

Former FDA Investigator Launches Global FDA Inspection Masterclass Video for Pharma and Biotech Compliance Teams

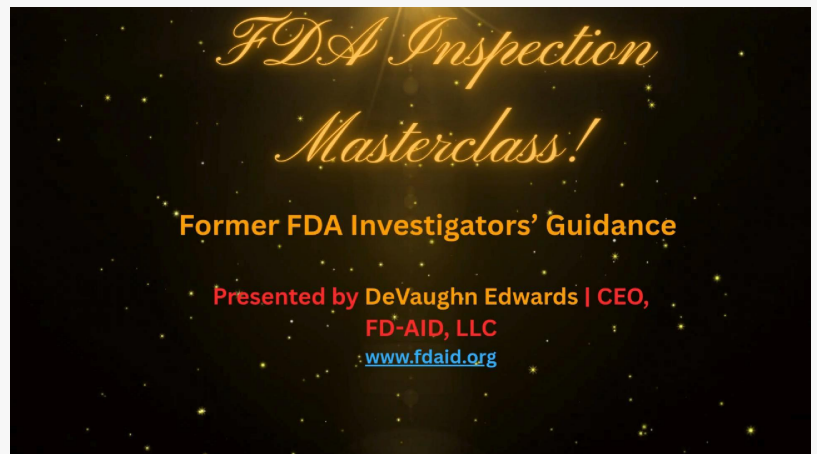
Global video course teaches inspection readiness strategies from former FDA Investigators—now available on-demand.

LUTZ, FL, UNITED STATES, May 13, 2025

/EINPresswire.com/ -- With FDA

inspections becoming increasingly rigorous and the announcement from the FDA that international international companies will no longer be given a heads-up of a pending inspection, a new video masterclass created by former FDA Investigators is now

available to help pharmaceutical and biotech companies worldwide better prepare for regulatory scrutiny.



FDA Inspection Master Class

[“FDA Inspection Masterclass: Former FDA Investigator’s Guidance”](#) is a comprehensive, on-demand training course developed by DeVaughn Edwards, Principal Consultant at FD-AID, LLC. Edwards, a former FDA Investigator with 14 years of field experience and over a decade in the pharmaceutical industry, collaborated with other regulatory experts to create an insider-driven guide to surviving — and succeeding — in FDA inspections.

“

“We created this masterclass because far too many companies wait until an FDA inspection is scheduled to prepare. That’s usually too late.”

DeVaughn Edwards, Principal Consultant, FD-AID, LLC

“This is the type of training I wish more companies had before we walked through the door,” said Edwards. “It covers what Investigators actually evaluate and how to avoid the most common and costly mistakes.”

The training includes eight modules covering:

- FDA inspection triggers

- PAI (Pre-Approval Inspection) strategy
- How to conduct internal mock inspections
- Interview techniques
- Real-world examples of Form 483 missteps
- Clinical and GMP-specific risks

Participants will receive:

- On-demand access to the full course
- Certificate of completion (optional)

An option for team-based licensing

With U.S. and global regulators increasing their oversight, especially of contract manufacturers and clinical operations, this video offers a critical opportunity to close compliance gaps before the FDA identifies them.

The course is now available internationally at:

<https://devaughnedwards.podia.com/fda-inspection-masterclass-former-fda-investigator-s-guidance>

About FD-AID, LLC

FD-AID, LLC is a pharmaceutical consulting firm led by former FDA Investigators and industry executives. The company provides mock FDA inspections, regulatory remediation, and GMP compliance training to clients around the world. More at www.fdaid.org.

Media Contact:

DeVaughn Edwards

Principal Consultant

FD-AID, LLC

devaughn@fdaid.org | +1 (732) 895-7831

5514 Garden Arbor Drive, Lutz, FL 33558

DeVaughn Edwards

FD-AID LLC

+ 17328957831

[email us here](#)

Visit us on social media:

[LinkedIn](#)

[Other](#)

This press release can be viewed online at: <https://www.einpresswire.com/article/811890875>

EIN Presswire's priority is source transparency. We do not allow opaque clients, and our editors try to be careful about weeding out false and misleading content. As a user, if you see something we have missed, please do bring it to our attention. Your help is welcome. EIN Presswire, Everyone's Internet News Presswire™, tries to define some of the boundaries that are reasonable in today's world. Please see our Editorial Guidelines for more information.

© 1995-2025 Newsmatics Inc. All Right Reserved.