

CURRICULUM VITAE

A.vamsi

✉ E-Mail: alurivamsi112@gmail.com

☎ Phone : +919010909422

Career Objective:-

To join health care/ Pharmaceuticals where I can utilize my knowledge and skills for the development of the company and where opportunities exist for the growth of my career and where I could come across the latest technology and to be part of the team that work dynamically towards the growth of the organization.

Educational Qualification:-

Course	Board/ University	Institute	Place	Year of Passing	Percentage of marks
B.Sc	A.N.U	N.N.S degree college	Chirala	2014	67%
Intermediate	Board of intermediate	Sri Vani J.R. College	Chirala	2011	50%
S.S.C.	S.S.C.	Vasavi High School	Chirala	2009	65%

Strengths:-

- Hard working with Positive attitude.
- Capability to mold myself to the situation.
- Sincere & Punctuality.
- Zeal to learn new things.
- Capability of being a good and efficient team player.

Area of interest:-

- Developing side.

Computer skills:-

Operating Skills : Windows Family.

Application Software's : Microsoft office.

Working Experience: -

Total experience- 10+ Years

Time period	Job Title-Dept.	Company Details	Company Profile
Till date To 05 Aug 2020	Executive-QA	M/s Sionc Pharmaceuticals Pvt. Ltd., (Unit-I), Parawada, Visakhapatnam, India	USFDA, EUGMP and KFDD certified Company & is a leading manufacturer of API's & Oncology Drugs in India.
26 Jul 2020 To 23 Oct 2017	Sr. Chemist-QA	M/s Rampex Labs Pvt. Ltd., (Unit-II), Parawada, Visakhapatnam, India	An ISO 9001: 2015 certified Company & is a leading manufacturer of API's & Intermediate's in India. Produced doc. To USFDA and Received FEI No.
22 Oct 2017 To 14 Apr 2014	Officer-QA To Chemist-QA	M/s KRS Pharmaceutical Pvt. Ltd., (Unit-II) Parawada, Visakhapatnam, India	An ISO 9001: 2008 certified Company & is a leading manufacturer of API's & Intermediate's in India.

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Core Competencies:-

- Preparation and Review of Batch production control record, Standard operating procedure's, Validation Protocol and Report, Agreements, Water and Environmental Trends & Annual Product Quality Review reports.
- Ability to handle validations like Process, Cleaning, Utility's and Equipment Qualifications, Trainings.
- Preparations of Technology transfer documents.
- Monitoring of Schedule & Validation Master Plan.
- Logging and handling of VMP, Change controls, Deviations, OOS, Complaints, Recalls, Return Goods, Initiation documents.
- Filling of Customer Vendor Questionnaires and Vendor management system
- Controlling all quality related documents for retrieval, issue and retention as Quality Document Management System.
- Releasing the API's & Intermediate's Product to Market as per the customer specifications / requirements, according to the Quality Norms (QA Release).
- Handling of Customer mail communications, Regulatory Updations and assisting in DMF Filing.
- Responsible for document retrieval and best documentation practices in timely manner.
- Online Audits (IPQA) based on the cGMP.
- Provide support to the various divisions for effective and efficient implementation of the Quality Management System.
- Upgrade Quality Control process. Co-ordinate with Production, QC, warehouse and Maintenance regarding non-conformance & corrective action for the same.
- To assist Head-QA in continuous system updation.



Achievements:-

- ❖ Successfully faced the audits EUGMP, KFDA, ISO 9001: 2008, Mylan, Cipla.



Personal Details:-

Father's Name : Mr. A Ramesh

Date of Birth : 05th May 1993

Languages Known : English, Telugu.

Marital Status : Married.

Present address : D.No. 11-2-5, kailsh Nagar, old gajuwaka, Visakhapatnam.

Permanent Address : Back side of dhanalakshmi Rice mill, Kothapeta, Chirala, Pin code : 523157
Prakasam (DIST), A.P.



Declaration:-

I hereby declare that all the above-mentioned information is true to best of my knowledge and belief.

Place : Visakhapatnam.

Date :

A.VAMSI