Life Cycle Manager (ESO LM)

Job ID
REQ-10001653
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India

Resumen

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About the Role

Life Cycle Manager

Location – Hyderabad #LI Hybrid

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Key Responsibilities:

- Lead projects for all Lifecycle events (LCE) in area of responsibility (DS, DP, FP for assigned Product(s). Successful implementation of Project/LCE in-time, budget, compliance
- Forms the project team during kick-off meeting in the initiation phase. Organizes and leads project meetings and arranges meeting minutes until hand-over to the Life Cycle Implementation Expert (LCI). Is responsible and accountable to develop together with the responsible functions in RA-CMC and RA the regulatory submission strategy and in conjunction with E2E-Supply Network Planning Manager, Sites, Markets and other stakeholder the implementation strategy.
- Identifies harmonization/bundling opportunities to reduce complexity. Is responsible to check the impact of the project on supply, possible interaction with other projects or brand related activities, ensures whenever needed the prioritization across Life Cycle Projects.
- Project Lead is accountable to ensure that the Master Data strategy is created by the responsible Master Data group. The Project Lead together with the LCI creates a first version of the changeover plan (COP) including affected assortment (provided by LCI), planned submission, planned approval and planned implementation timelines on SKU level
- Is responsible to distribute project information to sites and markets (incl. COP) to trigger initial local data set-up (planning BOM, material master data, sales forecast etc.). The project lead only does the 1st communication of the COP Version 1 to the stakeholder, thereafter the LCI is responsible.

- Is responsible for project reporting and update in the applicable Life Cycle (LC) reporting tool. Is responsible for communicating relevant information after submission to affiliates (Demand Managers) and affected manufacturing site(s) (SCM line functions).
- Is accountable to ensure that all implementation activities are completed according to the project plan and the regulatory approvals in order to ensure a regulatory compliant implementation and that any changes to the project (e.g. timeline shifts) will be communicated.

Essential Requirements:

- **Education:** University degree (technical, science, pharmaceutical, project management, supply chain management preferred) or other education with equivalent experience in the pharmaceutical industry.
- 12 years in the biopharmaceutical industry, specifically in Large Molecules. Extensive experience in Life cycle management of DP, DS and FS
- Experience in execution of product launch, transfer, change, divestment, pruning, new product development.
- Experience in biopharmaceutical manufacturing – aseptic/sterile manufacturing, in large molecules
- Preferred experience in Operations, Quality, Manufacturing Science, or related departments\
- Very good communication skills, Project Management techniques, Analytical & structured approach, Problem solving in complex environments.

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Job Type
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