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2. EU Medical Device Vigilance Reporting Requirements Under MEDDEV 2.12-1 Rev 7
   Speaker: Helen Colquhoun, Senior Vice President, Cromsource

3. How to Document and Implement an FDA-Ready CAPA System
   Speaker: Mark Perkins, Principal Consultant, Medical Device QA/RA Consulting

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   Speaker: Bill White, Senior Consultant, Quality System Strategies LLC

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   Speaker: Dr. Joy Frestedt, President & CEO, Frestedt Incorporated

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   Speakers: John Avellanet, Managing Director & Principal, Cerulean Associates LLC

9. Sample Size for Design Verification and Validation
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10. Medical Device Design Requirements and Considerations for Risk Management
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