



2010 DEA Pharmaceutical Training Seminars

Tentative Agenda *

Day 1

8:00 – 8:30	Registration
8:30 – 9:00	Welcoming Remarks and Introduction to the Seminars
9:00 – 9:45	Registration of Manufacturers and Importers (Section 303 Investigations)
9:45 – 10:00	Break
10:00 – 11:30	Inventories, Records, and Reports
11:30 – 1:00	Lunch
1:00 – 1:45	Quotas
1:45 – 2:15	Research vs. Manufacturing
2:15 – 3:00	Quota Applications
3:00 – 3:15	Break
3:15 – 4:00	Quota Applications (con't.)

Day 2

8:30 – 9:30	Import/Export
9:30 – 9:45	Break
9:45 – 10:45	Import/Export (con't.)
10:45 – 11:45	Diversion Investigator
11:45 – 1:00	Lunch
1:00 – 2:30	Year-End Report Generator
2:30 – 2:45	Break
2:45 – 4:45	ARCOS

2010 DEA Pharmaceutical Training Seminars

San Diego, California – April 13-16, 2010

Denver, Colorado – May 18-21, 2010

Introduction to the Pharmaceutical Training Seminars

- What is the Single Convention and the Psychotropic Convention
- Domestic Legislation in Relation to International Treaties
- Overview of Industry Responsibilities with Regard to International Treaties and Domestic Legislation
- DEA's utilization of information gathered from registrants for purposes of its reporting responsibilities to the United Nations

Registration of Manufacturers and Importers of Schedule Is and IIs

- What is a "303-investigation" and why is it performed
- How does DEA utilize additional questions in evaluating a manufacturer or import registration application

Inventories, Records, and Reports

- Role of the Diversion Control Program
- Registrant inventories, records and report requirements
- How industry can assist DEA to prevent diversion

Quotas

- Legislative History
- Combat Methamphetamine Epidemic Act
- Types and Purpose of Quotas
- Information and Procedure Used to Establish Quotas

Research vs. Manufacturing

- Registration Coincident Activities
- Research vs. Manufacturing Activities
- Historical Determinations

Quota Applications

- Explanation of Quota Applications
- Quotas Adjustments
- Authorized Destructions

Import/Export

- Legislation and General Information
- Explanation of the DEA-161, DEA-357 and DEA-236
- Transshipments
- Filing of Return Information

Diversion Investigator

- How DEA Approaches and Completes a Registrant Inspection

Year-End Report Generator

- Explanation of Year-end Reports
- YERs On line
- Common Problems Encountered

ARCOS

- What and How Manufacturers Report to ARCOS
- An Explanation of the ARCOS Manufacturing Codes
- Current Reporting Method and Future Reporting Requirements
- An Explanation of the ARCOS Edit Error Codes and How to Correct ARCOS Reports