Samir N. Chaudhary

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OBJECTIVE:

To excel myself in Pharmaceutical activities to work and acquire a post in an organization where good interpersonal skills are required.

STRENGTHS:

- Able to work in flexible schedule.
- Zeal to keep learning.
- Ability to work as a part of team.

AWARDING BODY	TITLE OF EXAMINATION	PERCENTAGE	INSTITUTION	AREA OF SPECIALIZATION	YEAR OF PASSING
R.U.H.S	**Master of Pharmacy**	**70.25%**	Bhupal Nobles' college of Pharmacy, Udaipur.	Industrial Pharmacy	2013
Pune University	**Bachelor of Pharmacy**	**61.45%**	Modern College of Pharmacy, Nigdi, Pune.	Pharmaceutical Sciences	2010
G.H.S.E.B Gandhinagar	**HSC**	**63.65%**	Shree Gattu Vidhyalaya, Ankleshwar.	Physics, Chemistry, Biology	2006
G.H.S.E.B Gandhinagar	**SSC**	**74.54%**	Shree Gattu Vidhyalaya, Ankleshwar.	Science, Mathematics	2004

Project during M.Pharm:

Controlled Release In situ forming Ciprofloxacin Hydrochloride Hydrogel for Ophthalmic Drug Delivery.

Study performed at **"Haffkine Bio-Pharmaceuticals Ltd, Govt of Maharashtra, Acharya Dhonde Marg, Parel, Mumbai Maharashtra.**

Computer Skill:

- Proficient in Microsoft word, Excel, Power Point, Internet Surfing operations.

- E-BRM, E-QAMS, SAP.

WORK EXPERIENCE:

1. Glenmark Pharmaceuticals, Goa.		
Plant approval	:	**MHRA, TGA, USFDA, ENVISA, MCC.**
Designation	:	**Quality Assurance Sr. Officer**
Duration	:	**From Nov 2013 to March 2016.**
Role and responsibility	:	➢ Routine Manufacturing compliances, in-process control for solid dosage facility. ➢ Communication to Superiors related to the Manufacturing Assurance Activities on Daily basis. ➢ Review and compliance of online BPR's, Manufacturing Process and SOP's. ➢ Implementation of GMP, Quality Systems and procedures during the Manufacturing Process. ➢ Involve in Cleaning Validation like swab sampling and its Documentation. ➢ Line Clearance of On-going Products for Dispensing, Manufacturing and Packing Operations. ➢ To Check and perform SAP related Activities and Compliances. ➢ Involve for Process Validation Activities and Data Compilation. ➢ Finished Product, Stability and Hold Time Study Sample Collection and submit to respective Department for Analysis.
2. Unique Pharmaceuticals Ltd. Panoli.		
Plant approval	:	**TGA, USFDA**
Designation	:	**Quality Assurance Executive**
Duration	:	**From March 2016 to November 2016.**
Role and responsibility		➢ Routine Manufacturing compliances & in-process activities. ➢ Communication to Superiors related to the Quality Assurance Activities on Daily basis. ➢ Review and compliance of online e BRM and SOP's. ➢ Implementation of GMP, Quality Systems and procedures during the Manufacturing Process. ➢ Involved in QMS Activities like change control, deviation & CAPA. ➢ Involved in Process Validation Activities and Data Compilation.

3. Intas Pharmaceuticals Ltd. Matoda.	
Plant approval	**USFDA, MHRA, TGA**
Designation	**Quality Assurance Executive**
Duration	**From November 2016 to till date.**
Role and responsibility	➢ Routine Manufacturing compliances & in-process activities. ➢ Communication to Superiors related to the Quality Assurance Activities on Daily basis. ➢ Review and compliance of online e BRM and SOP's. ➢ Implementation of GMP, Quality Systems and procedures during the Manufacturing Process. ➢ Involved in QMS Activities like change control, deviation & CAPA.

Audits faced:	US Food & Drug Administration (USFDA) ISO 14001 MHRA, TGA

Personal information:		
Mother's Name	:	ALKA NARENDRA CHAUDHARY
Father's Name	:	NARENDRA RAMDAS CHAUDHARY
Permanent address	:	C/4 Yagnavalkya soc. Plot no. 805, G.I.D.C., Ankleshwar - 393002, Gujarat.
Marital status	:	Married
Sex	:	Male
Linguistics	:	English, Hindi, Marathi, Gujarati

Declaration: The above details furnished are true & correct to the best of my Knowledge.

(Samir Chaudhary)