Associate, Automation Engineer

Job ID
REQ-10014731
Mar 04, 2025
Japan

Summary

Support design, implementation, and maintenance of automation systems in projects and site operations.

About the Role

Major Accountabilities

- Control of room temperature, humidity, and pressure.
- Manage of materials (shipping volume, usage fees)
- Support internal and external audits.
- Develop & implement Coding / Recipes (DCS / Scada / PLC's / Control Networks)
- Manage operations for (OT) systems, tools, and applications, ensuring their stability and integrity, while meeting customer service levels.
- Perform GMP risk assessments (incl. Sensors SR)

Requirements:

- At least 3 years' experience in GMP related work
- At least 3 years' experience in build-up facility system (any Industry)
- Must be proficient in global and local procedures for operations, maintenance and development.
- Experience with OQ, PQ, DQ, IQ, PV and other documentation.
- Communications skills essential.
- Skills in C&Q within the automation (CSV).
- Fluent level of Japanese and business level English language skills.
- Academic qualifications not required.
- Oversea business trip required few times annually
- Engineering experience in automation in the chemical or pharmaceutical industry desirable.

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- DCS Scada PLC
- (OT)
- GMP SRA

- GMP 3
- 3 ( )
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- OQ, PQ, DQ,IQ,PV
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- (CSV) C&Q
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Division
Operations
Location
Japan
Site
Sasayama
Company / Legal Entity
JP05 (FCRS = JP005) Novartis Pharma K.K.
Functional Area
Technical Operations
Job Type
Full time
Employment Type
Regular
Shift Work
No
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midcareer-r.japan@novartis.com

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