



# PHARMACEUTICAL INDUSTRY TRAINING

FDA • QA • CSV • REGULATORY COMPLIANCE

Fee: \$299 + tax per day, per person (Limited seating available)

*Professional Training for Regulated Pharmaceutical Industry*



**Date: May 14, 2026**

**Date: May 15, 2026**

**Speaker: Myriam Sosa, MSPH**  
President – The Gintegra Group (TGG)

**Speaker: José E. Matos, MS CpE, MS CIS**  
Senior Consultant / IT Leader-The Gintegra Group (TGG)

**8:00 AM – 12:00 PM**

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**Latest Trends in FDA Enforcement Actions & Significant Updates**

- CSV fundamentals (GxP systems)
- FDA & data integrity expectations
- Computer Systems Validation (CSV) and Data Integrity (DI)
- Risk-based validation approach

**12:00 PM – 1:00 PM /Lunch break**

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**1:00 PM – 5:00 PM**

**1:00 PM - 5:00 PM**

**Pharma Deviations & Investigation Fundamentals**

**What Do You Need to Know?**

- FDA enforcement expectations
- Root cause analysis & CAPA
- Inspection-ready investigations

- Global Information System Qualification and Validation
- Third Party Vendor or Supplier audits

**TARGET AUDIENCE:** QA / QC Departments, Compliance Auditor, Regulatory Affairs, CSV & IT Validation Teams-Manufacturing & Engineering. Information Technology (IT), Automation Engineering, Project Managers, Equipment and Process Validation professionals. Computer Science and/or Computer Engineering students.

**Registration & information:** [www.thegintegragroup.com/training](http://www.thegintegragroup.com/training)

**Email:** [D'Eduka@thegintegragroup.com](mailto:D'Eduka@thegintegragroup.com)

Limited seating available