

Pradeep P. Tamkhane.

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PRESENT ADDRESS:

Divyajyot Residency A-101, Plot No. - H-3164,
Opp. City Centre, 500 quarter road,
GIDC, Ankleshwar (GJ.) PIN-393001.

PERMANENT ADDRESS:

Palava City, Casa Rio, Antarctica B-0202,
Opp. Pawar Public School,
Kalyan Shil Road, Dombivali (MH), PIN- 421204.

OVERVIEW:

Prodigious expertise gained in the field of Production, Planning and operations with more than **20+ years** of experience. I have been involved in Production, Planning, Facility modification Project Management, and Operations Management & thorough activities in my endeavor. I have been very successful in mentoring the staff at each of my alienation due to my immense orchestration potential. I have gained experience as a Sr. Production manager, Dy. Manager, Assistant Manager, Team leader and a Team player who always gets the job done.

- ❖ Having more than 20+ Years of Experience in manufacturing and operations in Speciality chemicals industries and pharmaceuticals.
- ❖ Proficiency in Production Management and Operations Management in the Specialty chemical industry & Pharmaceuticals.
- ❖ Overseeing production of speciality chemicals, intermediates, key raw material and implementing plans to ensure timely accomplishment of production targets within the cost parameters.
- ❖ Well versed with kilo scale, pilot scale and commercial scale in Pharmaceutical industry and also experience in scale up techniques & accelerated process Improvement studies of products along with R&D scales.
- ❖ Overseeing all Manufacturing Blocks, effectively.
- ❖ Having the experience Lead-Operations for the site of 150 numbers of team and 300+ contractual manpower, where I am responsible for plant operations like production, engineering, EHS, projects, purchase and administration.

EDUCATION:

- ❖ B. Tech (Chemical Engineering) From UDCT, North Maharashtra University, Jalgaon, Maharashtra in 2003 with 67.62%.
- ❖ 10+2 from Nasik Board with 58.00%.
- ❖ 10th from Nasik Board (1994-95) with 81.42%.

CORE SKILLS:

- ❖ Easily getting familiar with any new system.
- ❖ Good interpersonal and interdepartmental communication skills.
- ❖ Analysis for Breakdown, losses and corrective actions in plants.
- ❖ Process Validation, scale up and Commercialization of existing New Products.
- ❖ Responsible for taking care of all audits (internal (quality and safety), external customer (quality and safety), regulatory like US FDA, EDQM, MHRA) in the plants all over the year.
- ❖ Production management from raw material to dispatch of Finished Goods.
- ❖ Experienced in various types of reactions.
- ❖ QMS system (Quality management system).
- ❖ Safety studies (HazOP, HIRA and PSSR).
- ❖ Mechanical Projects Monitoring.
- ❖ Ensuring the completion of projects as per project Scheduling, costing, planning and commissioning.
- ❖ Guiding team to fulfill daily production requirements.

Quality management System Implementation / Process Improvement

- ❖ Setting up facilities in the organization to spearhead quality management initiatives to improve the quality system in the plants.

- ❖ Establishing quality management systems across various processes to meet the pre-set quality standards and reduce rejection levels.
- ❖ Imparting training to all employees, pertaining to quality implementation and productivity improvement.

Operations Management

- ❖ Responsible for cGMP implementation, Safety and SAP purchase requests; preparing production reports and training records for the changes intended /implemented.
- ❖ Planning and effecting preventive maintenance schedules of various machinery and instruments to increase machine up time and equipment reliability.
- ❖ Responsible for the all Internal & external audits which includes US-FDA, regulatory and all customer audits.
- ❖ Responsible for the regulatory Audits like US-FDA, EDQM and etc.
- ❖ Good exposure to systems ISO 9001:2008, ISO 14001, ISO 18001:45001, EHS & cGMP.
- ❖ Exposures to the productivity enhancement tools like **MEP** (manufacturing excellence program) and **Operational excellence**.

Participation in the site social activities

- ❖ Active member of the ENCON (energy conservation) team on site.
- ❖ Active member of mock drill team.
- ❖ Active member of SAT (Safety Audit Team).

PROFESSIONAL EXPERIENCE:

Industrial Solvents and Chemicals Pvt. Ltd. Ankleshwar, (Gujrat).

Oct-2023 – till date

Role: Sr. Manager - Production

Responsibilities:

- ❖ Heading 05 commercial block, manufacturing organic as well as inorganic compounds, responsible for the production and all activities of the Production plant.
- ❖ Regularly monitoring the capacities of products across all the products running in the plant keeping in mind the Dynamics of Market.
- ❖ Executing the Production by following Safety and ISO norms.
- ❖ Prioritization of manufacturing based on throughput, to get best contribution.
- ❖ Co-coordination with dept. such as stores, Q.C., maintenance, safety, purchase and dispatch to streamline the operation activities.
- ❖ Responsible for all the documentation as per ISO standards.
- ❖ Maintaining the plant auditable by strictly maintaining the safety and ISO standards.
- ❖ Responsible for all deliveries in time.
- ❖ Regular follow-up of procurement, quality & dispatches.
- ❖ Responsible for the all Internal & external audits which includes ISO, regulatory and all vendor audits.
- ❖ Managing the team of 150 + employees in the current organization.

Aarti Pharmalabs Limited, Tarapur, Boisar.

June-2017 – Oct-2023

Role: Dy. Manager - Production

Responsibilities:

- ❖ Successfully monitored, completed the execution of API products commercial scale Validations
- ❖ Heading 1 commercial block, Steroid manufacturing, Responsible for the production and all activities of the Production plant.
- ❖ Regularly monitoring the capacities of products across all the Products running in the plant keeping in mind the Dynamics of Market.
- ❖ Executing the Production by following cGMP and Safety.
- ❖ Prioritization of manufacturing based on throughput, to get best contribution.
- ❖ Co-coordinating with dept. such as stores, Q.C., maintenance, safety, PD lab to streamline the operation activities.
- ❖ Involving in the activities from the development study of the new products.
- ❖ Responsible for all the documentation of the new products till commercial scale from the development scale.
- ❖ Maintaining the plant auditable by strictly maintaining the safety and cGMP.
- ❖ Responsible for all deliveries in time.
- ❖ Regular follow-up of procurement, quality & dispatches
- ❖ Responsible for the all Internal & external audits which includes US-FDA, regulatory and all customer audits.

- ❖ Managing the team of 55 + employees in the current organization.

Jubilant Life Sciences Limited, Ambarnath, Mumbai, Maharashtra. Mar 2010 to June 2017

Role: Assistant Manager - Production

Responsibilities:

Capacity Management-Mfg. Strategy:

- ❖ Maintained the Safety & GMP in all the plants.
- ❖ Successfully implemented ZLD, with the help of ETP (Effluent treatment Plant) with MEE (Multiple effective evaporator).
- ❖ Regularly monitoring the capacities of products across all the Products keeping in mind the Dynamics of Market.
- ❖ Managed the Unit of 01 production block and 60 employees.
- ❖ Setting up facilities in the organization to spearhead quality management initiatives. Implementation of QMS where a third party audit team conducts a half yearly check and submits a report to the management. Identified critical points because of this move and inculcated the quality mind set to employees of the Unit.
- ❖ Monitoring functioning of plant; implementing many recycle / re-use schemes for reducing pollution loads and improving recoveries.

Enaltec Labs Pvt. Ltd., Dombivali, Mumbai.

May-2009 to March-2010.

Role: Worked as Assistant Manager (Operations).

Responsibilities:

- ❖ Lead the site from planning to execution.
- ❖ As an Executive, I was fully responsible for the yields and a quality of various API's produced with firm fermenter.
- ❖ Responsible for operations of various pharmaceutical ingredients and final products.
- ❖ Managed people from different outsourcing units and established the products at commercial scale.

Glenmark Generics Limited, Solapur, (MH).

Sept. - 2004 to April-2009

Role: Worked as Sr. Officer (Production).

Responsibilities:

- ❖ Independent handling of shift duties.
- ❖ Ensuring smooth operation to maximize production and to meet preset targets.
- ❖ Maintaining good housekeeping.
- ❖ Handled various types of reactions and unit operations.
- ❖ Implementing operation policies lead down by mgmt.
- ❖ Cost reduction in one of the products by use of recovered solvent.
- ❖ Managed people from different outsourcing units and established the products at commercial scale.

Gujarat Themis Biosyn Limited, Vapi, (Guj).

Sept. - 2003 to Aug.-2004

Role: Worked as Officer (Production).

Responsibilities:

- ❖ Independent handling of shift duties.
- ❖ Troubleshooting process problems and maintaining parameters;
- ❖ Reporting to executive Operations.
- ❖ Ensuring smooth operation to maximize production and to meet preset targets.
- ❖ Maintaining good housekeeping.
- ❖ Implementing operation policies lead down by mgmt.
- ❖ Cost reduction in one of the products by use of recovered solvent.

Skill Set:-

- ❖ MS Office
- ❖ Well versed with SAP.

Personal Details:-

- ❖ Date Of Birth : 15 April1980
- ❖ Father's Name : Mr. P.D. Tamkhane
- ❖ Marital Status : Married
- ❖ Nationality : Indian
- ❖ Languages Known : English, Hindi, Marathi
- ❖ Passport no. : M5239677.

Declaration:-

I hereby declare that all the information provided by me in this application is factual and correct to the best of my knowledge and belief.