GPS CAPABILITIES PROFILE

validation

Your local, independent validation team

Computer, Process, Master Planning, IQ, OQ, PQ

Validation Excellence
Gavin Pharmaceutical Services offers the finest validation services available world wide. We blend a practical, how–to–do–it–right approach with a drive for excellence to deliver solid compliance. With rich experience dating back to the pharmaceutical industry development of validation techniques, Gavin's people are devoted to excellence, dedicated to performance, and skilled in making your validation project a real success.

Our Validation Team
Gavin Pharmaceutical Services employs scientists and engineers with academic training, validation training, cGMP training and experience to bring success to your project. We have expertise in Validation Master Plans, computer system validation, water systems, API's, liquids, solid dose, sterile products and more. If you require total project organization and management, or just need a little help, we are ready to assist your team.

Your Independent Contractor
Many of our competitors offer "integration" of validation with other critical services such as engineering design, construction, construction management, commissioning. We offer independent validation services. Independence is not required by FDA, but by common sense and good business practice. It means that we represent your interests, not our bottom line.

Your Local Validation Contractor
Tired of paying expensive airfare, fancy hotel, and rental car bills? Try our local service. That means no bills for the travel and living expenses of our people. In some cases, we may move to the project site, or we may travel. This way, you are not limited to only local contractors.

At a Glance

Service: *General Validation*
Compliance: *21CFR Part 211*
Capability: *Full Capability*
Best at: *High Quality Validation*
Project Scope: *Full or Partial*
Contact: *Blair Conley*
Sales: **1-800-700-5147**

Standard Operating Procedures
We write standard operating procedures to control equipment and processes without making compliance a nightmare. We are experts in writing SOP documents for compliance with cGMP.

Installation Qualification (IQ)
Our approach to IQ is comprehensive. One capability which distinguishes GPS is equipment design review. Our experience combines with GMP and analytical skills to evaluate the details of critical processing equipment.

Operational Qualification (OQ)
Our approach to OQ is practical. We cover the critical, process related operational issues with well planned and executed OQ studies. We like to present the results graphically for ease of interpretation and FDA review.

Performance Qualification (PQ)
We design meaningful, value added studies to prove robust performance of equipment and processes. Let us make your project a total success.

Act Today
Call today to discuss your project!

1-800-700-5147