Technical Standards & Regulatory Affairs Engineer

Location: Belgium (on-site)

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 **Technical Standards & Regulatory Affairs Engineer** to join our journey.

We are looking for a Technical Standards & Regulatory Affairs Engineer to support one of our clients active in the high-tech industry. The consultant will play a key role in ensuring that products and processes comply with applicable technical standards and regulatory requirements across relevant markets.

elmea
LIFE SCIENCES CONSULTING

- Identify, analyse, and monitor applicable technical standards and regulatory requirements (international and sector-specific)
- Support product development and engineering teams in integrating compliance requirements from early design stages
- Lead and coordinate regulatory submissions and technical documentation packages
- Maintain and update the regulatory affairs database and standards library
- Liaise with regulatory bodies, notified bodies, and standards organisations
- Provide internal training and guidance on regulatory and standards compliance
- Participate in audits and support corrective action processes

Required Profile

- Engineering degree (electromechanical, biomedical, physics, or equivalent)
- Experience in regulatory affairs or technical standards within a high-tech, medical device, or industrial environment
- Familiarity with standards such as IEC, ISO, CE marking, FDA requirements (an asset)
- Strong analytical and documentation skills
- Fluent in English; French and/or Dutch is a plus
- Ability to work in a structured, process-driven environment

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**.