

# Regulatory Science Symposium

## Principles of Global Clinical Research for Medical Devices

Friday, April 9, 2021 / 9am - 3pm PST



Asia-Pacific  
Economic Cooperation

USC School of Pharmacy  
International Center for Regulatory Science



9:00 AM PST	<b>Introduction</b> <i>Eunjoo Pacifici, PharmD, PhD</i> USC, SC-CTSI, School of Pharmacy I Chair & Associate Professor, Dept. of Reg. & Quality Sciences I Associate Director, DK Kim International Center for Regulatory Science
9:30 AM PST	<b>Clinical Investigation: Value and Key Aspects of the IMDRF Guidance Document</b> <i>Maria E. Donawa, MD</i> President, Donawa Lifescience
10:15 AM PST	<b>Break</b>
10:30 AM PST	<b>Good Clinical Practices and ISO 14155</b> <i>Danielle Giroud, RN, MBA</i> Founder, CEO, MD-Clinicals
11:15 AM PST	<b>General Discussion</b>
11:30 AM PST	<b>Lunch</b>
12:30 PM PST	<b>From Clinical Data to Clinical Evidence</b> <i>Cheryl Hergert, MPH</i> Principal Clinical Quality Specialist, Medtronic
1:15 PM PST	<b>Break</b>
1:30 PM PST	<b>Developing Clinical Evaluations</b> <i>Evangeline Loh, PhD, RAC (US, EU)</i> Global Regulatory Manager, Emergo by UL
2:15 PM PST	<b>Panel Discussion</b>
2:45 PM PST	<b>Wrap-Up</b> <i>Eunjoo Pacifici, PharmD, PhD</i> USC, SC-CTSI, School of Pharmacy I Chair & Associate Professor, Dept. of Reg. & Quality Sciences I Associate Director, DK Kim International Center for Regulatory Science



Please complete the course evaluation survey at the end of the symposium to receive a certificate of completion. Hours may be eligible for SoCRA and/or ACRP credit.

Series sponsored by The Greater LA CTSA Consortium



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