Long-Term Monitoring of Treatment Related Adverse Events in the Resource Limited Setting

The Forum for Collaborative HIV Research

Ben Cheng

www.hivforum.org

The Forum for Collaborative HIV Research is a public/private partnership including government agencies, industry, HIV researchers and clinicians, payors, foundations and the HIV patient advocacy community.

Our mission is to facilitate and enhance HIV research.

The Forum Executive Committee

- Government Agencies
  - US DHHS (NIH, CDC, FDA, HRSA), State Department (OGAC)
  - European Regulatory: EMEA
- Industries
  - Abbott, Bayer Diagnostic, Boehringer Ingelheim, Bristol-Myers Squibb, Gilead Sciences, GlaxoSmithKline, Monogram BioSciences, Roche Laboratories, Roche Molecular Systems, Pfizer, Schering-Plough, Tibotec, VIRxSYS
- Payors: Kaiser Permanente
- Academia
  - US and Europe
- Providers
- Patient Advocacy
  - US and Europe
- Foundations & Organizations (Gates, AmFAR, IAS)

Mechanisms & Outcomes

- Workshops
- Roundtable Discussions
- On-Going Working Groups
- Conference Activities
  - Satellite Symposia
  - Feedback of projects and reports
- Identification of gaps, recommendations, collaborations between programs, constituencies, etc; dissemination of information, publication and position papers, streamlined funding, etc

Planning Committee

Shabir Banoo	South African MCC/University of Witwatersrand
Ben Cheng	Forum for Collaborative HIV Research
David Cooper	NCHECR
Judith Currier	UCLA
Elly Katabira	Makerere University
Cissy Kityo	Joint Clinical Research Centre
Jens Lundgren	Copenhagen HIV Program/EuroSIDA
Veronica Miller	Forum for Collaborative HIV Research
David Pizzuti	Johnson & Johnson
Bill Powderly	Mater University Hospital
Ian Weller	Royal Free & University College Medical School

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  - Ipsita Das and Becky Griesse – project coordinators

Agenda

13:00 – 13:15	Welcome and Introductions	Bill Powderly and Ben Cheng
13:15 – 13:30	Long-term monitoring of treatment related adverse events in adults	Elly Katibira
13:30 – 13:45	Regulatory considerations for long-term monitoring of treatment related adverse events	Shabir Banoo
13:45 – 14:45	Discussion	
14:45 – 15:00	Break	
15:00 – 15:15	Considerations for data collection	Veronica Miller
15:15 – 15:30	Use of Information Technology for Data Collection	Pamela Johnson
15:30 – 16:45	Discussion	
16:45 – 17:00	Wrap up and next steps	Bill Powderly and Ben Cheng