CURRICULUM VITAE

ARUNKUMAR GUNDAIAH RAMESH

76 ST PAULS AVE
APT 2B, JERSEY CITY
NEWJERSEY, ZIPCODE: 07306,
: 201-423-2032
E. mail: apotex2014@gmail.com

OBJECTIVE

A career in pharmaceutical Product Development including Technology Transfer with emphasis on developing novel patentable concepts that are safe, cost effective and cater the present business needs of industry.

PROFESSIONAL EXPERIENCE

FENNY SPECALITY PHARMACY, JERSEY CITY, NEWJERSEY

PHARMACY INTERN FENNY SPECALITY PHARMACY from OCT 2009 till date

Technical Expertise

- WELL VERSED WITH PRIVATE INSURANCE BILLING SOFWARE.
- DISPENSING OF MEDICATIONS AND EDUCATING THE PATIENT ABOUT THE MEDICATIONS
- FORMULATION OF COMPOUNDED MEDICATIONS OF ORAL AND TOPICAL DOSAGE FORMS.
- BRAND AND GENERIC PRICE EVALUATION.

Apotex Research Private Limited, Bangalore (Apotex Inc.)

SENIOR FORMULATOR - Formulation Development, from FEB 2006 till JUL 2009.

Worked on development of generic equivalent of Oral dosage form such as Tablets and Capsules including Immediate-Release, Bi-layer, Modified-Release Dosage forms. Strategy finalization, patentable / non-infringing product development, exhibit batches and documentation in compliance with the current regulatory requirements for Solid oral dosage form like tablets and capsules for USFDA, European Union, Australia and Health Canada regulatory agencies.

Technical Expertise

- Development of novel, patentable and non-infringing Generic version of Solid dosage forms [viz., Immediate-Release, Extended-Release, Delayed-Release (Enteric coated)].
- Expertise in formulation development and scale up techniques involved in development of solid oral dosage form.
- Worked with a team of people to meet project and team objectives along with pilot plant.
- Well introduced with regulatory filing procedure, contents and regulatory requirements for ANDAs to USFDA, EMEA, TGA and Health Canada regulatory authorities.
- Dissolution, bioavailability & bioequivalence studies of newly developed Immediate-Release, Delayed / Controlled dosage forms.
- Well versed in the Roller compaction technique of dry granulation.
- Documentation of Product Development report, Batch Manufacturing Record, In-process & Finished product specification, Raw material specification, Licensing information, Dossier information and various protocols.

Dabur Research Foundation, Delhi

Research Scientist – Formulation Development & Generics, from DEC 2004 to JAN 2006

Worked on development of generic equivalent of Solid oral dosage forms such as immediate release Tablets and Bi-layer Tablets and Parenteral dosage forms. Involved in the development of products for USFDA, European Union, and for less Regulated market across the globe.

Product Development

- Development of Generic version of Solid dosage forms of Anti-cancer drugs for US & EU Market.
- Development of Generic version of parenteral dosage forms of Anti-cancer drugs for US & EU Market.
- Performing Pre-formulation studies of APIs and carrying out Excipient compatibility studies for Solids.
- Performing Pre-formulation studies of APIs and carrying out Ion-compatibility studies for Parenterals.
- Literature / Patent evaluation Reports.
- Patent Strategies & Non-infringement Reports.
- Documentation for Technology Transfer.
- Preparation of Development reports.

Zydus-Cadila Healthcare Limited, Bangalore

Research Pharmacist – Formulation Development, from MAY 2004 to OCT 2004

Product Development

- Development of Immediate-Release dosage forms of Tablets and Liquid orals and Pre-formulation studies.
- Compilation of Development reports
- Preparation of Standard Operating Procedures for all Equipments and executing the same.
- Preparation of IQ/OQ/PQ protocols for equipment and execution.
- Process improvement and Scale-up trials.

Strides Arcolab Limited, Bangalore

Research Executive – MAY 2003 and MAY 2004.

- Development of Immediate-Release dosage forms of Tablets and Liquid orals and Pre-formulation studies.
- Compilation of Development reports
- Preparation of Standard Operating Procedures for all Equipment's and executing the same.
- Preparation of IQ/OQ/PQ protocols for equipment and execution.
- Process improvement and Scale-up trials.

EDUCATIONAL QUALIFICATION

COURSE	BOARD / UNIVERSITY	YEAR	% OF MARKS	CLASS
S.S.L.C	Karnataka secondary education examination board	1992-93	**62.24**	**FIRST**
P.U.C	PRE-UNIVERSITY BOARD	1993-94	**52.33**	**SECOND**
BACHELOR IN PHARMACY	BANGALORE UNIVERSITY	1995-00	**67.92**	**FIRST**
MASTER IN PHARMACY	RAJIV GANDHI UNIVERSITY OF HEALTH SCIENCES	2001-03	**73.71**	**FIRST**

TECHNICAL COMPETENCY

Proficient in MS Word, MS Excel, MS PowerPoint and Design expert software.

ACHIEVEMENTS

- ✓ **Presented** my Research work "MUCOADHESIVE DOSAGEFORM OF CAPTOPRIL Formulation and its Stability studies" at 56th Indian Pharmaceutical Congress 2005, PUNE.
- ✓ Secured FIRST CLASS at the University Level in M. Pharm (Pharmaceutics) examination.
- ✓ Qualified **GATE 2001** with 89.12 (Percentile Score). All India Rank - **195.**

PERSONAL DETAILS

FATHERS NAME	:	G. RAMESH
SEX	:	Male
DATE OF BIRTH	:	8TH OCTOBER 1977
MARITAL STATUS	:	MARRIED

REFERENCES

1. **Dr. N. RAVI KUMAR**
   Head – Product Development
   Jubilant Life Science
   Noida – 201 301
   Mobile: +91 9880125998
   E.mail: ravikumarn.n@gmail.com

2. **RAJESHA B. C**
   Deputy General Manager – Formulation Development
   Apotex Research Private Limited
   Bangalore – 560 099
   Mobile: +91 95386 01400
   E.mail: brajesha@apotex.co.in

3. **S. SRINIVAS**
   General Manager – Technical
   Remedia Therapeutics Pvt. Ltd.,
   Mobile: +91 90084 11331.
   Email: srinivas_28123@rediffmail.com

4. **ANIL KUMAR**
   HEAD – FORMULATION DEVELOPMENT
   STRIDES ARCOLAB LTD
   Mobile: +91 9845241870