Health outcomes research in an era of cost containment

Improving efficiency of research: decreasing costs, increasing quality

PART 2: INTERVENTIONS
MARCH 12, 2015

Centre de Recherche Épidémiologie et Statistique Sorbonne Paris Cité (CRESS-UMR1153)

8:30 - 9:00 BREAKFAST AND INFORMAL GREETINGS

9:00 - 9:05 WELCOMING REMARKS

9:05 - 9:15 SYMPOSIUM OVERVIEW

9:15 - 10:45 PANEL 1: INNOVATIVE, MORE EFFICIENT TRIAL DESIGN

No of 1 trials
Sunita Vohra, MD, MSc, FRCPC, FCAHS

Direct to patient trials
Steve R. Cummings, MD

Public led online trials
Amy Price, PhD

10:45 - 11:00 COFFEE BREAK

11:00 - 12:30 PANEL 2: SIMPLIFYING TRIALS OR IMPROVING OBSERVATIONAL STUDIES ACCURACY

Changing trial metrics: The role of data management
Eric L. Eisenstein, DBA

Improving trial monitoring to reduce the costs
Tomasz Burzykowski, PhD

Analytic techniques to improve accuracy of observational studies
Jason Wright

12:30 - 13:30 LUNCH

13:30 - 15:00 PANEL 3: BLURRING THE LINES: USING OBSERVATIONAL STUDIES TO IMPROVE RECRUITMENTS IN RCTS

Registry-based RCTs: A disruptive technology
Ole Fröbert, MD, PhD

A new trial design to speed RCTs: The I-SPY 2 trial
Donald Berry, MD, PhD

Strategies to improve recruitment in RCTs
Jonathan Craig, MBChB, DipCH, FRACP, M Med (Clin Epi), PhD

15:00 - 16:30 PANEL 4: BETTER REPORTING AND DATA SHARING FOR BETTER TRIAL RETURN ON INVESTMENT

Interventions to improve the quality of reporting and posting of results
Isabelle Boutron, PhD

Sharing clinical trial data on patient level
Martin Posch

16:30 - 16:45 CLOSING REMARKS

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