# VIBHA VISHWAS RAUT

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## Summary

Experienced and resourceful analytical development research scientist having strong attention to detail specialized in method development, method validation, method transfer having knowledge in GMP, ICH guidelines and regulatory requirements for testing and validation procedures. Seeking Regulatory affairs position where 2.9 years of working in highly regulated and GMP compliant environment in leading pharmaceutical industry such as Sanofi Synthelabo Private limited will add value. Strongly reliable and focussed individual with excellent interpersonal , time management skills and have ability to grasp new concepts quickly who can work independently as well as in a team.

## Experience

- August 2017- July 2018
- Apprentice , Analytical Development at Sanofi–Synthelabo (India) Development centre, Verna-Goa, 403722
- August 2018 – July 2019.
- Research trainee , Analytical Development ,skill development, Yashaswi Academy, at Sanofi–Synthelabo (India) Development centre, Verna-Goa, 403722.
- August 2019-May 2020.
- Research Scientist-II, Analytical Development, Vergo Pharma Research Laboratory, Verna-Goa, 403722.

## Certifications

- Distance learning General course on Intellectual Property Rights and Introduction to the Patent Cooperation Treaty through World Intellectual Property Organization.

## Areas of Interest

- Regulatory affairs
- Quality Assurance

## Skills

- Basic knowledge of GLP, GMP, GDP, CTD modules and regulatory requirements for testing and validation procedures.
- Basic understanding of ICH guidelines.
- Basic knowledge of Intellectual Property Rights, Patents.
- Operational knowledge of analytical instruments like HPLC, Dissolution apparatus, UV spectrophotometer, Karl Fischer apparatus.
- Hands on Software's like Empower 3 and Chromeleon.

## Job Profile And Responsibilities

- Performed method development, method validation , method transfer.
- Performed qualitative and quantitative analysis such as assay, dissolution profile, moisture content, uniformity of dosage form, forced degradation studies, residue analysis,
- Performed calibration of analytical instruments.

## Educational Qualifications

- M Pharm in Quality Assurance from Goa College of Pharmacy in year 2017 .
- B Pharm from Goa College of Pharmacy in year 2015 .
- D Pharm from Goa College of Pharmacy in year 2012 .
- HSSC from Saraswat Vidyalaya in year 2010.
- SSC from Holy Family High School in year 2008.

## Project Work

- Development of analytical method for estimation of drugs in presence of its impurities by RP-HPLC.
- Validated absorption correction method for estimation of drugs in combined dosage form.

## Industrial Training

- One month training at Unichem Laboratories Ltd, Pilerne , Bardez Goa .

## Technical Activities

- APTICON 21$^{st}$Manipal University APTI Annual National Convention between 14$^{th}$- 16$^{th}$ October 2016,.
- Industrial visit to Indocco Remedies, Verna Industrial, Estate, Verna, Goa.

## Strengths

- Good communication skills.
- Ability to work as team as well as an individual .
- Eagerness to learn new things.
- Eager to take responsibilities.
- Focused and career oriented.
- Time management skills.
- Ability to grasp new concepts quickly.

## Languages

English, Hindi, Marathi, Konkani

## Declaration

I hereby declare that the above said information is proof read , refined, perfect and best of my knowledge.