

Step 1: Head to academy.greenlight.guru. The access code is right above the sign up fields for you to use if you haven't already signed up.

greenlight guru Academy

Use code *explore_engage_educate*
Partners and customers will have their codes communicated directly for special access.

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Step 2: Find the ER MDR & IVDR tile. Clicking on this tile will take you to all available sessions.

Level up your industry knowledge and explore the topics below

Welcome to Greenlight Guru Academy!
0 hr 6 min

Document Management
Learn about document management and get tips from medical device industry experts.
[Learn more →](#)

Design Controls
Learn why design controls are crucial for the medical device quality process and what the regulations require.
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Risk Management
Risk can be an overwhelming topic, but these courses can show you how to simplify and own the process.
[Learn more →](#)

Training Management
Learn about the regulations that require training and how you can solve for this in your quality process.
[Learn more →](#)

FDA 510(k) Overview
0 hr 34 min

EU MDR & IVDR
TRUE QUALITY SUMMIT SERIES
March 2021
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Step 3: Click the session you are looking to collect the certificate from. This will take you to a registration page.

MARCH 15-19TH

EU MDR & IVDR

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Whether you're looking to watch a session, or just want y

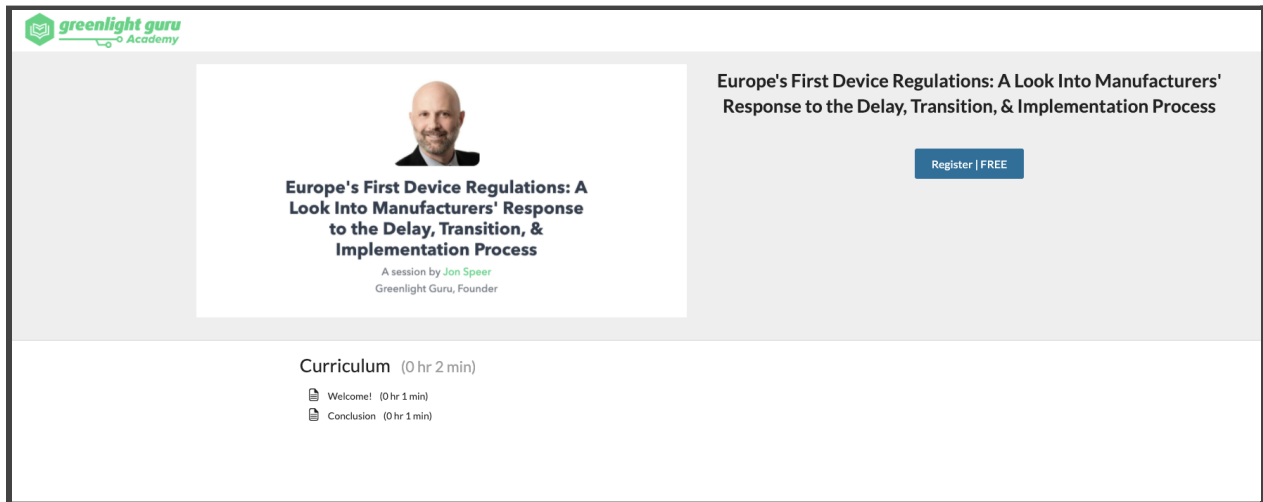


Complete



Europe's First Device Regulations: A Look Into Manufacturers' Response to the Delay, Transition, & Implementation Process
A session by Jon Speer
Greenlight Guru, Founder

0 hr 2 min

Step 4: Click the Register button, then view the “welcome” and “conclusion” lessons and the system will automatically send you a certificate.





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Academy


Europe's First Device Regulations: A Look Into Manufacturers' Response to the Delay, Transition, & Implementation Process
A session by **Jon Speer**
Greenlight Guru, Founder

Europe's First Device Regulations: A Look Into Manufacturers' Response to the Delay, Transition, & Implementation Process

[Register | FREE](#)

Curriculum (0 hr 2 min)

-  Welcome! (0 hr 1 min)
-  Conclusion (0 hr 1 min)

Step 5: Check your email! The certificate will be sent along with some easy instructions to follow for posting on LinkedIn.



Georgia Peach,

Way to go! You're hard work paid off and you should be proud of your accomplishment.

A certificate has been prepared for you and you can download it at this link:

<https://verify.skilljar.com/c/mxywyj226fqj>.

You should share this great milestone with your connections! Learn how to add this certificate to your LinkedIn profile using the button below. If you would be willing to tag Greenlight Guru in your post we would be very grateful. We want to share in your success and help give you the recognition you deserve.



Don't forget to share with us how these skills help you along in your medical device journey. Stories from professionals like you are the best part of our day and will help us continue to improve this experience for others!

Congratulations from all of us here at Greenlight Guru!

Certificate of Attendance

Georgia Peach

for the session

Europe's First Device Regulations: A Look Into Manufacturers' Response to the Delay, Transition, & Implementation Process

Issued: March 15, 2021
Certificate No: mxywyj226fqj

EU MDR & IVDR
TRUE QUALITY SUMMIT SERIES
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This [certificate](#) verifies that on March 15, 2021, **Georgia Peach** completed [Europe's First Device Regulations: A Look Into Manufacturers' Response to the Delay, Transition, & Implementation Process](#), as offered by **Greenlight Guru Academy**.

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