

QA Officer -Reputed Medical Device Industry -Delhi, NCR, India -3LPA -Kajal

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Company: Seven Consultancy

Location: Noida

Category: other-general

JOB DETAILS

- 1 Review & preparation of MES & other GMP Software Qualification Documents.
- 2 Preparation of URS, GxP Risk Assessment, Qualification protocols (OQ, PQ), RTM, VSR.
- 3 Review URS, RA, Design documents, qualification document.
- 4 Handling of post implementation changes documents.
- 5 Execute OQ and PQ.
- 6 To prepare compliance report for internal / external audit and Compliance of internal/external audit points.
- 7 To Initiation, investigation and evaluation of Change control, CAPA and deviation.
- 8 To prepare & review SOPs.
- 9 To initiate, investigate, lock and closing of TCC, CC, Deviation and Incident report and verify the implemented CAPA and its effectiveness.
- 10 To Prepare PQR/APR.
- 11 Investigation and report preparation of Market Complaints and product failures at shop floor.
- 12 Online review of GMP documents.

FUNCTIONAL AREA

QA Officer, QA Officer jobs in delhi, QA Officer jobs in NCR, Regulatory affairs, Regulatory affairs jobs in delhi, Regulatory affairs jobs in NCR, Quality control, Quality control jobs in delhi, Quality control jobs in NCR, Risk management, Risk management jobs in delhi, Risk management jobs in NCR, Auditor, Auditor jobs in delhi, Auditor jobs in NCR, Medical

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