

***Dissemination and Implementation in Embedded Pragmatic Trials:  
Getting the Timing Right in Real-World Research***

17<sup>th</sup> Annual Conference on the Science of Dissemination and Implementation in Health  
Co-hosted by AcademyHealth and National Institutes of Health  
“Moving Fast and Slow: Optimizing the Pace of Implementation”

Crystal Gateway Marriott, Arlington  
December 8, 2024

<b>DURATION</b>	<b>AGENDA TOPIC</b>	<b>SPEAKERS</b>	<b>GOALS</b>
10:00 – 10:10 a.m.	<b>Welcome Opening Remarks</b>	Emily O’Brien	<ul style="list-style-type: none"> <li>• Welcome and introduction of agenda, objectives, and Living Textbook</li> </ul>
10:10 – 10:40 a.m.	<b>What are Embedded Pragmatic Clinical Trials (ePCTs)?</b>	Beda Jean-Francois	<ul style="list-style-type: none"> <li>• Identify key considerations in the design and conduct of ePCTs and how they differ from explanatory trials</li> <li>• Learn about the advantages and disadvantages of ePCTs, when a pragmatic approach can be used to answer the research question</li> <li>• Q &amp; A with attendees</li> </ul>
10:40 – 11:10 a.m.	<b>Objectives and Trial Design: An Overview of Hybrid Designs</b>	Devon Check	<ul style="list-style-type: none"> <li>• Overview of the 3 types of effectiveness implementation hybrid trial designs and when they may be appropriate for ePCTs</li> <li>• Q &amp; A with attendees</li> </ul>

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11:10 – 11:40 a.m.	<b>Engaging with Health System and Community Partners</b>	Hayden Bosworth	<ul style="list-style-type: none"> <li>• Describe the breadth of individuals to engage as partners and approaches for engaging them through all phases of the study</li> <li>• Identify skills needed for a strong study team and consider the diversity of the team, including inclusive practices</li> <li>• Understand the real-world priorities and perspectives of healthcare system leaders and how to obtain their support</li> <li>• Identify engagement practices to obtain patient and community perspectives</li> <li>• Highlight challenges of partnering with diverse healthcare systems</li> <li>• Q &amp; A with attendees</li> </ul>
11:40 a.m. – 12:40 p.m.	<b>ePCTs in Context: Small Group Work Followed by Panel Discussion with NIH Collaboratory Trial PIs</b>	<b>Moderator:</b> Angelo Volandes  <b>Panel:</b> Andrea Cheville Julie Fritz Mike Ho Sebastian Tong	<ul style="list-style-type: none"> <li>• Introduce PIs of ongoing ePCTs and hear a brief overview of each trial</li> <li>• Have attendees work in small groups to discuss challenges faced by ongoing ePCTs</li> <li>• PIs discuss how they handled the challenges from attendees' discussion, reflect on the morning topics, and discuss lessons learned</li> <li>• Q &amp; A with attendees</li> </ul>
12:40 – 1:40 p.m.	<b>Lunch</b>		<ul style="list-style-type: none"> <li>• Networking among attendees and presenters</li> </ul>
1:40 – 2:00 p.m.	<b>Measuring Outcomes</b>	Angelo Volandes	<ul style="list-style-type: none"> <li>• Describe methods for measuring outcomes using data sources such as electronic health records (EHRs) and patient-reported outcomes (PROs)</li> <li>• Discuss the integration of a health equity lens in evaluating outcomes</li> <li>• Q &amp; A with attendees</li> </ul>
2:00 – 2:30 p.m.	<b>ePCT Design</b>	Jonathan Moyer	<ul style="list-style-type: none"> <li>• Learn about cluster randomized and stepped-wedge study designs</li> <li>• Q &amp; A with attendees</li> </ul>

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2:30 – 3:00 p.m.	<b>ePCT Analysis</b>	Jonathan Moyer	<ul style="list-style-type: none"> <li>Recognize the analytical challenges and trade-offs of pragmatic study designs, focusing on what principal investigators (PIs) need to know</li> <li>Q &amp; A with attendees</li> </ul>
3:00 – 3:10 p.m.	<b>Break</b>		<ul style="list-style-type: none"> <li>Networking among attendees and presenters</li> </ul>
3:10 – 3:40 p.m.	<b>Pilot &amp; Feasibility Testing</b>	Beda Jean-Francois	<ul style="list-style-type: none"> <li>Identify approaches to evaluating the capabilities of the partner healthcare system and testing key elements of various types of interventions</li> <li>Q &amp; A with attendees</li> </ul>
3:40 – 4:10 p.m.	<b>Ethical &amp; Regulatory Oversight Considerations</b>	Stephanie Morain	<ul style="list-style-type: none"> <li>Learn about the regulatory and ethical challenges of conducting ePCTs</li> <li>Discuss unique needs of historically underrepresented and mistreated groups</li> <li>Q &amp; A with attendees</li> </ul>
4:10 – 4:40 p.m.	<b>Writing a Compelling Grant Application</b>	Beda Jean-Francois	<ul style="list-style-type: none"> <li>Learn how to develop a compelling ePCT application</li> <li>Tips from Collaboratory PIs</li> <li>Q &amp; A with attendees</li> </ul>
4:40 – 5:40 p.m.	<b>ePCTs in Context: Small Group Work Followed by Panel Discussion with NIH Collaboratory Trial PIs</b>	<b>Moderator:</b> Stephanie Morain  <b>Panel:</b> Andrea Cheville Julie Fritz Mike Ho Sebastian Tong	<ul style="list-style-type: none"> <li>Have attendees work in small groups to discuss challenges faced by ongoing ePCTs</li> <li>PIs discuss how they handled the challenges from attendees' discussion, reflect on the afternoon topics, and discuss lessons learned</li> <li>Q &amp; A with attendees</li> </ul>
5:40 – 5:50 p.m.	<b>Closing Remarks</b>	Emily O'Brien	<ul style="list-style-type: none"> <li>Wrap-up including identifying sources for further learning</li> </ul>