

## Industrial Pharmacist

### About

Amaris is an independent and international Management and Technologies Consulting Group created in 2007. Present in more than 45 countries and 60 offices worldwide, it offers proximity support to its clients, wherever their locations. Amaris' added value lies in its teams' quality, in their attention to detail, and in their commitment to always deliver the most innovative solutions.

### Consulting

We are currently looking for a critical mind to join our expert team based in Belgium. Within Amaris, we are providing consulting to our partners. This means that we do not only deliver technical assistance, but we also excel at providing them with advice and innovative improvements in order to improve their business processes. Therefore we are looking for technical experts with great interpersonal skills in order to provide state of the art consulting.

### Your role

You will join Amaris as a consultant and be part of a team coordinated by your Manager. Together you will design technical and/or management solutions for our clients in pharmaceutical, medical devices and biotechnology industries. You will be present on client's plant to work on mid and long term project in Deviation and CAPA. As a CAPA & Deviation Coordinator you will be working in a team of consultants who are experts in the specific field of: Pharmaceutical, Medical Devices or Biotechnology; within these markets they have a specialization in: Quality Assurance, Deviation and CAPA management, Quality Management System, Aseptic manufacturing, GMP management.

Your role as a CAPA & Deviation consultant will be to suggest corrective actions and to identify the root causes of identified problems to prevent their recurrence. In this position you will be responsible for a variation of CAPA & Deviation aspects within the projects of our clients such as:

- Support business unit operations teams in their Deviation and CAPA management.
- Support operations teams in the deviation management: write deviations with the right appropriate level of quality, collect data to perform root cause analysis using appropriate tools, assess the deviation's impact and the associated risk.
- Identify efficient CAPA aligned with priority in order to fix problems permanently.
- Implement CAPA and check their effectiveness by performing audit.
- Implement continuous improvement projects linked with root causes of deviations: define solutions with operators, technicians and supervisors.
- Carry out trend analysis on recurring problems and propose a strategy of improvement.

A graphic consisting of two overlapping orange shapes: a larger solid orange trapezoid and a smaller white trapezoid with a thin orange outline, partially overlapping the bottom right of the larger one. The text "Sparkling Consulting, Smarter Innovation" is written in white on the larger orange shape.

Sparkling  
Consulting,  
Smarter  
Innovation

## Profile

You have received a university degree (Msc) or equivalent. You know the legal requirements for pharmaceutical industry (cGMP) and have a work experience in such environment. You are flexible, able to adapt yourself to some different process and to take into account the various priorities of your representatives. You have an analytical and creative mind and you are familiar with the root cause analysis tools (**5M method/Ishikawa diagram, 5 Why, process mapping**). You are proactive, have good communication skills, and are able to change people mindset in the way of efficient working. You are proficient in English (spoken and written) and French!

## What you can expect from this experience

As a first mission in consultancy, the Deviation & CAPA management project will provide you:

- A transversal knowledge of a major company's functional departments.
- The possibility to create your own network through the numerous interactions with experts and specific point-of-contact.
- A challenge to your problem solving abilities.

- An overview of a Quality Management System with a growing importance for health agencies.
- A knowledge of different manufacturing process for parenteral drugs.

## What we offer you

You will join an international consultancy group in continuous growth and with a strong presence in Europe, America and Asia. You will have the opportunity to develop your career as professional consultant, specialized in quality assurance, inside a sparkling company. A company which is able to provide you rapid career growth, by participation in projects that involves cutting-edge technologies for the main international pharmaceutical companies. You will become part of an international employee community across and within our offices, this will enable you to expand your professional network. You will be offered an industry competitive remuneration package, intensive training programs and an indefinite contract.

Apply at [pcools@amaris.com](mailto:pcools@amaris.com)