Getting it Right:

The Industry Representative Role in Device Advisory Committee Meetings

Complimentary Webinar

November 14, 2018 | 1:00 - 3:00pm EST

Going before an advisory committee can be a make-or-break moment for a device sponsor. Industry representatives (IRs) to these meetings play a critical role, helping to assure that these meetings focus on device safety and effectiveness and the issues under discussion.

In response, AdvaMed has updated its training document for IRs and developed this webinar, which joins industry and FDA device experts to talk about the importance of advisory committee meetings and the IR's key role. These experts will be available to answer your questions. **Register for the webinar here.**

WHO SHOULD ATTEND:

- Industry representatives to device advisory committees
- Prospective advisory committee members
- Regulatory affairs professionals with products that may come before advisory committees

KEY TAKEAWAYS:

- Essential elements of advisory committees: regulatory authorization, makeup, and conflict-of-interest review
- What happens at advisory committee meetings and which topics are on and off the table
- The IR's role and best practices before and during committee meeetings
- How consumer and patient representatives participate in committee meetings

WEBINAR SPEAKERS:

- James Swink, Director, Advisory Panel Program, CDRH, FDA
- Michael Pfleger, Vice President, Head External Affairs and Regulatory Affairs, Alcon
- Robert Durgin, Worldwide Vice President, Regulatory Affairs, DePuy Synthes Companies, Johnson & Johnson
- Sharon Starowicz, Director, Regulatory Policy Innovation, Global Orthopedics, Johnson & Johnson
- Jennifer Bolton, Regulatory Fellow, Boston Scientific

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Q&A session to follow speaker presentations

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