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**RECORDINGS:**

1. Requests for Off-label Information – The FDA Guidance and Its Implications  
   Speakers: Wendy Blackburn, Executive Vice President, Intouch Solutions, and Dr. Darshan Kulkarni, Principal Attorney, The Kulkarni Law Firm

2. Preparing for Upcoming Changes in EU Pharmacovigilance Requirements  
   Speaker: Dr. Jan Petracek, CEO & Chief Consultant, PharmInvent

3. Industry Update – Europe’s New Pharmacovigilance Regulations  
   Speaker: Stefan Blesse, Principal Consultant, Granzer Regulatory Consulting & Services

4. Effectively Managing an FDA Inspection – Opening Meeting to Rapid 483 Response  
   Speaker: John Avellanet, Managing Director & Principal, Cerulean Associates LLC

5. FDA Update – Responding to Unsolicited Requests for Off-Label Information  
   Speakers: Marian J. Lee, Partner, and Beverly H. Lorell, Senior Adviser, King & Spalding LLP

6. Centralized vs Onsite Monitoring – Applying FDA’s Risk-based Approach  
   Speaker: Sandra Maddock, President & CEO, IMARC Research, Inc

7. Off-label Promotion – What FDA Looks For & What You Need to Know  
   Speaker: Alan Minsk, Partner, Arnall Golden Gregory LLP

8. Expedited and Periodic Safety Reporting for Drug Trials  
   Speakers: Helen Colquhoun, Vice President, Cromsource

9. Following the 510(k) and 505(b)(2) Pathways to Regulatory Approval  
   Speaker: Dr. Joy Frestedt, President & CEO, Frestedt Incorporate

10. Successfully Responding to FDA 483s and Warning Letters  
    Speaker: Michael A. Swit, Special Counsel, FDA Practice, Duane Morris LLP
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