Mr. Kashiprasad Ramdayal Pardeshi

CV for Post of Chairman / Managing Director / Technical Director / Executive Director / Associate Director / CEO / President / GM / Plant Head / CQA Head / DQA Head / QA Head / RA Head / Lab Head / R&D Head / Project Head / Business Head / HR-Personnel-Admin Head etc.

Current Address:
17/4, Plot: 06, Tulip CHS, Bhawani Nagar Society, Marol, Andheri East, Mumbai, Maharashtra, India-400059.

Permanent Address:
At Post Lamkani, Tal. Dist. Dhule, Maharashtra, India-424307.

krpjbcpl@gmail.com
krpardeshi@zoho.com

9594640045 / 8369219571

linkedin.com/in/kashiprasad-pardeshi-095b79125/

PROFILE

- Corporate Quality Assurance
- Quality Assurance
- Regulatory Affairs
- Research & Development
- Projects
- Distribution & Supply Chain
- Contract Manufacturing
- Environment Health & Safety
- C&F/Logistics/4PL/3PL/2PL
- Business Development
- Consulting Services

EXPOSURE

- Pharma
- Biotech
- Cosmetics
- Nutraceuticals
- Ayush
- FMCG
- Medical Devices
- Diagnostics Kits
- Clinical Research
- Public Testing Laboratory

TOTAL WORK EXPERIENCE-21 YEARS

GENERAL MANAGER & PLANT HEAD (QUALITY ASSURANCE, REGULATORY AFFAIRS, R&D & PROJECTS)
TOWA PHARMA PVT. LTD., JALIYAMATH, DEHGAM, GANDHINAGAR, AHMEDABAD, GUJARAT — Oct 2024-Till Date

HEAD-PLANT, QUALITY ASSURANCE & REGULATORY AFFAIRS — Jun 2024-Sep 2024
SANJAR PHARMA LLP., DEDHROTA, HIMMATNAGAR, GUJARAT

HEAD-PLANT, QUALITY ASSURANCE & REGULATORY AFFAIRS — Jan 2024-Apr 2024
ELEGANT COSMED PVT. LTD., KUVADVA, RAJKOT, GUJARAT

SENIOR CONSULTANT
BOON MANAGEMENT CONSULTANTS PVT. LTD., THANE, MUMBAI — Oct 2023-Dec 2023

HEAD-LAB, QUALITY ASSURANCE, REGULATORY AFFAIRS & PROJECTS — Jul 2022-Aug 2023
JUBILANT PHARMA & CHEMICAL LAB. (OPC) PVT.LTD., NEW-PANVEL, NAVI-MUMBAI

MANAGER-QUALITY ASSURANCE & EHS (QEHS)
PATCHEMS PVT. LTD., BHIWANDI, MUMBAI — Apr 2022 - Jun 2022

EXECUTIVE-CORPORATE QUALITY ASSURANCE (QA/RA/R&D/PDD/SUPPLY CHAIN & DISTRIBUTION)
J.B.CHEMICALS & PHARMACEUTICALS LTD., WORLI-MUMBAI PRABHADEVI, THANE, BHIVANDI, DAMAN, PANOLI, ANKLESHWAR — Aug 2009 - Mar 2022

EXECUTIVE-CORPORATE QUALITY ASSURANCE & REGULATORY AFFAIRS — Jun 2008 - May 2009
SHARON BIO-MEDICINE LTD., VASHI, NAVI-MUMBAI

EXECUTIVE-REGULATORY AFFAIRS (QA/RA/R&D)
CIRON DRUGS & PHARMACEUTICALS PVT. LTD., BOISAR, TARAPUR — Dec 2006 - Jun 2008

OFFICER-QUALITY ASSURANCE — May 2006 - Nov 2006
BIOCHEM PHARMACEUTICAL INDUSTRIES LTD., DAMAN

PROJECT ASSOCIATE
DR. JAINS PVT. LTD., VASAI, MUMBAI — One Year

INDUSTRIAL TRAINING
SHREE SWAMI SAMARTH PVT. LTD., JALGOAN — One Year

EDUCATIONAL HISTORY

B.Pharm., M.Sc., M.Phil. MSC-IT

M.Phil. in Biotechnology/Bioinformatics — 2008 - 2009
TGOU (NAGALAND), INDIA — Grade: A/75%

M.Sc. in Biotechnology/Bioinformatics — 2004 - 2005
N.I.F.E.M (PUNE), SMU (KARNATAKA), INDIA. — Grade: A/76%

B.Pharm. — 2000 - 2003
COLLEGE OF PHARMACY SHAHADA, NMU (JALGOAN) — Grade: B/59%

H.S.C — 1998 - 1999
POOJYA SANE GURUJI VIDYA PRASARAK MANDAL SCHOOL LONKHEDA, SHAHADA, NASHIK BOARD, INDIA — Grade: A/70%

S.S.C — 1997
SHREE SATPUDA VIDYALAYA LONKHEDA, NASIK BOARD, INDIA — Grade: A/68%

CERTIFICATE

MSC-IT — 2003
MAHARASHTRA STATE CERTIFICATE IN INFORMATION & TECHNOLOGY, MITCON, PUNE, MAHARASHTRA, INDIA — Grade: A/62%

Management / Operational / Quality Skills:

- **Strong** Leadership
- Planning
- Scheduling
- Organizing
- Purchasing
- Controlling
- Automotive
- Corporate Social Responsibility
- Regulatory Compliance
- Project Management
- Quality / Assurance / Risk Management
- Inventory Management
- Material Management
- Training & Development
- Human Resource Management
- Daily Works Management
- Clean Workplace
- Occupational Safety
- Timely Delivery
- AMC (Maintenance, Calibration)
- Process Control
- Product Quality Safety
- Energy Management
- Environment Management
- Measurement & Analysis
- Supply Chain Management
- Business Development / Marketing / Sales Process Optimization
- Cost/Quote/BID/Agreement
- Innovative Strategies
- Productivity Enhancement
- Customer relationship management (CRM)
- Problem Solving
- Team Coordination
- Waste Management
- Technology Upgradation
- Idea Generation & Concept Development
- Trend Analysis
- Resource Conservation
- Internet of Things (IOT)
- Cloud Computing
- Artificial Intelligence (AI)
- LIMS, SAP, 5S, Six Sigma, Lean, Kaizen, 7 QC Tools etc.

ROLES & RESPONSIBILITIES: (PHARMA/BIOTECH/AYUSH/PUBLIC TESTING LABORATORY/MEDICAL DEVICES/DIAGNOSTIC KITS ETC.)

- Production Planning, Approval for Techno-Commercial Agreements, Tenders, Bids, Customers Orders, Purchase Orders, Work Orders, Quotations etc.
- Handling Audits, Inspections, Compliances for Local & Global GMP's.
- Attend Audits for new & periodic Registrations/licenses/renewals for Product Permissions.
- Implementation of Quality Management System (Change Control, Deviation, OOS, Incidence, RCA, CAPA, Market Complaint, Product Recall, Internal Audits, Self-Inspection, Vendor Audit, Pharmacovigilance, ADR etc.)
- R&D (Formulation/Method Development, Technology Transfer, Stability Studies, Packaging Development, IPQA)
- Responding to Customers/Clients Queries for Local & Global Regulatory, Loan, P2P, Contract etc.
- Approving of SOP's, SMF, VMP, Specification, Protocol, Report, Cleaning Validation, Data Integrity, GDP, Process Validation, Analytical Method Validation, Stability Studies, CSV, Quality Manual, Quality Risk Management.
- Coordination with Manufacturing Sites, RA, R&D, QA, QC, Purchase, Supply Chain, Distribution, C&F Agents, Marketing, Sales, Business Unit etc.
- Approving labelling for Artwork for Local & Global.
- Project works FAT, SAT, DQ, IQ, OQ, PQ, URS, UAT, etc.
- Master Vendor Audits with Planning, Monitoring.
- Information Management System (IMS) on Change Management & Projects Implementation Status.
- Pharmacopeial Change Management & Gap Analysis.
- Product Life Cycle Management, Risk Management.
- Approval for Project, Product File, CMC, CTD, Dossier.
- Exposure in Pharma Formulations (OSD-Tablets, Capsules, IV-Injectables-SVP, LVP, Lyophilize, ENT-Eye, Ear, Throat, Lozenges, Liquid Orals-Syrup, Suspension, Emulsion), Cosmetics/Topical/External/Personal Hygiene-Ointment, Cream, Gel, Paste, Soap, Diagnostic Kit, Medical Device, VET, AYUSH-Herbal, Powder, Nutraceutical, Aromatic oils, Flavors, Fragrances etc.
- Monitoring & update of National Projects gaps for (IPC, D&C Act, NDPS Act, DPCO, NPPA, PVPI, WHO-GMP, CDSCO, DCGI, DGHS, DCC, FSSAI, NABL, APEDA, MOFPI, BIS, Medical devices rules etc.).
- Monitoring and updating of International projects gaps for (USFDA, ANDA, UKMHRA, EUGMP, cGMP/GLP/GCP/GAMP/GXP, WHO, ICH, ISO, ISO-9001, 17025, 13485, 14001, 18001/OHSAS, 27001, FSMS, 21-CFR (Part-11, 210, 212, 820), EC/IEC/EU/EMA, TGA-Australia, SFDA-China, MHLW-Japan, HAS-Singapore, ANVISA-Brazil, INVIMA-Colombia, NAFDAC-Nigeria, MCC-South Africa, PIC'S, FDC, CIS, Row Markets, etc.)

PERSONAL INFO

- Name: Mr. Kashiprasad Ramdayal Pardeshi
- Nationality: Indian
- Domicile: Maharashtra
- Birthdate: 17/08/1981
- Gender: Male
- Marital Status: Single
- Languages: English, Hindi, Marathi, Gujarati
- Aadhar No.: 819889618013
- Pan No.: AMYPP5860H
- Pharmacist Reg. No.:73711
- Indian Passport No.: F6241233
- Educational Qualification: B.Pharm., Msc., M.phil.
- Certificate: MSC-IT

STANDING

- Total Experience: 21+ Years
- Current CTC: 12.00 LPA
- Expected CTC: 15-20 LPA (Negotiable)
- Notice Period: Immediate Joiner
- Relocate: Pan India / Overseas
- Job Location: Corporate / Factory / Warehouse / Travelling / WFH (Work from Home) / Part Time / Full Time / Freelance etc.
- PAN INDIA-Job Location Preference: Mumbai, Navi-Mumbai, Mumbai Suburban, Pune, Maharashtra, Palghar, Boisar, Tarapur, Daman, Vapi, Dadra, Nagar, Haveli, Gujarat, Goa, Rajasthan, Madhya Pradesh, Karnataka, Hyderabad, Delhi, Haryana, Uttrarakhand, Baddi, Himachal Pradesh etc.
- OVERSEAS-Job Location Preference: Canada, Australia, South Africa, USA, UK, Europe, Russia etc.

ROLES & RESPONSIBILITIES: (CFA/C&F/LOGISTICS/QHSE PROJECTS)

- SOP approval, implementation & compliance.
- CAPA assessment and its follow up.
- Communication with principal company-Commercial Quality & Business Quality related work.
- Handling the QA / EHS & Facility audits.
- Conducting Mock audit- twice in a year to all divisions and to monitor closure of audit observations.
- Validation activity (with Protocol/Report Approval etc.).
- Monitor WH cleanliness, maintenance and storage of records.
- Monitor Quality Compliance & following the requirements for temperature monitoring/mapping & recording devices.
- Auditing outsourcing services & Monitor AMC records.
- Monitor MRMs and closure of issues with given dates.
- Updating of Warehouse Internal / ISO audit related documents.
- Monitor disposal activity of expired & unsaleable stock.
- Recruitment of Employees, Housekeeping, Securities with Agreement and Liaisoning with agency.
- Monitoring & maintaining AMCs (ACs, WIF, DG, CCTV, Fire Alarm System, Goods Lift, Conveyor belts, Electric Motors, HVAC, AHU, Fire Extinguishers & Pest Control etc.) with service records.
- Monitor deviation and Change control.
- Monitor or organize quality awareness training for Quality related staff.
- Conduct MOCK audit prior to QA/ISO audit to minimize observations & risk.
- Attend FDA/Regulatory work related to new license/ Renewal or Competent Person (CP).
- QA / EHS audit observations received from respective divisions to be completed on or before due date time.
- Planning & Monitoring of Civil Projects like MEP (Mechanical, Electrical & Plumbing) for various locations.
- Implementation & Monitoring for Good Warehouse Practices (GWP), Good Distribution Practices (GDP), Good Supply Chain Practices (GSP), Quality Management System-QMS-ISO 9001, ISO 13485-Medical Devices, ISO-17025-Laboratory & Calibrations, EMS-ISO 14001-Environmental Management System, OHSAS-ISO 18001-Occupational Health & Safety, FSMS-ISO 22000-Food Safety Management System, ISMS-ISO 27001-Information Security Management Systems, IATF 16949-International Automotive Task Force etc.

DECLARATION: I HEREBY DECLARE THAT, THE ABOVE INFORMATION IS GENUINE TO MY KNOWLEDGE.

PLACE: MUMBAI
DATE:

YOUR'S SINCERELY;
Mr. KASHIPRASAD R. PARDESHI