US Agent and Imports

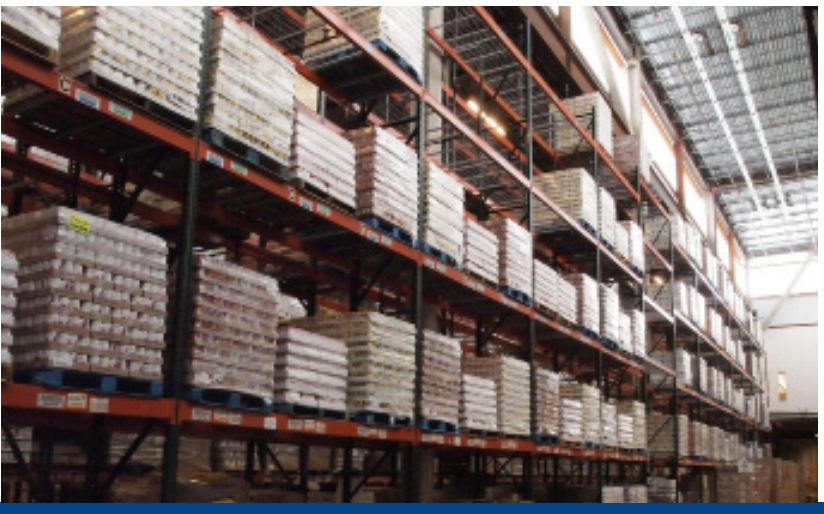
Specializing in FDA Regulatory Matters

EAS Consulting Group

A Certified Group Company

EAS Consulting Group provides registration and listing assistance as well as US Agent services to international clients in all FDA and USDA regulated areas. US Agents are required by FDA to act as an intermediary between foreign firms and FDA. By quickly responding to questions and concerns in a timely manner, EAS helps clients get their compliant products to the US marketplace as quickly as possible.

EAS is a leading provider of regulatory services to FDA and USDA regulated industries. We have over 50 years of experience assisting clients with developing regulatory strategies, implementing quality systems, filing regulatory submissions, and ensuring compliance with all FDA and USDA regulations. 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.



- ✓ FDA Import Requirements
- ✓ Establishment Registrations (Drug, Medical Device, Food, and Cosmetics)
- ✓ Drug Submissions (IND, DMF, ANDA, Amendments)
- ✓ Facilitation of eCTD submissions
- ✓ Medical Device Applications and Correspondence (510(k), 513(g), IDEs, De Novo, Pre-Submission)
- ✓ Process Filing for Acidified and Low-Acid Canned Foods (LACF)
- ✓ Foreign Color Manufacturers Batch Certifications
- ✓ Cosmetic product listings
- ✓ Preparation for FDA Inspections of Foreign Facilities
- ✓ Corrective Action Plans
- ✓ Assistance with Product Detention
- ✓ Seminars and Onsite Training

EAS Consulting Group
easconsultinggroup.com
(571) 447-5500