Based in Gosselies, Delphi Genetics is a biotech CDMO specialized in the production of GMP plasmid DNA to be used as starting material or drug substance in Cell and Gene Therapy.

In order to sustain its important growth Delphi Genetics is actively looking to recruit a

Qualified person

responsibilities:

As a Qualified Person, you supervise the QA department and are responsible of the production of plasmid DNA in compliance with GMP standards and you ensure the release of the batches. You report directly to the top management.

Your main tasks are:
- Release the GMP batches produced while respecting deadlines and quality standards.
- Overview the plasmid manufacturing activities.
- Manage the Quality Department and quality system in accordance with GMP standards.
- Ensure the proper functioning of maintenance, qualification, validation and service provisioning activities.
- Support the QA team in document writing and reviewing (on field when necessary)
- Participate in initial and ongoing training of staff.
- Participate in supplier audits and audits at other European production sites.

Profile:
- You have a Master's Degree in Pharmacy, an Advanced Master in Industrial Pharmacy and a QP number.
- You have proven experience in the biopharmaceutical industry. Candidates fresh out of studies are also welcome.
- You are respectful, trustworthy and enjoy teamwork.
- You are flexible and rigorous.
- You are dynamic and proactive and have good communication skills.
- You can speak English and French both orally and in writing.

Offer:
- A diversified function within a developing society.
- The opportunity to join a human-sized, dynamic and professional environment.
- A training course from the start and throughout your evolution.
- A contract of indefinite duration accompanied by an attractive salary package.

Interested in Joining us, send your application to Charline Savels: csavels@delphigenetics.com

Your application and related information will remain strictly confidential

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