

## APPLICATION AND CURRICULUM VITAE

To,  
The Head of Recruitment,

**Subject:** Willing to work with your organization in Quality Assurance.

**Applicant:** Mr. Rambir Singh

**Experience:** -10 Years and 02 Months

Respected Sir / Madam,

Please accept mine best wishes and regards.

Sir / Madam, I am well educated, experienced, sincere and hardworking, independently handled to assigned works and graduated in pharmaceutical.

I believe in loyalty, confidentiality and achievements.

I will utilize my knowledge and experience to meet to fulfill all stages of the finished products with quality aspects. I will effectively liaison with all your regulatory requirements.

Kindly confirm suitable time for meeting.

I hope, you will be given a positive response.

Thank you in anticipation.

Yours sincerely,

Rambir Singh

Bachelor of pharmacy

**Contact & whatsapp No:** +91-9355655529

**Email:** [rsrambir5529@gmail.com](mailto:rsrambir5529@gmail.com)

**Enclosure:** curriculum vitae.

## CURRICULUM VITAE

**RAMBIR SINGH**

**Contact & whatsapp No:** +91-9355655529

**Email:** [rsrambir5529@gmail.com](mailto:rsrambir5529@gmail.com)

### **CAREER AMBITIONS**

To work on Challenging assignments that provides benefits of the job satisfaction & a steady professional growth and to continuously aim towards innovation, perfection and excellence in pharmaceuticals with sound knowledge in Quality Assurance and achieve a position of Quality Assurance Specialist.

### **SKILLS**

- Leadership
- Interpersonal Skills
- Communication Skills
- Time Management
- Task oriented.
- Quality Management System (Change Control, Deviation, Incidents, NCR, Investigations & CAPA, cGMP & GAMP Compliance, Training, APR, Market Complaint, Internal / External Audit, OOT & OOS)
- Proactive approach during Qualification & Validation activities.
- Data Analytical Skills

### **WORK EXPERIENCE, ORGANIZATION DETAILS AND JOB PROFILE**

| ORGANIZATION  | LOCATION                    | FUNCTION             | FORMULATION  | PERIOD  |           |
|---|-----------------------------|----------------------|--|---------|-----------|
| <b>HIGLANCE LABORATORIES PVT. LTD.</b><br>(WHO GMP, PERU GMP & RUSSIAN GMP Certified) | <b>Greater Noida, U.P.</b>  | Assistant Manager QA | Tablets, Capsules, Oral Liquid, Soft gel capsules                                    | 04/2018 | Till Date |
| <b>RAVIAN LIFE SCIENCE PVT. LTD.</b> (WHO GMP Certified)                              | <b>Haridwar, Uttrakhand</b> | Sr. Executive QA     | Tablets, Capsules, Oral Liquid,  | 07/2016 | 04/2018   |
| <b>GRACURE PHARMACEUTICALS LTD.</b><br>(WHO GMP, TGA & EU-GMP Certified)              | <b>Bhiwadi, Rajasthan</b>   | Sr. Executive QA     | Tablets, Capsules, Oral Liquid, Dry syrup, Ointment,                                 | 04/2015 | 07/2016   |
| <b>AKUMS DRUGS &amp; PHARMACEUTICALS LTD.</b><br>(WHO GMP Certified)                  | <b>Haridwar, Uttrakhand</b> | Officer QA           | Tablets, Capsules, Dry syrup   | 01/2014 | 04/2015   |
| <b>BRAWN LABORATORIES LTD.</b><br>(WHO GMP Certified)                                 | <b>Faridabad, Haryana</b>   | QA Chemist           | Tablets, Capsules, Oral Liquid, Dry injection, Liquid Injection, Ointment, Dry syrup | 06/2012 | 01/2014   |

### **AUDIT FACED**

- Successfully faced multiple time WHO audit in different organization like Higlance Laboratories, Ravian Life science and Gracure Pharmaceuticals.
- Successfully completion of following regulatory audit like Peru, Russia, Uganda, Sri Lanka & Kenya.
- Successfully faced different P2P customer audit like Wockhardt, Alkem, Abbott, Cadila, Lincoln, etc.

### **EDUCATIONAL QUALIFICATION**

Graduation in “Bachelor of Pharmacy” from “Maharishi Dyanand University, Rohtak” (2011).

## JOB RESPONSIBILITIES FOLLOWED

- **Audit preparation and Overall compliance of regulatory requirements and maintaining cGMP system in the organization.**
- **Utility validations (Purified Water System, Compressed air System, HVAC system with ISO-14644).**
- Preparation & implementation of the first level documents like Site Master File, Validation Master Plan, Quality Manual as per the cGMP requirements.
- Gap assessment of the SOP, revision of SOP, implementation of the revised SOP in the existing system effectively.
- **Coordination with cross functional team for timely execution, completion of the qualification and validation activities as well as QMS activities.**
- Manage the team members (IPQA) and Managed all in process activities with the help of IPQA persons and documentation work as per their capabilities and allot their responsibilities.
- **Qualification of equipment's and systems with GAMP-5 in atomization equipment's and software systems.**  
(*Quality control: Autoclave, stability chamber, LAF, Dynamic Pass box, BOD incubator, Cooling chamber etc).*  
(*Production: RMG, Octagonal Blender, Compression machine, Blister packing machine etc.*)
- Preparation of Hold time study Protocols & Execution for Hold time study (*For products, cleaned equipment's, unclean equipment's, volumetric solutions, purified water, staging of material, bulk, compressed tablets, coated tablets, coating solution, binder solution, filled capsules).*
- Preparation of protocols and Execution of Process Validation.
- Investigation of root causes, implementation of corrective and preventive actions. Handling of Deviations, Non-conformances, Change Controls, Market Complaints and CAPA Tracking.
- Preparation of Training syllabus, Training needs, Training modules & training calendars. Imparting training on shop floor.
- **Review of documents** (SOPs, Master & executed Batch Records (BMR & BPR), COA of bulk, semi finish, finish, STS & STP, analytical method validation & analytical method verification, environment monitoring data, analyst validation, calibration of the instruments, OOS & OOT etc.
- Preparation of Stability Study protocols & bracketing for the stability studies.
- Participating in Internal Audit team & in inter departmental audit program.
- Active participation during Technology Transfer from one site to other sites and product development to commercial scale up.
- **Prepare and performed risk assessment with severity occurrence and detection, comparative study data respect to material & equipment equivalence, handled, update, track, completion and trending of the QMS tools (Deviation, Change Control, Market Complaint, and Incident) on yearly basis.**

## ACHIEVEMENTS

- Participated in Three days training programme on “Qualification & validation” presented by Dr. Rao at Akums drugs & pharmaceuticals.

- Participated in training programme on “Market complaint & investigation, CAPA” presented by Mr. Prabhakar Patil, CQA head, Alkem laboratories.
- **Get the “Best productivity award with QMS system for the 2017” in the Ravian Life Science Private Limited.**

### CAPABILITIES

- Capable of making good decisions and able to solve problems / issues pertaining to people, processes or systems without compromising on the quality.
- Capable of taking higher responsibilities to contribute towards the growth of the organization.
- Capable of leading the team by example, to complete the assigned responsibilities within the timelines.
- Capable of motivating people and receiving the best efforts out of them.

### STRENGTHS

- Fully conversant with all relevant Quality Assurance procedures.
- Ability to evaluate audit findings and implement appropriate corrective actions
- Ability to work independently and in a team environment. Work well in an interdependent team environment, and promote positive, respectful professional relationships
- Strong proficiency in common office applications (e.g., Word, Excel, PowerPoint, Outlook, Adobe Acrobat, etc.) and demonstrated adaptability and rapid fluency to new relevant applications.
- **Hard working and motivated Positive ‘can do’ attitude.**

### REFERENCE

- Mr. Teeka Ram Sharma, Sr. G.M. Quality Assurance in Kwaliti Pharmaceuticals Amritsar. (9694726387)
- Mr. Sanjeev Yadav, Head Quality Assurance in Unicare India Pvt. Ltd. Greater Noida. (9908330503)
- Mr. Sanjeev Kumar Singh, Head Quality Assurance in Shree Shyam Life Science Baddi. (9045498780)

### SALARY AND OTHER DETAILS

|   |   |
|---|---|
| <b>Current In-hand salary:</b> 6.99 lac per Annum | <b>Notice period:</b> 45 days                           |
| <b>Gross Salary:</b> 7.51 lac per Annum           | <b>Expectation:</b> 25 percent of current package       |
| <b>Cost to Company:</b> 7.73 lac per Annum        | <b>Post applied:</b> Deputy Manager / Assistant Manager |

### PERSONAL DETAILS

|                        |   |                   |  |
|------------------------|---|-------------------|--|
| <b>Father’s Name</b>   | : | Sh. Chhotey Lal   | <b><u>Address:</u></b>   |
| <b>Date of Birth</b>   | : | 30 Nov. 1988      | <b><u>Current address:</u></b> H.No.143, Delta-I, Greater      |
| <b>Sex</b>             | : | Male              | Noida, U.P. (201308).  |
| <b>Nationality</b>     | : | Indian            | <b><u>Permanent address:</u></b> Village-Ronija, Post-Asawata, |
| <b>Religion</b>        | : | Hindu             | Tehsil-Palwal, Distt.-Palwal, Haryana (121102)                 |
| <b>Marital Status</b>  | : | Married           |  |
| <b>Languages Known</b> | : | English and Hindi |  |

### DECLARATION

I hereby declare that the information given above is true to the best of my knowledge.

Place:\_\_\_\_\_

[RAMBIR SINGH]