# REGULATORY AFFAIRS SPECIALIST

*Brussels, Belgium*

monia@elemed.eu

## THE COMPANY

Do you want to be part of one of the world's biggest healthcare companies? With an extremely dynamic growth strategy and great future prospects, this is definitely a company where you won't get bored!

This is a great opportunity to join their well-established regulatory department and work on globally leading products. Come and join an exciting company whose new ideas are transforming the future! With 75 thousand employees across the globe, they are committed to improving safety, efficiency, and better diagnosis of diseases.

Dedication and passion are the driving forces to face the challenges of developing innovative solutions for global issues and you could be a part of it!

# THE OPPORTUNITY

As a Regulatory Affairs Specialist, you will support the Benelux team in regulatory activities for different product categories. You will be part of France and Benelux Regulatory Affairs reporting to the RA Manager.

By joining the company in this role you will gain access to training and development opportunities and the ability to gain excellent exposure to medical device and healthcare career options.

This will be a great way to further your career development in a fulfilling role!


# YOUR RESPONSIBILITIES:

- Updating of regulatory documents and internal databases
- Handling customers' questions
- Monitoring Regulatory changes in Benelux
- Attend internal meetings with EU Regulatory Affairs team or project teams
- Support cross-functional teams for products launches
- Reviewing promotional materials in French and Dutch

# YOUR QUALIFICATIONS:

- Minimum of 2 years experience in Regulatory Affairs
- Good communication skills verbal and written in French, Dutch, and English

# INTERESTED TO EXPLORE THIS FURTHER?

If you are interested in this exciting role, please send your application directly to **monia@elemed.eu**

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