

LOGESHWARAN YOUVARADJE

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Experience in Engineering quality assurance complaints handling, track wise. Knowledge of Good manufacturing practices, QMS, Training, regulatory, industry and corporate guideline requirement.

SKILLS

- 5+ Years of experience at Engineering Quality assurance –Audit handling, customer- complaints, validation & Qualification of Equipments- URS, DQ,FAT,IQ,OQ and PQ
- 1+Years of experience in Quality inspector the concepts – Approval of final materials, quality events, internal quality audit etc.

WORK EXPERIENCE

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| • Executive in Kawman Pharma | MAR 2026 – Till now |
| • Officer at Hikal | APR 2025 – FEB 2026 |
| • Executive at Solara Active Pharma Science | AUG 2021– APR 2025 |
| • Quality Assurance inspector in Hi-tech Engineers | AUG 2017 – JUN 2019 |

EDUCATION

- B. Tech Mechanical engineering, Manakula vinayagar institute of technology, Puducherry India 2013-2017.

CERTIFICATE

- 5S & KAIZEN practices
- Lean manufacturing

WORK EXPERIENCE

Kawman Pharma

Key Responsibilities

- Handling of new equipment Commissioning & qualification URS, DQ, FAT, IQ, OQ and PQ preparation, review and approval.
- Prepare and review of requalification, temperature mapping and revalidation activities of equipment's.
- Review and follow up of preventive maintenance checklist are completed within the timeline and equipment breakdown analysis.
- Tracking of project validation planner for the new facility and projects.
- Assessment and initiating of change control.
- Closing change control related action items and task within the timeline.
- Handling deviation, CAPA and verification of effectiveness check CAPA.
- Performing risk assessment for equipment, process and facility.
- Preparation and Review of CPV and annual product quality review documents.

- Handling of customer compliant and return goods.
- Preparation and review of periodical revision SOP.
- Handling internal audits and customer giving response to the observation.
- Completed Japanese customer audit and CDSCO in April -2026.
- Giving training to the employees as per the annual calendar planner along with CFT.
- Preparation of monthly report related to quality events.
- Providing support to the vendor qualification team.
- Issuance and archival of documents.

Hikal Ltd.

Key Responsibilities

- Responsible for monitoring and ensuring compliance in line with cGMP w.r.t assigned departments
- Review of preventive maintenance planner, procedure, breakdown analysis, root cause of equipment failure
- Preparation and review of PM checklist along with CFT.
- Monitoring of calibration and preventive maintenance activities
- Initiating change control in track wise, salesforce.
- Preparation and review of Quality Assurance procedures.
- Review of procedures related to Production, Warehouse, Engineering and other support functions.
- Handling of deviations, change controls, incidents & Market complaints.
- Monitoring of CAPA and necessarily follow up for closure.
- Co-ordination for internal audits and customer audits.
- To monitor equipment qualification, requalification, temperature mapping and revalidation activities.
- Preparation and Review of annual product quality review documents.
- Preparation of internal audits and customer audit response
- Issuance and archival of documents.
- To handle any other tasks, other than above, assigned by Head Quality Assurance.

Solara Active Pharma Science

Key Responsibilities

- Intake- Gathering information of the complaint like – situation, date of aware, writing description, writing complaint investigation summary.
- Initiating change control regarding quality events through track wise.
- Finding Investigation Root cause, drafting.
- Follow-ups for additional information and product returns, Completes Investigations within required timelines and support to close the complaint within the specified timeline.

- Pre-Approval and approval routine Periodic Re-Qualification and Qualification of Equipments and HVAC systems.
- Monitoring preventive maintenance performing within due date as per planner.
- Review of executed preventive maintenance checklist and verification status label.
- Review and approval of SOP.
- Review and approval of layouts and updating in SMF.
- Issuance of equipment log books and archival of records.
- Revising IQ, OQ, PQ and RQ protocol as per change control.
- Approval of executed documents as per GDP and GEP practice.
- Preparation of yearly planner for equipment's and temperature mapping.
- Pre-Approval and approval of temperature mapping protocol.
- Completing daily task, CAPA and effectiveness check of change control.
- Preparation and reviewing of master lists.
- Tracking of project validation planner.
- Handling customer audits monthly.
- Handled European union good manufacturing practice and Japan good manufacturing practice regulatory audits.

Quality assurance inspector

Key Responsibilities

- Monitoring the overall quality of processes, identifying improvement areas.
- CAPA for failure.
- Verification of manufacturing materials and final approval.
- Solving the problem in the customer end.
- Quality events.
- Preparing yearly trend.

Personal details

Date of Birth : 17/08/1995

Languages : English, Tamil, French