

Clinical Trials in the Framework of MDR and FDA

Date:

14-Oct-2025

10.00 am - 12.00 pm

Venue:

online

Costs:

free of charge

Type:

Webinar

Organiser:

CERES GmbH evaluation & research

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Language:

English

Links:

[🔗 to the event and registration](#)

The webinar on clinical trials in the framework of MDR and FDA will encompass presentations on the FDA approval versus CE conformity, how to proceed after obtaining FDA approval and the set up of clinical trials for efficient data collection for both markets. The invited guest speaker has vast knowledge of both markets and will talk about the pearls and the pitfalls of accessing the US market, while our in-house experts will deliver the presentations on the planning, set up and execution of the clinical trials. In order to instigate a lively discussion, the focus after the presentations will be on answering your questions. Therefore, please insert one required question in the registration form, for our thorough preparation ahead of the event. Should your question go beyond the framework of the webinar, we will be happy to contact you 1:1 to clarify your open point. All slides and presentations will be in English. Please note that for quality purposes, the webinar can be recorded.

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