**Curriculum Vitae**

**Name**  : Raghavendra R

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Professional Summary

* 9.5 years of experience in the diversified field of Pharma.
* Knowledge on stability Program Management, Laboratory Testing, Documentation, Quality Management System, Safety and Good Laboratory Practices.
* Responsible for day-to-day operational activities like Analysis, Review, Compilation, Incubation of Stability samples.
* Having good understanding of Business Requirement Documents, Functional and System Requirement Documents.
* Strong Analytical and problem solving skills.
* Efficient in interacting with business requirements, teams and refining the requirements for clarity and simplicity.
* Handling LIMS

Experience Summary

* Currently working as a Sr.**Associate** for Stability at Mylan, Bangalore from July 2010 to till dated.
* Worked as **Quality Assurance Officer** for Sequent Research Limited, Mangalore from September 2008 to July 2010.
* Worked as **Chemical Analysis Chemist in Quality Control** for Remidex Pharmaceuticals Pvt.Ltd

**Qualification:**

* **Bachelor of Science (58%)** - Adhichunchanagiri College, Channarayapatna.
* **Pre University Course (PCMB 49%) -** Government PU college, Mangalore.
* **SSLC (56%) -** Navodhaya School, Channarayapatna

**Career Profile:**

**Agila Mylan Company: – Currently Working as Associate since July-2010**

**Employer** : Mylan Company

**My Role** **:** Sr.Associate

**Responsibilities / Contributions:**

* Work allotment and handling the team.
* Working on **LIMS (Lab Information Management System )**
* **Stability Program Management**: Aware of guidelines, stability testing requirements perform and conduct appropriate stability tests, record results in order to ensure product robustness. Ensure the raw data and results.

1. Incubation of Stability samples as per commitment with customer and regulatory requirements by coordinating with plant QA, RAD and customers.
2. Review of stability chamber calibration and mapping reports received from HTA
3. Ensured the stability chamber cleanliness and surrounding area clean.
4. Reviewed the stability temperature and humidity record data.
5. Preparing for audits during audit in all the plants.
6. Prepared qualification protocol (Design Protocol) and reviewed Installation qualification, Operational qualification, Performance qualification protocols and reports.
7. Relevant SOP’s prepared.
8. Reviewed qualification documents on (HSF) stability.
9. Providing stability protocols and specification and also STP per schedule.

* **Laboratory Testing**: Receive and analyze product samples during various stages of product development and line testing for external locations. Compare and analyze results against expectations, record and submit results in order to provide timely information.
* **Documentation**: Prepare documentation of specifications and test procedures, review pharmacopeias and update documents accordingly, seek approval and submit to user departments in order to ensure proper documentation and technology transfer to product location.
* **Quality Management System**: Ensure the compliance of the documents, instruments. Maintain the calibration schedule, calibration record and history cards. Ensure the testing that has to be performed as per the Standard Test Procedures.
* **Safety and Good Laboratory Practices**: Aware of laboratory safety and follow good working practices in daily activities in the laboratory towards the implementation of GLP.

Sequent Research Limited: –Worked as Quality Assurance officer September 2008 to July 2010

**Employer** : Sequent Scientific- Research Ltd.

**My Role** **:** Quality Assurance – QA Officer

**Responsibilities / Contributions:**

* Work allotment, Handling and troubleshooting of Analytical issues.
* Investigation and closure of Out Of Specification results, Investigation and Closure of Incidents.
* Analysis of stability samples related to out of specification and Review of analytical documents and Review of Monthly Instrument Calibration Reports.
* Taken sole responsibility in GLP (Good laboratory practices)
* Internal Audit, Issuance of Analytical Raw data sheets.
* Raising requisitions for Material requirement and planning trainings for the new joinees.

Remidex Pharmaceuticals Pvt.Ltd: – Worked as Chemist in Quality Control from Sept-2006 to Sept-2008

**Employer** : Remidex Pharmaceuticals Pvt.Ltd.

**My Role** **:** Chemical Analysis Chemist in Quality Control

**Responsibilities / Contributions:**

* In - process and finished products.

Audits Faced:

* GSK
* MERCK
* FDA
* WHO
* Internal Audits
* ANVISA Audits

**Instruments Handled:**

* Melting point Apparatus, Disintegration Tester, Dissolution Tester, FTIR Spectrometer, Tablet friability Tester, UV – Visible Spectrophotometer & HPLC,
* Automatic KF Titrator, Auto Titrator Polari meter and pH Meter.

Personal Profile:

* Name : Raghavendra. R
* Date of birth : 26th October, 1981
* Sex : Male
* Father’s Name : Rudrappa. S
* Marital Status : Married
* Nationality : Indian
* Languages known : English, Hindi, Kannada, Tulu.

Declaration:

I, **Raghavendra R** do here by declare that the information given above is true and correct to best my knowledge.

I also understand that any false declaration or willful suppression of information shall amount to misconduct and may result in appropriate disciplinary action.

Place : Bangalore

Date : **Raghavendra R**