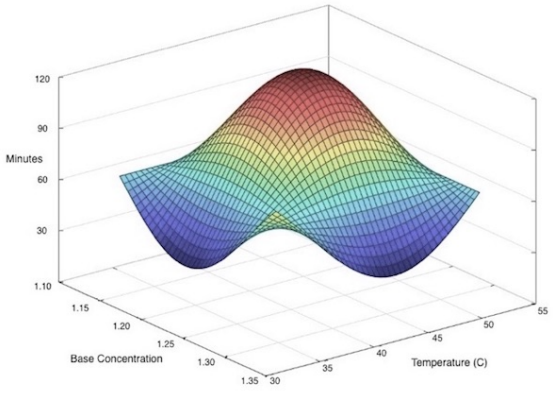
JARVIS PHARMACHEM

Welcome to Jarvis PharmaChem Consulting, a sole proprietor limited liability company, headquartered in Selma, NC, that is committed to optimizing pharmaceutical, chemical and food companies' processes and procedures to maintain compliance with today's regulatory standards.

Jarvis PharmaChem offers a wide range of services including equipment qualification, process/cleaning validation, process safety management, EHS auditing, business and chemical process optimization/ troubleshooting, GMP remediation/improvement, deviation investigation/CAPA Implementation, and batch record standardization.

Our goal is to improve your company systems and processes to allow for safety, efficiency, regulatory compliance and cost savings. Here at Jarvis PharmaChem Consulting, we don't reinvent the wheel. Instead, we help reduce friction to keep companies moving forward.

Optimization is defined as, "the action of making the best or most effective use of a situation or resource." Jarvis Pharmachem Consulting works with companies and its employees to ensure cohesion between company processes and best practices of the industry thus improving performance and quality. We review your company's critical processes and operation parameters to solve existing manufacturing or service problems and remove the defects and variations.



In manufacturing, We employ Design of Experiments (DOE) or cause -and-effect relationships to identify existing issues. DOE offers a practical approach for exploring multi-factor analysis. This requires the use if statistical tools and experimentation concepts to establish validity, reliability and replicability.

- Process Optimization/ Troubleshooting
- Process Safety / EHS Auditing
- GMP Remediation / Improvement
- Equipment Qualification
- Deviation Investigation / CAPA Implementation
- Batch Record Generation
- Process / Cleaning Validation