# **Shaik Azmath**

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### **OBJECTIVE**

Aspiring professional passionate about Organic Chemistry, seeking opportunities in Regulatory Affairs, R&D and QA. Proficient in synthesis, sustainability, and regulatory compliance. Eager to contribute expertise to drive innovation and ensure quality standards in the pharmaceutical industry.

### **EDUCATION**

A.V. College of arts, science & commerce (OU)

M.Sc. (Organic Chemistry) |

Gauthami Degree College (TU)

B.Sc. MPC (Maths, Physics, Chemistry) | GPA: 9.1/10.0

Intermediate MPC (Maths, Physics, Chemistry)

Board of intermediate Education, TS | GPA: 8.0/10.0

**Secondary School Certificate** 

Board of Secondary Education, TS | GPA: 8.8/10.0

Hyderabad, TS

Graduation Date: Oct2023

Nizamabad, TS

Graduation Date: Aug2021

Nizamabad, TS

Graduation Date: Apr2018

Nizamabad, TS

Graduation Date: Mar2016

#### **SKILLS**

ChemDraw, ChemOffice

Microsoft Office (Excel, Word & PowerPoint)

Soft skills: Communication | Time management | Collaboration | Adaptability | Problem solving

### **PROJECTS**

# **Dummy DMF (Drug Master File)** | S.G Pharma | <u>Link</u>

Nov2023

- Crafted a comprehensive dummy Drug Master File (DMF) as a pivotal assignment during coursework, showcasing adeptness in understanding DMF formatting, structure, and compliance with regulations without disclosing confidential information.
- Developed a simulated DMF project, emphasizing adherence to formatting **guidelines** and industry standards, demonstrating practical application of theoretical knowledge acquired during the Regulatory Affairs course.
- Created a dummy DMF project, focusing on meticulous **structuring and adherence to regulatory guidelines**, highlighting a proactive approach to gaining hands-on experience in the realm of regulatory affairs within a controlled educational environment.

# **Dummy Technical Package** | S.G Pharma | <u>Link</u>

Oct2023

- Constructed a comprehensive dummy technical package as part of practical training, showcasing an understanding
  of technical documentation requirements without disclosing proprietary information.
- Engaged in the creation of a simulated technical package, focusing on the structure, organization, and key elements necessary for compliant documentation, demonstrating adherence to **industry guidelines**.
- Developed a simulated technical package, demonstrating meticulous attention to detail and knowledge of essential components, contributing to a foundational understanding of technical documentation principles within the industry

# **CERTIFICATION**

# **Drug Regulatory Affairs** | S.G Pharma | Link

Oct2023-Nov2023

- Successfully completed a drug regulatory affairs certification program, expanding my knowledge of pharmaceutical regulations and approval procedures.
- Proficiency in preparing regulatory documents and ensuring compliance was honed during the certification.
- Gained a heightened awareness of the importance of maintaining product quality and safety in line with industry standards.

# **LANGUAGES**

English, Hindi, Telugu, Urdu, Arabic (Basic)

### **DECLARATION**

I hereby that the above-mentioned information is correct up to my knowledge and I bear the responsibility for the correctness of the above-mentioned particulars.