



# REGULATORY SCIENCE SERIES

**Tuesdays, 12:05–12:55 p.m.**

(plus one Thursday and two Friday panel discussions)

**Tucson and Phoenix locations connect by videoconference**

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DATE	SPEAKER	TITLE
<b>FEB. 6</b>	<b>Prova Ahmed and Ibrahim Garba</b> Regulatory Science Fellows	<b><i>mHealth Applications and Regulatory Process</i></b> Tucson: AHSC 3230 Phoenix: Building 2, T-Health 2309
<b>FEB. 8</b>	<b>Patrick O'Brien</b> General Counsel, Arrowhead Research Corporation	<b><i>Labeling</i></b> Tucson: AHSC Library 4150A Phoenix: Building 2, T-Health 2309
<b>FEB. 13</b>	<b>Brooke Courtney</b> Senior Regulatory Counsel, FDA Office of Counterterrorism and Emerging Threats in the Office of the Commissioner	<b><i>Emergency Use Authorizations for Medical Countermeasures</i></b> Tucson: AHSC 3230 Phoenix: Building 2, T-Health 2309
<b>FEB. 27</b>	<b>Andi Mitchell</b> Officer, UA Institutional Animal Care & Use Committee Program	<b><i>Animal Use in Research</i></b> Tucson: Nursing 470 Phoenix: Building 2, T-Health 2309
<b>MAR. 13</b>	<b>Mariette Marsh</b> Director, Human Subjects Protection Program	<b><i>Human Subjects Research and Updates to the Common Rule</i></b> Tucson: College of Medicine 3230 Phoenix: Building 2, T-Health 2309
<b>MAR. 20</b>	<b>Mabel Crescioni</b> Director, Electronic Patient-Reported Outcome Consortium, Critical Path Institute	<b><i>Conflicts of Interests and General Introduction to C-PATH</i></b> Tucson: College of Medicine 3230 Phoenix: Building 2, T-Health 2309
<b>MAR. 23</b>	<b>PANEL: Joseph Ross and Martha Brumfield</b> Martha Brumfield, President and CEO of Critical Path Institute; Joseph Ross, Associate Professor of Medicine (General Medicine), Yale University of Medicine	<b><i>Implications of Amgen v. HHS</i></b> Tucson: Nursing 470 Phoenix: Building 2, T-Health 2309
<b>MAR. 27</b>	<b>Leslie Boyer</b> VIPER Institute, UA College of Medicine	<b><i>Antivenom in the USA</i></b> Tucson: College of Medicine 3230 Phoenix: Building 2, T-Health 2309
<b>APR. 20</b>	<b>PANEL: Maureen Dreher and Heather Vander Ploeg</b> Maureen Dreher, Policy Analyst, Early Feasibility Study Program, Breakthrough Devices Program, FDA; Heather Vander Ploeg, Clinical Operations Manager, Regeneris Biomedical Inc.	<b><i>Medical Devices</i></b> Tucson: Kiewit Auditorium, Cancer Center Phoenix: T-Health Amphitheater 2306



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