

## **-: CURRICULUM VITAE: -**

**NAME: Mr. MRUGESHKUMAR.H. PANDYA.**

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### **CAREER OBJECTIVE: -**

- A pharmaceutical professional of experience having Intend to build a career with leading corporate in Hi-tech environment with committed and dedicated peoples. And to do challenging work in pharmaceutical field.
- To prove myself best for the organization and for creativity teamwork and leadership.

### **ACADEMIC QUALIFICATIONS.**

- |  |              |
|--|--------------|
| ➤ Bachelor of Science (chemistry): Gujarat University, India | October-2017 |
| ➤ Higher Secondary Certificate (Science stream): H.S.E.      | March-2013   |
| ➤ Secondary School Certificate (General): S.S.C.             | March-2011   |

### **PROFESSIONAL EXPERIENCE DETAILS:**

- ❖ **Current Working:** Presently Working in **TORRENT PHARMACEUTICAL LTD. Chhatral - Ahmedabad (Gujarat)** since February-2024 to till date as a Technical Assistant in QA Department.
- ❖ **Previous Organization:** Working in **Tatva Chintan pharma chem Ltd. Dahej – Bharuch (Gujarat)** since September -2022 to February-2024 as a Jr. Officer in QA Department.
- ❖ **Previous Organization:** Working in **Shiva Pharmachem Limited. Dahej – Bharuch (Gujarat)** February-2021 to September-2022 as an **Assistant -Quality Assurance** in QA Department.

### **JOB PROFILE & RESPONSIBILITIES:**

#### **Responsibility:**

- ❖ To review Batch Manufacturing Records for Batch release and online BMR.
- ❖ Preparation & review of Annual product Quality Review Report.
- ❖ Preparation of validation Protocols & monitoring process validation activities.
- ❖ Review of Master BMR & ECR.
- ❖ Handling of change control activity.
- ❖ Preparation and review of Standard Operating Procedure.
- ❖ Perform the sampling of Finish product/intermediate and storage of reserve sample
- ❖ Review of Qualification and Validation documents.
- ❖ Handling of SAP.
- ❖ Review, Tracking & retrieval of BMR, SOPs, Formats, logbook, & Test data sheets.

## **-. CURRICULUM VITAE: -**

- ❖ Complete training on relevant SOPs and develop an understanding of the activities to be conducted prior to undertaking any task.
- ❖ To conduct Q.A. rounds as per established schedule and carry out follow up audit to ensure the compliance of finding.
- ❖ To do online operation checking and ensure online entries in Batch Manufacturing Records log books & related documents as per Standard Operating Procedure (SOPs).

### **KEY ROLE: -**

- Review of Validation and Qualifications documents.
- Review of Master BMR and ECR
- Batch release activity through SAP (Intermediate).

### **ACHIEVEMENTS / EXTRA-CURRICULAR ACTIVITIES:**

- **Suggestions** related to System Improvement
- **Software** handle like SAP, SCADA, C-DAS, DMS, SABA, QMS.

### **AREA OF INTEREST: -**

- QMS & Validation related documents.
- Documentation & IPQA Work.

### **STRENGTH: -**

- Always interested in challenging work. & try to solve problem with win case.
- Positive attitude,
- Self-motivated,
- Excellent grasping power.

### **PERSONAL PROFILE: -**

➤ Name	:	Pandya Mrugeshkumar.
➤ Father Name	:	Pandya Harishbhai.
➤ Gender	:	Male
➤ Marital Status	:	Married
➤ Date of Birth	:	15/12/1995
➤ Nationality	:	Indian
➤ Language Known	:	English, Hindi, Gujarati

### **DECLARATION:**

“I hereby declare that all the details mentioned above are in accordance with the truth and fact as per my knowledge and I hold the responsibility for the correctness of the above-mentioned particulars.”

**Place:**

**Date:**

**Mr. Mrugeshkumar.H. Pandya**