PARIS, FRANCE

Regulatory Affairs & Quality Specialist

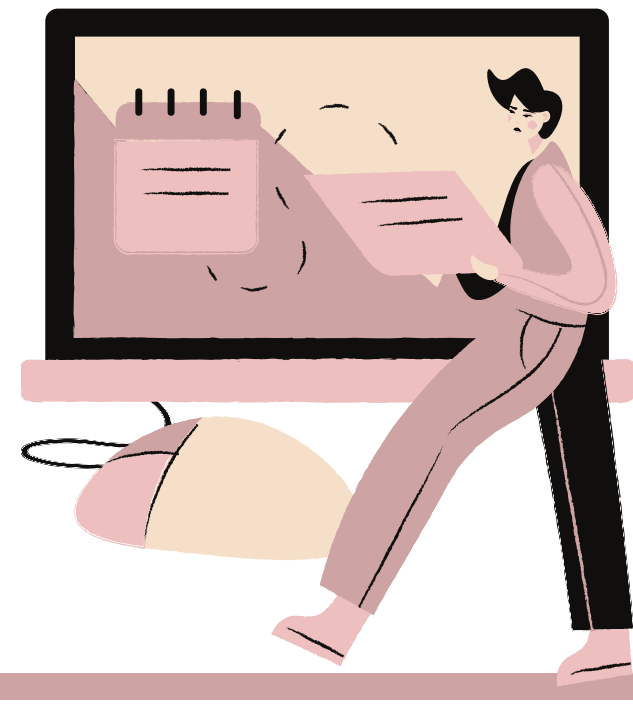
CLARISSE@ELEMED.EU

elemed

COMPANY

We will consider candidates with either Regulatory Affairs or Quality experience.

Join a human-size consulting agency at the heart of the health industry. This company is known for their expertise, dynamism and reactivity, and offers regulatory, compliance, QMS, market access and training solutions. If you enjoy project management and working in a versatile environment this role is for you.

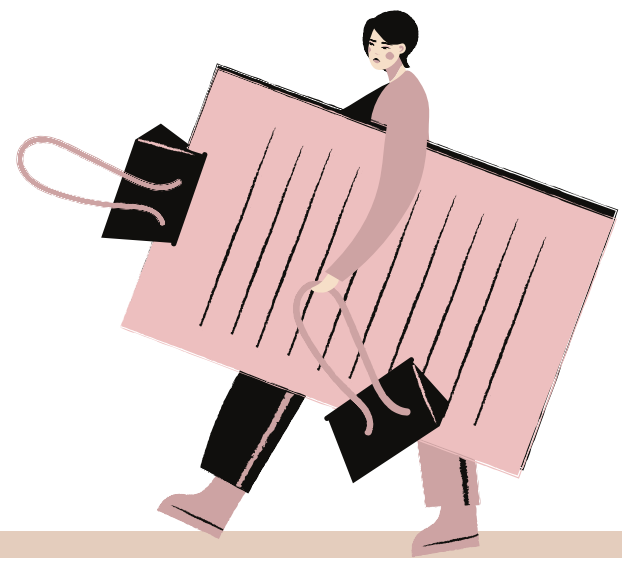


OPPORTUNITY

You will have the chance to join a team that values solidarity and teamwork.
You will also benefit from an attractive remuneration and social advantages (31 days of holidays, PEE, PERCO, and 2 yearly seminars (winter and summer).
And you will have the opportunity of working in the incredible Intra Muros offices in Paris with a gym, cantine, rest zone and much more at your disposal.

RESPONSIBILITIES

- Developing regulatory strategy project, quality and clinical
- Managing regulatory projects (Technical documentation, CE marking, 510k (USA)
- Managing and set up quality system ISO 13485
- Managing audit (ISO, MDSAP, distributor, manufacturer)
- Developing and/or delivering training.
- Participate in business strategies to develop and impact the growth of the businesses



REQUIREMENTS

- *A degree in pharmaceutical, Ingenieur or science*
- *Experience in regulatory, quality or/ and clinical*
- *Fluent in English and full professional capacity (read, write and speaking)*
- *3 years experience*

INTERESTED IN FURTHER CONVERSATION?

PLEASE SEND YOUR CV TO CLARISSE@ELEMED.EU TO ARRANGE A CONFIDENTIAL CAREER DISCUSSION.