Regulatory Science Series

Medical Device Regulation

Friday, April 20, 12:00–2:00 PM

Maureen Dreher

Heather Vander Ploeg

Join us for a panel on Friday, April 20, from 12:00 p.m. to 2:00 p.m. with **Maureen Dreher**, PhD, Policy Analyst in the Clinical Trials Program at the FDA's Center for Devices and Radiological Health (CDRH), and **Heather Vander Ploeg**, MBA, CCRA, Clinical Operations Manager at Regenesis Biomedical. Dr. Dreher is a biomedical engineer and has been with the FDA since 2007 as a researcher and subject matter expert on medical devices. Ms. Vander Ploeg brings extensive industry experience in medical device development, including clinical trial management at Medicis Pharmaceutical prior to joining Regenesis.

Please join us for an engaging discussion featuring FDA and industry perspectives on the regulation of medical devices. Lunch is included with your RSVP!

To RSVP, please visit:
https://uarizona.co1.qualtrics.com/jfe/form/SV_cUy62lUXMP3kuPz

LOCATIONS

Tucson Campus (Videoconference)
Kiewit Auditorium
Arizona Cancer Center
1515 N. Campbell Ave.
Tucson, AZ 85724

Phoenix Campus (Live)
T-Health Amphitheater
Building 2, Room 2306
550 E. Van Buren St.
Phoenix, AZ 85004

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JAMES E. ROGERS COLLEGE OF LAW
Regulatory Science