GxP Industry Services

Specializing in FDA Regulatory Matters



EAS Consulting Group provides a variety of domestic and international regulatory services to industries under FDA oversight including in-depth expertise and guidance in the areas of Good Manufacturing Practices (GMPs), Good Laboratory Practices (GLPs), Good Clinical Practices (GCPs), and Good Documentation Practices (GDPs) and compliance with the Food Safety Modernization Act (FSMA) for all FDA regulated industries.

Our seamless approach helps organizations navigate compliance with quality management and system development. Whether your firm is involved with the manufacturing, packaging, distribution of products; ingredients and additives; or is a testing or research laboratory facility; EAS is committed to helping you understand, meet, and sustain compliance with the FDA's numerous technical requirements.

Choose EAS for:

- ✓ GMPs, GLPs, GCPs, GDPs, FSMA
- ✓ Quality System Development
- ✓ Corrective and Preventative Action (CAPA) Plans and Implementation
- ✓ Gap Analysis
- ✓ SOP Writing and Training
- ✓ Quality Agreement Preparation
- ✓ Managing FDA Inspections
- ✓ Investigatory Systems
- ✓ Laboratory Operations:
 - » Method Development, Validation and Transfer
 - » Specification Development
 - » Stability Program Development
 - » Out of Specification (OOS) Procedures
- ✓ Regulatory Remediation
 - » 483 and Warning Letter Responses
 - » Consent Decree Remediation
 - » Meetings with FDA
- ✓ Expert Witness Services
- ✓ Seminars and Onsite Training



