CURRICULUM VITAE

SATYAJEET M. BHAGWAT
Assistant Manager – QA
208, Crown Residency,
303 Silicon City, Rau,
Indore. 452012
Mob. No: 9028241863
satyajeetbhagwat@gmail.com

PROFESSIONAL OBJECTIVE:

To join a firm & team, have good vision & best policy. I am willing to perform best in favor of growth of organization by optimizing cost, increasing productivity with full energetic talent & achieve the quality standards & goals that appreciate team efforts.

SUMMARY:

A competent professional with **11+** years of experience in Pharmaceutical Quality Assurance in Oral Solid Dosage formulation which covers the ***Manufacturing, Packaging and Warehouse of Pharmaceuticals***. Presently employed as Assistant Manager – Quality Assurance with Mylan (Viatris) Pharmaceutical ltd., Pithampur (Indore).

AREA OF INTREST:

Corporate Quality Assurance (CQA), Quality Assurance, Regulatory Affairs (RA), Research and Development

ACADEMIC QUALIFICATION:

Degree/ Certificate	University/Board	Year of Passing	Marks obtained (%)
Diploma in Pharmacy (D. Pharm.)	MSBTE	2007	66.00 %
Bachelor of Pharmacy (B. Pharm.)	University of Pune	2010	55.33 %
Master of Pharmacy (M.Pharm.)	North Maharashtra University, Jalgaon (NMU)	2012	64.76 %

AUDIT FACED:

Exposure and faced the National & International Audits like USFDA, EU, MHRA (UK), MCC, ANVISA, WHO(Geneva), Customer Audits and response to the observation made by auditors.

PROFESSIONAL EXPERIENCE:

Firm/ Company	Period		Designation
	From	To	
Mylan Laboratories Limited, (Viatris) Pithampur, Indore	July - 2021	Till date	Assistant Manager-QA
Mylan Laboratories Limited, (Viatris) Waluj, Aurangabad	August - 2015	July - 2021	Executive - QA
Cipla Ltd. Verna, Goa	February -2014	August - 2015	Officer - QA
Marksans Pharma, Verna, Goa	January - 2013	January-2014	Management Trainee - QA

EXPERIENCE/WORK & JOB ROLE PERFORMED:

Sr.No.	Firm/ Company	Job role performed
1	Mylan	▪ Responsible for Batch production and control record (BPCR) review. (Documentation In charge) ▪ Maintain all Time Audit Readiness & Compliance. ▪ Preparation & Review of SOP & its Related Documents. ▪ Self-Inspection/Internal Audit of cross-functional departments. ▪ Active participation in Audit, Response and Compliance of observations from Regulatory and customer audits at site. ▪ To complete Assign Trainings, Impart Trainings, and associated training related tasks. ▪ Ensure the implementation of current Good manufacturing practices at all levels of functioning in all plant activities. ▪ Compilation and Review of APQR. ▪ To perform Effectiveness check and task completion assigned Intrackwise for change control in timely manner. ▪ Responsible for investigations and CAPA implementation of various QMS like Deviations, Market complaints, Out of specification, non-conformances in Quality Audits. ▪ Perform GAP analysis of various activities against regulatory guidance through Internal Audits & recommends corrective actions. ▪ Handling of Software - SAP, Trackwise, MA-Management-My Portfolio,Share Point, D2-Documentum, etc. ▪ To impart training to subordinates and new joiners.

Sr.No.	Firm/ Company	Job role performed
2	Cipla	▪ Handling in-Process checks & issuing of line clearance of various dosage forms like tablet, capsule at manufacturing & Packing. ▪ Handling of control sample along with review of documents. ▪ Responsible for implementation & development of Quality system on shop floor. ▪ To perform sampling of in process, finished bulk, Hold time and finished product as per the product BPCR or approved protocol. ▪ Handling of sampling activities for process validation, swab / rinse sampling, BUA Sampling (Blend Uniformity Analysis Sampling) in-process sampling & hold time study at manufacturing and packaging stage. ▪ BMR, BPR Review & Compliance. ▪ AQL Inspection at manufacturing Stages. ▪ Calibration of IPQA laboratory Instruments as Analytical balance, DT, Friability, Hardness Tester etc.
3	Marksans Pharma Ltd.	▪ To certify line clearance. ▪ To perform in process testing. ▪ Environmental monitoring of ware house and process area. ▪ To perform swab sampling. ▪ To review and completion of BPCR. ▪ To perform IPQA activities in various departments, in order to ensure cGMP compliance. ▪ Preparation, and closing of NCR (Non Conformance Report) if any non-conformance found in tablet manufacturing and Packing.

EXTRA CURRICULAR ACTIVITIES:

- Playing Cricket.
- Cooking Food.
- Singing

STRENGTH:

- Very flexible and adaptable to change.
- Responsible, operate with integrity and team work.
- Learning attitude and logical approach.

PERSONAL DETAILS:

- Date of Birth : 4th Jan 1988
- Gender : Male
- Nationality : Indian
- Religion : Hindu
- Marital status : Married
- Language known : English, Hindi, Marathi.

NOTICE PERIOD:

- 3 Months

DECLARATION:

I hereby to declare that the details furnished above are true to the best of my knowledge.

Satyajeet M. Bhagwat
(Sign/Date)