KAMALESH KUMAR
UDAYAGIRI Tower flat no 206 2nd floor,
Near to Mustafa Junction iti road, Kurmannpalem
Visakhapatnam-530046 (A.P) India.
Contact No. 09696116581, 08894979542
Email ID : **kamalesh.prajapati326@gmail.com**

To,

Respected Sir/Madam,

Sub: - Application for the suitable position in your Organization.

I take this opportunity to introduce myself as **Kamalesh Kumar Prajapati** did my **B**. **Pharm** in **2010** and currently working as **Sr. Executive Quality Assurance in Pfizer Healthcare India Pvt. Ltd. Visakhapatnam A.P.**

I am interested to work in your organization with the knowledge and practical skills gained during my studies and occupation, I am sure that I am competent enough to fulfill the needs and demands of your esteemed organization. I will be very grateful if you have to consider my application.
I am enclosing my Resume for your kind consideration and favor. Looking to hear you soon with a positive response.

Thanking You in Anticipation

Yours sincerely,

Kamalesh Kumar Prajapati

Quality assurance Sr. Executive

Pfizer Healthcare India Pvt. Ltd. Visakhapatnam (A.P.)

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CAREER OBJECTIVE: -

To work in an organization where I can get ample opportunities to explore and enhance my skills there by contributing to organizational and self-growth.

Education: -

B. Pharm in 2010 from Saroj institute of technology and management Lucknow. Affiliated with Uttar Pradesh Technical University.

Professional Experience: - 10 Years.

- Currently Working with **Pfizer Healthcare India Pvt. Ltd. Visakhapatnam**, A.P. as Quality assurance Sr. Executive in Injectable section from 6th April 2015 to till date.
- Worked with **Theon Pharmaceuticals Ltd**. Nalagarh (H.P.) as Sr. Officer QA in Injectable section from 22 April 2013 to March 2015.
- Worked with **Akums Drugs and Pharmaceuticals Ltd**. Haridwar, U.K. as QA Officer in Injectable section from 1th October 2012 to 20 April 2013.
- Worked with **Aqua Vitoe Laboratories, Jharmajri, Baddi**, **H.P.** as Q.A. Chemist in Injectable section from August 2010 to 28 September 2012.

Current C.T.C.: - 7.7 Lac per Year.
Expected C.T.C.:- As per Company Norms.

Working with **Pfizer Healthcare India Pvt. Ltd. Visakhapatnam**, A.P. as Quality assurance Sr. Executive in Injectable section from 6th April 2015 to till date

Key Responsibilities:

- Ability to supervise area related activities in (Downstream) Terminal sterilization, Visual inspection and Packing area.
- Ensuring the products meet cGMP requirements
- Area monitoring during visual inspection and packing activity.
- Line clearance of autoclave for Terminal sterilization of product as per procedure specification (loading and unloading).
- Execution of process instruction sheet at various stages line clearance, area and equipment cleaning details, availability of equipment for specific area etc.
- Online review of batch reports at various stages (Terminal sterilizer, and packing).
- Initial impact assessment and closure of quality notification within specified duration.
- Review of Master packaging records at various stages (TS, VI and Packing).
- Review of ECN for various stages (TS, VI and Packing).

- Quarterly review of recipe and active user list for the following equipment Terminal sterilizer, Visual inspection machine (Bosch and Eisai), Leak testing (Wilco and Nikka) and semi-automatic inspection machine (Maschinpex).
- Review of process instruction sheet during batch and after completion of batch activity for attachments and other discrepancies if any.
- Review of Audit trial of the equipment which is having Audit trial.
- Review of Change control.
- Review of batch records and closing exceptions in AMPS (Rockwell Pharma Suite)

Theon Pharmaceuticals Ltd.

Worked with **Theon Pharmaceuticals Ltd**. Nalagarh (H.P.) as Sr. Officer QA in Injectable section from 22 April 2013 to March 2015

Key Responsibilities:
Validation:

- Co – ordination during scheduled validation of equipment's, process validation and media fill.
- Sample collection at various stages during Validation activity.
- Participation of equipment Validation like:
- Autoclave
- Tunnel
- HVAC

. **Documentation:**

- Prepare the area monitoring record.
- To check the Artwork of products.
- Preparation of BMR and BPR
- Handling Deviations.
- Handling the change control.

Production and Packing:

- Line clearance at various stages of manufacturing and packing activity.
- Line clearance of autoclave for various cycles such as rubber stoppers, m/c parts, and garments.
- Check the all parameters of Tunnel and washing machine.
- Balance calibration and verification.
- Coordination with QC and Production departments.
- To ensure the availability of current documents in each department.
- Calibration of IPQA Instruments/apparatus.
- Completion and review of online BMR/BPR.
- Sample collection at various stages of manufacturing and packing activity.
- Implementation of GMP on Floor.
- Area monitoring of aseptic area.

Akums Drugs and Pharmaceuticals Ltd

Worked with **Akums Drugs and Pharmaceuticals Ltd**. Haridwar, U.K. as a QA Officer in Injectable section from 1st October 2012 to 20 April 2013.

Key Responsibilities: -

Production and Packing:

- To Check the in-process (Manufacturing of batch, Filling & Packing) activities as per batch record.
- Sample collection at various stages of manufacturing.
- Filled volume/weight checking during filling operation.
- Maintain online documents, B.M.R., B.P.R., and in process control record.
- Review of batch manufacturing Packaging and Analytical records, equipment logs etc. before batch release.
- Balance calibration and verification.
- Monitoring issuance & review of BMR & BPR.
- Analysis of Secondary Packing Material.
- Collect Control & Stability Sample as per requirement.
- Coordination with QC and Production departments.
- To ensure the availability of current documents in each department.
- Calibration of IPQA Instruments/apparatus.
- Destruction of secondary packing materials.

Aqua Vitoe Laboratories, Jharmajri, Baddi, H.P.

Worked with **Aqua Vitoe Laboratories, Jharmajri, Baddi**, **H.P.** as Q.A. Chemist in Injectable section from August 2010 to 28 September 2012.

Key Responsibilities: -

Production and Packing:

- Good knowledge of dry powder antibiotics manufacturing process in Carbapenem, Penicillin, Non beta lactam and Cephalosporin.
- In process Q.A. related activities.
- Line clearance at various stages of manufacturing and packing activity.
- Checking the batch details printed on label, carton and shipper.
- Filled volume/weight checking during filling operation.
- In process checking during various stages of manufacturing and packing activity.
- Balance calibration and verification.
- Completion and review of online BMR/BPR.
- Sample collection at various stages of manufacturing and packing activity.

Personal Details:

Date of Birth	:	11 Sept. 1984
Father's Name	:	Mr. Manaraj Prajapati
Nationality	:	Indian
Language Known	:	Hindi & English
Attitude	:	Think Positive, Punctual, Honesty & Smart Working

Declaration:

I hereby declare that the information furnished above is true to the best of my knowledge.

Date:

Place: (**Kamalesh Kumar)**