

Global System Integration Expert, Global DQC CoE

Job ID
REQ-10027504
Oct 28, 2024
India

About the Role

Job Purpose:

The Global System Integration Expert supports the efforts to create system integrations via TetraScience Data platform

Experience Required: Minimum 10 years of laboratory experience in a pharma industry.

Major Accountabilities:

- Act as technical expert in creation of pipelines programmed in DataWeave programming language.
- Creates pipelines (custom code) to enable data from various laboratory systems to be transferred to the current LIMS system
- Supports establishment and maintenance of global documentation related to the systems in scope
- Identifies and anticipates site needs, determines what features should be implemented, and support prioritization
- Supports establishment of release timelines, content of each release, oversee the application development stages and supports completion of each release in accordance with the approved plan.
- Supports Business screening, PQ scripting and PQ execution.
- Provides required periodic progress reports, milestone activities and communications to the program management.
- Supports establishment and maintenance of global documentation related to the systems in scope (e.g. SOPs, WIs, user guides, etc)
- Contribute to Laboratory Operations Quality System in defining and implementation of strategy and defined activities.
- Adheres to all applicable procedures, cGMPs, company policies and any other quality or regulatory requirements.

Key Performance Indicators:

- Metrics according to target
- Individual project completion
- Achieves agreed targets and objectives in terms of quality, time and cost
- Supports departmental objectives to implement systems according to overall program plans

Minimum Requirements:

Education:

University degree in Pharmacy, Engineering, Chemistry or equivalent Discipline

Experience:

Thorough knowledge of cGMP requirements:

- * The profile must have strong knowledge in Data Weave programming language.
 - * Computer System Validation experience is key expectation, similarly coding experience
 - * Strong understanding of regulatory requirements for commercial products.
 - * Technical understanding of laboratory business processes and enterprise data expertise
 - * Experience with Labware LIMS and/or TetraScience data platform
 - * Strong understanding of risk assessment and risk management fundamentals/tools.
 - * Team and consensus builder, with definitive and authoritative decision-making ability.
- Critical Negotiations.
 - Functional Breadth.
 - Project Management.
 - People Leadership.
 - Collaborating across boundaries.
 - Operations Management and Execution.

Why Novartis: Our purpose is to reimagine medicine to improve and extend people's lives and our vision is to become the most valued and trusted medicines company in the world. How can we achieve this? With our people. It is our associates that drive us each day to reach our ambitions. Be a part of this mission and join us! Learn more here: <https://www.novartis.com/about/strategy/people-and-culture>

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Role Requirements

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Division

Operations

Business Unit

Innovative Medicines

Location

India

Site

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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