Quality Assurance Specialist

We are looking for a highly skilled Quality Assurance Specialist to join our Quality Assurance team in Denmark.

About us

BioPorto is an in vitro diagnostics company that provides tests and antibodies to clinicians and researchers around the world. We use our antibody and assay expertise to transform novel research tools into clinically actionable biomarkers that can make a difference in patients' lives. BioPorto is headquartered in Hellerup, Denmark and is listed on the NASDAQ Copenhagen stock exchange.

Our people

BioPorto has approximately 35 dedicated employees globally, most of whom work in Hellerup. Our organization can be characterized as energetic, passionate, and committed to providing a supportive work environment. If you wish to be part of our organization, we are eager to hear from you.

Your role

We are looking for a dedicated employee who will help ensure the quality of our products. You will be part of a team consisting of three committed professionals who, in collaboration with sales, marketing, R&D and production, will handle quality assurance and registration tasks within production and sales of antibody products and in vitro diagnostic (IVD) equipment. If you like to have an impact on quality, have a clear ability to maintain overview but at the same time a sense of detail and a wish to make a difference in a small company, it maybe you that we are looking for.

The position is for occupation no later than the 1st of December.

Key responsibilities

- Continuous optimization and standardization of the department's and the company's processes and existing procedures
- Processing of complaints, product, and process deviations as well as corrective actions
- Review and approval of validation protocols and reports
- Release of products for packaging and sale
- Continuous monitoring of the company's maintenance of development / production equipment
- Internal and external audits as well as supplier management
- Assist the company in maintaining and developing a proper quality spirit through training of colleagues
- Participation in the company's transition to new European rules
- Be part of the implementation of a new IT tool for the QMS.

Your qualifications

- Academic background (engineer, cand. Pharm., Cand. Scient. Or similar) and preferably knowledge of immunochemistry
- Several years of experience from a QA function or alternatively has good knowledge of production and QC processes in a quality-assured company, preferably ISO 13485 knowledge and FDA 21 CFR part 820
- Has experience with validation, handling of deviations, suppliers and / or audits

- Ability to write, read, and speaking in English at a professional level.
- Can work independently while maintaining a high quality and attention to detail.
- Ability to work in a dynamic environment where effective teamwork is critical to succeed.
- Can stay focused and meet timelines while managing multiple activities.

As a person, you are structured, thorough, and responsible as well as good at meeting deadlines. The position also requires that you a capable of having influence and are not afraid of being "hands-on". Finally, you are used to communicating constructively and entering into collaborations with a focus on finding the right solutions. You are able to simplify complex situations and are good at stakeholder management.

Our offer

We offer a wide range of professional, social and financial employee benefits in addition to exciting and dynamic position, with great opportunities for personal and professional development. You will work in a small and informal team, with interface to other departments (Production, R&D, etc.), where your expertise and opinion will be valued.

Additional Information

If you have any questions, please contact Head of QA, Asger Dalgaard (ada@bioporto.com or +45 45 29 00 00)

Please send your application and CV **in English** using the link below no later than **25th October 2021**.

Your application will be treated with confidentiality.

Interviews will be performed in parallel to the application period. We reserve the rights to proceed with the employment process if the right candidate is identified during this period.