PDC*line pharma
ADVANCED CANCER VACCINES

PDC*line Pharma (www.pdc-line-pharma.com) is a clinical-stage spin-off of the French Blood Bank (EFS), that develops a new class of potent and off-the-shelf therapeutic cancer vaccines based on a proprietary cell line of Plasmacytoid Dendritic Cells (PDC*line).
Based on a robust preclinical package and a first-in-human Phase Ib in melanoma, PDC*line Pharma has initiated a clinical development in lung cancer with a new candidate (PDC*lung) and neoantigens (PDC*neo).

PDC*line Pharma comprises a team of 27 persons based in Belgium (Liège, headquarters) and France (Grenoble). The company has raised nearly €52 million in non-dilutive equity. The last round of the financing has been led by the Asian leading Venture Capital company KIP (Korean Investment Partners).

In order to strengthen its team located in Liege (Belgium), we are actively looking for several

Quality Control Technician

In relation to the activities carried out by PDC*line Pharma, the person occupying the position of Quality Control Technician has the following characteristics:

Responsibilities of the Function

The responsibilities of the QC Technician position are as follows:

In context of material resources and operation management

- Participate in utilities stock management;
- Participate in equipment qualification and maintenance/requalification;
- Perform requests for quotation, orders and reception of incoming materials;
- Perform QC tests for incoming materials release;
- Participate in the QC stock management: inventory and monitoring of laboratory materials;
- Participate in routine tasks related to the good management of the laboratory (shared responsibility between all operators);
- Provide support for cleaning/storage of QC areas and filling relative checklists, including logbooks of equipment;
- Participate in the development and validation of QC analytical methods;
- Perform QC tests related to IMP clinical batches: manipulation, analysis and interpretation of results, drafting of reports;
- Perform immuno-monitoring activities (blood processing, enumeration and immuno-phenotyping);
- Participate in the stability studies of IMP;
- Record samples in QC sample library;

- Participate in the creation and edition of MBR in SmartReg, create and execute BR, update of SmartReg database.

In context of human resources management:

- Training of trainees

In context of continuous improvement management:

- Initiate and follow all non-conformities (deficiencies/deviations/OOS/OOT/derogations) observed from the established and approved rules/specifications, and immediately inform the QA;
- Follow up Change Controls;
- Participate in the internal audits;
- Ensure CAPA follow-up and implement corrective and preventive actions;
- Comply with applicable GxP regulations.

Skills

- Scientific education related to biotechnologies or equivalent experience;
- Knowledge of molecular biology;
- Theoretical and practical knowledge of the techniques used within the company;
- Knowledge of basic GMP guidelines;
- Knowledge of the activities of PDC*line;
- Knowledge of English (comprehension of English text).

We offer :

- A diversified position with responsibilities within a fast-growing start-up.
- The opportunity to join a human-sized, dynamic and professional environment.
- A permanent contract and an attractive salary package in line with your experience.

Interested?

Please apply by sending your CV and cover letter via e-mail to the attention of Valérie Donolato - HR&Admin.
v.donolato@pdc-line-pharma.com

Your application will be treated as confidential.