



Universität
Basel

Juristische
Fakultät



EU Clinical Trial Regulation New Legal Framework for Multicenter Clinical Trials

Friday, December 16, 2016, 1 pm – 5 pm

WWZ Auditorium

University of Basel, Law Faculty, Peter Merian-Weg 8, CH-4002 Basel

Preliminary Program:

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|---------------|---|
| 13.00 – 13.05 | Welcome
Prof. Dr. Dipl.-Biol. Herbert Zech, University Basel, Faculty of Law |
| 13:05 – 13.15 | Introduction
Prof. Dr. Claudia Seitz, M.A., University Basel, Faculty of Law
Dr. Alexander Meier, Novartis Pharma AG |
| 13.15 – 13.45 | EU Clinical Trial Regulation and the Swiss Human Research Act
Prof. Dr. Claudia Seitz, M.A., University Basel, Faculty of Law |
| 13.45 – 14.15 | EU Clinical Trial Regulation – A View from Legal and Regulatory
Dr. Alexander Meier, Novartis Pharma AG |
| 14.15 – 14.45 | Coffee Break |
| 14.45 – 15.30 | EU Clinical Trial Regulation – A View from the Regulator
(tbc) |
| 15.30 – 16.00 | EU Clinical Trial Regulation – A View from the Industry
Judith Creba, Senior Global Program Director, Novartis Pharma AG |
| 16.00 – 16.30 | EU Clinical Trial Regulation – A View from Data Privacy
Noemi Alonso Calvo, Data Privacy Officer, Sandoz International GmbH |
| 16.30 – 17.00 | Discussion |
| 17.00 | Apéro |

No registration fee, please register by email to zlsr-ius@unibas.ch by Thursday, December 8, 2016

Organized by: Dr. Alexander Meier, Novartis Pharma AG, Head Legal Global & Pharma Drug Development, Basel

Prof. Dr. Claudia Seitz, M.A., Center for Life Sciences Law (CLSL), University Basel, Faculty of Law