AMSTERDAM, THE NETHERLANDS

RA/QA SPECIALIST

ELENA@ELEMED.EU

COMPANY

This is a fantastic opportunity to join an innovative, high-tech startup working with groundbreaking software.

You will have a real impact on the company and support their creativity, whilst having the chance to really own their regulatory compliance strategy. They are passionate about improving and developing their software in their own specific industry.

As a RA /QA Specialist, you will be part of their project to expand internationally and lead their transition from MDD to MDR.

There will be the chance for you to develop your own method of work and take on a huge learning opportunity through training and new projects.

This is also a unique opportunity to fast forward your career and create your own network by leading monthly meetups.

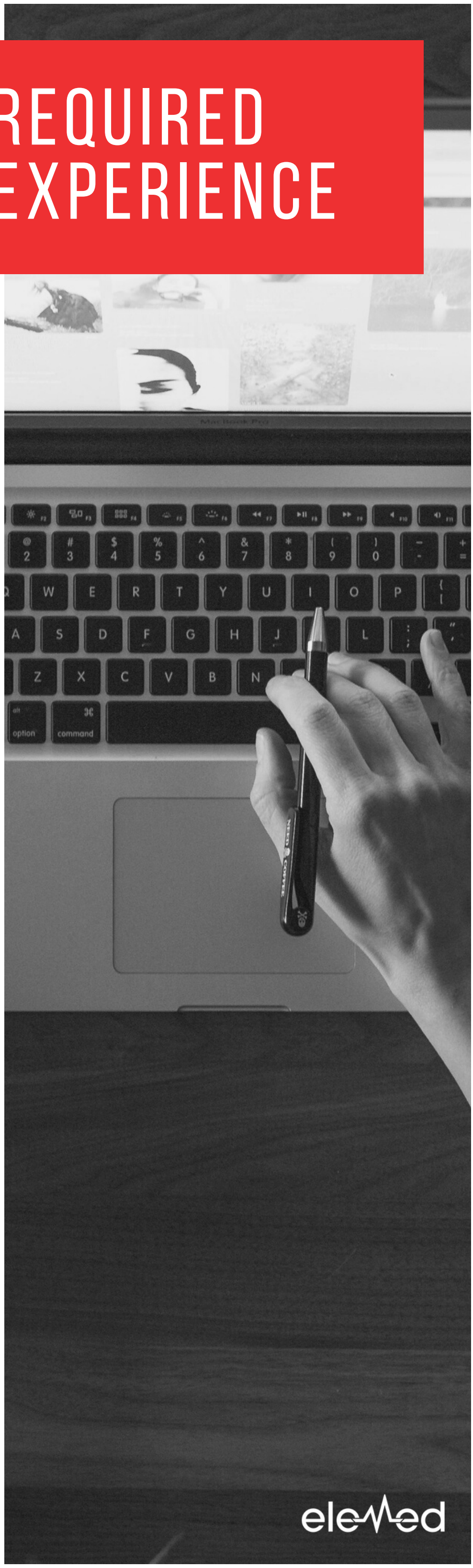
Still looking for more reasons to join this company? Well, you will receive daily lunch, free drinks, and fresh fruit!

RESPONSIBILITIES

As RA/QA Specialist, your responsibilities will be to:

- Take charge of developing and delivering the company's regulatory strategy whilst also championing quality and compliance within the business
- Take ownership of the company's QMS
- Work closely with the senior leadership team
- Work closely with internal teams to create and document cross-functional processes
- Lead regular audits of company procedures and processes - both internally and externally
- Lead the medical software certification process and oversee continuous compliance (MDR)

REQUIRED EXPERIENCE



Do you have the following experience? If so, we want to hear from you!

2-5 YEARS
EXPERIENCE

in Quality and Regulatory in the Medical Devices Industry

MDD TO MDR
KNOWLEDGE

Do you have knowledge of the transition from MDD to MDR?

ISO 13485
EXPERIENCE

Do you have experience with QMS (ISO 13485)?

ENGLISH
SPEAKER

Are you fluent in English?

ARE YOU INTERESTED IN THIS POSITION?

SEND YOUR APPLICATION TO ELENA@ELEMED.EU