To support our team in Hamburg
we are seeking a qualified

QM Manager (m/f/d)

***We are** a physician-led, integrated global oncology company. Our goal is to unveil the complex mechanisms of cancer, in order to support precision oncology.*
Through our three divisions, IndivuServ, IndivuType, and IndivuTest, we offer specialized products and services that support our customers in biomarker and target discovery, drug development, clinical trials, individualized therapy, and more.
With more than 200 employees across Europe, North America, and Asia, we are an internationally minded, future-oriented company with an ambitious growth strategy and a strong international reputation. You will work with an innovative team of leading laboratory and bioinformatics scientists, product managers, and clinicians. Become part of our community, united by the goal of curing cancer.

Your Responsibilities

A QM Manager at Indivumed delivers all quality services needed to enable and enhance the full set of mainly GCP/GLP laboratory processes while ensuring the ISO 9001:2015 certified state, setting the stage for future development and growth of the laboratory work environment of Indivumed. Maintenance, support and further development of the functional QM system are an integral part of this function, as well as the close collaboration with the Senior Director Quality Management for daily QM content.

- Responsibility of all quality process within one business unit of Indivumed, focusing on laboratory requirements under ISO 9001:2015 as well as GCP/GLP-compliance (GxP guidelines)
- Maintenance and further development of functional QM system, taking into account the laboratory needs set within the global context of the company quality strategy and policy
- Creation and update of quality processes under ISO certification and GxP guidelines for a functional laboratory working environment
- Support (management and writing) in all quality-related questions from the laboratory due to GCP/GLP requirements
- Tracking of CAPA measures, derived from OOS, deviations, audits, supplier qualifications
- Change Control, Handling of deviations, trainings management, computer system validation (CSV) within the GCP/GLP laboratory work environment
- On-site management in case of customer audits and regulatory inspections (international) for (re-)certification(s): preparation, performance and follow-up

Our Requirements

- Successful completed studies in the area of natural sciences, pharmacy or equivalent
- Min. 3 years of professional experience in a regulated quality laboratory environment, ISO or GxP-certified
- Delivering results relying on full personal responsibility but always focusing on enabling and enhancing the GCP/GLP laboratory processes
- Prominent IT affinity is definitely an advantage
- Fluent in German and English (oral & written)
- Passion for analysing complex issues to streamline and simplify them by applying ISO/GxP quality tools
- Highly communicative with an open mind-setting related to common sense to support the laboratory functions in terms of quality services by QM
- Loves being a team-player bringing all social skills to the table

What We Offer

- Demanding area of responsibility in a fast-growing company
- Creative freedom
- A collegial team and a good working atmosphere
- Continuing education and training
- Grant for the HVV ProfiTicket
- Subsidy for sports membership and events
- Fresh fruits and drinks
- Christmas and summer party
- Flexible working hours
- German and English courses
- The 24th and 31st of December are holidays and won´t be deducted from your annual holidays.

Would You Like to Contribute to the Success of Our Company?

Please apply online via jobs@indivumed.com and include your salary expectations and current notice period.

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For more information about
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