Experienced QC Scientist for Production Team

We are looking for an experienced and skilled QC Scientist to join our Production 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. We are 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 and responsibilities

Our Production team produces antibodies and assays (ELISA, turbidimetric immunoassays and lateral flow assays). As a QC Scientist you will be involved in various daily operational tasks, in a dynamic and challenging environment. Your main responsibilities will be to assist with:

- Maintenance of the department's documentation and equipment
- QC, data processing and Certificates of Analysis (CoAs)
- Validation of new equipment, as well as novel procedure for both manufacturing and testing
- Analytical method validation, stability reports and protocols
- Writing CAPAs, non-conformities, and customer support
- Annual review of suppliers
- Updating lab inventories for both manufacturing and QC testing

Your profile

You have 3+ years of QC and validation experience in the medical device or diagnostics industry. It is also a requirement for this position to have a relevant academic background within pharmacy, biotechnology, biology, chemistry or similar. It is of utmost importance to have a self-motivating personality with a pragmatic attitude and approach to the daily work — without compromising quality. The environment is dynamic and challenging, which requires the ability to find solutions, deliver on promises and give our customers a positive experience. Therefore, as a person you are flexible and able to meet timelines while managing multiple activities. You can work independently, ensuring high quality standards and attention to detail. You enjoy a dynamic working environment where effective teamwork is critical to succeed.

Moreover, you have experience with several of the following areas:

- Production of diagnostic assays or other IVD Medical Devices and working in a GMP-regulated environment
- ELISA, clinical biochemistry analysers, and/or lateral flow assay
- Recombinant protein expression in *E. coli* and protein purification, specifically usage of ÄKTA purification systems
- Cell culturing and monoclonal antibody expression
- Antibody conjugation/modification
- Knowledge of Master Control

BioPorto has international customers, and our company language is English. It is therefore a required ability that communication in both written and spoken English is fluent.

Our offer

We offer you an 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 (R& D and QA/RA, etc.), where your expertise and opinion will be valued.

Additional information

If you have any questions, please contact the Head of HR, Karen Stendal (ks@bioporto.com, +45 45 29 00 00).

Please send your application and CV **in English** using the link below no later than **13 December 2021**.
www.bioporto.com/careers/

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.