

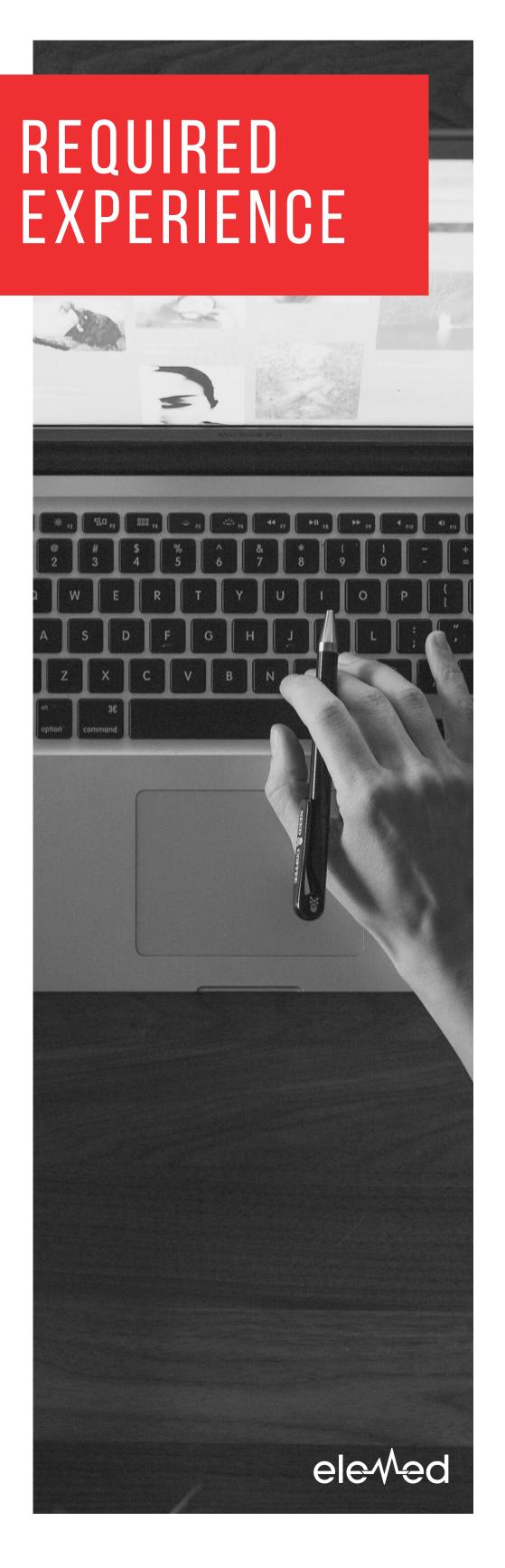


This is a fantastic opportunity to join an innovative start up with groundbreaking software. You will have a real impact in the company and support their creativity. They are passionate about improving and developing their software in their own specific industry.

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

That will be the chance for you to develop your own method of work, and will acquire further knowledge through training.

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 a daily lunch, free drinks, and fresh fruits!



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?

