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Grand Rounds Special Series:
Ethics and Regulatory Issues in Pragmatic Clinical Trials

Grand Rounds is a weekly webinar hosted by the NIH Health Care Systems Research Collaboratory and PCORnet. Starting in September 2015, one Grand Rounds per month will feature a topic from the *Clinical Trials* 12-article special issue on ethical and regulatory challenges in pragmatic clinical trials. Each session will include a presentation by the author(s) followed by time for discussion. For details and archived presentations, visit https://www.nihcollaboratory.org/Pages/Grand-Rounds-Hub.aspx.

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Date	Title / Presenter
9/25/2015	*Ethical and Regulatory Issues in Pragmatic Clinical Trials: Introducing a Special Series in Clinical Trials* Jeremy Sugarman – Johns Hopkins
10/16/2015	*Data Monitoring Committees for Pragmatic Clinical Trials* Susan Ellenberg – University of Pennsylvania
11/20/2015	*Gatekeepers for Pragmatic Clinical Trials* Danielle M. Whicher – PCORI
12/18/2015	*The Ethics and Regulatory Landscape of Including Vulnerable Populations in Pragmatic Clinical Trials* Mary Jane Welch – Rush University
1/15/2016	*Harmonization and Streamlining of Research Oversight for Pragmatic Clinical Trials* Pearl O'Rourke – Massachusetts General John Lantos – Children's Mercy Hospital
2/19/2016	*Harms, Benefits, and the Nature of Interventions in Pragmatic Clinical Trials* Joe Ali – Johns Hopkins
3/18/2016	*Ethical Responsibilities Toward Indirect and Collateral Participants in Pragmatic Clinical Trials* Jaye Bea Smalley – PCORI
4/15/2016	*Considerations in the Evaluation and Determination of Minimal Risk in Pragmatic Clinical Trials* John Lantos – Children's Mercy Hospital
5/20/2016	*Use of Altered Informed Consent in Pragmatic Clinical Trials* Ross McKinney – Duke University
6/17/2016	*Oversight on the Borderline: Quality Improvement and Pragmatic Research* John Finkelstein – Boston Children's
7/15/2016	*The Food and Drug Administration and Pragmatic Clinical Trials of Marketed Medical Products* Monique Anderson – Duke University
8/19/2016	*Privacy and Confidentiality in Pragmatic Clinical Trials* Deven McGraw – HHS OCR Alan Rubel – University of Wisconsin

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