

Speakers
Part 15 Hearing – Adverse Event Reporting to IRB’s
Docket 2005n-0038
March 21, 2005

All presenters have requested 20 minutes or less

Dr. Juhana Idanpaan-Heikkila (1)
Secretary General
Council for International Organizations of Medical Sciences

Yvonne Higgins (2)
Associate Director, Human Research
University of Pennsylvania

Michael Susko (3)
President
Citizens for Responsible Care and Research

Dr. Sandra Alfano (4)
Vice Chair, Human Investigations Committee
Yale University

Maureen Hardwick (5)
Partner
Garner, Carton, & Douglas on behalf of IRB Sponsor Roundtable

Dr. Gary Chadwick (6)
Associate Provost
University of Rochester

Dr. Owen Reese (7)
Executive Director
Western Institutional Review Board

Dr. Howard Dickler (8)
Senior Consultant for Research
Association of American Medical Colleges

Paul Covington (9)
Executive Vice President of Development
PPD Development

Dr. William Hendee (10)
Senior Associate Dean and Vice President
Medical College of Wisconsin

Dr. Vish Watkins (11)
Project Leader
Eli Lilly and Company

John Isidor (12)
Schulman Associates IRB, Inc.
Representing Consortium of Independent Review Boards

Dr. Jean –Louis Saillot (13)
Schering Plough
Representing PhRMA

Dr. P. Pearl O'Rourke (14) (did not attend meeting- David Borasky read statement)
Chair
PRIM&R Board of Directors

David Borasky (15)
Immediate Past President
Applied Research Ethics National Association (ARENA)

Dr. Wendy Stephenson (16)
Co-Chair
Council for International Organizations of Medical Sciences VI Working Group

Thomas Adams (17)
President and Chief Executive Officer
Association of Clinical Research Professionals

Dr. Greg Koski (18)
Massachusetts General Hospital
Representing Association of Clinical Research Professionals

Dr. Sorell Schwartz (19)
Chair, IRB Subcommittee on Adverse Events
Georgetown University Medical Center