



# CURRICULUM VITAE

**SHYAM KUMAR CHOUDHARY**

**Approved Manufacturing Chemist**  
Approved in Tablet, Capsule & Dry Syrup from  
**Food & Drugs Administration Haryana**

## OBJECTIVE:-

My objective is to embark upon a carrier, which satisfies my creativity and is associated with a company where my growth is boosted along with the company's growth.

## CONTACT INFORMATION:-

SHYAM KUMAR CHOUDHARY  
Address:- F426/23(old) H no.10029 Gali  
no.317 Sanjay Colony, Sec-23, Faridabad  
Haryana 121005

E Mail [sh44shyam@gmail.com](mailto:sh44shyam@gmail.com)  
Mobile No. 9560084937

## PERSONAL INFORMATION:-

Father's Name  
Date of Birth  
Sex  
Marital Status  
Nationality  
Languages known

Maha Kant Choudhary  
21-Nov-1976  
Male  
Married  
Indian  
English and Hindi

## HOBBIES:-

Reading Books, Amiable and discussion with technical matters and reading newspapers.

## STRENGTH:-

Punctual, Self motivation, Hard working and Positive attitude, Intriguing personality.

## WEAKNESS:-

I Can't compromise with quality of work assigned to me.

PROFESSIONAL DETAILS:-	
Work Experience  Skills Industry Type Role	More than 13 years experience in Pharmaceutical company Formulation Manufacturing Department. Pharmaceuticals (Production) Tablet, Capsule, Dry Syrup and oral liquid.
ACADEMIC QUALIFICATIONS:-	
B.sc Passed From L.N.M.U Darbhanga Bihar in 1998. Intermediate From Bihar Intermediate Council Patna in the year 1993. High School From Bihar Board Patna in the year 1991.	
EXPERIENCE SUMMARY:-	
<p>I am an experienced, trained, and qualified Manufacturing personal possesses good communication, Interpersonal &amp; manufacturing skills and fulfill with the spirit of teamwork.</p> <p>I am also having hands-on experience in developing, reviewing, and auditing SOPs, specifications, validation protocols, reports, batch documents, and document control in the light of a great understanding of the latest trends of developed regulatory market requirements in the present scenario.</p>	
Successfully faced various Customers audit and Third-party audits such as Vietnam, Ethiopia, Philippines, Myanmar, Ukraine, Kenya, Iraq, Cambodia, Uzbekistan, Sri Lanka, Afghanistan, Nigeria, etc.	
CURRENT ASSIGNMENT AND JOB RESPONSIBILITIES:-	
Presently working as <b>Assistant Manager Production</b> in <b>M/s Brawn Laboratories Ltd.</b> situated at Plot No. 13, N.I.T Faridabad Haryana 121001 dated from <b>09.10.2015 to till date</b> . Looking after Projects, Manufacturing Process, Production Planning, and Inventory Control, Administration for their Tablets, Capsule, Dry Syrup, Liquid orals product, etc.	
CAREER AFFILIATION:-	
I have worked as a <b>Sr. Production chemist</b> in <b>M/s ZEE Laboratories Ltd.</b> situated at Paonta Sahib in Himachal Pradesh dated from <b>01.06.2013 to 30.09.2015</b> looked after the manufacturing process, production planning, inventory control, administration for their Tablets, Capsule, oral liquids and dry syrups products.	
I have worked as <b>Asst. Production chemist</b> with the Companies <b>M/s Arbro Pharmaceutical Ltd, M/s Anrose Pharma, M/s Oyster Pharmaceutical Ltd.</b> During period <b>03.06.2006 to 26.04.2013</b> . which produced many formulations.	

<b>RESPONSIBILITIES:-</b>
To Co-Ordinate with QA/QC department in controlling their process and product at every stage of manufacturing to meet the established.
Maintenance of Manufacturing records and BPRS for each batch manufactured.
Routine cGMP auditing at manufacturing and related area.
Development and implementation at the cGMP system to ensure international standard requirements.
To suggest and organize a training program for the development of technical and administrative skills of all the employees to meet with cGMP. Regulations on a continual basis which shall be done by Co-coordinating with corporate quality division and GMP Committee.
Development and implementation of the cGMP system to ensure international regulatory requirements.
<b>JOB PROFILE:-</b>
Overseeing the production process, drawing up a production schedule.
Ensuring that the production is cost-effective.
Making sure that products are produced on time and are of good quality.
Working out the human and material resources needed.
Estimating costs and setting the quality standards.
Monitoring the production process and adjusting schedules as needs.
Being responsible for the selection and maintenance of equipment.
Monitoring products standards and implementing quality control programmes.
Liaising among different departments, e.g. Suppliers, Managers.
Working with managers to implement the company's policies and goals.
Supervising and motivating a team of workers.
Reviewing the performance of subordinates.
Identifying training needs.
Ensuring that health and safety guidelines are followed.
<b>DECLARATION:-</b>
<p>I hereby declare that the information given about are true to the best of my knowledge and belief.</p> <p style="text-align: center;"><b>SHYAM KUMAR CHOUDHARY</b></p> <p style="text-align: center;">(Since the document is prepared electronically. Hence no signature is required)</p>