SAIDUL ISLAM

BACHELOR OF PHARMACY

From-KANPUR INSTITUTE OF TECHNOLOGY AND PHARMACY

SPECIALISATION:-
Executive QUALITY ASSURANCE

ADDRESS:

Permanent Address:-
S/O- LT Abdulla Seikh,
Vill- Hazipur, PO- Begunia ,
PS-Mayureswar, Birbhum,
731245,(West Bengal),Indian.

Mobile: +91-8170038090

Correspondence Address:-

Vill- Hazipur, PO- Begunia , PS-Mayureswar, Birbhum,
731245,(WB),Indian.

Mobile: **+91 8170038090**

E-MAIL:-
Saidulislamahm90@gmail.com

PERSONALPROFILE:-
Father's Name: LT Abdulla Seikh
Date of Birth: 22 Nov, 1995
Sex: Male
Nationality: Indian
Language Known: - Bangla, Hindi, English

HOBBIS:-
- Travelling.
- Playing Pool.
- Internet Surfing.
- Work in a team Efficiently.
- Disciplined.
- Punctual.

OBJECTIVE:-

"To obtain a creative and challenging opportunity where I can exercise my pharmaceuticals and interpersonal skills which will enable me with an ability to learn new things and explore future career prospects."

EXPERIENCE:-

- 25 July 2018 to 30 Augest 2019 experience in the field of Quality Assurance Officer(IPQA) in **Unicure India limited (**Raipur, Bhagwanpur**)** UTTRAKHAND. Worked in Tablet Section(Granulation,Compression,Coating) Capsule, Dispensing,Primary and Secondary Packing.
- 1 Sep. 2019 to 28 Feb. 2024 experience in the field of Executive Quality Assurance(IPQA) in **SURAKSHA PHARMA Pvt. Ltd.** (Karoundi,Roorkee) UTTRAKHAND. Worked in Tablet Section (Granulation,Compression,Coating), Capsule, Primary and Secondary Packing.
- 1 Mar. 2024 to Till date experience in the field of Executive Quality Assurance (IPQA) in **DenMark PHARMACEUTICALS Pvt. Ltd**. (Karoundi, Roorkee) UTTRAKHAND. Worked in Tablet Section (Granulation , Compression and Coating).

SUCCESFUL AUDIT FACED:-

- Faced WHO-GMP audit.
- Bihar, Kerala Government.
- NHP Canada Government.

WORK RESPOSIBILITY:-

- Line clearance on shop floor.

- Monitoring of Environment conditions as per the requirement. To check and ensure the proper use of input materials.
- To check and ensure the proper segregation of the materials as per the status. To give the Line clearance for batch to batch as well as product-to-product as per The requirement.
- To carry out the In-process checking stages of manufacturing and packaging operations.
- To review the batch record up to the relevant stage and ensuring the yield is within the standard yield mentioned.
- To collect the finished product control samples as per the requirement.To collect and send the stability samples as per the requirement.
- To Perform Daily and monthly Balance Calibration & Verification.Issue and verification of BMR, BPR and stereo.
- To review and retrieval of BMR & BPR after completion of manufacturing and Packing of batch. To perform Process validation batch sampling.
- Ensuring Quality of product in tablet, capsule as per specification.
- Handling and storage of Documentation of all QA records.Proper storage & Retrieval of BMRs and BPRs.
- Raising change control/deviation control for any deviating manufacturing process and keeping the record of the same.
- Process Validation all activity Flow.
- Art work Check in Foil and Carton Before printing.
- Product dispatch inspection and approval.
- To report the daily activity to QA Manager and DGM.

COMPUTER AND TECHNICAL SKILL

- **Basic knowledge of computer** : Operating System - Windows10
- **Microsoft Office**: Microsoft Word, Excel

Date:

Place: (**SAIDUL ISLAM)**