

# PRODUCT QUALITY MANUFACTURING & COMPLIANCE

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The FDA's continued emphasis on proactive quality management and compliance with current Good Manufacturing Practices (cGMP) presents new challenges and opportunities for the private sector, making it essential for companies to build quality, safety, and integrity into their products.

Greenleaf's in-depth knowledge and understanding of the U.S. Food and Drug Administration (FDA) provides clients with the technical and strategic support needed to align their approach with the FDA's quality, safety, and compliance expectations.

## STRATEGIC & TECHNICAL CAPABILITIES

Greenleaf's Product Quality, Manufacturing, and Compliance Team has a proven track record of achieving success and providing services that are recognized as best in class by companies seeking to strengthen their quality management systems.

Greenleaf professionals work as teams specializing in product quality, manufacturing, and compliance; medical devices and combination products; and drug and biological products. Greenleaf's comprehensive approach provides a full-service engagement that ensures clients can count on expert direction as they encounter complex regulatory challenges.

Greenleaf's compliance experts specialize in corporate quality and compliance systems; FDA inspections, compliance, and enforcement processes; FDA organization and structure as they relate to compliance functions and decisions; FDA communications, including enforcement letters and facility and supply chain audit reports; and the complete spectrum of compliance- and enforcement-related actions.

Experts from Greenleaf's Product Quality, Manufacturing, and Compliance Team also provide advisory services that include extensive research and due diligence for investors engaged in potential mergers and acquisitions that require regulatory risk analyses before and after decisions and transactions.



In today's dynamic regulatory environment, keeping pace is not enough. Success is achieved by staying one step ahead of the pack. That's easier said than done, which is why successful companies recognize the need for a trusted regulatory partner to help navigate the challenging FDA landscape.

## ABOUT GREENLEAF

Greenleaf Health is a leading FDA regulatory consulting firm that provides strategic and technical guidance to pharmaceutical, biotechnology, and medical device companies researching, developing, and manufacturing innovative solutions to pressing global public health challenges.

Greenleaf is committed to serving our clients' needs with extensive expertise, unwavering integrity, and strategic insight in a manner that supports availability of safe, effective, and high-quality drugs, biologics, and devices.

## UNMATCHED EXPERTISE

Greenleaf's team is comprised of experts with a combined total of more than 250 years of FDA experience. The firm includes former leaders and regulatory professionals from the FDA, Capitol Hill, top global pharmaceutical and medical device companies, leading law firms, and the top U.S. biotechnology trade organization.

With decades of experience in senior positions at the FDA and throughout industry, Greenleaf's team of respected professionals brings unmatched expertise that companies need when working directly with the FDA and when navigating today's evolving regulatory environment.

## COMPREHENSIVE SERVICES

The Greenleaf Product Quality, Manufacturing, and Compliance Team uses its combined depth of knowledge and breadth of skill to provide the following services:



### Compliance Assessments

Greenleaf works with life science entities to evaluate and strengthen compliance functions with an eye toward optimizing processes, mitigating risks, and creating a culture of compliance, while continuing to meet business objectives.



### Compliance Remediation

Greenleaf experts bring an unmatched level of credibility and trust when interacting with the FDA. The team has the experience and insight to successfully guide companies along the remediation pathway.



### Consultation, Training & Regulatory Guidance

Greenleaf partners with companies to deliver insight and guidance that helps achieve business and regulatory objectives. The team also develops and delivers training on a variety of regulatory subjects.



### Inspection Readiness

An FDA inspection is a noteworthy moment for any regulated company. Greenleaf experts strengthen clients' readiness for FDA preapproval and surveillance inspections through consulting, training, audits, and mock inspections.



### FDA Communications

Greenleaf's best-in-class experts bring value to any communication with the FDA, including formal regulatory communication, in-person meetings, and responses to compliance actions and regulatory correspondence.



### GCP Services

Greenleaf specialists use a risk-based methodology to develop and improve clinical quality systems. Because there is no one-size-fits-all approach to GCP compliance, each solution is built to fit the targeted needs of the client.

## LEADERSHIP

Members of the Greenleaf Product Quality, Manufacturing & Compliance Team include:



### JOHN TAYLOR

Principal,  
Compliance & Regulatory Affairs

Former FDA senior official held many high-profile positions at the Agency, as well as senior leadership roles within industry.



### CYNTHIA SCHNEDAR

Executive Vice President,  
Regulatory Compliance

25-year compliance career, including serving as director of the FDA's drug compliance office.



### DAVID ELDER

Executive Vice President,  
Regulatory Compliance

23-year veteran of the FDA with prominent roles in domestic and foreign inspections, recalls and emergencies, and compliance actions.



### KALAH AUCHINCLOSS

Senior Vice President,  
Regulatory Compliance

10+ years of experience on Capitol Hill, in the private sector, and at the FDA, including role as Deputy Chief of Staff.



### KRISTEN GRUMET

Senior Vice President,  
Regulatory Compliance

25-year compliance career, including role as an FDA field investigator specializing in medical devices.



### LIZ OESTREICH

Vice President,  
Regulatory Compliance

Diverse background and knowledge of legal, public policy, and non-profit sectors.

## ADDITIONAL TEAM MEMBERS

BRITTANY MILBY, MADELEINE GIAQUINTO, LAURA BARTEE