**RAHUL BHARDWAJ**

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**Permanent Address**: H.NO-32, Sihi Gate, Rajwara, Ballabgarh-Faridabad (121004)

Haryana

**CAREER OBJECTIVE:**

**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.**

**PROFESSIONAL SUMMARY**

**Approx. 11 years working experience in the field of QA Compliance, Validation, Qualification, & QMS. (HVAC Validation)**

* Effectively co-ordinate with various departments and effectively handed for customers and regulatory audits.
* Good understanding of c**GMP, WHO**, **ISO 9001, CQA** etc.
* Excellent Communication, Functional / Presentation / **Technical writing & time Management** skills.
* Excellent Leadership, **Team Facilitator, Decision Making**& Organizational People development Skills.

**EDUCATIONAL QUALIFICATION:**

* Completed **“Bachelor of Pharmacy”** from “**M.D.University, Rohtak”**.(2011)

**TRAINING & CERTIFICATIONS:**

* Done 45 days training in Brawn Laboratories Ltd. Faridabad.

**AUDIT FACED:**

* WHO, EU-GMP,UGANDA, KENYA, UKRAINE, VIETNAM, MYANMAR, ETHOPIA, US Labs., UNILAB, AVENTIS, CLARIS LIFESCIENCE, CIPLA, SUN PHARMA, JUBLLIENT LIFE SCIENCES & TORRENT, ABBOTT, BLUECROSS.

**SOFTWARE SKILLS:**

* Windows operating systems MS Office, Word, Excel, Power Point Presentations& Internet skills.
* Practical knowledge of software such as **SAP, ERP and BIMS (Batch Issuance Management System).**

**ORGANIZATIONAL EXPERIENCE:**

1. **Arbro Pharmaceuticals Pvt. Ltd. (May-2021 to Till )**

**DESIGNATION- Manager QA**

**Arbro Pharmaceuticals -** is involved in the manufacture of a wide range of pharmaceutical products. The Plant located at Kirti Nagar Industrial Area, New Delhi is a state-of-the-art facility with critical equipments and latest technology for the manufacture of Tablets, Capsules, Liquid Orals, Herbal and Cosmetic having **WHO-GMP and ISO-9001-2015** approvals and having Export for various country like Kenya, Cambodia, Afghanistan etc.

**CURRENT JOB RESPONSIBILITY IN DAFFODILLS:**

* cGMP Compliance.
* Responsible for artwork approvals
* Vendor Management.
* Initiation , review and trending of change control, Deviation, Investigation , CAPA.
* Management of Product Quality Complaints.
* Review of Documents Like Batch Production Record, Validation Protocols.
* Management of SOPs including distribution and archival.
* Preparation of Process Validation protocol & Compilation of report.
* Handling all approvals related to QMS activity.

1. **Daffodills Pharmaceuticals Limited (Nov-2017 to April-2021)**

**DESIGNATION- Asst. Manager QA**

**Daffodills Pharma** is involved in the manufacture of a wide range of pharmaceutical products. The Plant located at Rohta Road, Jawahar Nagar, Meerut (UP), is a state-of-the-art facility with critical equipments and latest technology for the manufacture of Tablet, Capsules, Liquid Orals, and Injection having **WHO-GMP and ISO-9001-2015** approvals and having Export for various country like Cameroon , Philippines and Sri Lanka, Kenya, Uganda etc.

**CURRENT JOB RESPONSIBILITY IN DAFFODILLS:**

* cGMP Compliance.
* Responsible for artwork approvals.
* Vendor Management.
* Handling all approvals related to QMS activity.
* Handling QA/QC Activity.
* Management of Product Quality Complaints.
* Review of Documents Like Batch Production Record, Validation Protocols.
* Initiation , review and trending of change control, Deviation, Investigation , CAPA.
* Management of SOPs including distribution and archival.
* Preparation of Process Validation protocol & Compilation of report.
* Handling all approvals related to QMS activity.

1. **AKUMS DRUGS AND PHARMACEUTICAL LTD.(AKUMS GROUP) (08 AUG. 2016 TO OCT-2017)**

**DESIGNATION- Sr. EXECUTIVE**

Akums group is involved in the manufacture of wide range of pharmaceutical product. Akums group has Twelve manufacturing plant having wide range of manufacturing of Tablet, Capsule, Liquid, Ointment, Sachet, Dry Syrups, Liquid injection, Dry injection, Large Volume Parenterals and Akums group pure and Cure healthcare Pvt. Ltd. Haridwar plant having Asia largest third party manufacturing plant in whole asia.

**JOB RESPONSIBILITY IN AKUMS:**

* Preparation, issuance & retrieval of SOPs.
* Actively Involved in Project of Two Large line.
* Preparation of Process Validation protocol & Compilation of report.
* HVAC Validation.
* Qualification of Equipments.
* Review of BMR and BPR.
* Initiation , review and trending of change control, Deviation, Investigation , CAPA

1. **AHLCON PARENTERALS INDIA LTD.(A B/BRAUN GROUP COMPANY) (16 APRIL 2014 TO 25 AUG. 2016)**

**DESIGNATION-QA EXECUTIVE**

Ahlcon Parenterals (India) Limited which is now a B/Braun group Company(German MNC) and is involved in the manufacture of Parenterals preparations (LVP and SVP) by employing a highly sophisticated production process; The dosage are clear solution which is produced in the conditions using state of the art FFS Technology.

**JOB RESPONSIBILITY IN AHLCON:**

* Working in Project stage organization to helping the project & German engineering team to perform the activity as per the regulatory norms.
* Preparation & Review of documents such as Equipment / Instruments SOP (Operational / cleaning & Calibration).
* Responsible for distribution, revision, retrieval, archival & destruction of controlled documents.
* Prepare & Review of protocols & review of **Qualification / Validation** / **Re-Qualification** and Commissioning of Equipments of Manufacturing as per the Regulatory requirements.
* Prepare the protocol and report of HVAC system (**URS, DQ, IQ, OQ**, and **PQ**) Process Validation, Hold Time Study.
* Review audit observations to ensure compliance & Conduct internal audit (self inspection) to ensure compliances on the deviations.

1. **BRAWN LABORATORIES LTD.( 03 May 2011 TO March 2014)**

**DESIGNATION- QA Executive**

**BRAWN LABS-** is involved in the manufacture of a wide range of pharmaceutical products. The Plant located at FCA-13, NIT FARIDABAD, (HR) Industrial area is a state-of-the-art facility with critical equipments and latest technology for the manufacture of Tablet, Capsules, Liquid Orals, and Injection having **WHO-GMP and ISO-9001-2008** approvals and having Export for various country like Kenya, Myanmar, Ukraine, Vietnam and Uganda etc.

**JOB RESPONSIBILITY IN BRAWN:**

* Manage all in process activities with the help of IPQA persons.
* Compilation of documents like change control, market complaint, deviation &non compliance report.
* BMR & BPR issuance, receiving & reviewing.
* Preparation and Approval of Standard operating Procedures.
* Do all the in-process activity in the area of dispensing & material receiving of the raw material and packing material, granulation, blending, compression, primary packing & secondary packing.
* To Check Instruments Calibration:
* Disintegration Apparatus
* Friability Apparatus
* Weighing Balance.
* Preparation of APQR.
* Ensuring that all Log sheets & Records are updated on-line with the operation.
* Monitoring of Temperature, Humidity, Cleaning, and Sanitation of all area.

**DECLARATION:**

I, undersigned, confirm that to the best of my knowledge, this CV correctly describes my experience, my qualifications and me.

**REFERENCES**:

My working experience and my dedication towards my responsibilities and work.

**PERSONALDETAILS:**

* Name : Rahul Bhardwaj
* Father’s Name : Sh.Ved Prakash
* Date of birth : 15th Mar. 1987.
* Gender : Male
* Nationality : Indian
* Marital status : Married
* Languages known : English, Hindi

Place: Faridabad (Rahul Bhardwaj)

Date**:**