FDA WORKSHOP

UCONN

Join us for an interactive and informative workshop on the FDA process led by three accomplished FDA experts and consultants. Gain a basic understanding of the FDA approval and licensing program. Sign up during registration for a 1:1 meeting with one of the experts to get answers to your critical questions.

WHO SHOULD ATTEND:

Anyone involved in a startup looking for insight into the FDA process. Primary focus is on medical devices or diagnostics, with limited discussion on cell, tissue and pharma. The workshop is open to the public and a light lunch will be provided.

PANELISTS:

Susan Alpert, former SVP and Chief Regulatory Officer, Medtronic; former Director of the Office of Device Evaluation at FDA.

Rosemary Harry, President, Device Foundations, Inc.

Pam Weagraff, Director and Group Lead Medical Devices and Diagnostics Regulatory, Quintiles Advisory Services.

WEDNESDAY 10/14/15
10:00 A.M. – 3:00 P.M.

UCONN HEALTH CENTER
CELL AND GENOME
SCIENCES BUILDING

400 FARMINGTON, CT 06030

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