Inspection Management Quality Lead

Location : **Braine-l'Alleud, Belgium**
Working model : **Hybrid – approximately 2 days/week on-site**
Contract : **Full-time**

Are you ready to take your career to the next level?
Do you thrive in environments that challenge and inspire you?
Would you like to grow within a team that values both excellence and enjoyment at work?

ABOUT ELMEA CONSULTING

Elmea Consulting, a brand of Taleo Consulting, is a life sciences consulting firm supporting healthcare companies from development to distribution.

DEVELOPMENT

From proof-of-concept to licensing, we guide our clients through each stage of development, ensuring efficient, science-driven, and regulatory-compliant pathways.

PRODUCTION

We help healthcare organizations accelerate market access while maintaining the highest standards of quality, safety, and regulatory compliance.

What Will You Do?

At **Elmea Consulting**, we believe that advancing healthcare starts with **investing in people**. When our consultants grow, feel empowered, and are supported, they can make a real impact for our clients and, ultimately, for patients.

Our teams are made up of curious, committed, and passionate professionals who thrive in a collaborative environment. By developing their expertise and careers, they actively contribute to the success of healthcare and life sciences organizations at every stage of innovation.

Our success is built on the **talent, collaboration, and commitment** of our people and we're now looking for a **Pharmacovigilance Quality Lead** to join our journey.

Mission

As a Pharmacovigilance Quality Lead, you will join an international pharmaceutical company based in Braine-l'Alleud and support global pharmacovigilance quality, regulatory compliance, and inspection readiness.

This role is ideal for a senior professional with strong expertise in PV Quality Assurance, GVP compliance, auditing, and regulatory inspection management.

Challenges we trust you with

· Lead and coordinate pharmacovigilance inspection readiness activities, including preparation for and hosting of regulatory authority inspections.

· Ensure compliance with global pharmacovigilance regulations and guidelines, including GVP, GCP, FDA, EMA, and MHRA requirements.

· Oversee and continuously improve PV quality management systems, processes, and procedures.

· Coordinate PV audits, inspection documentation, findings, and follow-up activities.

· Manage CAPAs, ensuring appropriate corrective and preventive actions are implemented and tracked.

· Collaborate with global cross-functional teams and external service providers to maintain quality standards and regulatory compliance.

· Support continuous improvement initiatives and promote a strong quality and compliance culture across the organisation.

What we value

You have:

· 8+ years of experience in Pharmacovigilance Quality Assurance, PV Compliance, GVP, or PV Auditing.

· Proven experience managing and/or hosting pharmacovigilance inspections (mandatory).

· Strong knowledge of global PV regulations, particularly GVP, and applicable regulatory requirements.

· Experience with PV quality systems, audits, CAPAs, inspection preparation, and documentation management.

· Experience working within the pharmaceutical or biotechnology industry.

· Ability to collaborate effectively with international stakeholders and manage quality-related activities across teams.

· Excellent written and spoken English (mandatory).

Nice to have:

· Experience with Veeva and/or Argus.

· Experience overseeing external vendors or service providers.

· Previous experience in a global, cross-functional environment.

· French language skills.

Our Vision at Elmea Consulting

We aim to build a leading healthcare and life sciences consulting firm driven by expertise, collaboration, and impact. By attracting top talent and fostering the right environment, we empower our teams to grow and deliver meaningful value to healthcare innovation.

Why Join Elmea Consulting?

Joining **Elmea Consulting**, a brand of **Taleo Consulting**, means becoming part of an international and dynamic team. We offer real career growth, continuous learning, and the opportunity to work on high-impact healthcare projects.

If you're ready to grow with a firm that values excellence and collaboration, **send us your CV**.