

## CURRICULUM VITAE

T.M.S.Reddy.  
S/o T.Venkat Reddy,  
Plot No 43,  
Tulasaivanam colony,  
Hayathnagar,  
Hyderabad,  
Telangana.  
INDIA. 501511

Email: msreddy9666749333@gmail.com  
(+) 9182989167

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### **OBJECTIVE:**

To seek a challenging career in Quality Assurance Department in a growth oriented leading organization, which provides opportunities for continuous growth and learning.

### **Total Experience:**

Around 19 years 0 months in Quality Assurance Department in Reputed Pharmaceutical Organizations.

### **PRESENT EMPLOYMENT:**

Presently working as Manager in Quality Assurance Department at Chemic Life sciences private limited (**ISO 9001:2015, GMP certified facility**), Choutuppal facility from June-2020 to till date.

### **PREVIOUS EMPLOYMENT:**

Worked as Manager in Quality Assurance Department for Lucent Drugs Private Limited API facility(**ISO 9001:2015, WHO GMP certified facility**), Kazipally facility from June-2017 to May-2020.

Worked as Manager in Quality Assurance Department around 2.5 years for M/s Tagoor chemicals (**An ISO 9001:2015 certified facility**). Jeedimetla.

Worked as In-charge in Quality Assurance Department around 3 years for M/s Shree Jaya Laboratories Pvt Limited (**An USFDA certified facility**). Choutuppal, Nalgonda district.

Worked as Executive in Quality Assurance Department for 3 years 6 months in M/s Vasudha Pharma chem. Limited (**An USFDA,KFDA,MHRA and ISO 9001:2008 certified facility**), Jeedimetla.

Worked as Chemist in Quality Assurance Department for 3 Year in Vasudha Pharma chem. Limited (**An USFDA, KFDA, MHRA and ISO 9001:2008 certified facility**), Jeedimetla.

### **ACADEMIC QUALIFICATIONS:**

- ❖ B.Sc from Osmania University 2003.
- ❖ Board of Intermediate 2000.
- ❖ SSC from Board of Secondary Education, Andhra Pradesh 1998.

### **Summary of Work Experience:**

- Involved in Vendor Qualification and Evaluate of new Vendors for Raw Materials, Packing Materials and continuous monitoring of supply chain approval and disapproval of vendors with annual trending.
- Preparing/Reviewing of Process validation protocols and reports, Monitoring of its effectiveness.
- Preparing/Reviewing of Cleaning validation protocols and reports related to Product change over, Batch to Batch, Periodical Cleaning Validations as per the Regulatory requirements.
- Participation in Line-clearance activities.
- Stability studies/ Holding Studies initiation and monitoring of results in various conditions.
- Preparation & Review of Annual Product Quality Review Report (APQR) by covering all the aspects.
- Handling of Change control management.
- Handling of Deviations, OOS, OOT, and Return Goods & Complaints by participating in investigations for identifying of Root cause.
- Responsible for Preparation of Audits presentations various Customers full fill the requirements as per audit agenda.

- Responsible for preparation of Audit response for the observation made during the audits as per the CAPA obtained from the internal departments.
- Participate in self-inspections & identifying non conformities and verifying its effectiveness.
- Preparing & Updating of Site master File (SMF) & Validation Master Plan (VMP).
- Review of Executed BPCRs, ECRs along with Analytical Quality Data.
- Planning and monitoring of product dispatches as per customer requirement and committed timelines.
- Reviewing of Qualification documents of Equipment's, Facility, water systems.
- Preparation/Reviews of the Performance Qualification Protocols & Reports for equipment's.
- Reviewing and follow up the calibration, preventive maintenances of all Departments.
- Responsible for investigation of Out of calibration results.
- Conducting the cGMP trainings and on job trainings periodically.
- Participation in safety committee meetings, Management Review Meetings.
- Provide support to the various divisions for effective and efficient implementation of the QMS.
- Manufacturing Licenses and GMP certificates filing and Responsible for Documents archival.
- Giving guidance to colleagues for quality systems and cleanliness of the work places.
- Responsible for implementation & Inspection of respective area operations periodically and check on the implementation of Standard Operating Procedures (SOP) during the operations.
- Ensuring that the changes to the system or process are being implemented with the valid approach as per the established norms/systems for effecting the changes with due approvals from the concerned departments and customers and regulatory authorities, where defined.

**Audit faced:** More than 50 audits Like KRKA, Dr. Reddys, TEVA, GVK Life sciences, Mylan laboratories, Aurobindo Pharma, MSN laboratories, Cadila pharma, Jubliant Life sciences, Morepen labs, Indi swift pharma, Sun pharma, Mankind pharma, Lee pharma, Raks pharma, Hetero drugs, Smilax laboratories, Venkata narayana API, Vasudha pharma chem. Limited, Metrochem API, Rakshith drugs, IPCA laboratories, Fleming laboratories etc...

**Marital Status: Married.**

**Date of Birth: 28/08/1983.**