

eCompliance Manager

Job ID
REQ-10061249

10月 09, 2025

India

摘要

The eCompliance Manager is responsible for providing Quality Assurance oversight and guidance with regard to computerized systems validation (CSV), operating within the framework of regulations (GxP, 21CFR11, etc.) and requirements defined in the Novartis Quality Manual and global procedures.

eCompliance Manager provides the needed operational support such as approving the GxP impacted changes, Periodic Review Reports, deviations etc., Provides the guidance to the project and operations team on the CSV related topics and related information. Reviews and/or approves the global Computerized Systems key validation deliverables as a part of the eCompliance support to the GxP projects.

About the Role

Key Responsibilities:

- Quality oversight of operational activities of GxP systems (e.g., Changes, Periodic Reviews, Deviations etc.,)
 2. Provide needed support to meet the applicable Novartis and regulatory requirements for GxP regulated computerized systems projects.
 3. Point of Contact for all CSV related matters for GxP Computerized Systems and act as an interface between IT and Business for eCompliance topics in relation to GxP classified Computer Systems promoting a Quality Culture.
 4. Review and approve project related documents for GxP relevant systems including determination of GxP applicability for all GxP and non-GxP relevant systems.
 5. Establish trusted partnership with assigned IT Function with understanding of business drivers, and provide the needed day to day operational support.
 6. Review and approve the GxP Changes and the associated deliverables.
 7. Review and approve the GxP impacted deviations, ensure appropriate CAPA are implemented.
 8. Contribute for the preparation of VMP and execute the plan for the systems associated with the respective functions.
 9. Review and approve the Periodic Review Reports for the GxP computerized systems and the associated gaps within CAPA Management System.
 10. Perform supplier qualification assessment activities.
 11. Provide Audit support as assigned and in case of CAPAs, provide the required Quality support.

Essential Requirements:

- 1. GxP relevant computerized systems are developed, implemented and maintained according to the Novartis requirements.
- 2. On time review and approval of changes, deviations & periodic review reports for the GxP computerized systems.
- 2. Documentation supporting eCompliance and CSV requirements is in place, maintained up-to-date and can be presented during audits and inspections without delays and issues.
- 3. Gaps in eCompliance and CSV activities are proactively identified, escalated and the development of mitigation plans supported.
- 4. Client/stakeholder satisfaction and corresponding feedback.

Desirable Requirements:

Work Experience:

- its related technologies to support the operation
 - Good understanding in system application management, its Quality support approach and industry best practices (ITIL, ITSM, etc.)
 - Experience in the development, implementation and lifecycle management of key computerized systems in the Pharmaceutical Development, Manufacturing, Quality, Commercial and Infrastructure space (e.g. ERP/SAP, MES, LIMS, CRM, IAM, etc.)
 - Successful cross-divisional/functional work with complex international teams
 - Experience in GxP supplier qualification activities
 - Proven ability to adjust to multiple demands, shifting priorities and unexpected events while maintaining a positive work attitude
 - Ability to effectively interact and present to Management

- Ability to manage the key stakeholders and build trust
- Proven ability to influence without hierarchical authority and build trusted partnerships
- Proven self-starter with experience in initiating and delivering projects and processes
- Excellent communication, negotiation, facilitation, and interpersonal skills

Languages :

- English.

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Operations

Business Unit
Universal Hierarchy Node

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站点
Hyderabad (Office)

Company / Legal Entity
IN10 (FCRS = IN010) Novartis Healthcare Private Limited

Functional Area
Quality

Job Type
Full time

Employment Type
Regular

Shift Work
No

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