

(U)HPLC and/or Electrophoresis Laboratory Professional

Scientific Operations

SCOPE

Quality Assistance is a leading analytical CRO based in Thuin, Belgium.

We assist our clients through non-clinical and clinical studies to marketing authorisation, using our state-of-the-art, product-dedicated expertise in analytical sciences.

For each customer and each project, **we design customised solutions**, define analytical protocols, develop and validate specific new analytical methods and perform characterisation, stability, pharmacokinetic, biomarker and immunogenicity studies as well as batch release testing, **in order to evaluate the Quality, Safety and Efficacy of the given drugs.**

You will be part of a team involved in projects outsourced by our clients.

Those projects are related to the development, validation and/or application of analytical methods based on a protocol for the characterisation and quality control of biomolecules.

YOUR MISSION

You will be part of a team of 5/6 people and report to the Technical Leader Chromatography.

You will work on the bench and you will perform daily laboratory activities.

Depending on your level of expertise, you will work on projects related to either application, development and/ or validation of analytical methods.

This implies:

- Analysing samples according to specifications and ensuring traceability of every analysis,
- Processing data,
- Writing reports and associated supporting documents,
- Presenting/discussing results.

Depending on your level of autonomy, you will be requested to ensure the following tasks:

- Validating raw data,
- Training team members and others on technical skills,
- Taking part in the writing/preparation of protocols,
- Supporting audits and investigations.

PROFILE

Scientific background ideally with knowledge of biologics (antibodies, peptides, proteins, ADCs...).

You have hands-on technical expertise in either:

- HPLC / (U)HPLC,
- Capillary and/or gel Electrophoresis (cGE, cIEF, SDS-Page).

You have work experience in a regulated environment ideally in the pharmaceutical industry.

You are experienced **working with software** such as Empower, iCE3, 32 KARAT. Quantity one is a plus.

You are fluent in **French** and have a good command of **English** (spoken and written).

Why join *Quality Assistance*?

Do you want to thrive in a professional setting that still maintains a human touch? Are you looking for a working environment based on mutual respect, communication and support, where it is good to live and work?

Apply now to join our analytical CRO! We are pursuing a common goal: to accelerate access to new medicines.

You will benefit from a permanent employment contract with a **competitive compensation package** in line with the industry, including **many fringe benefits** (meal vouchers, hospitalisation and outpatient care insurance, group insurance, bonuses, and for certain positions, a company car and petrol card).

As soon as you start your job, you will follow a comprehensive **training programme adapted to your profile and role.**

Did you know that in 2021 we welcomed, and trained 45 new colleagues? We also promoted 23 team members (vertical mobility). 13 positions were filled by internal candidates. We provided 1,275 general and technical training sessions as well as 696 practical training courses.

We offer **multiple opportunities** so that you can integrate yourself into your new work environment and get to know your new colleagues (after-works, sports and recreational activities, team building, department dinners, end-of-year parties, BBQs, events for families, etc.). **We pamper our team members and take care of them:** free sports lessons, free fruit and sugar-free drinks, daily delivery of lunches and bread, free car wash, ironing service via service vouchers, books and board games available, and much more...

You will join a company that listens to your needs and your suggestions!

About *Quality Assistance*

Quality Assistance is a leading **European Contract Research Organisation (CRO)** providing the pharmaceutical industry with all the analytical services required by **EMA** and **FDA** regulations for the development and marketing of innovative human medicinal products.

From candidate selection, through non-clinical and clinical studies, to marketing authorisation, *Quality Assistance* provides **customised solutions** for its clients:

- We define **analytical protocols**;
- We develop and validate **specific new analytical methods**;
- We perform **characterisation, stability, pharmacokinetic, biomarker and immunogenicity studies as well as batch release testing**.

These tests are performed in order to evaluate the **Quality, Safety and Efficacy** of the given drugs.

With **40 years' expertise** at the forefront of analytical sciences, *Quality Assistance* holds a **unique place on the market thanks to:**

- **All of its laboratories located on one site** (Donstiennes, Belgium);
- **250 highly qualified professionals**;
- A **wide range** of analytical methods and state-of-the-art equipment.

The *Quality Assistance* environment is **GMP, GLP and GCLP/GCP compliant**.

Visit <https://www.quality-assistance.com/quality-assistance/leading-analytical-cro> to learn more.

How can you apply?

Send your application (**JOB186.2**) now to Mrs Isabelle Lebrun, Talent Acquisition Manager, to recrutement@quality-assistance.be or consult the Careers page on our website <http://www.quality-assistance.com/careers>.

Address: Technoparc de Thudinie 2, 6536 Donstiennes, Belgium