RESUME

✉mm_murugan1982@yahoo.co.in
M.ELAMURUGAN ☎ +919840822740 & 7397375061

Career Objective

Looking for a responsible and challenging position with opportunities to enrich my knowledge and skills while contributing my best and devote myself towards the growth of the Organization.

Employment History - Present

M/S. NATCO PHARMA LIMITED (USFDA Approved Facility)

Position Title **: Manager – Quality control**
Industry : Bulk drug Manufacturing (Generics)
Duration : **Nov 2009 – Till date**

Job Responsibilities:-

- Ensure effective functioning of the department activities and Manpower planning for regular shift activities.
- Assigning job responsibilities to overall QC persons.
- Ensuring Quality of Raw materials, intermediates and finished products and timely release to meet management goals.
- Ensuring proper investigation of deviations, complaints, stability failures and out of specifications.
- Review and Training with respect to GMP, GLP, Protocols, SOPs, STPs and SPECs.
- Instruments handled like HPLC(WATERS, SHIMADZU,AGILENT), GC(AGILENT, PERKIN ELMER, VARIAN), FTIR(PERKIN ELMER, NICOLET), UV(AGILENT, VARIAN, PERKIN ELMER), IC (METRHOM), TOC (SHIMADZU), etc.,
- Having knowledge about data integrity (manually created data and system generated data) and 21 CFR Part 11 and training to QC personnel for the same.
- Training to Quality control employees related to network based Empower3 (FR2) and other standalone softwares, Stability studies, Analytical method validation, etc.,
- Preparation of training matrix (Yearly, monthly and weekly) and conduct class room training weekly and covered 35 employees in a month.
- Evaluation for production related SOPs, BPCRs and BCRs.
- Coordination with QA to prepare site master file, process, drying, cleaning validation protocol and report.
- Coordination with QA for new document initiation, deviation, change control, vendor audit and OOS investigation.
- Customer and regulatory audit preparation.
- General lab compliance, Document review & release and Facing internal and external audits, OOS investigation.

Employment History - Previous

M/S. ACTAVIS PHARMA MANUFACTURING PVT LTD

(USFDA Approved Facility)

Position Title : **Executive - Quality Control**

Industry : Bulk drug Manufacturing (Generics)

Duration : **Apr 2008 – Nov 2009 (1.7 Years)**

Employment History - Previous

M/S. COVALENT LABORATORIES PRIVATE LIMITED

Position Title : **Executive - Quality Control**

Specialization : Shift in charge

Industry : Bulk drug Manufacturing (Cephalosporins, Non Cephalosporins)

Duration : **APR 2003 – SEP 2008 (4.6 Yrs)**

Educational Background

Master of Science (M. Sc)

Field of Study : Chemistry

Major : General chemistry

University : Kovempu University, Karnataka-India

Bachelor of Science (B. Sc)

Field of Study : Chemistry

University : Madras University, Tamilnadu-India

Personal Strengths:

- Ambitious, Committed
- Adaptability
- Leadership skills to work/lead along with other members
- Analytical and problem solving
- Ability to relate to co-workers in a close environment
- Deal with people, problems and situations with honesty and respect

Personal Particulars

Date of Birth	: 25.05.1982	Age	: 36 years
Gender	: Male / Married	Nationality	: Indian
Address	: 1353, LIG-1,Mathur, Chennai – 600 068. Tamilnadu.		

References

1. Mr V.Venkata ramanujam
Head – QC
Natco PharmaLtd,
Chennai – Tamil Nadu
Mobile : +919444686410

2. Mr. M.Vinayagamurthi
GM – QC
Chennai – Tamil Nadu
Mobile : +919840722079

Yours Truly,

Station : Chennai

Date :

(M.Elamurugan)