

JOB DESCRIPTION

Clinical Supply Manager

Service	Supply Chain	
Direct Supervisor	Sr Director, Manufacturing, Sourcing & Pharmaceutical	
	Development	
Management of Team	N/A	

Position Summary

The incumbent will be responsible for establishing and managing all the supply chain related activities for the clinical projects at Theratechnologies Inc. <u>Please note that this position may be</u> <u>based in Dublin, Ireland, or Montreal, Quebec, Canada, depending on the location of the</u> <u>successful candidate.</u>

Key Responsibilities

- Planning the manufacturing (including required quantities to fill requests from clinical team) to ensure continuity of clinical supplies
- Monitor inventories
- > Manage logistics for product shipments from CMO's to clinical distributor
- Manage clinical distributor (shipment to clinical sites)
- > Manage logistics/shipments for clinical test kits from testing labs to clinical sites
- Manage logistics/shipments of clinical samples (ex. blood and biopsies) from clinical sites to testing labs
- Monitor budget for all the above, take care of contracts, POs, invoice approvals, change orders, etc.
- Develop and/or procure clinical secondary and tertiary packaging (e.g., carton for vials, and shipper boxes) and ensure adequacy for given shipment conditions (e.g., dry ice)
- Determine permit requirements, perform permit application/renewal, and maintain valid permit or ensure that CMO has valid permit at all times
- > Manage EU importation and release for distribution
- > Develop and implement contracts with an EU warehouse/clinical distributor
- > Performs any other duties required by the management.

External Customers

- > CMO's
- ➢ TPL's
- Analytical Laboratories

Qualifications Required

- > Bachelor's degree in science or equivalent
- ➢ 5 years of team management experience

- > 5 years of relevant experience in a supply chain role
- > 5 years of experience in a pharmaceutical or biotechnology company
- Good knowledge of European guidelines, laws, regulations, and regulatory practices (Good Manufacturing Practices (GMP), Good Distribution Practices (GDP))
- > Excellent communicator, and well-organized with superior presentation skills
- > Understand the quality mechanisms for releasing batches of manufactured drugs.
- Knowledge of Word, Excel, Power Point, Outlook, and Project

Personal Qualities for all Staff

- Ability to set goals and meet deadlines.
- > Ability to prioritize
- Ability to generate ideas and find solutions
- Ability to work both alone and in a team
- > Team spirit and tendency to help each other
- Versatility and flexibility

Personal Qualities Specific to Position

- Leadership and Accountability
- Results-oriented
- Ability to manage and train staff
- Ability to negotiate and to influence others
- Capable to establish priorities
- Ability to multi-task with ease
- > Ability to learn quickly and demonstrate curiosity
- > Able to manage and control stress in the workplace
- > Excellent attention to detail and quality in all matters
- Fluency in written and oral expression
- Discretion
- > Creativity
- Rigor in tasks
- Available for business travel

Current incumbent:

Validated	by:	
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_____ Date: _____

Signature of direct supervisor: ______