Senior Quality Assurance consultants to Stockholm or Malmö

Sofus has grown to become the leading life science consultancy firm in the Nordics. We are now expanding our services in GMP and medical device and are looking for experienced QA-consultants to our offices in Stockholm or Malmö.

We are now part of Xendo, an international, growing consulting company that is well established in regulatory affairs, product development and GMP/GDP, as well as pharmacovigilance for both (bio)pharmaceuticals and medical devices.

We are big enough to have built a good reputation after working with more than 250 pharmaceutical and biotechnology companies worldwide. Nevertheless, we are small enough to ensure that all employees are recognized and their contributions valued and appreciated.

With us, you will work with a broad variety of customers from the early stage of development to mature products. As a QA-consultant will you support our customers with services such as strategic advice, project management, auditing, QA-agreements or as acting QP/RP.

Your background
We are looking for someone that has worked with QA in a GMP environment for at least 3-5 years and meets requirement for QP according to LVFS 2004:7. You are looking for new opportunities and want to combine your knowledge and skills with the latest in drug development and manufacturing. Experience from validation/IT-validation, medical device, QA-supplier management, auditing, IMPs, GLP, biologicals or acting QP/RP is an advantage.

We offer an independent role with challenging and varied tasks, customer contacts and with great opportunities to actively participate in the company's continued development.

For more information, contact Anders Nyholm, tel: +46 8 56 00 20 36.
Please send us your CV and a personal letter as soon as possible, but at the latest by 8th January 2018, to: recruitments@sofus.se

We will hold interviews during the application period.

Sofus offers services in European regulatory affairs. We believe that a reliable expertise in regulatory affairs is the most important factor in reaching the market successfully. Our driving force is to use our expertise to reduce regulatory risks in all parts of the product life cycle.

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