MaSTherCell

The global cell and gene therapy CDMO partner

We are currently looking for an:

Analytical Development Expert

Full-time | CDI | Gosselies

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

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JOB OFFER

Analytical Development Expert

RESPONSABILITIES

The analytical development expert main role is to develop or optimize analytical methods associated to one project. In order to fulfill his/her role, the analytical development expert:

- Is an active member of analytical development team
- Maintains open and effective communication with all other QC team member and with other departments (Prod, QA, Project Management, technical services) when required in his/her daily function
- Respects MTC's values, QC department vision and mission, and QC chart
- Participates actively to the weekly QC team meeting
- Follows the needed training on due time
- Works to keep department KPI in green
- Is involved in technology transfer from the client, including meetings and observation of production/QC at the customer's premises
- Keeps up-to-date with scientific literature related to the project
- Is involved in the project follow-up as regards the development phase of the project including contact with the clients, results reporting and presentation
- Suggests new approaches and brings ideas to be discussed internally before communication with the client
- Drafts or reviews protocols, batch records and development/ tech transfer reports of analytical methods
- Supervises and performs new QC tests implementation or improvement on due time and reports progress and issues to the QC supervisor/project leader/project manager
- Interprets the obtained results and informs immediately the QC project leader in case of unusual event/result
- Documents his/her work according to current MTC's procedures and GDP rules
- May write validation protocols and reports of analytical methods under the supervision

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of the QC project leader/Head of QC.

- Is responsible for ordering adequate material and ensures its availability in due time
- Double checks data associated to analytical development generated by other operators
- Performs his/her duties according to current procedures
- Respects safety rules in working area
- Informs immediately the QC Supervisor/ QC project leader/ the head of QC in case of problem or failure or security issue
- Participates to daily maintenance of the laboratory (bench and floor cleaning, waste evacuation, general aspect,...) and assists also the QC support in this task
- Assists the QC support to perform on due time routine and planned maintenance, calibration, qualification of equipment in the QC Department
- Suggests new approaches and brings ideas to be discussed internally before communication with the client
- Finds adequate subcontractors, if necessary
- May participate to audits
- Is involved in the transfer of the developed test to the QC specialists, in due time according to the project plan
- Drafts QC documents (INS, PRO, FORM, GSPE)
- Initiates deviation, CHC linked to his/her work and performs the investigation when potential impact on GMP activities
- Implements CAPA when applicable (impact on GMP activities)

QUALIFICATIONS & EXPERIENCE

- At least Bioengineering degree or Master degree in chemistry, biochemistry, biology, pharmacy, etc.
- At least 5 years experience in culture and analytics of cells from human origin
- Experience in cGMP environment is a plus
- Fluency in French and English is a must
- Knowledge of a third language is a plus

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SKILLS & COMPETENCIES

- Organization skills, multitasking
- Quality minded, autonomous, rigorous, attentive to details, persevering and pro-active
- Customer oriented
- Teamwork and teaching skills
- Good communicator, cheerful

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

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