Agenda

8:00 – 8:45 AM, May 9th, Registration

8:45 – 9:15	**IND Development Plans to Enable First-in-Human Studies** ***Pushpa Singh, Ph.D. and Mauricio Futran, Ph.D.***
9:15 – 9:45	**Getting your Drug to Market - Drug Development 101** ***Carol S. Auletta, M.B.A., D.A.B.T., R.A.C.***
9:45 – 10:15	**Deciphering the CMC Regulatory Code: GCCGCTCMCFORTHEINDGCAGCG....** ***Frederick (Simon) Golec, Jr., Ph.D.***
10:15 – 10:45	**Drug Product: The Other Critical Part of Pre-Clinical Development** ***Denita Winstead, Ph.D.***
10:45 – 11:00	***Break***
11:00 – 11:30	**Toxicology Considerations in Developing Biologics: A Case Study** ***Robert M. Parker, Ph.D., D.A.B.T.***
11:30 – 12:00	**IND-Enabling Study Requirements and Technical Considerations** ***Aidan Curran, Ph.D.***

Register for this free symposium at: http://pddi.ticketleap.com/
or by contacting Rhoda Joseph at: rjoseph@padrugdiscovery.org