Telix Pharmaceuticals Limited
ACN 616 620 369
Suite 401, 55 Flemington Road
North Melbourne
Victoria, 3051
Australia

TLX-JD
Job Description : Bioprocess Production Manager (2 roles)
Type : Full-time
Location : Melbourne
Date : Immediate

About Telix Pharmaceuticals Limited

Telix Pharmaceuticals Limited ("Telix", the "Company") is an Australian public Company (ASX:TLX) headquartered in Melbourne with operations in Europe, the US and Japan. Our mission is to be a leading, global biopharmaceutical Company in the field of "theranostic" radiopharmaceuticals and the Company is currently developing a mid-to-late stage pipeline of products in prostate, kidney and brain cancer.

Description

To contribute to the achievement of the company's strategic goals by managing all technical bioprocess activities for a late-phase biological candidate program in support of Telix's development and commercial aims

Responsibilities

- Planning, coordination and active management of all biologics manufacturing activities conducted by external vendors for a late-phase program such as analytical method qualifications, antibody production, bioconjugate manufacture and fill/finish activities.
- Biologics manufacturing strategy development including phase-appropriate vendor selection in line with technical, regulatory & quality requirements to suit clinical and commercial needs.
- Effective management of outsourced manufacturing activities and tracking of expenditure to ensure delivery on time and budget.
- Contributing to regular project meetings and providing verbal or written feedback to inform the team or management of progress.
- Providing technical support to other TLX projects (or sourcing appropriate advice) to support radiochemistry process development and regulatory filings.

Qualifications, Skills and Experience

- At least 10 years of relevant Biologic drug production experience with at least 2 years in commercial drug development.
- Post-graduate qualifications strongly preferred (with a minimum of an honours degree in a relevant scientific subject plus demonstrated relevant experience)
- Thorough knowledge of GMP / Commercial Process development & validation.

- Hands on experience with quality management systems used in the manufacturing environment including control of raw material and product specifications.

- Formal regulatory/quality training is not required, but useful.

Scientific & Pharmaceutical Knowledge Areas

- A solid science background, with qualifications in biochemistry/ pharmacology/ chemistry preferred.

- Direct experience in biologics manufacturing processes including cell line development, upstream & downstream development, viral clearance studies, biologics-relevant assay qualification, bioconjugation and fill/finish.

- Knowledge of a range of scientific techniques from relevant subject areas such as biochemistry, pharmacology, proteomics, immunology.

- Understanding of regulatory requirements for biologics including preparation of documentation describing manufacture activities for regulatory filings and clinical study submissions.

Contact

For further details and information pertaining to compensation for the role, as well as expressions of interest, please contact Telix human resources at employment@telixpharma.com or visit our careers page at www.telixpharma.com .