

MaSTherCell

The global cell and gene therapy CDMO partner



We are currently looking for a:

QC Specialist

Full-time

CDI

Gosselies

Interested to work with us? Send your resume and a cover letter at job@masthercell.com

RESPONSIBILITIES

- Writing validation/transfer, protocols of analytical methods, certificates of analyses, stability protocols
- Interpretation of results and writing of analytical methods validation/transfer reports
- Taking part to discussion with clients and subcontractors regarding analytical methods
- Performing QC tests on due time, according to cGMP, current procedures and the QC Manager's instructions
- Performing his/her duties and documenting his/her work according to current procedures and GMP
- Performing on due time : routine and planned maintenance, calibration, qualification of equipment in the QC Department
- Checking the "conform" status of an equipment prior the test
- Checking the expiration date of reagents/solutions prior the test
- Daily maintenance of the laboratory (bench and floor cleaning, waste evacuation, general aspect, etc.)
- Raw data collection, CAPA implementation, Drafting QC documents
- Set up of new QC tests
- Declaring and initiating deviation, OOS linked to his/her results
- Taking part to audits when necessary
- Taking part in technology transfer from the client (may include meetings and observation of QC test on the customer's premises)



QUALIFICATIONS & EXPERIENCE

- At least Graduate level in Sciences
- Experience of minimum 1 year in Quality Control, preferably in a biopharmaceutical company
- Previous experience in cGMP environment

SKILLS & COMPETENCIES

- Fluency in French and good reading comprehension and writing skills in English
- Microsoft office, ERP system
- Rigorous & conscientious
- Autonomous and organized

WE OFFER

- Full-time position (40h/week)
- An Indeterminate contract
- The opportunity to take part in a growing dynamic biotech company
- A human-sized working environment with a convivial atmosphere