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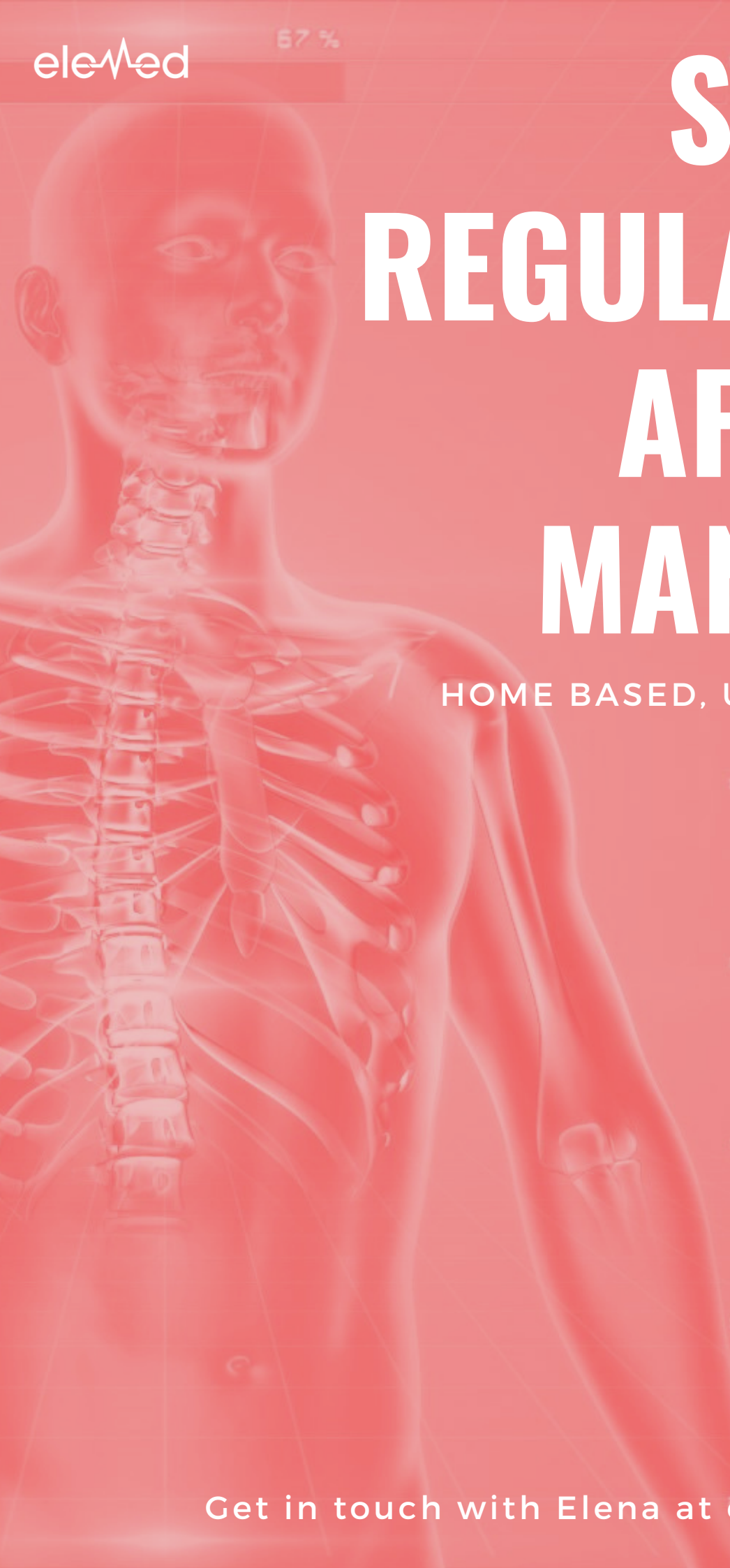
SENIOR REGULATORY AFFAIRS MANAGER

HOME BASED, UNITED KINGDOM



Get in touch with Elena at elena@elemed.eu

THE COMPANY

Come and join a global leader in cardiovascular medical devices as their new Senior Regulatory Affairs Manager!

This is an opportunity to join a stable company with mature processes and structure but the pace and agility of a start-up. With 15,000 employees worldwide and a strong company brand perception, they are known all globally and are really at the forefront of the most cutting edge technology in the cardiovascular field. But it doesn't stop there...This company is continuing to invest in its people and R&D and has plans to grow even further over the next 5 years. With that growth comes your opportunity to evolve, develop and advance your professional career alongside some of the best professionals in the industry.

THE OPPORTUNITY

This is a high-visibility, advisory position as Senior Regulatory Affairs Manager, reporting to the Senior Director Regulatory Affairs. You will sit at the table of directors, interface with commercial VPs and really have the opportunity to make an impact within the business. You will navigate complex regulatory challenges such as MDR and BREXIT for the two business units and advise the key stakeholders on strategy and technical elements building trust internally and externally.

YOUR QUALIFICATIONS:

- 6+ years experience working in Regulatory Affairs for medical devices
- Experience with Class III OR minimum Class IIb implantable medical devices
- Previous project OR people management experience
- Fluent speaking, reading and writing in English

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YOUR RESPONSIBILITIES:

- Develop, lead and implement complex regulatory affairs strategies in core countries globally
- Interface, liaise and build relationships with internal and external parties such as senior stakeholders, notified bodies, competent authorities
- Advise key stakeholders from a technical and regulatory perspective to help achieve current and future business goals
- Lead and develop one RA specialist that will report directly to you and support you in your activities
- Navigate complex regulatory issues such as BREXIT and MDR and ensure compliance with the relevant regulations
- Liaise with commercial VPs and ensure that the European voice is heard to the wider global teams

INTERESTED TO EXPLORE THIS FURTHER?

Please send your CV to elena@elemed.eu to arrange a confidential career discussion.