Pharmaceutical Services

Specializing in FDA Regulatory Matters

EAS Consulting Group

A Certified Group Company

EAS Consulting Group provides a variety of FDA-based regulatory services to the global Rx and OTC pharmaceutical industry. We offer a seamless approach to help your organization navigate the regulatory intricacies associated with product development, submissions, claims and labeling, manufacturing and packaging of commercial products, as well as facility and product registrations and listings.

Our team of pharmaceutical regulatory experts provide assistance with quality management system development and strategies, GMP/GLP/GCP assessments, Mock-FDA inspections, regulatory submissions, and more, all to help your organization succeed in this complex and competitive regulatory environment.

Employing a unique team of former FDA officials and industry experts, many with more than 30 years of experience, EAS has unparalleled expertise to assist clients with all of their consulting needs.

EAS is committed to helping the domestic and international pharmaceutical market understand, meet and sustain compliance with the FDA's numerous requirements in a manner that is respectful of your time and business drivers.

- ✓ Regulatory Strategy Development
- ✓ DMF, NDA, ANDA, NADA
- ✓ Regulatory Compliance Support
- ✓ Auditing GMPs/Quality Systems
- ✓ Labeling and Claims
- ✓ Cosmetic to OTC Switch
- ✓ Homeopathic
- ✓ Allopathic
- ✓ Quality Management System Support
- ✓ Registration and Drug Product Listing
- ✓ US Agent Services
- ✓ FDA Detention Assistance
- ✓ Seminars and Onsite Trainings

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