Tobacco Services

Specializing in FDA Regulatory Matters

EAS Consulting Group

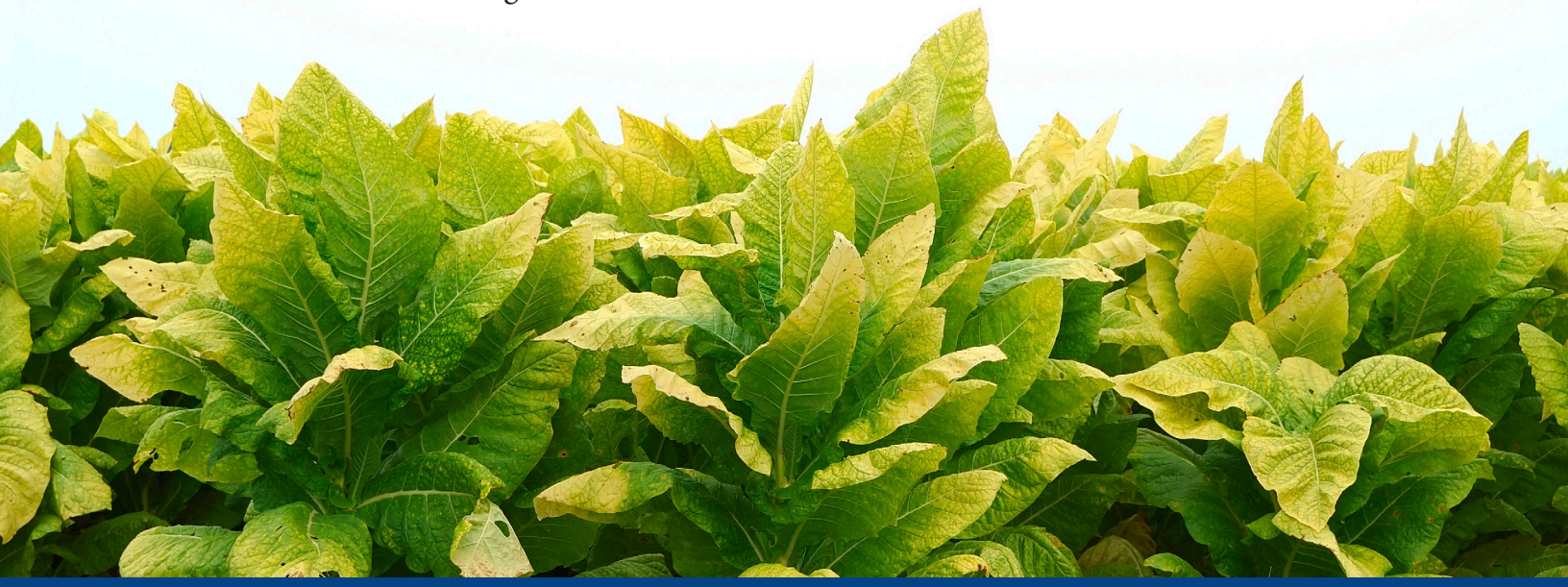
A Certified Company

EAS Consulting Group, LLC is a leading provider of regulatory consulting services to the tobacco industry, including newly deemed products such as ENDS products, cigars and pipe tobacco. The firm has extensive experience assisting clients in implementing systems and preparing for compliance with the FDA tobacco TPMPs, the Family Smoking Prevention and Tobacco Control Act and the May 2016 Deeming Rule and Regulations.

Originally founded in 1960, EAS has over 50 years of experience assisting clients in developing regulatory strategies, implementing quality systems, filing regulatory submissions, and ensuring compliance with applicable regulations. Employing a unique team of former FDA, USDA and state agencies 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.

Choose EAS For:

- ✓ Regulatory Consulting
  - » Preparing for Tobacco Product Manufacturing Practices (TPMP)
  - » Compliance with Deeming Rule and its requirements
  - » Implementing FDA Quality Systems
- ✓ Registration and Listing Assistance
- ✓ Mock Audits
- ✓ Onsite Training



Employing a unique team of former Food and Drug Administration (FDA) officials and industry experts, EAS offers unparalleled expertise with many consultants having more than 30 years regulatory experience and quality compliance.

EAS Consulting Group LLC provides tobacco consultation services and assistance with registrations and listings of tobacco products, TPMP compliance, manufacturing site audits and regulatory compliance training.

Regulatory Compliance

EAS offers assistance with the development of quality systems, compliance assessments of manufacturing facilities, filing of regulatory submissions and more to ensure your facility, product and labeling all meet FDA's stringent requirements for the tobacco industry. Whether ENDS products, cigars, pipe tobacco or c-cigarettes, cigars, certain dissolvables that are not "smokeless tobacco," gels, and waterpipe tobacco, EAS Independent Consultants offer the expert answers you need no matter the regulatory question.

EAS Independent Consultants are well versed in the details of The Family Smoking Prevention and Tobacco Control Act, supplemental FDA guidance and the May 2016 Deeming Rule. We monitor FDA tobacco activities and regulation development, keeping clients informed of the latest updates and thorough assessments of FDA guidance.

Registrations and Listings

EAS provides assistance with facility registration, product listings, and ingredient listings.

Mock Audits & Compliance Assessments

EAS Independent Consultants provide "compliance assessment" and "mock FDA-inspection" services to both large and small tobacco and e-cig/e-liquid companies, providing detailed analysis and reports on how your systems and procedures align with current FDA expectations and what corrective actions and enhancements you may want to consider implementing in preparation for future TPMP requirements.

Onsite Trainings

EAS takes pride in our ability to conduct training courses tailored to meet the unique needs of your organization. Our trainers, former high level FDA officials and industry executives, travel to your facility, providing customized training and a thorough analysis and review of current industry regulations. Students who have attended our programs have found EAS trainings to be unique in that that they provide an opportunity to ask former regulators questions that simply cannot be addressed by other training organizations.

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