Ram Wandhekar

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CURRICULUM VITAE

CAREER OBJECTIVE:

To expand my knowledge in the field of Quality and to excel in my area of activity by taking up challenging assignments and contribute to the organization by working as a key player in challenging & creative environment.

PROFESSIONAL PROFILE:

- **Total 13.5 Years experience with exposure of Oral liquid, Tablets and Capsules, API, MDI-nasal spray and Eye/Ear drops, Sterile.**
- Presently working as **Deputy Manager QC, in Piramal pharma Limited. Pithampur, Indore** in QC department (As Investigator-Handling QMS-Event, Incident, OOS, OOT, Change control and Deviation of FG and Stability section) The Piramal pharma is Manufacturer of Solid dosage forms (Tablets, Capsules) and Ophthalmic, oral liquid, eye and Ear drops.
  Period: From 17th April 2023 to till date.
- Worked as **Assistant Manager – in FDC ltd. Waluj Aurangabad** in QC department **(As Stability section head)** The FDC Ltd is Manufacturer of Ophthalmic, Oral liquid, Eye and Ear drops. Period: From 29th August 2022 to 11th April 2023.
- Worked as **Assistant Manager-** in **Zydus cadila healthcare Ltd.** Ahmedabad in QC department **(As a Reviewer Exhibit FP Section)**, The Zydus cadila is manufacturer of solid dosage forms (Tablets, Capsules). Period: From: 09th October 2017 to 21st August 2022.
- Worked as **Sr Executive** in **Cipla Ltd. Pithampur, Indore** in QC Finished Product section
  (As an analyst), which one is manufacturer of solid dosage form, (Tablets, Capsules, Ophthalmic, Oral liquid, Eye drops and Nasal spray.)
  Period: From 29th August 2016 to 4th October 2017.
- Worked as **QC Officer- in Sun Pharmaceutical industries Ltd Ahmednagar** which one is manufacturer of API's. **(As an Analyst)**
  Period: From 11th July 2013 to 23rd August 2016
- Worked as **Trainee Officer-**in QC department, Brassica Pharma Pvt Ltd at Boisar, Mumbai which one is manufacturer of solid dosage form, (Tablets and Capsules). Period: From 3rd July 2012 to 3rd July 2013.

JOB PROFILE:

- Handling QMS i.e. Event, Incidents, OOS, OOT, Change control, Deviation in FG and Stability section. Implementation of CAPA against root cause.
- Leading Stability Section, Planning and execution of stability samples analysis and release within timeline. Verifying stability chamber related activities, QMS related activities i.e., Incidences, OOS, OOT, Change control and Deviations regarding stability section.
- Overall responsible for day-to-day activities in QC lab, managing Stability releases and maintaining GLP and GDP in laboratory.
- Leading Finish Exhibit section, planning and execution of Finish products analysis and release.
- Investigation and report Preparation/Review of laboratory incidents, OOS and OOT's, online review and take necessary actions for daily audit trail/ Chromeleone message center related concerns.
- Process validation and cleaning validation of new products.
- Pharmacopeia gap assessments and implementation activities related to revised Pharmacopeia monographs and general chapters.
- Analytical method transfers on new projects.
- Analytical method verifications of revised Pharmacopeia monographs.
- SOP's gap identification and accordingly updating SOPs of laboratory using Ensure software.
- Initiation of Change controls in Trackwise software.
- Actively involvement in departmental safety activities like, leading safety training, awareness about use of lab. PPE's, HIRA preparations and review of laboratory instruments.

HANDS ON EXPIRIENCE:

- Detailed investigation by six sigma for OOS and OOT, Implementation of CAPA.
- Master document preparation like Analytical Method Verification protocol, Analytical Cleaning Method Validation protocol, Analytical Method transfer protocol, SOP, STP, Specification preparations and review.
- Handling and Coordination of SAP, implementation of TDS in SAP.
- Handling of Calber LIMS (3.2 and 3.4), Thermoscientific LIMS (12.1).

INSTRUMENTS HANDLED:

- Handled and very well familiar with Hardware of HPLC systems (Makes: Waters alliance, Agilent, Shimadzu).
- Handled and very well familiar with Software of HPLC systems (**Chromeleone 6.8 and 7.2**).

- Handled and very well familiar with G.C. instruments with headspace (Make: Perkin Elmer and Shimadzu).
- Handled and very well familiar with UV Spectrophotometer, F.T.I.R. (Makes: Shimadzu, Perkin Elmer).
- Handled and very well familiar with Dissolution Apparatus (Make: Electrolab).

EDUCATIONAL PROFILE:

- Bachelor of Science : From Pune university with secured Distinction.
- Master of Science (Organic Chemistry) : From Pune University, Ahilyanagar Secured Higher Second Class.

AUDITS FACED:

- Faced GMP audits from USFDA, MHRA, ANVISA -Brazil, MCC – South Africa,
- Faced USFDA audit at Piramal pharma, Pithampur Indore Madya Pradesh.

COMPUTER SKILLS:

- Certificate Course in Computer Concepts (MSCIT), Work with MS Office, Open office and Internet applications.

PERSONAL DETAILS:

Name	**:**	Ram R. Wandhekar
Date of Birth	**:**	20th Dec 1988
Permanent Address	**:**	At-Gunjale Post-Katrad, Ta- Rahuri Dist- Ahmednagar, Maharashtra (India) Pin:413705
Gender	**:**	Male
Marital Status	**:**	Married
Languages Known	**:**	English, Hindi and Marathi.

DECLARATION:

I hereby declare that the above written particulars are true to the best of my knowledge and belief.

Date :_____
Place :_____