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Webinar on

What You Should Know About Export Requirements For FDA Regulated Products From The US To Foreign Countries

Areas Covered

- *Laws and Regulations, FDA Guidance, Definitions*
- *FDA Process for Issuing Export Certificates, Enforcement Actions: Case Studies*
- *Export and Documentation Requirements, PASS-IT Recommendations: Best Practices*
- *Important Information for Exporting FDA Regulated Products: Food, Cosmetics, Animal Drugs, Drugs, and Medical Devices*
- *Regulatory Requirements Exported from the US,Prior Notice Process*
- *How to Deal with Refusal to Issue Export Certificates*
- *Common Misunderstanding and Key Considerations*

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This training will provide an in-depth explanation of FDA-Regulated Products that were exported from the United States to Foreign Countries.

PRESENTED BY:

Dr. Gerald is an Academic Professor and Founder of RVG International Consulting Firm, LLC. He has over twenty-five years of business experience in, strategic management, marketing analysis, and supply chain management. He also earned a Doctor of Business Administration in International Business and Advanced Professional Business Certificate in Marketing from Argosy University College of Business.

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On-Demand Webinar

Duration : 60 Minutes

Price: $200

Webinar Description

This training will provide an in-depth explanation of FDA-Regulated Products that were exported from the United States to Foreign Countries. Also, it will address what export documentation must be used in the transportation of FDA-regulated products. Then, it also will discuss the FDA procedures that exporter must comply to in order to meet the Modernization Act agreement. Lastly, it explains what local government and private resources that will have an impact on the exporter's transaction process when it comes tariff costs in an FDA trading environment.

Who Should Attend ?

Regulatory Affairs

Quality Professionals

Product Development Professionals

CROs

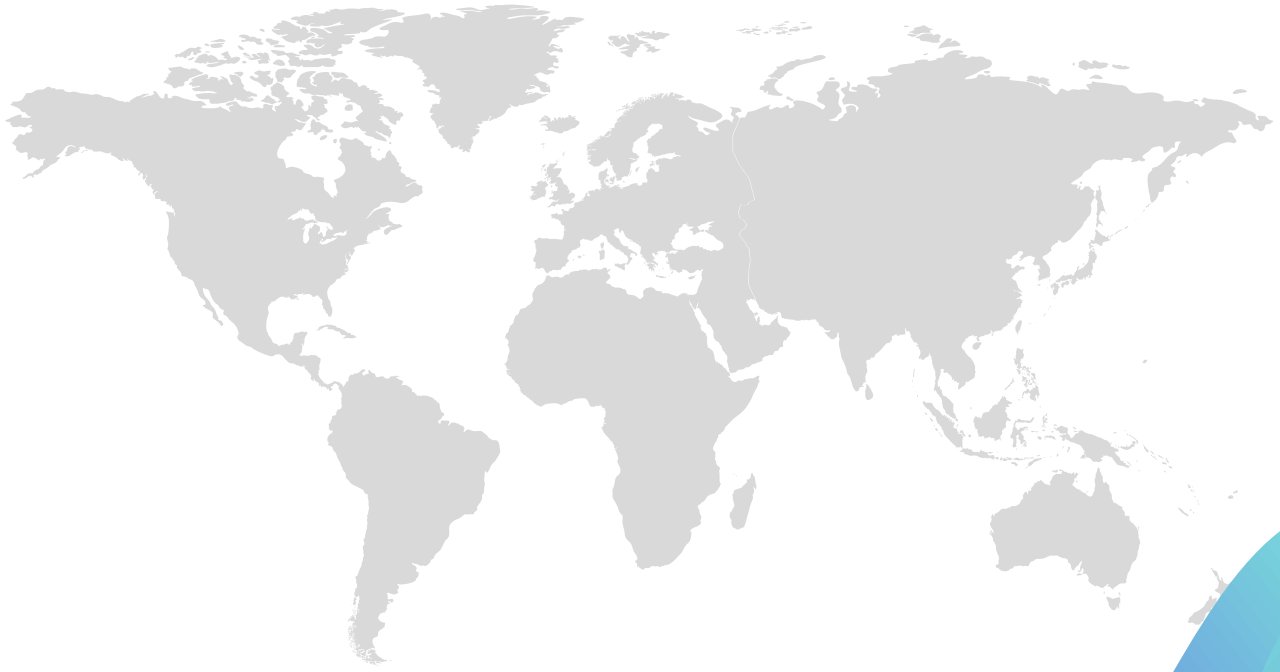
Consultants

Senior Management

Contractors and Subcontractors

Anyone interested in the subject

R&D Scientists, Engineers, Managers, and Directors



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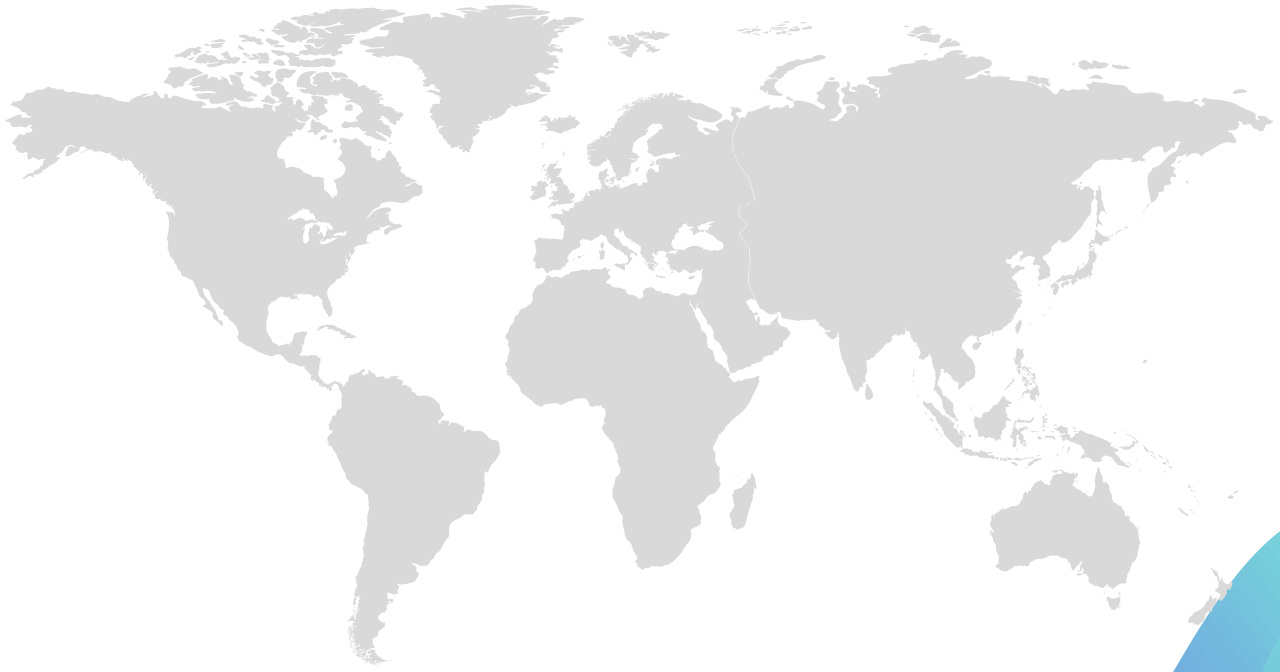
Why Should Attend ?

As exporters, when exporting outside of U.S., then they must consider the Food and Drug Administration (FDA) provisions because it helps them to understand what documents and procedures are needed in coordination transaction process. The Food and Drug Administration (FDA) measures will spell out the transportation documentation that is required for exporting to a foreign port. This Food and Drug Administration (FDA) has various multipart that contains compliance documentation and procedures that are necessary for this international business in order to facilitate the entry of goods.



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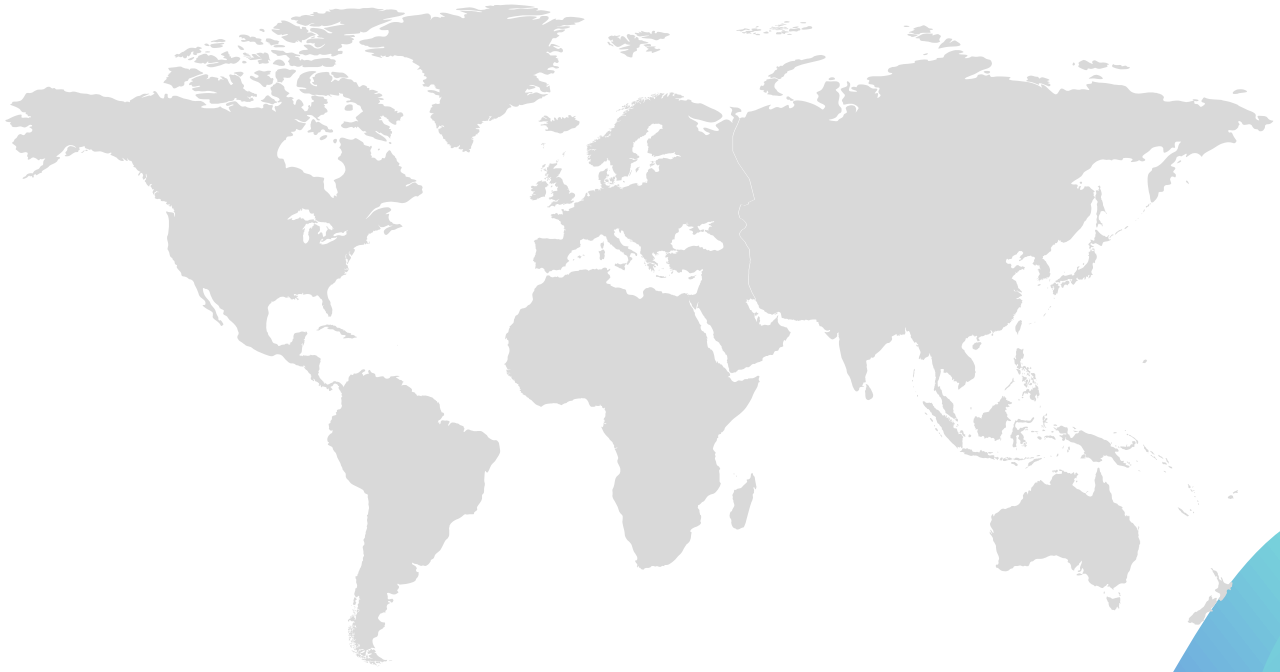
Exporters, we must understand what their exporting roles to determine the export certificate for FDA-regulated products are. Also, exporters must address what is their responsibility when it comes to distribution and selling of their goods. Next, exporter needs to know the determining factors for the origination of goods under the FDA Regulations. For this reason, exporters must be aware of their roles when it comes to generating the documentation and what procedures that must be followed to meet the duty compliance processes, which could lead to legal challenges and delay of the exportation process in this marketplace.



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