

## TMF SWAT Manager

Job ID  
REQ-10045337

4月 29, 2025

United Kingdom

### 摘要

This is a unique opportunity to join Clinical Development Document and Governance Management team as a TMF SWAT Manager!

### About the Role

This is a hybrid role and can be based in London, Dublin or Hyderabad offices. The expectation is to be in the office 12 days/month

Major accountabilities:

- Act as CDGM point of contact for assigned portfolio of In-Licensing / Out-Licensing / Acquisition / Divestment Projects, collaborating with key stakeholders with CDGM teams, Development Informatics, Legal, Development Quality Assurance and Global Project Teams.

- Lead and/or Contribute to the development of TMF Transition Plans and ensure the successful transitions of TMF (paper and electronic) documentation outside of Novartis in support of out-licensing and divestment projects, and into Novartis in support of in-licensing and acquisition projects.
- Develop and maintain paper and electronic document processes & standards relating to BD&L projects and Out licensing activities, in compliance with internal and external requirements & regulations
- Identify and communicate risks/trends/patterns relating to TMF, BD&L projects, Out licensing activities and work with key stakeholders to define and implement pragmatic remediations.
- Executes vendor oversight plan, monitors service metrics and identifies opportunities for improvement to the operating model. Acts as point of escalation for issues.
- Serves as Subject Matter Expert on TMF transition related training materials, formal and informal processes and tracking tools for TMF transition oversight activities in collaboration with CDM Process team and other key stakeholders
- Provides support for inspections/audits, contributes to root cause analysis identification and creation/delivery of CAPAs.
- Identify and implement improvements to document management processes to improve quality of TMF integration activities. May act as business lead for innovation projects to enhance integration processes.
- Supports the TMF Integration Lead with respect to forecasting and planning of B&L projects.

#### Relevant Experience:

- Minimum of 5 years working in clinical research and development in the pharmaceutical industry (and/or Contract Research Organisations) with specific experience in clinical documentation and/or records & information management.
- Demonstrated success in planning and executing cross functional projects.
- Strong influencing and presentation skills. Ability to communicate effectively at all levels.
- High organisational awareness, including experience working in multi-disciplinary teams, across cultures and geographies.
- Good negotiation, problem solving and conflict resolution skills; experience establishing trusted relationships with internal and external stakeholders.

#### Languages :

- English.

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reasonable accommodation for any part of the recruitment process, or in order to receive more detailed information about the essential functions of a position, please send an e-mail to [diversity.inclusionch@novartis.com](mailto:diversity.inclusionch@novartis.com) and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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部门  
Development

Business Unit  
Universal Hierarchy Node

地点  
United Kingdom

站点  
London (The Westworks)

Company / Legal Entity  
GB16 (FCRS = GB016) Novartis Pharmaceuticals UK Ltd.

Alternative Location 1

Dublin (NOCC), Ireland

Alternative Location 2  
Hyderabad (Office), India

Functional Area  
Research & Development

Job Type  
Full time

Employment Type  
Regular

Shift Work  
No

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