CQV Engineer

Do you like to do what you are best at every day - and do you prefer innovative challenges over admin routines?

If you also have a profound experience with commissioning and validation within the pharmaceutical industry, and you can use it together with your interpersonal skills to ensure the right level of documentation in the complex equipment projects in the pharmaceutical industry, you should continue reading.

JOB BRIEF

We are a fast-growing market leading pharmaceutical consultancy specialized in IT and automation systems. And we consider it our main cause to deliver state-of-the-art technical solutions that enable the world´s best pharmaceutical companies to produce lifesaving pharmaceuticals for patients all over the world.

As a CQV Engineer with Trust GMP, you will work with implementation of the latest technologies and be part of a dynamic growth company with a flat organization and high degree of trust and independence.

THE POSITION

You will be the CQV lead in high priority projects defining and ensuring the right level of documentation according to the latest cGMP and our customers QMS systems and use your experience to implement systems ready for the operational phase of the equipment life cycle.

Your role will be to act as the glue that ensures successful documentation of systems, equipment, processes, and procedures and to always create the most value for our customers. In other words, you will be connecting the dots. Your responsibilities will be:

- Specifying or giving input to validation strategies
- CQV review of project documents such as:
  - Validation planning and reporting
  - Requirement and engineering specifications
  - Equipment design
  - Design review/DQ
  - FAT/SAT
  - Commissioning and validation
- Manage and follow up on documentation in project

QUALIFICATIONS

Technical qualifications

- You most likely hold a B.Sc. or M.Sc. in engineering or similar
- Extensive experience with pharmaceutical equipment and processes
- GMP
- Validation

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CVR: 39 75 62 77

- Fluent in English and Danish - both spoken and written

Personal qualifications:

- You have a high-quality mindset
- You have a welcoming, proactive, and solution-oriented approach and like to share knowledge with your team
- You make decisions by having one eye on the details and the other on the broader surroundings and long-term goals
- You have a natural urge to always look for smarter ways to do things
- Various assignment and project environments keep you sharp
- Team events, small talk and social activities give you energy
- You generally thrive in a hectic everyday life with more questions than answers and in finding possibilities in the unknown

WE OFFER

At Trust GMP we offer a dynamic, flexible, and ambitious work environment and a fun and friendly work spirit. We want to do things differently, so we have eliminated all unnecessary bureaucracy and let the management team take care of pitching and admin work, so you instead can take care of what you are best at.

We have a solid high-quality market brand and are known for nurturing long term costumer relations based on trust with world leading pharmaceutical companies. And we can therefore guarantee you great both professional and personal development opportunities. The daily work is based at our costumer or at home, which allows us to spend the office expenses on team and social family events, something we prioritize highly.

In Trust GMP we work as a team, and you can always count on having passionate, skilled, and knowledgeable colleagues to spare with.

Trust is a fundamental value to us. We build our business on trust in our colleagues, customers, and business partners just as we always strive to enable everyone to trust in us.

CONTACT AND APPLY

Would you like to hear more about the position or submit your application and CV, please contact:
Mads Nielsen, +45 6171 7139, mdni@trustgmp.com.

"We believe in making complicated things simple."

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