MaSTherCell

The global cell and gene therapy CDMO partner

We are currently looking for a:

CELL CULTURE EXPERT

Full-time | Webster, TX - USA

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

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JOB OFFER

Cell Culture Expert

RESPONSIBILITIES

- May be involved in all stages of production process maturity (from development or acquisition to continuous commercial Good Manufacturing Practice (GMP) production) and follow up of closed projects.
- Manage and master critical parameters of production process.
- Must deal with equipment and technology.
- Responsible for management of direct reports.
- Create, finalize and/or approve instructions/forms/protocols related to specific production or generic documentation which could be approved by a Supervisor and relevant stakeholder prior to production start.
- Compile key performance indicators (KPIs) of your team and encourage continuous improvement.
- Respect environmental health and safety (EHS) rules in working area.
- Plan, coordinate and perform daily production.
- Organize production campaigns.
- Must be highly qualified/trained to work in A/B environment and able to train colleagues.
- Must dedicate appropriate time in production area to ensure all duties to produce are met and perform production activities to support Cell Culture Specialists.
- Create, finalize and/or approve GMP documentation related to performed production for final approval by hierarchy/certified reviewer.
- Upon request, must represent the project you manage at internal/external meetings.
- May be responsible for one (1) or several production areas out of dedicated project (daily double checks of area parameters).
- Plan (in partnership with Shared Service Team) and guarantee realization of grades A/B major cleaning (monthly, quarterly, yearly).
- Coordinate (in partnership with Shared Service Team/Technical Service) vaporized hydrogen peroxide (VHP) realization.
- Ensure current Good Manufacturing Practice (cGMP) implementation through Masthercell's quality system and encourage continuous quality improvement.
- Continuously train direct reports working in aseptic environment and manage all

A/B area monitoring.

- Finalize Deviations and Curative and Preventive Actions (CAPA) for hierarchy approval.
- Create product investigations related to out of specifications (OOS) opened by Quality Control (QC).
- Prepare and execute audits/inspections.

SKILLS/QUALIFICATIONS

- Expert in Cell Culture and GMP Aseptic Environment.
- Cell Therapy Specific Context.
- Quality-Minded.
- Rigorous.
- Customer Oriented.
- Continuous Improvement Driven.
- Demonstrated Team Management Capabilities.
- Cross-Functional Team Spirit.
- Well Organized.
- Able to Multi-Task.
- Flexible.
- Autonomous.
- Good Communicator.

EDUCATION/EXPERIENCE

- Master's Degree in Biology, Pharmacy, Clinical Chemistry, Bioengineering or relevant experience in the fields of Biotech/Pharma/Cell Therapy.
- Minimum of two (2) years confirmed experience in GMP environment or equivalent experience in University non-GMP CORE training or training facility.
- Minimum of two (2) years confirmed experience in cell culture production and/or human cells cultivation.

REPORTING RELATIONSHIP

Direct line reporting to the Production Supervisor.

- APPLY TO: jobs-us@masthercell.com.
- Please be sure to include your salary requirements.
- NO TELEPHONE CALLS PLEASE!