



AdvaMed

Advanced Medical Technology Association
Events & Education

Investigational Device Exemption (IDE) Submissions Workshop

AdvaMed Office
701 Pennsylvania Ave. N.W. Suite 800 Washington, D.C.
October 9, 2019

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|-------------------------|---|
| 8:30 – 9:00 am | Registration Check-In and Continental Breakfast |
| 9:00 – 9:05 am | Welcome and Introductions |
| 9:05 – 9:45 am | What Is an IDE?
<i>Douglas Dumont, FDA</i> <ul style="list-style-type: none">• The purpose of an IDE exemption• Different types of IDEs• What an IDE does (and does not) permit• When manufacturers or physicians should seek an IDE• Roles of IRBs, investigators, and sponsors |
| 9:45 – 10:00 am | Break |
| 10:00 – 11:30 am | Preparing the Technical & Functional Aspects of an IDE
<i>Pam Weagraff, IQVIA</i> <ul style="list-style-type: none">• Elements of an IDE: Intro/Background, Manufacturing/Device, Labeling, Reference/Other• Avoiding common errors and deficiencies• The role of risk analysis in an IDE• Managing planned or unplanned device or study changes |
| 11:30 – 12:15 pm | Regulatory Compliance During Study Conduct
<i>Brian Malkin, Arent Fox</i> <ul style="list-style-type: none">• Monitoring• Consenting of patients• Enrollment requirements• Adverse event reporting• Sponsor records and reports• Investigator records and reports• Protocol deviations |
| 12:15 – 1:15 | Networking Lunch |

Important Notice

The information provided in this course represents the personal opinions of the instructors and does not necessarily represent the opinions of AdvaMed staff. Companies relying on the information do so at their own risk and assume the risk of any subsequent liability that results from relying on the information. The information does not constitute legal advice.



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1:15 – 2:00 pm

Reporting Results

Pam Weagraff, IQVIA

- Clinical study reports (interim and final)
- Dissemination to the medical community and to regulators
- Incorporation into pre-market submissions
- Assessment of impact to product labeling
- Requirements for registering trials on CT.gov

2:00 – 2:30 pm

Optimizing the Pre-PMA Meeting

Kristin Davenport, Covington & Burling LLP

Christina Kuhn, Covington & Burling LLP

- Purpose and value of the meeting
- Requesting a Pre-PMA meeting
- Identifying discussion questions
- Team preparation and rehearsals

2:30 – 2:45 pm

Break

2:45 – 3:45 pm

Developing an IDE Strategy

Kristin Davenport, Covington & Burling LLP

Christina Kuhn, Covington & Burling LLP

- What to consider and when
- Preclinical testing before human studies
- Making the best use of pre-submission meetings
- Using foreign data in a US submission
- Characteristics of a successful IDE submission

3:45 – 4:45 pm

BIMO Audits

Christopher Gioffre, FDA

- The purpose of a BIMO inspection
- When and how a BIMO inspection occurs
- Preventing findings and responding to findings
- Typical and atypical observations – cautionary tales from the field

4:45 – 5:15 pm

Speaker Q&A

5:15 pm

Adjournment

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