CURRICULUM VITTAE

NAME:

AKSHAY SHIVNATH PAWAR.

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+918600243315,
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FATHERS NAME:

SHIVNATH AMBADAS PAWAR.

DATE OF BIRTH: ***29/10/1990.***

SEX: ***MALE.***

MARITAL STATUS: ***MARRIED.***

LANGUAGES KNOWN:

ENGLISH, HINDI, MARATHI.

NATIONALITY: ***INDIAN.***

PERMANENT ADDRESS:

AT.POST: DEOGAON (BHAVANINAGAR), TAL: NIPHAD, DIST: NASHIK, STATE: MAHARASHTRA, (INDIA), PIN CODE: 422305.

APPLYING FOR:

Suitable position in your esteemed organization.

CAREER OBJECTIVE:

Looking forward to associating myself with an organization where there is an opportunity to share, contribute and upgrade my knowledge for the development of self and organization served.

EXPERIANCE DETAILS:

Overall Experience: 9 years, 2 months in Pharmaceutical Industry – Oral Solid Dosage Formulation

Department - Quality Assurance & Quality Control

Current and Previous Employers:

Mylan Laboratories Limited, A Viatris Company, Nashik Maharashtra Designation: Assistant Manager Duration: From 14th of Aug. 2016 to Till Date
Alembic Pharmaceuticals Ltd. Baroda, Gujarat. Designation: Executive Duration: From 10th Of Nov. 2014 To 08th of Aug. 2016
Cipla Ltd. Verna, Goa. DESIGNATION: QA Officer: Under Management Staff From 11th of February 2013 to 22nd of October 2014.

JOB PROFILE:

• Customer Complaints Responsibilities –

- Preparation of Market complaints investigation reports (Quality Complaints & Drug Efficacy Complaints) for all Markets,
- Preparation of Product Quality Defect Trend Assessments,
- To perform Root Cause Analysis, Risk assessments,
- To monitor planned /unplanned deviation and initiate & monitor Corrective and Preventive Action (CAPA),
- To monitor Change Control activities
- Preparation of monthly, quarterly, and annual reports of market complaints.
- Participate in various project meetings internally or externally (vendor & customer) and represent senior management.

AUDIT EXPOSED:

W.H.O.

USFDA

ANVISA

MHRA

TGA

AND OTHER CUSTOMER AUDITS.

PERSONAL TRAITS:

ABILITY TO ADAPT WITH DIFFERENT CULTURAL AND WORKING ENVIRONMENT.

POSITIVE APPROACHES TOWARDS ALL SITUATIONS

QUALITY LEADERSHIP, ENERGETIC, CONFIDENCE.

GOOD COMMUNICATION SKILL.

- Performing any other important tasks that management may assign from time to time. Work closely with Head of Quality, Manufacturing and Regulatory to ensure the compliances.

• Databases Handled –

- **TrackWise** - for Market Complaints / Other investigations
- **Tableau** – for normalized complaint rate
- **LIMS** - for Analytical data
- **SAP** - for Manufacturing, Packaging data
- **CARA** - for documentation

• Computer System Validation –

- CSV Preparation, Execution & Review of documents like URS, FRS, SDS, Impact & Risk Assessment, IQ, OQ, PQ as per 21 CFR Part 11, GAMP5, EU Annex 11, etc. and Management, Co-ordination, and support for CSV / Equipment Qualification activities.

- **Also performed the tasks as:**

- To monitor quality systems and procedures are followed in accordance with cGMP.
- Conduct Unit Operation.
- APQR entry and APQR review.
- To review process performance qualification protocols and reports
- Responsible for carrying out validation activities according to the validation protocol, reporting for the validation activities and review of the same in manufacturing as well at packaging process.
- Initiation in modification of sampling activities and tracking for the same to avoid mishandling and errors.
- To carry out the Hold time activities for intermediate and bulk products.
- Review of master BMR, BPR and checking of draft documents.
- Performed online process checks in tablet, and capsule sections like appearance, DT, Friability, thickness, hardness, weight variation, etc.,
- To perform AQL inspection and monitor manufacturing processes.
- Responsible for packing line proof checking.
- To ensure cleanliness of areas.
- Monitoring machinery logbooks and temperature, humidity charts.
- Responsible for giving line clearance, Bulk checking, Log review, Daily rounds.
- To ensure online documentation and timely entries of all operations and activities.

ADDITIONAL EDUCATIONAL QUALIFICATION:

MAHARASHTRA STATE CERTIFICATE COURSE IN INFORMATION TECHNOLOGY. (MSCIT)

INDUSTRIAL TRAINING PERFORMED DURING GRADUATION:

AT "REVE PHARMA"

WORKS: PLOT NO.78, STICE, MUSALGAON, TAL: SINNAR, NASHIK 103. TELEFAX: 02551 240127.

TRAINING COMPLETED IN:

PRODUCTION DEPARTMENT ABOUT; STANDARD OPERATING PROCEDURES TO BE FOLLOWED, GMP, cGMP, TABLET COMPRESSION MACHINE, FBD, MASS MIXER, RMG, BLENDER MACHINE, BLISTER MACHINE, OINTMENT FILLING MACHINE, AND RESPECTIVE PACKAGINGS.

- Responsible for handling of data logger activities.
- Responsible for handling of packing material rejection.

- **Also functioned as Laboratory Quality Assurance Auditor/ warehouse auditor**

- To monitor standards as per cGLP.
- To carry out inspections in quality control laboratory.
- To investigate OOS/OOT and Analytical incidence.
- To ensure as the safety being used.
- To ensure online reporting and to check the calibration reports.
- To check audit trails of instruments like UV, IR, KF, FTIR, POLARIMETER and other softwares being used for printouts.
- To ensure location charts, stock, and proper storage of materials according to their storage conditions in warehouse area.

ACADEMIC CREDIANTIALS:

University/Board: Pune (Maharashtra).

Graduation Qualification: Bachelor of Pharmacy

QUALIFICATION	COLLEGE/ SCHOOL	YEAR OF PASSING	%MARKS
B. Pharm	S.N.D. College of Pharmacy, Yeola.	2011-12	63.83 % **First class**
H.S.C.	Karmveer BhauraoPatil Junior College Of Science, Vinchur.	2007-08	50.00 %
S.S.C.	Gautam Public School, Gautamnagar, Kolpewadi. (English Medium)	2005-06	52.00 %

DECLARATION:

I hereby declare that the information provided above is true to the best of my knowledge. I assure you, if i get a chance, i will execute my work to the fullest satisfaction of my superiors.

PLACE: Nashik, Maharashtra **YOURS TRULY,**

DATE: **AKSHAY S. PAWAR.**