

More than a job. We make the DIFFERENCE.

JOIN CSM !



To strengthen our Quality Unit, our fast growing company is currently seeking a highly motivated and enthusiastic

## QUALIFIED PERSON (QP)

This position is based in Mont-Saint-Guibert (Belgium)

### Main responsibilities

- Batch certification & release of medicinal products & maintain a batch certification register EU import of IMP.
- Verify/evaluate GMP manufacturing documentation.
- If applicable, perform GMP audits to proof that the medicinal products have been manufactured and tested according to EU GMP.
- Ensure that CSM services are delivered on time, within budget and according to the agreed quality standards.
- Establish the technical/quality agreements between CSM – customers and CSM – subcontractors.
- Take part in the overall improvements of internal processes and systems.
- Contribute to the quality department's activities to ensure continuity of operation. Coordinate information collection via "quality questionnaires" sent to suppliers and subcontractors.

### Skills

- Qualification as Qualified Person mandatory
- Experience as a QP of 2-5 years
- IMP experience of 2 years mandatory
- Experience in manufacturing, packaging & labelling and/or supply chain.
- GMP-GDP-GCP knowledge.
- French & fluent in English.
- Good computer skills : Word, Excel, Outlook, Powerpoint...
- Assertiveness and diplomacy.
- High organizational and planning skills.
- Excellent verbal and written communication skills.
- Team worker and team spirit.
- Customer oriented.
- Eagle eye for details.

### Offer

At CSM, we understand our work within the clinical trial process could not happen without our dedicated employees. That's why we are striving to provide our employees with a challenging working environment which still promotes a work-life balance. We provide competitive compensation and a complete benefits package.

If you have the desire to play your part in making a difference in the lives of patients participating in clinical trials, we want you to apply today by submitting your resume to: [HREurope@csmondemand.com](mailto:HREurope@csmondemand.com)

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*CSM, a Clinigen company, is providing innovative clinical trial supplies and biological sample management services since 1997. With two facilities in Post-Brexit EU and three in the US, CSM is committed to ensuring clinical trials are a success, regardless of size or scope, from Phase I to Phase IV projects.*

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