

MEDICAL DEVICE CLINICAL STUDIES

Understand How to Conduct Pre and Post Market Medical Device Clinical Studies

WEBCAST

COURSE LEADERS

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- ▶ **The Regulatory Framework Applicable to Medical Device Studies**
- ▶ **Standards and Guidelines**
- ▶ **Clinical Evaluation ~ Literature Review**
- ▶ **Designing Pre Market and Post Market Studies**
- ▶ **Ethical Considerations Throughout Europe for Pre and Post Market Studies**
- ▶ **Regulatory Clinical Investigation Notifications**
- ▶ **Site Assessment and Initiation Visits**
- ▶ **How to Write a Final Study Report**
- ▶ **Current Key Issues Affecting Medical Device Studies**
- ▶ **How to Write a Final Study Report**

Reference Documents are available with this webcast