



THRIVE IT QUALIFY IT



Qualify IT Solutions a Startup company in engineering & design. Having successfully executed CSV & CQV projects in Infra, Pharma, Biotechnology and software industries. Qualify IT Solutions provides expert delivery of world class Pharma Facilities. Our team of CSV & CQV engineers is skilled at taking your project from concept to operation, providing local support with global subject matter expertise. Qualify IT solutions will deliver a cost-effective design and to meet regulatory requirement.

Qualify IT solutions provides services for the validation of the process equipment's, utilities, control systems and the facility determined to be GXP and Non-GXP system used in pharmaceutical manufacturing facilities.

COMPUTER SYSTEM VALIDATION

- Upgradation of Legacy Systems
- Automation Support
- PLC and SCADA Validation as per GAMP Requirements
- Application Software Validation as per GAMP and EU ANX 11 and 21 CFR Part 11
- EMS/BMS Qualification
- Excel Sheet Validation
- Audit Trail Review Assessments
- Periodic Review Assessments
- ERES Assessments
- GAP Assessments for All types Computerized Systems

Computer System Validations



COMMISSIONING & QUALIFICATION



Concept & Designing for Pharmaceutical Green field & Brown Field projects

- Process Engineering
- Procurement Management
- Commissioning & Qualification
- HVAC Design Support
- Process Equipment's Automation Support
- Facility, Black Utility, Clean Utility, Process Equipment's P&ID Concept review
- CIP/SIP Studies Designs
- AMC Services for All type of Process Equipment's (Vial Filling Line, BFS, Ampule Line, Tube Filling Line and water system)



THERMAL VALIDATION

- Steam Sterilizers
- Depyrogenation tunnel
- Dry Heat Sterilizers
- Mixing Vessels
- BOD Incubators
- Cold chambers
- Refrigerators
- Clean Rooms Mapping
- Thermal mapping of Warehouse, Cold Room & Retention Room
- Lyophilizer Validation
- Ultra-low Temperature Equipment and cold chain services



Regulatory & Customer Audit Support & Trainings provided by professionals

- GMP Training Program
- Data Integrity
- Good Documentation Practices
- Quality Management Systems
- Audit Behavioural Training
- Soft skills Training
- Safety Training
- CSV Training Program
- Risk Assessment
- Digital Training Module





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