**CURICULUM VITAE**

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| **Objective** | Looking forward to associate myself with an organization, where there is an opportunity to continue and upgrade my knowledge for self and organization served. |

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| **Personal Skills** | Having 12+ Years of experience with comprehensive Problem Solving abilities, Excellent verbal and written communication skills, ability to deal with people, willingness to learn, team facilitator. |

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| https://lh4.googleusercontent.com/FWuFexnGcKAtASh02pWdjN2QAzip37zp8CIQBYm5kuPQY2YrfPq65VdR9gpB-iczGitboIKXQXnWlOm49-YSBDBfvciIMh0Hm3qra4-eDnostvPyt4E  **Highlight of Professional Career** | Excellent exposure to product **Manufacturing and Packing**, independently.  Coordination with **Regulatory Affair, Corporate Quality assurance, and Packing Development Dept**.  Key role-played in regulated market approvals of **USFDA, ANVISA, WHO-GMP, TGA, IMB AND MHRA, EU-GMP AUDITS, TFDA & ISO 14001**.  Review of Equipment Qualifications (**URS, DQ, IQ, OQ, PQ**). Preparation of Equipment Qualifications  Having Excellent knowledge of **TRACKWISE** used for the Deviation, change control, OOS, OOT and for the PQC(product quality complaint)  Independently handle the market complaint and share the interim report within given time line.  PQC (Market Complaint) will be closed by performing the investigation with proper root cause and CAPA. Implementing major cost savings strategies**,** productivity enhancements. |

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| **Education** | |  |  |  | | --- | --- | --- | | **Examination** | **School/College/University** | **Year** | | **M. Pharmacy** | **Hamirpur Tech. University** | **2014** | | **Bachelor in Pharmacy** | **U P Tech. University** | **2006** | | **12th** | **Uttar Pradesh Board** | **2000** | | **10th** | **Uttar Pradesh Board** | **1998** | |
| **Professional**  **Experience** | ***AUROBINDO PHARMACEUTICAL LTD.***  **(FEBURARY 2016 to TILL DATE)**   * Working as **Manager Production-Packing.** Responsible for the packing activity and produces 950 mil every month. * Leading the Team of 300 person including Deputy manager, Assistant manager, Executive and operators. * Support operation support team Investigates and conducts trouble-shooting of unexpended trends or results related to product quality – to include root cause analysis and CAPA determination. * Specific responsibility for meeting Production schedules with efficient and optimized use of resources and machine time. * Interfaces with Supply Chain to develop mid- and long-range Master Production Schedules. * Coaches sub-ordinates to build capabilities, skills and technical knowledge. * Ensures adequate resources are available to meet current and projected production volumes. * Collaborates with Validation Department in qualification / validation of equipment and processes. * Prepare and review the SOP, investigation reports, performance appraisals and technical reports. * Strong problem-solving capabilities derived from extensive equipment knowledge, demonstrated track-record of continuous improvement yielding tangible business results * Strong leadership track-record with  high levels of ownership, accountability and technical competency at all levels within the organization. * Has extensive experience in managing cGMP compliance needs, deployment of Quality Management Systems and relations with external auditors and regulators and USFDA audit exposure.   ***JAGSONPAL PHARMACEUTICAL LTD.***  **(SEPTEMBER 2014 to FEBURARY 2016)**  Working as **Assistant Manager (Head of Department)** Production & packing and responsible for the production as well as for the packing activity for the 02 Block ( General Block and Oral contraceptive Block).  **Company profile**: Jagsonpal is among India’s premiere pharmaceutical companies. Founded in 1964 the company specializes in developing and manufacturing bulk drugs and pharmaceutical formulations. Over the past 40 years the firm has grown by leaps and bound.  ***RANBAXY LABORATORES LTD.***  **(DECEMBER 2008 to SEPTEMBER 2014)**  Working as **Production Executive**  **Company profile**: The organization shares the top position in all dosage forms in global market and largest manufacturer of solid products in worldwide. In Dec 2013 successfully faced the multiagency Audit at Paonta Sahib Plant and gets approval. It was first plant in INDIA who faced the multiagency audit.  ***NECTAR LIFESCIENCES LTD****.*  **(AUGUST 2007- NOVEMBER 2008)**  Worked as **Officer Production**  ***JUPITER REMEDIES***  **(JULY 2006 –JULY 2007)** |
| **Job Profile in Companies** | To handle the regular activity and planning of Manufacturing and Packing section with their good OEE Standards.  Having experience of project set up in Ranbaxy as well as in Nectar life sciences.  To schedule the production and packing on the basis of weekly, monthly and Yearly plan.  To execute the production and packing schedule and co-ordination with the QA, QC and warehouse Dep’t for effective team coordination and communication.  Also having good knowledge of Warehousing and scheduling.  To have effective control and unitization of manpower on daily basis.  To ensure online documentation of production activity for timely dispatch.  To take training programmers periodically to ensure that production takes place as per cGMP Guidelines.  To guide the production and packing operators and Trouble Shooting in case problems or issues on day to day activity for monitoring as schedule delivery of products.  To ensure strict adherence to SOP’s during all stages of production and packing.  Implementation of new concept for improving quality, productivity and safety such as **KAIZAN and PULL PUSH CONCEPT.**  To trouble shoot any technical or equipment problems in both formulation and packing.  To ensure the well maintenance and up keeping of the production premises.  To raise purchase requisition and to maintain inventory of miscellaneous requirement.  To Raise the **Purchase request, REVEX and CAPEX** for the machine and their spare parts.  To work under the SAP and ERP system in the coordition with the other departments. Having good knowledge of **SAP(Version ECC 6.0)**  To work under the guidelines of cGMP & to fulfill the requirement as per requirement.  Perform root-cause analysis of defects while developing & improving the processing by taking corrective actions to prevent any reoccurrences and defective issues.  Map the process while conducting the gap analysis, documenting all work flows and review of Impact Analysis Document for further processing, ensuring action plan as per requirement.  Having Excellent knowledge Deviation, change control, OOS, OOT.  Independently handle the market complaint and share the interim report within given time line.  PQC will be closed by performing the investigation with proper root cause and CAPA.  Having Excellent knowledge of **LMS** (Learning management solution) used for conducting the training. |
| **Audit Exposure** | **USFDA, ANVISA, WHO-GMP, TGA, IMB AND MHRA, EU-GMP AUDITS & MCC**. |

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| **Computer Skill** | Ability to work on MS words, Window 98, 2000,2007, Command on word, Power Point, Excel Sheet & the entire general program. Experience on SAP, ERP, LMS and TRACKWISE Application. |

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| https://lh6.googleusercontent.com/PCqpgbRg6xcM_f6xNFwPAa6NG_UXXHUSphkzwoCZ1D1RC_N0FWdJ8vg6Kb6gjIR9lbQcXZrWBHWxgG7Gq_H8UOeMHk4EoKIynwCd4-l5xd-cYrkJ2Tg  **Personal Details** | FATHERS NAME: Shri Kumud Kumar.  PERMANENT ADDRESS: H.No.111  Vill- Purna, Post- Amilaudha, Dist- Sonbhadra  Uttar Pradesh- 231210  SEX: Male  MARITAL STATUS:  Married  Mail ID : [viveksingh.kks@gmail.com](mailto:viveksingh.kks@gmail.com)  Mobile No. : 07060777277 & 07060777377 |