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:

Technical Service Equipment Expert

Responsabilities

- Support of facility in terms of preventive and curative maintenance, alarm management and utilities management;
- Support of all new equipment and their installation in the facility. This covers periodic requalification and problem solving with current equipment/process and requires entries in clean rooms if needed;
- Support for the PROD and QC department in facilitating their day to day operations with equipment;
- Interface for Environmental Monitoring System;
- Writing and updating documents associated to equipment management (ie: URS, PQ protocols);
- Ensuring Follow-up of sub-contractors: he/she is coordinating their planning of interventions with users in Logistic, Production & QC(ie: installation, requalification, preventative and curative maintenance);
- Being first line contact point in Technical Service for subject matter experts (SME) ;
- For his/her direct responsibility, he/she ensures cGMP implementation through MTC's quality system and feeds continuous quality improvement;
- He/she Documents his/her work according to current procedures and GMP;
- He/She is involved in preparation and execution of any audits/inspection;
- Supports change control preparation;

- He/She Initiates and finalizes Deviations (Owner role) and CAPA for hierarchy approval. When transversal (on top of the specific field) actions to close Deviations or CAPA are identified, the expert is able to evaluate impacts across MTC on top of its own responsibility field;
- He/She is trained for required activities and is providing data for relevant KPI;
- He/she maintains opened and effective communication with all other departments;
- He/she operates in safe conditions and following EHS rules in place.

Qualifications & Experience

- Master/Engineer in Science (e.g.: in biology, pharmacy, clinical chemistry, bioengineering) or relevant experience in a similar position in the sector of Biotech/Pharma/Cell therapy.
- At least 2 years of experience in GMP environment and
- Aseptic production is a plus
- Fluency in French.
- Good reading comprehension and writing skills in English.
- Computer skills: Microsoft office, use of Data Base systems

Skills & Competencies

- Technical oriented
- Quality minded, rigorous, customer oriented and continuous improvement driven
- Demonstrated cross functional team spirit.
- Organization skills, multitasking, flexibility and autonomous
- Good communicator

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

Interested?

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