

Email ID: mukeshr5184@gmail.com

Mukesh Chaudhari

C-108, Shivsagar Tenements, Nr. Saint Marry School Naroda, Ahmedabad-382330 Gujarat, India.

Career Objective:

Intend to build a career in leading pharmaceutical company with committed & dedicated people, which will help me to explore myself fully and realize my potential. Willing to work as a key player in Challenging & Creative Environment.

Professional Experience: 14 years in IPQA, Validation, QMS, Qualification in Parenteral & OSD

Sr. No.	Company	Designation	Duration	Dosage Forms
01	Intas Pharmaceuticals Ltd. Matoda, Ahmedabad.	Senior Executive-Q.A.	March-2016 & Onwards	Solid Dosage form
02	Mylan Laboratories Ltd. Pharmasez, Ahmedabad.	Executive-Q.A.	May-2014 to March-2016	Parenteral & Solid dosage form
03	Torrent Pharmaceuticals Ltd. Indrad, Mehsana	S.T.AQ.A.	Nov-2010 to May-2014	Solid dosage form
04	Lincoln Pharmaceuticals Ltd. Khatraj, Gandhinagar.	Officer-Q.A.	Aug-2009 to Oct-2010	Parenteral dosage form

Current Employer and Job Profile:

1. Intas Pharmaceuticals Limited, Matoda Ahmedabad

(Oral Solid Dosage)

- Monitoring of shop floor activities like line clearance, IPQA, sampling.
- Review of executed manufacturing & packing BPCR, release of batches.
- > Process validation activities like sampling during process validation, monitoring of validation batches.
- Preparation, execution and review of process validation protocol and reports and its compilation.
- Preparation and review of APQR (Annual Product quality review).
- ➤ To review of executed electronic mfg. & packing BPCR of oncology solid and Parenteral.
- Review and submit documents to Regulatory Affairs department like batch records, process validation protocol and reports, stability studies reports, specification and MOA, AMV protocols and reports etc.
- > Provide batch release & other related documents to client prior to batch release.
- ➤ Handling of query received from client, regulatory department and compliance them.

Previous Employer and Job Profile:

2. Mylan laboratory Limited.

(Parenteral and OSD)

- ➤ Handling, review & evaluation of change control related to documents, process, product, equipment etc.
- ➤ Handling and investigating Deviations, OOS, Incidents & Market Complaints.
- ➤ Handling & review of CAPA i.e. Implementation / Effectiveness review / Closure.
- Conducting Risk assessment and Impact evaluation for QMS activities and events.
- > Participated in self inspection & observation compliance activity.

3. Torrent Pharmaceuticals Limited.

(Oral Solid Dosage)

- > Dispensing activity, Line clearance activities in manufacturing and packing department.
- > IPQA and sampling activity in manufacturing, compression and packing department.
- ➤ Review of GMP documents like bmr-bpr, sops, area log books, balance calibration records, environmental monitoring records.
- Process validation activity such as monitoring of process validation and exhibit batches, sampling at various stage i.e. blend uniformity sampling, stratified sampling.
- Cleaning validation sample collection, hold time study samples, finished, retain & stability samples.
- > Preparation & review of process validation, protocols and reports.
- Execution of CSV validation activity as per VMP, GAMP5 and 21 CFR part 11 compliance.

4. Lincoln Pharmaceuticals Limited.

(Parenteral)

- Monitor and ensure that all activities involved in Manufacturing, filling, visual inspection, labelling and packing functions are in compliance with GMP and Standard operating procedures
- Line clearance & IPQA activity in Dispensing, Vial washing and depyrogenation, manufacturing, filtration, filling, Lypholization, sealing, visual inspection, labelling and packing operations.
- Sampling activity for In-process, Finished Product, Retained/control and Stability sample for analysis and other samples as per protocols (Process Validation, Cleaning Validation etc.).
- > Participated in Media fill activity, Area qualification, AHU Qualification, Water system etc.
- > Review of executed batch documents of manufacturing and packing.

Audits exposure:

➤ I have successfully faced USFDA-US, MHRA-UK, TGA-Australia, ANVISA-Brazil, WHO, Ukraine, Tanzania, Uganda, Ghana & various party audits during my working tenure.

Computer Proficiency:

- Microsoft word/ Excel
- > SAP
- Qedge Software (QMS)
- Process Xe (Electronic- BMR)
- > LIMS
- Minitab
- > FTS
- > TMS

Educational Qualification:

Qualification	Institute	University	Year	Percentage
B.Pharma	Bapuji Pharmacy College, Davangere	R.G.U.H.S., Bangalore. Karnataka 2007 70.12	2008	70.12%
D. Pharma.	Smt. R.D.Gardi Diploma Pharmacy College, Rajkot	Saurashtra University, Rajkot	2004	58.11%
H.S.C.	Raghunath School Ahmadabad	G.H.S.E.B., Gandhinagar	2002	66.67%
S.S.C.	Raghunath School Ahmadabad	G.S.E.B., Gandhinagar	2000	81.29%

Personal Details:

Father's name : Late Shree Rajaram P. Chaudhari

➤ Date of Birth : 5th January, 1984

Marital statusMarried

Language Proficiency: English, Hindi, Gujarati.

Expected Salary : Negotiable

➤ Hobbies : Watching movies, listening music, Playing snooker.

Strength : Positive attitude, hardworking, quick learner, leadership abilities, effective

communication skills, analytical skills, readiness to explore & learn.

Reference:

Available on demands.

Declaration:

I hereby declare that, the above mentioned details furnished by me are true & correct to the best of my knowledge & belief.

Place: Ahmedabad

Date:

(Mukesh Chaudhari)