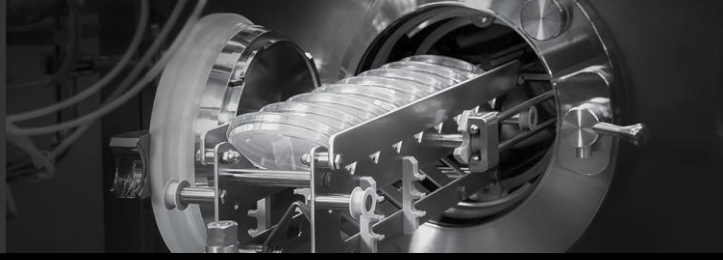
castus®



GMP and Validation Expert, QA (m/f/d)

Join castus on the way to the top!

The castus group is a leading global provider of systems for sterile material transfer in aseptic pharmaceutical production, including transfer systems, ports, and equipment. With castus Single Use Bags (CSU), our sterile single-use transfer bag, we are building a new, rapidly growing business segment.

Around 80 brilliant minds already work for us – and now we are looking for **you**! Are you a GMP expert with a passion for validation, quality, and aseptic processes? Then join our team, take charge of the validation of our sterile single-use transfer bags (CSU), and contribute your GMP expertise across the entire company!

Your Responsibilities – Where You'll Grow

- You will oversee the validation of our CSU products and coordinate testing with external laboratories, such as for extractables and leachables, particles, bioburden, shelf life, leak testing, and sterilization.
- You will create and maintain validation dossiers as well as technical customer documentation.
- You will support customers in the qualification of our products, address technical and GMP-related requirements, and assist with customer audits.
- You will handle nonconformities, complaints, and CAPA actions, and be responsible for change control as well as the qualification of relevant suppliers.
- You will advise various internal departments—such as Development, Sales, and project teams—on GMP requirements for our entire product portfolio, particularly regarding EU GMP Annex 1, contamination control, and equipment qualification.
- You will train employees on GMP topics and current regulatory requirements.
- You will monitor regulatory developments and strengthen our technical positioning through case studies, technical presentations, and technical articles.

Your Profile – How You'll Move Us Forward

- You have a degree in pharmacy, biotechnology, chemistry, microbiology, process engineering, or pharmaceutical engineering, or a comparable qualification.
- You have at least 5 years of professional experience in quality assurance or validation, ideally in the pharmaceutical industry or aseptic manufacturing.
- You have experience with GMP, EU-GMP Annex 1, validation, sterilization, and ideally extractables and leachables, as well as sterile packaging.
- You are able to provide technical guidance to testing laboratories, evaluate test results, and support customer audits.
- You work in a structured and independent way, communicate complex topics clearly, and bring strong expertise.
- You are fluent in German and English and are willing to travel for client meetings and conferences.

Why castus is the Right Fit for You

- You want to create, not just execute: At castus your ideas count and you'll have the freedom to make a real impact.
- You thrive on variety: Every day is different here – your skills will be in constant demand.
- You don't want to be just a number: With us, you'll be part of a dedicated team with flat hierarchies and a personal, collaborative atmosphere!



Top Company 2026 2025 2024 kununu

Ready to take the next step?
We look forward to receiving your application!

Ms. Michaela Schuler · Phone: +49 7352 92187-190
E-Mail: karriere@castus.pro · Website: www.castus.pro

castus GmbH & Co. KG
Kolpingstraße 34 · 88416 Ochsenhausen