PHARMQ MAVEN

Your Trust....Our Commitment....

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About Us

PHARMQ MAVEN
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PharmQ Maven is a trusted consultancy service provider of comprehensive solutions for the Pharmaceutical, Nutraceutical, Ayurvedic and Cosmetic Industries. Although established recently, our team brings over 15 years of industry experience in pharmaceutical infrastructure, compliance, and project execution. Under one roof, clients can avail all the services from designing to documentation and assurance to get the desired regulatory approvals.

Headquartered in Zirakpur (Tricity), Punjab.We cater primarily to North India as well as clients across the rest of India.

We have an experienced team of skilled professionals with extensive industry knowledge and practical expertise, enabling us to deliver reliable and effective solutions that support manufacturers in developing efficient, compliant, and high-quality facilities aligned with global regulatory standards.

Our approach focuses on innovation, precision, and regulatory compliance, ensuring excellence in every project we deliver and supporting our clients in all aspects according to their needs.

OUR CORE SERVICES

Documentation

It's a challenging task now a days for many organizations to prepare any document which could represent the actual practice and at the same time comply with the regulatory standards. Our Eminent Quality individuals are providing end to end support to our clients in case of documentation preparation and implementation as per the organization practice and policies.

Standard Operating Procedures	Process Validation Protocol and Report
Risk Assessment : New & Existing Facility, Product, Facility Contamination etc.	Cleaning Validation Protocol and Report and matrix
Hold time Validation Protocol and Report	Media Fill Validation Protocols and Reports
Site Master File	Validation Master Plan
Quality Manual	Onsite Emergency Plan
Business Continuity Plan	Analytical Method Validation & Transfer
MFR/BMR/BPR	Specifications

Commissioning , Qualification & Validation (CQV)

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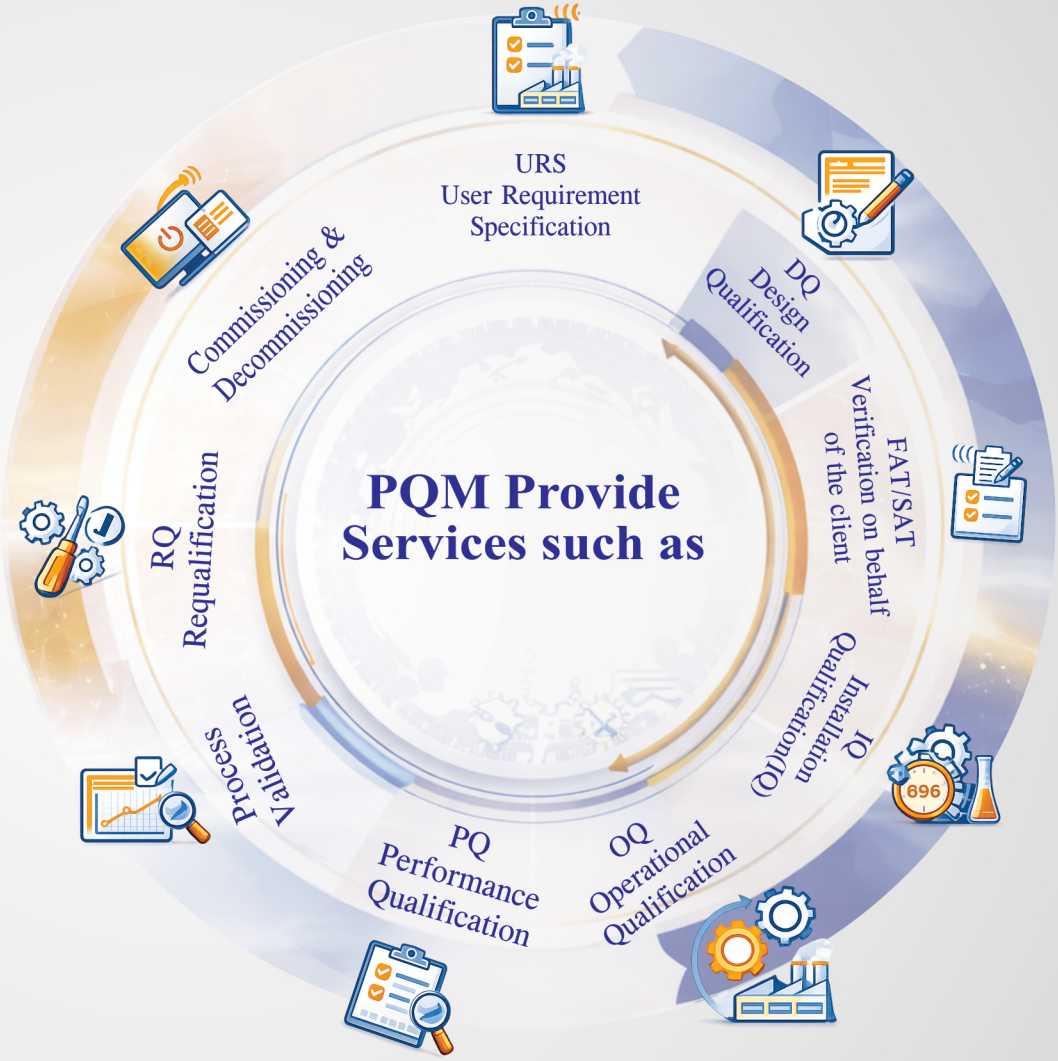
PQM provide end to end support from selection of Equipment/Instrument till it's ready for use and Ensure compliance with Schedule M & regulatory bodies standards.



OUR SERVICES

PROJECT CONSULTANCY

- Greenfield & Brownfield Project Execution
- Facility Design & Layout Optimization
- Equipment Selection & Vendor Finalization
- New Production Line

DOCUMENTATION

- Apex Documents
- SOP Development & Review
- CAPA Response
- Gap Analysis & Remediation Support

VALIDATION SERVICES

- Process Validation (PV)
- Cleaning Validation (CV)
- Engineering & Media Fill Batches
- Hold Time Studies (HTS)

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Cleaning Validation

Cleaning validation is a method that validates cleaning procedures. This methodology ensures the effectiveness and uniformity of a specific cleaning process in plants and equipment to remove residues and contaminants.

PQM provides end to end services for cleaning validation such as

Regulatory Affairs

PQM regulatory team prepares CTD dossiers, ACTD dossiers for pharmaceutical, Nutraceutical, veterinary & herbal products in the African, ASEAN, Middle East, CIS & LATAM markets i.e. ROW markets.

The rich industry experience of our team in every department including Quality Assurance, Quality control, production, packaging & regulatory affairs helps us understand the grassroot level problems in Dossier preparation or documentation & do the needful. Thus, we deliver practical & logical solutions to you.

PQM regulatory team can help you in all technical aspects of queries & documentation till your product is registered.

Traning & Technical Guidance

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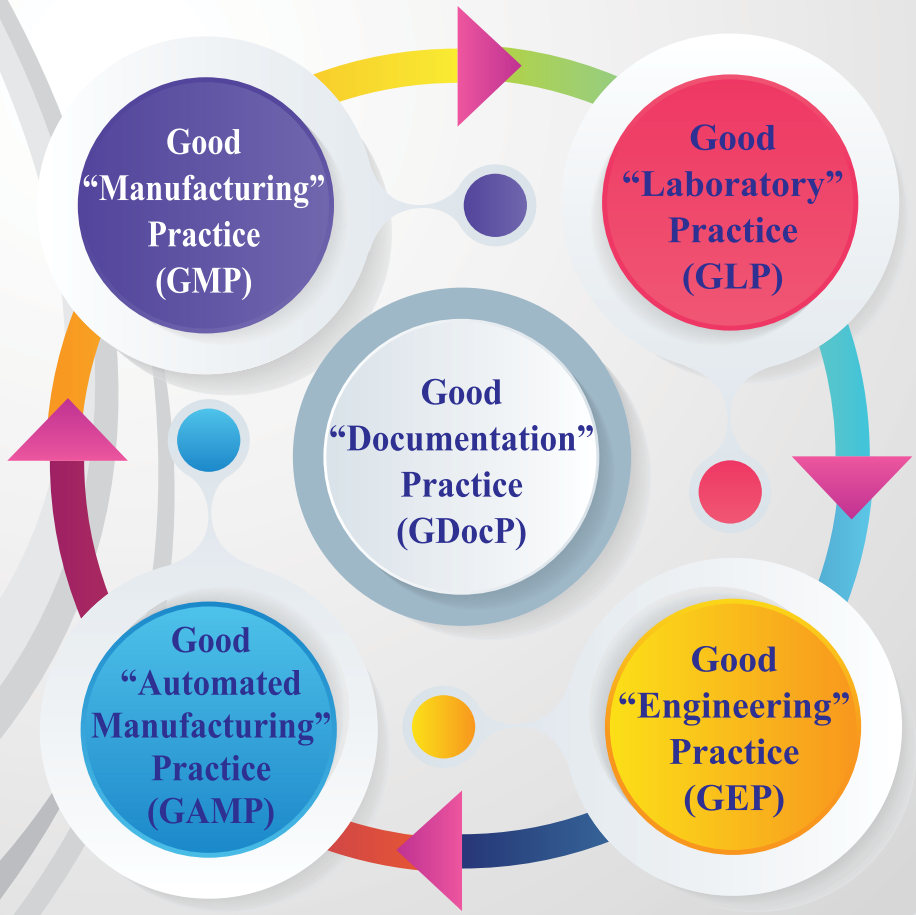
Training is the foundation that every organization wants to get it strong and effective. Organizations attempt to provide high-quality, meaningful training, a variety of obstacles often stand in the way, such as limited time from subject matter experts (SME's) or inability to access the proper training environments.

Data Integrity, QMS (Quality Management System) and GxP are the most vulnerable sides of the industry which need to be taken care of to uphold the organization practice and culture.

GxP is the general abbreviation for "good practice" quality standards and guidelines. The "x" represents the various areas it can be applied to. The term GxP is often used to refer to a collection of quality requirements.

There are 5 P's that we are focusing on as



PQM provide effective tailored training solutions to upkeep the Industry Practices and empower the workforce to do the right thing at all time and all situations, which can help organization to grow as intended.

Investigation Analysis

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Deviation, Incident, OOS, OOT etc. are integral part of the Manufacturing Industry which could occur anytime, anywhere intended or unintended. Investigation of those Non-Conformances are need to be performed accurately and precisely with logical reasoning and evidences and if not then it could lead to the critical observations during any regulatory and non-regulatory inspections and also impact the product integrity.

PQM expert QMS Team can provide or guide you to an effective and concrete Investigation Analysis report based on scientific evidences and back up data which could withstand all the queries from any regulatory and non-regulatory inspections.

Audit & Compliance

Our PQM expert auditors help you to ensure that you comply with international & Domestic regulations & standards and stringent market requirements recognised in the pharmaceutical industry in order to bring safe drugs to the market.

Below are the types of audits PQM offered

Turnkey Projects

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PQM is associated with trusted and reliable service providers for cleanroom partitions and equipment, HVAC, water systems, epoxy, electrical works, ETP/ STP and process equipment manufacturers (OEMs), ensuring end-to-end excellence in turnkey project execution.

PQM and Our Strategic Partner provides below services in affordable way such as

01 Detailed Project Report

02 Master Planning

03 Plant Layout Design

04 Architectural, Civil & Structural Layout

05 Cleanroom Design

06 HVAC Design

07 Electrical and Lighting Engineering

08 Plant Utilities Design

& Customizable Services

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