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| **Sahel****Sahel.371382@2freemail.com** **QA / QC Professional**  |  deede |
| **PROFILE SUMMARY**To obtain a Senior or Intermediate position in Pharmaceutical Quality Control or Quality Assurance that will utilize my skills in regulatory laboratory operations, knowledge and application of compliance requirements interpersonal communication, leadership and technical competence to seek areas of improvement, increase efficiencies to meet organizational objectives. |
| **STRENGTHS** |
| * Technical skills related to Analytical testing and QC lab operations.
 | * Compliance auditing skills (21CFR 110,210,211,820 & HACCP systems).
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| * Skilled in report writing & presentation
 | * Familiar with ISO Environment
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| * Data recording & record keeping management
 | * Can work independently with keen eye for details
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| * Able to work under pressure & tight timescales
* Well-developed time management skills
 | * Has positive approach to teamwork
* Planning and Organizing
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|  | *EDUCATION* |  |
| **H. S.S.C. (Science) 1999-00**Guardian angel High school, curchorem Goa, Board of education Goa, India. |  |
| **Bachelor of science (B.Sc.)**Government college Quepem, Goa university, Goa, India. |  **2000-03** |
| **Master of Business administration with specialization in Six sigma green belt , Pharmaceutical Management and Event management** Institute of Business management, Delhi, India  |  **2011-13** |
|  | *ACHIEVEMENTS* |  |
| * Successfully Faced Audits :-

 * US FDA ( Thrice)
* MHRA (Thrice)
* SANDOZ
* WHO (Twice).
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|  | *WORK EXPERIENCE* |  |
| **QA/QC Chemist -** MicroLabs Verna Goa **Nov 2012 – July 2015**Responsibility:* Working in Stability section
1. Analysis of Exhibit and Commercial Stability samples.
2. Review of Specification, Standard test procedure and Test Data sheet for Stability Analysis.
3. Handling of Stability samples for charging and summary sheet preparation for samples.
4. Preparation of method transfer protocol and report of submission products.
5. Handling instruments like Dissolution autosampler (Electrolab) and HPLC (Waters)
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| **QA/QC Chemist -** Lupin pharmaceuticals Verna Goa | **Aug 2007 – Oct 2012** |
| Responsibility:* Worked in raw material section
1. Analysis of excipient and API, Sampling of raw materials
2. Review of Specification, Standard test procedure and Test Data sheet for raw material. Analysis of water.
* Prepared BMR and various documentation as per USFDA guidelines and CGMP norms.
* Worked extensively on Dissolution Apparatus I & II and Disintegration Apparatus for Quality Control Testing of Solid Dosage Forms.
* Prepared Solutions and Dilutions as per the test requirements by Senior Chemist to perform HPLC analysis.
* Responsible for analyzing compounds as given and interpretation of those results.
* Pursued Training for different Analytical Techniques like IR Infrared Spectroscopy, NMR Nuclear Magnetic Resonance, HPLC, FTIR Fourier Transform IR Spectroscopy etc.
* Performed Calibration of analytical instruments like UVVis Spectrometer, pH meter.
* Handled projects like reduced testing for excipient and API
* Handled project for common specification where 25 common specifications for excipients prepared
* Handled all RA requirement and fulfilling all requirements for FILING OF ANDA PRODUCTS related to RAW MATERIAL and PACKING MATERIAL
* Knowledge of SAP related to RM
* Handling other QA and HORA QUERRIES
* Worked in PMQC and handled all packing QC responsibility like RA submission, release of material thru SAP, sampling and approval of material.
* Carried out reduced testing and implemented in various packaging material like HDPE Bottles, inserts, medication guides, closures, labels, silica gel etc.
* Carried out reduced testing and implemented in various API and excipients of raw material for various tests.
* Worked on a project of shelf life updation and maintained all the stock of raw material by updation of shelf life for all the available material in line with vendor COA, SPECIFICATION and SAP.
* Knowledge of SAP related to RM and PM
* Handled NIR and knowledge of library creation and validation for raw

 Materials in NIR.  |  |
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|  **QC chemist** - Cipla Pharmaceuticals Ltd Verna Goa | **Mar 2005 – May 2006** |
| Knowledge of work: {In Cipla}* Worked in non-routine section for 8 months
* Preparation, Maintenance of record and standardization of all volumetric solutions
* Preparation and Maintenance record of reagents and indicators
* Daily Standardization of Karl Fischer and diluted KF record maintenance and calibration
* Handled all potentiometric titrations by auto-titrator.
* Worked in documentation area for 4 months.
* Master preparation of finish product, stability, and raw material.
* Software knowledge of TQC, Validation of masters.
* Daily, weekly, Monthly Balance calibration.
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|  | *AREAS OF EXPERTISE* |  |

* Analytical skills, API, Balance, calibration, documentation, filling, FTIR, materials, Perkin Elmer, QA, SAP software knowledge regarding QC, Specification, UV, Validation.

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|  | *COURSE -TRAINING* |  |
|  * Diploma in computer through NIIT obtaining 75 %
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|  | *TECHNICAL PROFICIENCY* |  |
|  * Knowledge of High Performance Liquid Chromatography (Shimadzu).
* FTIR Spectrophotometer (Perkin Elmer).
* UV-VISIBLE Spectrophotometer. (Perkin Elmer).
* pH meter and Conductivity Meter.
* Refractometer and Viscometer.
* Melting Point Apparatus.
* Karl Fischer Apparatus.
* Particle Size Analyzer.
* Auto-titrator
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|  | *IT SKILLS* |  |
|  Proficient in computer skills, e.g. Word, Excel, etc. Problem solving and organizational skills. |

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|  | *PERSONAL DETAILS* |  |
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| Nationality | : | Indian |
| Date of Birth | : | 2nd July 1981 |
| Marital Status | : | Single |
| Visa Status | : | Indian Citizen |
|  Languages Willing to relocate  | :: | English, Hindi, Marathi & KonkaniYes |