研究開発・メディカルアフェアーズ統括本部 Pharmaceutical Project Management（PPM）/P…

> **日本イーライリリー株式会社での募集です。 臨床開発リーダー・臨床開発プロジェク…**

募集職種

人材紹介会社
株式会社ジェイ エイ シー リクルートメント

採用企業名
日本イーライリリー株式会社

求人ID
1576844

業種
医薬品

会社の種類
外資系企業

雇用形態
正社員

勤務地
兵庫県

給与
600万円 ~ 経験考慮の上、応相談

勤務時間
08:45 〜 17:30

休日・休暇
【有給休暇】有給休暇は入社後2ヶ月目から付与されます 初年度 10日 2か月目から 【休日】完全週休二日制 夏季休暇 年末年始…

更新日
2026年04月02日 04:00

応募必要条件

キャリアレベル
中途経験者レベル

英語レベル
ビジネス会話レベル

日本語レベル
ネイティブ

最終学歴
大学卒： 学士号

現在のビザ
日本での就労許可が必要です

募集要項

【求人No NJB2355427】
Pharmaceutical Project Management （PPM）
PURPOSE:

The purpose of Pharmaceutical Project Management （PPM） role is to have overall responsibilities for working with the

leadership of the Global counterparts and the corresponding Affiliate Business Unit functions to assure development and implementation of appropriate pharmaceutical development strategies and plans from early phase・ to end of the lifecycle of the molecule on time and on budget in the relevant therapeutic area.・ The key deliverables are full portfolio coverage simultaneous development and launches and creating value across the entire product life stage.・ The role will be assigned to one or multiple Therapeutic Areas （Immunology Neuroscience DOCTA/CMH or Oncology）.

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PRIMARY RESPONSIBILITIES:
1.・ Align the Portfolio Development to BU Strategies
・Engage Medical Clinical Regulatory CMC/Manufacturing Marketing Safety Functions and other functions to deliver the portfolio and align drug development strategies with BU strategies.
・Provide leadership for development of Product Lifecycle Plan in alignment with the Therapeutic Area・development and commercial strategies.
・Deliver necessary information and data about the new/ongoing projects for the portfolio and business plan management.

2.・ Lead Value Maximization Through Product Lifecycle
・Lead to develop robust Japan Project Plan that fortifies linkages to Asset Profile Value proposition MMFA （Manufactured Marketed Forms Agreement） and scope assures quality and consistency of content and translates into operational objectives.
・Translate Japan marketing medical and regulatory needs for the molecule/new indication/line extension （including new formulation / delivery solution） into the global integrated plan.
・Lead overall launch readiness and co lead the affiliate overall launch readiness in collaboration with respective commercial leader.
・Ensure post marketing activities including issue management are managed well by providing leadership to cross functional collaboration.

3.・ Deliver Results
・Lead the team to meet or exceed overall project goals （completion of clinical studies CMC deliverables and NDA/device preparation submission and approval timing etc.）
・Deliver project milestones on time on budget within scope and to high quality standards by holding functions accountable for achievement of key project deliverables in alignment with the agreed to project plan and in compliance with Japan quality standards.
・Identify areas for where process improvement will enable better results and work with discipline leaders to impact change.

4.・ Apply Project Management Knowledge and Processes・・・・・・・・・・・
・Work with Japan governance to authorize project scope and plans. Scope and plans should be based on broad scenario planning robust risk management planning and reliable execution （timeline FTE and budget） planning which ensures all functional requirements are appropriately reflected.
・Develop and implement stakeholder global and team communication plan to ensure alignment with medical affairs strategy and product development strategy.・・Lead external communication strategy related to development （e.g. data read outs regulatory outcomes）.・ Provide rapid and quality responses to external customers.
・Monitor and control performance of the overall project that utilizes appropriate tools and techniques （project systems budget: plan vs. actual FTE: plan vs. actual etc.） and ensure change management process is in place to manage scope and plan.

5.・ Build Winning Culture Through Effective People Management
・Coach team members by providing timely constructive feedback and recognition to good results and behaviors.
・Create an environment that encourages intelligent risk taking and fosters wining culture within the team.
・Support the growth of project management expertise in Lilly Japan through participation in shared learning forums staff meetings and training programs.

スキル・資格

※以下はManager/Sr. Manager/Sr .Group Manager Project Mgmtの要件です

QUALIFICATION REQUIREMENTS:
Knowledge and Skills
・Strong leadership skills.
・Excellent interpersonal and communications skills both in Japanese （native fluent or proficient） and in English （native fluent or proficient） communicate clearly with diverse interpersonal styles and able to develop credibility cross functionally.
・Ability to influence others positively negotiate effectively and manage conflicts effectively.
・Ability to lead and deliver in the volatile uncertain complex and ambiguous environment.
・Project Management Expertise
・Strong strategic and logical thinking skills.
・Strong conceptual skills.
・Understanding of Project Management concepts tools and skills.
・Business acumen.
・
Education and Work Experience Desirable to Perform Role:
・A degree in healthcare science or business discipline （BS/MS/PhD MD MBA）.
・Minimum 3 years of experience in R D function（s）.
・English language proficiency required or substantial experience as an active member or a leader of a global team.

Additional Preferences
・Working experience in pharmaceutical industry.
・Drug development knowledge and expertise.
・Qualification in project management professional （PMP）

会社説明

医療用医薬品の輸入・製造・販売