To,

R/Sir,

As a quality conscious personality with over 11 years’ experience in a pharmaceutical quality control and I think, I can contribute effectively to your organization.

Now I am working for **Mylan Laboratories Ltd**., SEZ Ahmedabad (Gujrat) as an **Executive.**

I have experience in analysis, review of Stability, In-process & Finished products, Analytical method transfer, Analytical instruments calibration etc.

I have experience in documentation as per **MHRA & USFDA requirements & also faced MHRA, USFDA & Other customer audits.**

Hope I can get the opportunity to meet you during the Interview.

Thanks & regards,

**RAKESH KUMAR**

**A 602, Gulmohar Nirvana**

**Opp. Pleasure Club, Bopal Ghuma Road**

**Ghuma, Ahmedabad**

**Gujrat - 380058**

**Mob.- 7679505042**

**Rakesh Kumar**

Village: Chainpur Hata

Dist. : Gopalganj

State : Bihar

**Contact No : 7679505042**

**E-mail: rakshahi@gmail.com**

**OBJECTIVE**: To work in a Technology driven organization that will offer strong challenges, career advancement & opportunities to use & expand my knowledge& capabilities, thereby contributing to the enhancement of organization & personal value.

**TECHNICAL QUALIFICATIONS**:

* M.sc Biotechnology from University of Pune.
* B.sc from Ganga Singh college (J P University, Chapra).

**EXPERIENCE (Over 11 years)**

* Worked with **Ankur Drugs & Pharma Limited, Baddi** from June 2009 to March 2010 as a **Microbiologist** of pharmaceutical & formulations like, Tablets, Capsules, Ointment & Oral liquid.
* Worked with **IPCA Laboratories Ltd, Sikkim** as a **QC Chemist** from Sep. 2011 to June 2014.
* Worked with **SUN Pharmaceuticals Ltd, Sikkim** as an **Officer QC** from Sep. 2014 to Jan. 2016.
* Worked with **Concord Biotech Ltd, Ahmedabad, Gujrat** as a **QC Officer** from May 2016 to June 2017.
* Worked with **Cadila Pharmaceuticals Limited, Ahmedabad, Gujarat** as an **Executive QC** from June 2017 to April 2018.

**AT PRESENT:**

**Mylan Laboratories Ltd(a Viatris Company), SEZ, Ahmedabad (Gujrat)** from April 2018 to till date.

**JOB PROFILE-**

* Analytical Data review and compilation of In-process, Finish Product & Stability data as per requirement of various regulatory agencies.
* Review of reports and protocol of the Analytical method transfer, Method equivalency & method verification documents.
* Review of reports and validation protocol for Process validation and Cleaning validation of commercial and exhibit products.
* Pertaining GMP training, Functional training and on job training to the analyst.
* Coordination with Research & Development for technology transfer and review of related documents.
* Reviewing the **Out of Specification (OOS) and Out of Trend (OOT)** results observed during analysis and monitoring during investigation. Monitoring Out of trend results before releasing the finished products.
* To check & verify all the test reports, Instruments logbooks and documents and ensure compliance as well as data integrity.

**INSTRUMENTS KNOWLEDGE: -**

* **HPLC**  I. WATERS- Alliance - Empower-3 software

II.SHIMADZU SYSTEM - Chromolean software.

III. Dionex (Ultimate 3000) -Chromolean Software

* DISSOLUTION TEST APPARATUS (Lab India & Electrolab)

And all QC relevant Instruments

**STRENGTH:**

* Willing to work as a key player as a Team leader and able to work independently with Working commitment and Positive Attitude.

**COMPUTATIONAL SKILLS:**

* Microsoft office (Word, Excel, Power Point)
* One Year Diploma in MIACT (Master in Advance Computer Technology).

**PERSONAL DATA**:

* Father’s name: Mr. Bhuvneshwar Shahi
* Date of Birth: 23rd January 1983
* Hobbies: Traveling & Reading
* Marital status: Married
* Sex: Male
* Nationality: India

**CURRENT CTC: 9.22 lacs**

**EXPECTED CTC: Negotiable**

**DECLARATION:**

I hereby declare that all above information given by me is correct to the best of my knowledge.

**PLACE:**

**(Rakesh Kumar)**

**DATE:**