Senior Expert - QC

Job ID
REQ-10067376

11月 27, 2025

Singapore

摘要

Primary responsibilities for this position include performing tasks associated with release testing and reviewing laboratory data, coordinating the team/lab under its responsibility. Communicating with internal & external partners for the Quality Control organization. Supports site as technical expert in related field

About the Role

Key Responsibilities:

- Act as SME in the area of equipment calibration and qualification in the QC laboratory ensuring compliance with local and global procedures while minimising the impact on the business
- Coach trainers to ensure effective training of QA/QC analysts in area of laboratory equipment and qualification by developing and advising standardized content, materials and assessment

tools.
- Act as SPOC for major QC equipment projects and liaise closely with end users and other departments in relation to cross-functional projects
- Management of QC investment budget/QC asset management and utilisation/QC CapEx projects.
- Troubleshoot and resolve challenging issues relating to modifications of current quality compliance processes and systems within and across various functions to ensure all health, safety or training procedures meet requirements.
- Lead cross functional project teams to drive team results including Lab Excellence projects
- Initiate and supervise the Strive to constantly continuous improvement of systems and practices on site with respect to achieve compliance, reducing waste and improving efficiencies
- Stand in for other team members in the QC Support Services team in their absence.

Essential Requirement:

- BS: 6+ years related experience with 3+ years in management
- Related experience should be in GMP-regulated industries in Quality Control.
- Must have a working knowledge of health authority and regulatory requirements as well as industry quality management tools, standards, and quality systems.
- Must have an understanding of pharmaceutical industry trends and practices.
- Broad cGMP experience is required with knowledge and understanding of manufacturing, quality control, and validation requirements and activities.

Desirable Requirement:

- English fluent in speaking and writing. Mandarin Chinese a plus

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部门
Operations

Business Unit
Quality

地点
Singapore

站点
Tuas South Avenue

Company / Legal Entity
SG12 (FCRS = SG012) Novartis Singapore Pharmaceutical Manufacturing Pte Ltd

Functional Area
Quality

Job Type
Full time

Employment Type
Regular

Shift Work
No

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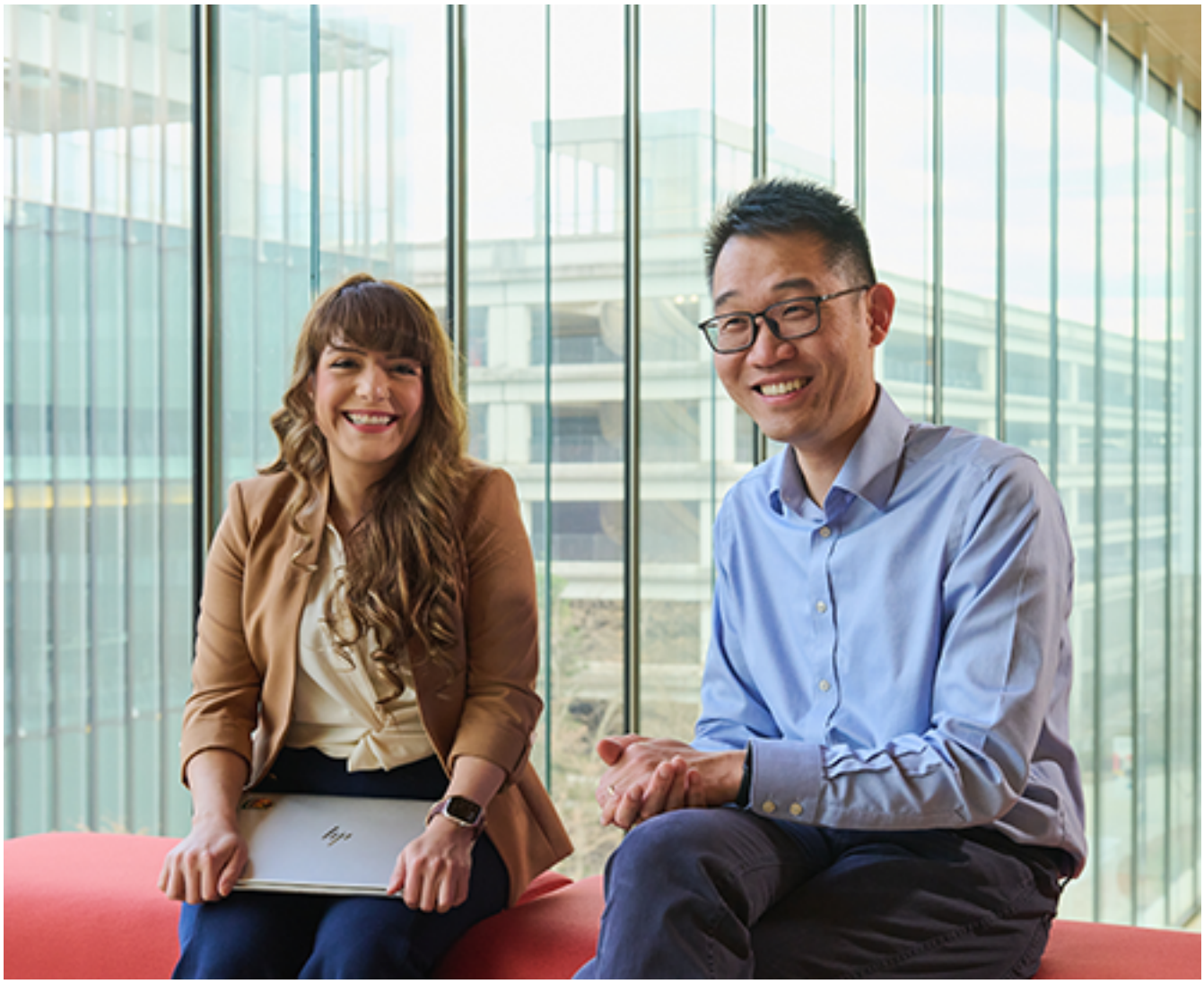

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