MaSTherCell is a dynamic and global Contract Development and Manufacturing Organization (CDMO) on a mission to deliver optimized process industrialization capacities to cell therapy organizations, and speed up the arrival of their therapies onto the market. From technology selection to business modeling, through GMP manufacturing, process development, quality services, MaSTherCell's teams are fully committed to helping their clients fulfill their objective of providing sustainable and affordable therapies to their patients.

To lead its increasing number of development and manufacturing projects, MaSTherCell is looking for a highly motivated:

QC Expert (m/f)

Responsabilities

The QC Expert P00 main role is to develop, optimize and perform generic activities P00 in the Quality Control department. In order to fulfill his/her role, the Quality Control Expert P00:

- Is an active member of back-end team;
- Takes part in analytical method validation under the supervision of QC Project Leader/ QC Supervisor (computerized validation);
- Works to keep department KPI in green;
- Follows the needed training on due time;
- Performs his/her duties and documents his/her work according to MTC's current procedures and GMP rules;
- Drafts generic QC documents (INS, PRO, FORM, and SPEC);
- Initiates Deviation, OOS, CHC linked to his/her work, performs investigations and implements CAPA;
- Takes part in new tests implementation/improvement under the supervision of QC Project Leader/ QC Supervisor;
- Double checks data generated by other operators, for which she/he is allowed to;
- Takes part to internal and/or external audits, if necessary;
- Is the back-up of QC Support (if needed).

Outsourcing activities:

- Finds adequate subcontractors, if necessary;
- Drafts documentation associated to subcontractor validation.

Equipment:

- Drafts documentation associated to installation, use and qualification of QC equipment;
- Contacts suppliers and manufacturers.

Qualifications & Experience

- At least Graduate level in chemistry, biochemistry, clinical chemistry, biology, etc.;
- 3-5 years experience in cGMP environment;
- Experience in quality control of cells from human tissue origin is an asset;
- Experience in microbiological testing is a plus.

Skills & Competencies

- Organization skills, multitasking;
- Quality minded, autonomous, rigorous, attentive to details, persevering and pro-active;
- Technical expertise;
- Teamwork and teaching skills;
- Good communication skills and cheerful;
- Good reading comprehension and writing skills in English.

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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Interested?

Please send your detailed CV and your application letter to Mrs. Elodie Noël, HR Manager: job@masthercell.com