

Graduate Trainee Program in Drug Regulatory Affairs (2 years) – Emerging Markets

Switzerland, Basel-Town, Basel

Job Facts

As a Graduate Trainee within the Pharma Development Drug Regulatory Affairs Group in Basel, you will gain hands-on experience with growing responsibility on the work of Regulatory Affairs professionals who are responsible for providing the Regulatory input and leadership from drug development through to lifecycle maintenance of the product for International countries.

The trainee program is a two-year program that offers you the opportunity to get hands on experience working in the regulatory affairs department of one of the largest Pharmaceutical companies in the world. Your assignments may include both activities in early development and marketed products with your own discrete tasks and responsibilities.

You will be responsible for supporting products and countries within the International Region, working closely with the International Product Partners, Regional Managers and Affiliates. Your tasks may cover activities across the whole development value chain from clinical trial applications through to license applications and maintenance.

As example you may:

- Prepare regulatory submissions in collaboration with other disciplines in the department
- Participate in and support the development and implementation of the regulatory strategies in collaboration with Global and Local colleagues
- Acquire and maintain an overview of relevant regulatory requirements
- Assist in the planning and preparation of regulatory documents
- Become a strong partner to affiliates, finding solutions and challenging the status quo

As part of your assignment you may have the chance to spend some time in another regulatory function as well. And to broaden your experience across the development value chain you will also have the possibility to spend up to six-month period on rotation in another Pharma Development function.

Who you are

You're someone who wants to influence your own development. You're looking for a company where you have the opportunity to pursue your interests across functions and geographies. Where a job title is not considered the final definition of who you are, but the starting point.

We are looking for a highly motivated person who has recently received hers/his Master or PhD degree in life or pharmaceutical sciences (preferably with focus in regulatory affairs) within the last 12 months or is about to receive it very soon.

Furthermore:

- You have already been exposed to the Pharmaceutical Industry (ideally to Regulatory Affairs) during an internship
- You are very collaborative and strong team-work is one of your strengths
- You can handle stressful situations well and you have the ability to follow deadlines
- As we are working in a very dynamic environment, you are a person that demonstrates flexibility, good coordination as well as strong communication skills
- Finally, you are someone who learns quickly and has very good written and spoken English skills

Who we are

At Roche, 94,000 people across 100 countries are pushing back the frontiers of healthcare. Working together, we've become one of the world's leading research-focused healthcare groups. Our success is built on innovation, curiosity and diversity.

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