

CURRICULUM VITAE

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1. OBJECTIVE :

To pursue a challenging career in **Quality Control**, which enables me to actively participate an ample scope to enhance my knowledge and skills, and be recognized for the service.

2. EXPERIENCE DETAILS :

- ⇒ 14+ years of extensive experience in Quality Control Department,
- ⇒ Now working as, a Dy. Manager in Shreyasvi Life sciences Pvt limited, Hyderabad from Nov-2021 to till date.
- ⇒ Worked in Aurobindo Pharma Ltd (Unit-XV) as Sr. Executive, from September-2017 to March-2020 in QC.
- ⇒ Worked in Quality control as a Sr. Executive, in HETERO DRUGS LTD UNIT-IX, Nakkapalli (Vizag) from July-2013 to Sep-2017.
- ⇒ Worked as a Quality Control Sr. Chemist, in Auctus Pharma Ltd, Parawada, Visakhapatnam from May-2011 to June-2013
- ⇒ Worked as a Quality Control Chemist in Sarvotham care Ltd, Baddi, HP. from Aug-2008 to May-2011.

3. EDUCATIONAL QUALIFICATIONS :

M.Sc (Organic Chemistry) From Acharya Nagarjuna University, Guntur.

B.Sc (Chemistry, Physics and Matches) from Bhasker Degree college, Parvathipuram.

4. KEY RESULT AREAS :

- ⇒ Work Planning and Audit readiness.
- ⇒ Having Exposure on EMPOWER, LIMS and Audit trail verification.
- ⇒ Handling of OOS, LIRs and Change controls

5. JOB RESPONSIBILITIES :

❖ Lab Management:

- ⇒ Development, implementation and monitoring of quality systems.
- ⇒ Ensure data integrity in the laboratory.
- ⇒ Review of daily activities in laboratory and resolving the issues.
- ⇒ Ensuring compliance to Pharmacopoeia monographs Investigation of Laboratory incidents, OOT and OOS.
- ⇒ Analytical Method Transfers.

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❖ Summary of Major Responsibilities:

- ⇒ Manage, plan and direct the raw material section to ensure quality support for manufacturing and to ensure the testing is executed in a timely and compliant manner.
- ⇒ Review, plan and direct the Method Transfer and Process validation to ensure quality support for manufacturing and to ensure the testing is executed in a timely and compliant manner.
- ⇒ Review of analytical method performance through life cycle management of analytical methods and improving the test methods.
- ⇒ Planning and work allocation, assurance the completeness of the activity.
- ⇒ Review of analytical documents and compilation.
- ⇒ Handling of laboratory incidents and deviation in (QMS).
- ⇒ Review of all analytical reports like Finished product and Raw material.
- ⇒ Daily GLP rounds in Quality control area to reduce the non-conformances.
- ⇒ Documents Review in Caliber E-LIMS Software.
- ⇒ Handling of OOS and OOT investigations.
- ⇒ Analysis of in-process, finished products and Validation samples.
- ⇒ Documentation of relevant analysis by using LIMS software.
- ⇒ Monitoring preventive maintenance & Calibration activities in the QC.
- ⇒ Preparation of the stability protocols and stability data sheets, review of stability data.
- ⇒ Review of audit trails.

❖ Instruments & Soft wares Handled:

- ⇒ HPLC (Waters2489,2487, Empower3 software)
- ⇒ HPLC (Agilent Technologies 1200 series) HPLC (Shimadzu 2030 with cromeleon sr-14)
- ⇒ GC (Agilent 7890A) with EMPOWER-3
- ⇒ Shimadzu GC2012 with Lab Solutions
- ⇒ U.V-VISIBLE Spectroscopy (Schimadzu)
- ⇒ KARLFISHER Titrator (Metrohm)

6. ACHIEVEMENTS :

- ⇒ Involved as a review team member in audits of USFDA, EDQM, WHO, KFDA, and ANVISA

7. PERSONAL DETAILS :

Date of Birth : 17/07/1985
Fathers Name : Lakshmana Rao
Permanent Address : Chollapadam, Komarada (M), AP.
Marital Status : Married
Languages Known : Telugu, English and Hindi

8. DECLARATION :

I hereby declare that the above furnished details are true and correct to the best of my knowledge.

Place: Hyderabad

Date :

Signature:

(Ch.Venkateswara Rao)