New England Chapter
(Boston)

Advanced Topics in Medical Device Regulation

Program Agenda:
4:00 – 4:30 pm – Arrival & Networking
4:30 – 6:00 pm – Presentation by Chris Cain with Q&A after

Program Description:

2016 has been a big year for FDA and the release of Guidance Documents (20+ Draft Guidances). Many of which will have a direct impact on clinical and regulatory strategy. During this presentation three Guidances will be discussed in detail.

Upon completion of this Webinar, attendees should be able to:

1. Understand how recent FDA (Draft) Guidances may impact clinical and regulatory strategy.
2. Identify three key elements from each (Draft) Guidance.
3. Learn practical ways of implementing the Guidance through real-world experience

Target Audience: Open to all Clinical Research Professionals

Further Information: Contact Denise Clarke at nec.acrp@gmail.com or call 617.306.5514.

Refund Policy: N/A

Level: Open to all clinical research personnel

Date: November 30, 2016

Time: 4:00 pm – 6:00 pm

Location:
Constant Contact
3rd Floor, Great Room
1601 Trapelo Rd,
Waltham, MA 02451

Speaker: Chris Cain
VP, Clinical and Regulatory Affairs,
Corindus, Inc.

Registration Cost: Free for all to attend.

Contact hours: 1.5 Contact hours have been applied for through ACRP. Membership is not required for online registration of contact hours.

To receive contact hours:
Purchase the contact hours, sign in at the registration desk, and attend the program. Log on to the ACRP website then "ACRP Learning Portal" to complete the evaluation no later than 30 days following the event and obtain the online certificate.

Contact Hour Cost:
$0 Chapter Members
$15 ACRP Members
$30 Non-Members