POSITION: Project Manager, Chemistry, Manufacturing & Controls (CMC)

Date: 29/11/2024

Type of contract: Permanent

Field: Biotech / Health / Antibiotic resistance

Working time: CDI Full-time

Location: Lyon Gerland, France

Workplace type: Hybrid

Ref: ATX24-09

Start date: As soon as possible

Presentation of the company:

AUROBAC THERAPEUTICS is a biopharmaceutical company created in 2022 by 3 renowned life sciences innovation companies, *Boehringer Ingelheim, Evotec* and *bioMérieux,* to become a global leader in the fight against bacterial infections, Antimicrobial Resistance (AMR) and their consequences in acute hospital settings.

We focus our efforts to develop the next generation of antimicrobial and acute care products, following a targeted precision medicine approach, i.e. associated with rapid diagnostics to support product use.

With strong growth ahead, within an international environment, the company is now looking for a **Project Manager CMC.**

Description:

As a Project Manager CMC at AUROBAC, you will manage the development of products in our pipeline and contribute to the CMC department transversal activities.

You will report to Head of CMC and work closely with Project Management, Clinical Operations, Quality Operations & Regulatory Affairs teams.

Missions:

- **Manage process development, analytical and stability activities conducted by external subcontractors**
  - ✓ Monitor development plan execution, timelines & deliverables, identify and escalate risks
  - ✓ Review technical documentation, analyze and interpret results
  - ✓ Lead weekly project meetings with **subcontractors**, produce/review agendas & minutes
- **Interact & collaborate with project team members & external experts**
  - ✓ Work closely with Clinical Operations team to coordinate clinical trial supplies
  - ✓ Work closely with Quality Operations team to manage quality events (deviations, OOS/OOT, change controls) and ensure Good Manufacturing Practices (GMP) compliance
  - ✓ As needed request & prepare meetings with external experts, draft and distribute agendas & minutes
- **Ensure communication with project and executive teams**
  - ✓ Share progress of development activities during monthly project meetings
  - ✓ Complete trackers for project activities follow-up
  - ✓ Produce material to support decision making

- **Contribute to CMC department transversal activities**
  - ✓ Supply & consolidate development trackers (clinical batch history, manufacturing parameters & results, stability timepoints & results)
  - ✓ Manage stock, supply & dispatch activities

- **Carry out literature research & technological/regulatory intelligence in accordance with project (excipients, formulation techniques, guidelines…)**

Skills:

- Strong knowledge of formulation, process and analytical development
- Good knowledge of Good Manufacturing Practices
- Experience in subcontractor management is required
- Experience in parenteral forms pharmaceutical development is a plus
- The candidate is rigorous, involved and demonstrates good organizational and communication skills
- Fluency in English is a must-have, French language skills would be a major plus
- Used to working in an international environment

Training and experience:

- PharmD, Master of science in biotechnology or chemistry or chemical engineering
- At least 2 years experience as scientist in the pharmaceutical industry

To apply, please send your **CV and cover letter** application to jobs@aurobac-tx.com mentioning the reference **ATX24-09**

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