Global Regulatory Intelligence Manager

Zug, Switzerland

Are you an experienced Regulatory or Quality professional wanting to “do something new”, but not sure what you can do with your experience?

We have a great opportunity which is something quite different - We are looking for somebody who understands the medical laws, standards and guidance concept and supports to understand and implement regulations.

THE COMPANY

This role is all about having your “finger on the pulse” of regulatory changes worldwide.

Located in the company’s headquarters you will be sharing the corridors with the top people and be reporting to the Global Regulatory Affairs Director which will offer you great exposure!
In this unique role you will be responsible for monitoring relevant trends, regulations and hot topics globally by attending networking events and participating in working groups to improve the company’s internal processes and guide it into the future. In this role you’ll work independently, learn about new markets and participate in company special projects i.e. transition to MDR.

This is a corporate role across the whole company and a great opportunity for someone who wants to directly impact the business and work autonomously using their experience in a completely different context.

As Global Regulatory Intelligence Manager you will:

- Guide and consult teams throughout product development projects and product lifecycle on standards and questions related to risk management/product safety
- Monitor and analyse worldwide registrations, trends and hot topics that may impact the company in terms of products and processes across all portfolios
- Prioritise and distribute collected information to necessary internal partners (Quality, Regulatory and Marketing departments) and work cross functionally to review and prepare for new/updated laws
- Attend and participate in networking groups, working groups and medtech events to establish and maintain external relationships and exchange information
- Establish and maintain all relevant laws, regulations, guidelines, directives for all products in a system
- Participate in special projects such as the transition to MDR

ARE YOU ...

1. A regulatory OR quality professional with 5+ years experience in the medical technology industry?
2. Strong experience with CE marking OR 21CFR/820 and ISO 13485:2016
3. Strong English skills

Please contact hello@elemed.eu for a confidential career discussion.