Virtual Consulting Services

Specializing in FDA Regulatory Matters

EAS Consulting Group

A Certified Group Company

EAS offers the unique capability of leveraging our expansive cGMP auditing and mock-FDA inspection expertise to support our clients with a virtual audit option that is tailored to fit your needs. Our detailed suite of virtual consulting techniques is utilized to assess and provide feedback on a firm's current operational programs benchmarked against applicable regulatory requirements and industry "Best Practices."

These virtual EAS services cover assessments of Good Manufacturing Practices (cGMPs); environmental, facility, equipment and worker hygiene; laboratory operations and compliance; recall, traceability and food defense utilizing your written documents and records as well as available video technology to provide an interactive experience. Through virtual auditing, EAS delivers an effective level of feedback, gap identification and recommendations from our expert consulting team on identifying and implementing effective corrective actions.

From extensive document reviews, staff interviews with key personnel, to a remote/virtual walk-through of your facility using video and conferencing technologies, EAS can provide your company with valuable feedback and actionable CAPAs to strengthen your company's compliance with applicable FDA, State and International regulations such as cGMPs.

Additionally, EAS offers many other regulatory compliance services remotely. This includes FDA, USDA and EPA registration and submission services, training, building FSVP written program and the review of product labels. Regardless of your needs, trust EAS to provide the level of service and expertise you need from your regulatory partner.

When in-person consulting is not practical, utilizing EAS's virtual options provide a ready alternative. Contact EAS to learn more about our client regulatory partnerships.

EAS Virtual Consulting Services Include:

Compliance Training

- ✓ GMP/GLP/GCP/GDP
- ✓ Managing Virtual and In-Person FDA Inspections
- ✓ Investigatory Systems
- ✓ FDA and USDA Labeling & Ingredient Compliance
- ✓ OSHA Worker's Safety
- ✓ Food Safety Modernization Act (FSMA) Regulations
- ✓ Laboratory Operations
- ✓ Facility, Equipment and Staff Environmental Hygiene
- ✓ Food Defense/Intentional Adulteration
- Imports & Country Specific Exports

Virtual Audits

- ✓ Audits Utilizing Focused Remote Facility Site Tours or Pre-recorded Video Reviews of Processes and Equipment
- ✓ Remote Operational Document and Supporting Record Reviews,
- ✓ OSHA Compliance Requirements
- ✓ FDA & USDA Compliance Mock Audits
- ✓ Process and Processing Equipment Assessments
- ✓ Corrective Action and Regulatory Remediation
- ✓ Drug Development CMC Reviews
- ✓ Toxicology Assessments GRAS, NDI Submissions
- ✓ Expert Opinions for Legal Matters

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