Belgium Jobs Expertini®

Analytical Scientist

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Company: UCB

Location: Braine-l'Alleud

Category: computer-and-mathematical

Make your mark for patients

We are looking for an **Analytical Scientist** to join us in our Pharma science organization based in **Braine-l'Alleud, Belgium**

About the role:

As the Analytical Scientist, you will lead and deliver on all analytical development activities to ensure a sufficient knowledge of the drug candidate in its pre-clinical, clinical and commercial phases.

The development stage covers products emerging from R&D function, entering pre-clinical and clinical pipeline and extends to lifecycle management of UCB's major marketed products, including transfer exercise to QC's worldwide.

What you will do:

You will be primarily responsible for implementing, developing, and leveraging the analytical technologies, platforms, processes, and knowledge management required to efficiently solve existing and future analytical development challenges.

You will implement/develop/maintain technological analytical platform(s) based on spectroscopy/spectrometry/physical measurement of Drug Substances (DS) and Drug Products (DP)

You will support the mass spectrometry activities by developing and improving the characterization, quantification and structural identification of compounds in DS and DP.

You will support the development of the API and DP processes by providing relevant platform, technology, analysis and reporting.

You will be responsible for coaching the analysts/scientists involved on specific analytical platforms /technologies /processes.

You will be actively involved in method and technology transfer and regulatory submissions preparing with internal and external business partners.

You will actively participate to the development roadmap and guidance, technical implementation, scientific and operational workflow definition, automation of the process/technology you will be responsible for.

You will implement digital solutions (equipment, process and data) that deliver value in terms of efficiency and compliance.

You will be responsible for the compliance with relevant regulations (GMP, HSE, ISO140001..) and SOPs

For this position you'll need the following education, experience and skills:
You have knowledge in pharmaceutical DS and DP analytical development and technologies for NCE.

You have proficiency with QbD development and data processing/management tools.

You have a detailed understanding of regulatory and GMP quality constraints.

You have in-depth knowledge in key analytical techniques including liquid and gas chromatography mass spectrometry with multiple ionization modes, … (working on Thermo Scientific instruments (Orbitrap) and knowledge of Chromeleon is a plus).

You have demonstrated capacity to work in a matrix organization in different technical areas.

You have demonstrated capacity for developing and managing projects with outsourced resources.

You have demonstrated good interpersonal, verbal, and written communication skills with ability to balance multiple priorities.

You have key technical backgrounds (depending on scientist key accountability(ies): EU, US and JP regulations, small-molecule chromatography (HPLC/GC) and detection techniques (UV/Vis). You'll need to demonstrate the ability to work effectively in a matrix organization and manage projects with outsourced resources. MS); PAT and chemometrics; NMR and MS spectrometry; dissolution techniques, USP, EP, and JP pharmacopeias.

You are able to find technical solutions and activate all appropriate internal /external experts to solve the issue.

You are able to efficiently translate acquired data into knowledge answering the project needs.

You are able to independently manage scientific tasks in order to meet deadlines within given constraints in resources, e.g. headcount, equipment, time, etc.

Are you ready to ‘go beyond’ to create value and make your mark for patients? If this sounds like you, then we would love to hear from you!

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