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Webinar on

Answering and Replying to an FDA 483

Areas Covered

- *GMP Compliance*
- *Regulatory requirements*
- *History*
- *Application*
- *Do’s and don’ts*

If you are new to the topic or need to brush up on the expectations or how to react to 483s consider attending.

PRESENTED BY:

Danielle DeLucy, MS, is the owner of ASA Training and Consulting, LLC which provides Pharmaceutical and Biologics based companies with training and quality systems assistance in order to meet regulatory compliance.

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On-Demand Webinar

Duration : 60 Minutes

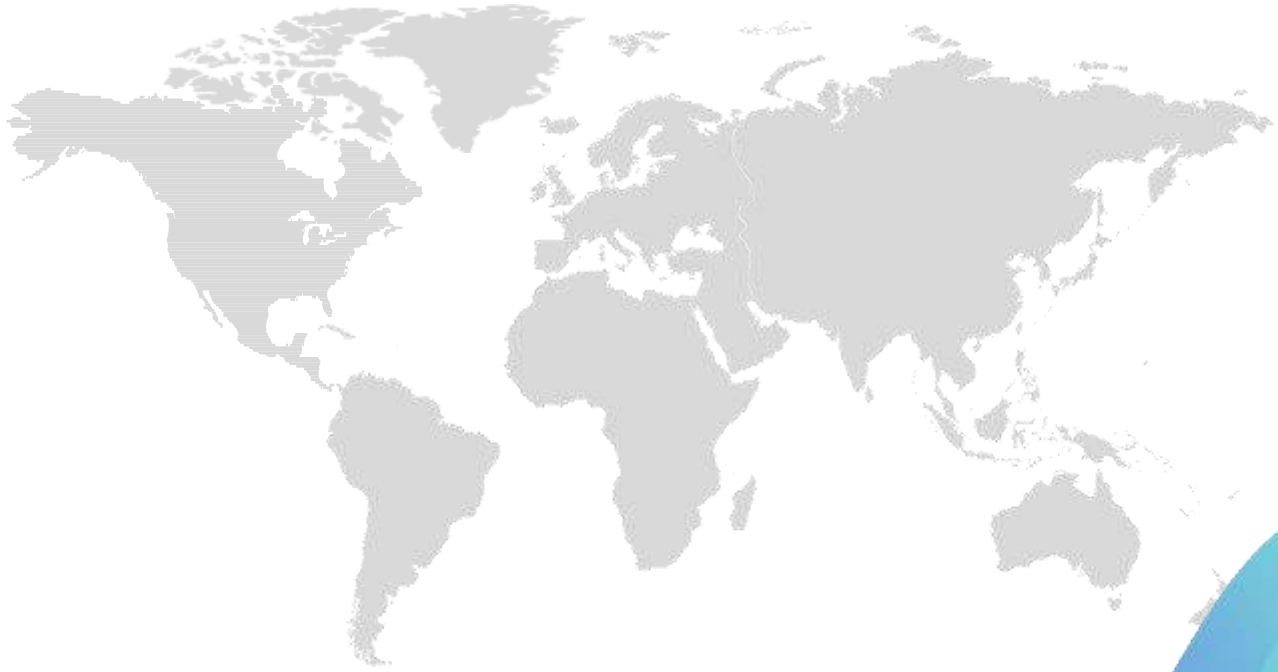
Price: $200

Webinar Description

A brief introduction into the FDA 483 and its use. Responding and responding appropriately is extremely important. This is not an advanced course on 483 response but a recommended approach to best position your company to prevent even worse FDA regulatory measures. All pharmaceutical and medical device facilities who need to know how to react to FDA 483s. Properly responding to FDA observations will make or break your organization and its standing with the regulators.

Who Should Attend ?

Those new to the practice and those who would like an update. Experienced as well as inexperienced



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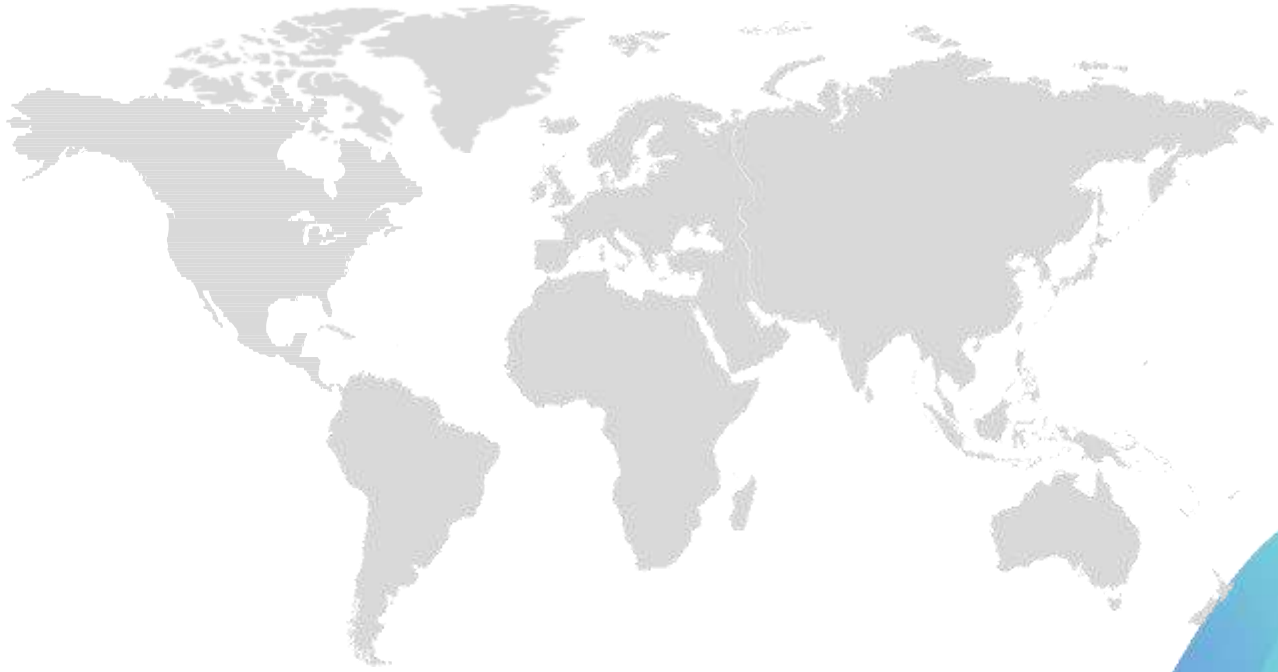

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