinary project team membership, preparation and review of regulatory documentation, study reports and manuscripts, participation in toxicology and pharmacology initiatives, and other active cross-functional collaborations. Additional responsibilities include investigations into mechanisms of toxicity, proactive management of potential safety liabilities and communication of impact to teams and governance committees. Experience with cross-functional teams and capability to build productive cross-functional collaborations both within and external to Genentech are desired.

### **Who You Are**

A PhD and post-doctorate or equivalent experience in toxicology or a relevant field along with a minimum of 3 to 5 years relevant experience in the pharmaceutical/biotechnology/CRO industry related to toxicology. Experience with biotherapeutics is required. Board certification in toxicology is desirable. The successful candidate must also demonstrate strong decision-making, complex problem-solving, critical data analysis and interpretation, excellent written and verbal communication skills; and the ability to build productive cross-functional collaborations both within and external to Genentech.

Genentech is an Equal Opportunity Employer.