Due Diligence and M&A Regulatory Support

Specializing in FDA Regulatory Matters



EAS is a leading consulting firm assisting with due diligence assessments and third-party audits of FDA and USDA regulated companies. We identify regulatory risks and compliance gaps and can determine whether appropriate steps and corrective actions have been taken. Our assessments offer an unbiased and clear picture of regulatory risk.

EAS provides a rapid and flexible response to short notice due diligence inqueries with customizable services including an overall assessment of a firm's regulatory history, past FDA, client and ISO audits, quality systems, GMP compliance, documentation, complaints, adverse events, CAPAs, facility condition and staff training/expertise. Post assessment you will receive a detailed report of findings and summary of observations, including criticality, as well as insights into recommended potential corrective actions.

Our due diligence services are customizable to client needs and conducted to provide a high-level assessment of a firm's overall state of regulatory and quality compliance. It can include:

- ✓ Remote Assessment of site's Quality Management System (QMS) and Regulatory history.
- ✓ On-Site (1-3 days) Due Diligence Assessment.
- ✓ Detailed Report of Findings. Consisting of submitting a detailed report that summarizes our observations, the criticality of each observation and insight into recommended potential corrective actions

Through our strategic alignment with Certified Laboratories, we can also ensure your testing requirements and challenges are met through a robust scope of sophisticated testing services.

No matter your FDA or USDA regulated business, EAS and Certified Laboratories can assist you with a reliable, rapid, flexible, comprehensive, accurate and customizable portfolio of due diligence services.

Choose EAS for your due diligence assessments of:

- ✓ Human Food Facilities
- ✓ Animal Feed/Food Facilities
- ✓ Dietary Supplement Facilities

- ✓ Pharmaceutical Facilities
- ✓ Medical Device Facilities
- ✓ Labeling and Marketing Compliance