

**Long-Acting/Extended Release (LA/ER) Antiretroviral Research Resource Program (LEAP)  
Investigator Meeting and Annual Workshop 2022  
Hyatt Regency Denver  
AGENDA**

**SATURDAY, February 12 | 1:00 AM – 5:30 PM | PLENARY SESSION | TBD**

**12:30 – 1:00 PM      ONSITE REGISTRATION AND LUNCH**

- 1:00-1:05 PM      **Welcome:** Carl Dieffenbach, Director, Division of AIDS, NIH
- 1:05-1:15 PM      **Where are we LEAPing next?!** – Charles Flexner, Johns Hopkins University
- Overview of Workshop** – Charles Flexner, Johns Hopkins University

**Plenary Session I – Charles Flexner, Chair**

- 1:15-2:35 PM      **Plenary Session I: Current Status of Existing Technologies:  
10-Minute Presentations followed by Q & A**
- 1:15-1:25 PM — “Update on Transcutaneous Microneedles for Antiretroviral Drug Delivery” –  
                         **Ryan Donnelly, Queens University Belfast**
- 1:25-1:35 PM — “Current status of LA/ER Cabotegravir and Rilpivirine including pipeline report  
on novel CBT formulations” – **William Spreen, ViiV**
- 1:35-1:45 PM — “Current Status of the Merck LA/ER Pipeline” –  
                         **Jay Grobler, Merck**
- 1:45-1:55 PM — “Current Status of the Gilead LA/ER Pipeline” – **Martin Rhee, Gilead Sciences**
- 1:55-2:05 PM — “Open discussion on pharmaceutical pipeline” – **ViiV, Merck, Gilead**
- 2:05-2:15 PM — “Update from the LEAP Modeling and Simulation Core” –  
                         **Andrew Owen, University of Liverpool**
- 2:15-2:25 PM — “LEAP Systematic Reviews of LA formulation use in pediatrics and pregnancy” –  
                         **Lynn Bertagnolli, Johns Hopkins University**
- 2:25-2:35 PM — “DAIDS/NIH resources to support preclinical and clinical development of ARVs and  
related anti-infective drugs” – **Keith Crawford, Division of AIDS, NIH**

**2:35 PM-2:45      10 Minute Break**

**TIME (EST)      Plenary Session 2 – Andrew Owen, Chair**

- 2:45-4:15 PM      **Plenary Session 2: Novel approaches to LA/ER Drug Delivery:  
15-Minute Presentations followed by Q & A**
- 2:45-3:00 PM — “Tenofovir implants and local toxicity: What have we learned to date?” -  
                         **Marc Baum, Oak Crest Institute**
- 3:00-3:15 PM — “Developing a long-acting formulation for Hepatitis B virus treatment and prevention”  
                         **A. Chatterjee, Calibr, Scripps Institute, San Diego**
- 3:15-3:30 AM — “Novel long-acting protides of approved ARVs” -  
                         **Benson Odagwa, University of Nebraska**
- 3:30-3:45 PM — “Update from LONGEVITY: Novel long-acting formulations for Tuberculosis,  
Malaria, and Hepatitis C Virus” – **Andrew Owen, University of Liverpool**
- 3:45-4:00 PM — “Update from the Targeted, Long-acting and Combination Anti-Retroviral Therapy  
(TLC-ART) program” – **Rodney Ho, University of Washington**
- 4:00-4:15 PM — “CADO 4/PADO 5: Approach to delivery of LA ARVs for HIV Treatment and  
Prevention in LMICs, report from a WHO-sponsored meeting” –  
                         **David Ripin, Clinton Health Access Initiative**

**4:15-4:25 PM      10 Minute Break**

**Focus Group Summary Reports – Sue Swindells, Chair**

4:25-5:25 PM

10 mins for each report

**4:25-4:35 PM Group 1:** “Developing LA formulations for treatment and prevention of HBV and HDV: Where are we now and where do we need to go?” –

**Andy Kaytes, UCSD**

**4:35-4:45 PM Group 2:** “Long-acting oral drugs and formulations: how do they stack up compared to other routes of drug delivery” –

**Polly Clayden, iBase**

**4:45-4:55 PM Group 3:** “Best practices for conversion of immediate release approved ARVs: To prodrug or not to prodrug?” –

**Paul Domanico, CHAI**

**4:55-5:05 PM Group 4:** “Challenges in developing combination LA products and regimens?” –

**Bob Bollinger, JHU**

**5:05-5:25 PM**

Discussion of Focus Group Reports and Next Steps – ALL

**5:30 PM**

**Close of Workshop**