

Clinical Document Management - Business Migrations Manager

Job ID
REQ-10039157

Feb 10, 2025

India

Summary

Clinical Document Governance Management (CDGM) is accountable for strategy and delivery of clinical document management (CDM) systems, processes, standards and operations of CDM services (including Trial Master File management (TMF), clinical submission readiness, record retention and archiving, Good Documentation Practice capability build) across Novartis globally. In addition, CDGM is driving the transformation of TMF at Novartis, through the introduction and adoption of new technologies, processes and ways of working.

About the Role

The CDM Business Migrations Manager is responsible for effective planning and delivery of migrations to and from of Novartis enterprise-wide clinical electronic document management systems (eDMS) aligned to Novartis business needs, partnering with technical vendors, internal IT and business stakeholders.

- Drives implementation of CDGM initiatives, projects and process improvement activities to enhance the planning and execution of migrations to and from enterprise clinical eDMS at Novartis.
- Act as CDGM point of contact for partnering with CDGM, IT (internal and external) and business stakeholders to plan and execute migrations to and from eDMS, in line with Novartis business, compliance and operational requirements.
- Partner with CDGM and business stakeholders, especially those planning in-licensing and out-licensing activities to identify and agree migration business requirements, understand source and target system capabilities and develop/maintain a future migration roadmap.
- Contribute to activities to ensure efficient processes & integrations of systems with eDMS based on strong understanding of Novartis enterprise systems landscape and in line with compliance and business priorities. Serves as Subject Matter Expert for training materials, formal and informal processes and tracking tools for eDMS migration activities, in collaboration with CDM Process team and other key stakeholders
- Plan and contribute to agile working methodologies being applied during development cycles to prepare for migration and during post migration hyper care period.
- Owner or Contributor of activities related to migration related Incident Management, Change Management and ongoing operations of the eDMS.
- Support forecasting of internal resource allocations and vendor provided activities as part of eDMS migration roadmap management.
- Executes vendor oversight plan, monitors service metrics and identifies opportunities for improvement to the operating model in relation to migration. Acts as point of escalation for issues.
- Provides support for inspections/audits, contributes to root cause analysis identification and creation/delivery of CAPAs.

Minimum Requirements:

- Advanced degree or combination of Bachelor ' s degree in information or life-sciences/healthcare and relevant industry experience. Minimum of 6 years working in Pharmaceuticals, Lifesciences and Clinical Research with specific experience in contributing and leading of clinical document management, TMF and/or records & information management.
- Minimum of 5 years of full-scale migrations
- Minimum 2 medium to major Veeva related hands-on and provable experience in leading and planning of migrations.
- Prior experience in Electronic Document Management systems, specifically in Clinical and Regulatory highly desired.
- Business relevant technical and working experience of eDMS systems like Veeva Clinical vault, RIM, Documentum D2LS or similar
- Knowledge of industrywide Electronic and Clinical Document Management systems and features
- Deep knowledge of Agile way of working with cross functional teams for releases
- Strong influencing and presentation skills. Ability to communicate effectively at all levels

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部門

Development

部門

Innovative Medicines

国

India

勤務地

Hyderabad (Office)

Company / Legal Entity

IN10 (FCRS = IN010) Novartis Healthcare Private Limited

Functional Area

Research & Development

職種

Full time

雇用形態

Regular

Shift Work
No

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Accessibility and accommodation

Novartis is committed to working with and providing reasonable accommodation to individuals with disabilities. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to perform the essential functions of a position, please send an e-mail to diversityandincl.india@novartis.com and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

Novartis is committed to building an outstanding, inclusive work environment and diverse teams' representative of the patients and communities we serve.



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