WE ARE HIRING

QC VALIDATION SUPERVISOR

First, did you know that AKKA Belgium is going to be AKKODIS soon? Because of the the merger with Modis.

AKKODIS is a global leader in the engineering and R&D market that is leveraging the power of connected data to accelerate innovation and digital transformation.
As part of the development of the activities at one of our clients we are looking for a QC VALIDATION SUPERVISOR

So what do you do all day?

As QC Validation Supervisor, you will be able to perform project management, while keeping a job with scientific thinking.

It's a hybrid job that will allow you to combine work at home and on customer site. You will participate to regular teleconferences and meetings with customer and internal team as appropriate. But you will also be required to go to the laboratory when necessary (without performing any manipulation).

CONTACT US jobs.belgium@akka.eu - www.akkodis.com

We are Akkodis

Your responsibilities :

- Participate to the reflection of the validation design and the treatment of experimental data in collaboration with QC technicians, test supervisor and QC manager
- Write validation analytical documents (validation protocols/reports, qualification protocols/reports, Rational documents, Comparability documents...)
- Participate to transversal interactions with other teams in QC Department and different other stakeholders
- Participate to the investigation phases in case of acceptance criteria unmet;
- Participate to the administrative tasks of the team (Continuous improvement, complete the follow-up files, etc..)
- Participate to the management of quality element (CAPA, eCC, Deviation, etc) in relation to validation projects
- Guarantee the application and maintenance of adequate cGMP rules in his work area.

What skills do you bring to the table?

- You are graduated as a bachelor's and/or master's degree in Chemistry, Biochemistry, Bioengineer or Pharmaceutical. Or, you have a significant experience in a similar role in pharmaceutical industry.
- You have a good expertise in some analytical methods, like: ELISA, Western blot, Liquid chromatography, Spectrophotometric assay, Kjeldahl, Karl Fischer, bioburden test, rapid diagnostic tests, etc. (Knowledges in laboratory skills is a plus).
- You have also good knowledge in validation of analytical methods and cGMP environment
- You are interested by the biochemical, physiochemical, virology and/or microbiology
- French fluent and knowledge in English
- You master the Microsoft Office software.
- You enjoy meeting and speaking with people, presenting your results, and advising customers,
- You are proactive, solution-oriented, open-minded and flexible

What's the offer you can't refuse?

As an AKKODIS Team member, you will be:

o In charge of diverse transversal analytical validation projects in a cGMP environment,
o Part of a dynamic and collaborative team of validation supervisors under the direction of a Service Manager,
o Onboarded in your position via a buddy process,
o Supported in your career by your Business Manager,
o Actor of your training plan and your personal and professional development,
o Benefiting from a permanent contract,

Benefiting from a competitive salary package including several extra-legal benefits

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