6th isirv-AVG Conference Advances in Respiratory Virus Therapeutics

Tuesday 13th - Thursday 15th November 2018 Hilton Washington DC/ Rockville Hotel, Washington DC, USA

Preliminary Programme

Tuesday 13 November	
08:30-08:45	Welcome & Opening
08:45-09:30	Session 1: Keynote Lecture: BARDA Perspectives on Advancing Respiratory Virus Therapeutics <i>Rick Bright, BARDA, Washington, D.C., USA</i>
09:30-11:00	Session 2 : Symposium - Preclinical Topics Eritoran, a TLR 4 Antagonist Stefanie Vogel, University of Maryland, Baltimore, MD, USA Repurposing of drugs as novel influenza (and other respiratory viruses) inhibitors from clinical gene expression infection signatures Andres Pizzorno, Centre International de Recherche en Infectiologie (CIRI- Team VirPath) & Signia Therapeutics, Lyon, France Repurposing Host Targets for Influenza Therapy Kevin Harrod, University of Alabama, Birmingham, AL, USA HA Minibinders Proof-of-Concept in Mice and Ferrets Deborah Fuller, University of Washington, Seattle, WA, USA
11:00-11:30	Coffee Break
11:30-13:00	Session 3: Oral Abstract Session 1- Preclinical Papers
13:00-14:00	Sponsored Lunchtime Seminar - TBC
14:00-15:00	Session 4: Symposium- Clinical Trial Design Issues Clinical Pharmacology Considerations for Influenza and RSV Trials FDA Speaker Considerations of use of PROs in SARI and Hospitalized Influenza Studies FDA COA Speaker
15:00-15:30	Coffee Break
15:30-17:00	Session 5: Oral Abstracts Session 2- Preclinical and Phase 1 Clinical Papers
17:00-18:00	Welcome Buffet Reception

Wednesday 14 November	
08:30-10:35	 Session 6: Symposium - Updates on Clinical Trials Evaluation of Anti-HA Stem Monoclonal Antibody FluA in Two Phase 2 Influenza Studies Melicent Peck, Genentech, South San Francisco, California, USA Baloxavir Marboxil, a Cap-Dependent Endonuclease Inhibitor - Development Updates Takeki Uehara, Shionogi & Co., Ltd, Osaka, Japan RSV Antiviral Treatment for HCT patients: Results from Recent Phase 2 Studies for Presatovir Michael Boeckh, University of Washington, Seattle, WA, USA Overview of RSV and Influenza Programmes James Witek, Janssen Research and Development, Titusville, New Jersey, USA
	Randomized Phase 2 Study Evaluating Nitazoxanide Versus Placebo in Hospitalized Subjects with Severe Acute Respiratory Illness <i>John Beigel, NIAID, Bethesda, MD, USA</i>
10:35-11:00	Coffee Break
11:00-13:00	 Session 7: Symposium- Clinical Trial and Regulatory Issues FDA Considerations for Influenza Drug Development Trial Designs for Serious Influenza Therapeutic Combinations (Antivirals, Adjunctive Therapies) Pediatric Considerations FDA Speaker Statistical Considerations/Endpoints LaRee Tracy, FDA, Silver Spring, MD, USA EMA Perspective Dr Radu Botgros, EMA, London, UK Investigator Perspective (Influenza) Michael Ison, Northwestern University, Chicago, Illinois, USA Discussion on RSV - FDA Guidance for Industry and Trial Designs/Endpoint FDA Speaker Investigator Perspective (RSV) John DeVincenzo, University of Tennessee, Memphis, Tennessee, USA
13:00-14:00	Sponsored Lunchtime Seminar - TBC
14:00-15:00	Session 8: Roundtable Discussion- Clinical Trial and Regulatory Issues Academic, Regulatory, and Industry representatives
15:00-15:30	Coffee Break
15:30-17:00	Session 9: Oral Abstract Session 3- Clinical Papers (Including Disease Pathogenesis, Biomarkers, Diagnostics)

17:00-18:00	Poster Session
Thursday 15 November	
08:30-10:00	Session 10: Oral Abstract Session 4- Clinical Trial Papers
10:00-10:30	Coffee Break
10:30-12:30	Session 11: Hot Topics and Late-Breakers Antiviral Resistance Monitoring Strategies Aeron Hurt, WHO Collaborating Centre, Melbourne, Australia Novel Coronaviruses Ralph Baric, University of North Carolina, Chapel Hill, NC, USA Adjunctive Therapies Nelson Lee, University of Alberta, Edmonton, Canada
12:30-13:00	Summary and Close of Conference
	Lunch and Depart