Perusing Ph.D in chemistry

from Venkateshwara

University.

Name:

Aviral Sharma

Permanent Address

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Personal

Information

Name – Aviral sharma S/O – Sh. Ram sewak sharma Date of Birth: Oct 30, 1990

Sex - Male

Nationality - Indian Marital Status - Single

Hobbies & Interests

- Surfing the Internet
- Learning new things.
- Blogging
- Solving Sudoku.
- Listening to music

<u>Strength</u>

- Self Confidence
- Honest
- Truthfulness
- Leadership
- Positive attitude
- Optimistic

CURRICULUM VITAE

Career Objective

To enhance my skills set by working in a challenging and competitive environment while at the same time contributing to the growth and progress of the organization and to put the best efforts towards mutual growth.

Academic Qualifications

DEGREE/EXAM	INSTITUTE	UNIVERSITY/BOARD
Ph.D. (Start From 2016)	Shri Venkateshwara University	Shri Venkateshwara University
M.Sc.(chemistry)	Monad University	Monad University
B.Sc. (P.C.M)	Bundelkhand University	Bundelkhand University
10+2	M.B.I.C	U.P Board
Matriculation	Prabhat Public higher secondary school	U.P Board

Author and developer of www.phamadocument.com

CURRENT EMPLOYER:

Zifam Pyrex Myanmar Co. Ltd. (Yangon, Myanmar).

Working as section Head (Aspiring to TGA)

(Tablet, Capsule, Dry Injection, Dry Syrup)

Since June 2017

- ➤ Handling of change control, OOS, OOT, Deviations & investigations (Quality Management System) QMS.
- Review of Analytical Method Validation Protocol, Reports and documents for accuracy and Compliance.
- Resource management, planning, monitoring and control of manpower, instruments and equipment's usage, to accomplish workload requirements.
- ➤ To Coordinate with QA and Regulatory Affair Department for data filing and submitting.

- > To participate in internal audits and regulatory inspections at the site.
- ➤ To prepare Analytical Method Validation Protocols.
- ➤ To update documents as per pharmacopoeia changes and change in regulatory guidelines.
- ➤ Preparation, training of SOPs and execution of activities to ensure compliance as per standard operating procedures. To prepare the URS for instruments.
- To prepare STP, GTP of Products as per regulatory requirements.
- ➤ To develop internal training modules for staff and workers to enhance their abilities and to update them w.r.t. latest requirements of GLP and GMP.
- > To provide training to the analysts for operation of lab Instruments and trouble shooting.

PREVIOUS CARRIER SUMMARY:

Ph.D. Course Work at University Campus:

Since: October 2016 to June 2017

Previous Employer

Company: Nectar life Science (Baddi) Himachal Pradesh

Manufacturing unit for dosage forms (Tablets and Injectable). The unit has the approvals of ANVISA, EU (EUROPEAN UNION) and USFDA.

Post: Executive

Since: JUNE 2014 to June 2016

JOB PROFILE:

- > Preparation of Stability Protocols as per ICH Guidelines or any country specific requirements.
- Preparation of Stability schedule as per Stability Protocols for Stability products.
- > Management of Stability chambers and monitoring their temperature and Humidity records.
- Preparation of Working Standards.
- > Preparation of Stability Summary Reports for tabulated and graphical presentations.
- Review of Stability data for accuracy and compliance.
- Review of calibration and Qualification documents for their accuracy and completeness.
- ➤ Handling of HPLC and GC Column.
- Handling of Capex.
- Analysis of Stability samples (Tablets, Capsule, and Syrups)
- Analysis of in process and finished Product samples.
- Preparation of Analytical Reports as per GLP.

Previous Employer

Company: Glenmak Pharmaceutical (Baddi) Himachal Pradesh

The unit has the approvals of ANVISA, EU (EUROPEAN UNION) and USFDA.

Manufacturing unit for Injection, Capsule and Tablets. Post: Officer

Since: December 2011 to May 2014

JOB PROFILE:

- Analysis and sampling of Raw Material
- Analysis and sampling of Packing Material
- Analysis of Stability Samples.
- Calibration of Instrument.
- Review of Stability, Raw Material and GLP Data.
- Analysis of in process and finished Product samples.
- Preparation of Analytical Reports as per GLP.
- Preparation of Working Standards.

Handling of following Instruments:

- Operation and Calibration of High Performance Liquid Chromatography.
- Operation and Calibration of Gas chromatography.
- > Operation and Calibration of Atomic Absorption Spectrophotometer.
- Operation and calibration of U.V Spectrophotometer.
- Operation and calibration of FTIR
- Operation and calibration of Dissolution.
- Operation and calibration of viscometer
- Operation and calibration of Auto-Titrator.
- Operation and calibration of Karl fisher
- Operation and calibration of DT Apparatus.
- Operation and calibration of Friability Test Apparatus
- Operation and calibration of Refractometer.
- Operation and calibration of Melting Point Apparatus.

Technical Skills:

- Proficiency in MS-Office (MS-Word, Excel etc.)
- Result Recording and usage decision in System application and Products in data processing

I hereby, declare that all the information's provided here are correct to the best of my knowledge.

Place:	
Date:	Aviral Sharma