



## Collocation of Buprenorphine with HCV Treatment to Improve Adherence and Reduce Harm in PWID with HCV: Preliminary Data from the ANCHOR Investigation

*Elana Rosenthal MD, Kristi Hill BA, Laura Nussdorf BA, Poonam Mathur DO MPH, Chloe Gross RN, Rachel Silk RN MPH, Elizabeth Akoth RN MPH, David Sternberg BA, Nadeera Siddique BA, Chloe Chaudhury BS, Benjamin Emmanuel MPH, Henry Masur MD, Shyamasundaran Kottillil MD PhD, Sarah Kattakuzhy MD*

*Institute of Human Virology, University of Maryland School of Medicine, Baltimore, MD, USA and NIH DC Partnership for HIV/AIDS Progress, Bethesda, MD, USA*



### Disclosures

- Sarah Kattakuzhy has received institutional grants from Gilead Sciences
- Elana Rosenthal has received institutional grants from Gilead Sciences and Merck
- Poonam Mathur has received institutional grants from Merck

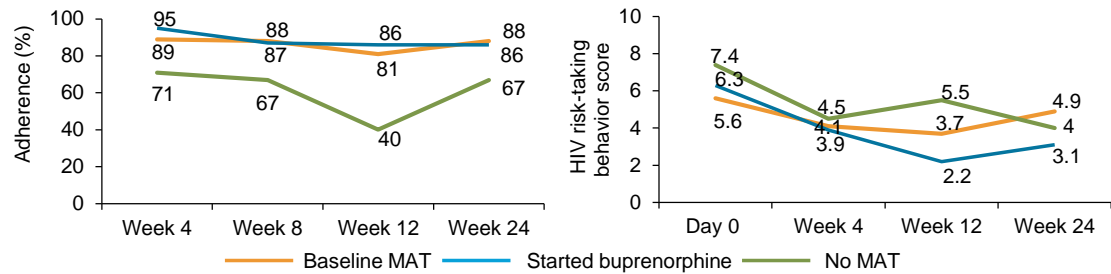
## Background/Aims

- Studies modeling HCV treatment as prevention in PWID highlight need for MAT
- Limited real-world data to evaluate
  - effect of MAT on success of HCV treatment
  - the optimal timing of MAT relative to initiating DAA therapy
- **To evaluate the impact of colocated buprenorphine with HCV treatment in PWID with active IDU on HCV related outcomes and IDU associated harms**

## Methods

- Single centre study conducted at an embedded clinic within in a harm reduction organization drop-in center in Washington, DC
- N=100
  - Opioid Use Disorder
  - Injection of an opioid within 3 months
- Treatment
  - Sofosbuvir/Velpatasvir for 12 weeks
  - Optional buprenorphine
- Endpoints
  - Adherence to medication and visits
  - Risk taking behaviour
  - SVR & Reinfection

## Results



### Compared with patients not on medication-assisted treatment (MAT) for OUD, those who started buprenorphine:

- Were more likely to attend treatment visits: Week 4 (95% vs. 71%,  $p=0.02$ ); Week 12 (86% vs. 40%,  $p=0.007$ )
- Were more likely to receive the second bottle of SOF/VEL (100% vs. 82%,  $p=0.02$ ), and were more likely to receive the second bottle at a clinic visit ( $p=0.01$ )
- Had a significant decline in the Darke HIV Risk-Taking Behaviour Scale during and after HCV treatment (-2.2 at Week 4,  $p=0.003$ ; -4.4 at Week 12,  $p=0.001$ ; -3.2 at Week 24,  $p=0.003$ )

Of the 45 patients who attended the Week 24 visit to date, the overall SVR was 93%

## Conclusions/Implications

- Preliminary results of the ANCHOR study support that active PWID not on MAT can be successfully initiated on buprenorphine during the course of HCV treatment
- Initiation of colocated buprenorphine
  - Improves adherence to medical visits and medication pick-up
  - Decreases resources necessary to dispense medication
  - Decreases risk-taking behaviour during and after HCV treatment, a result not seen in those on baseline MAT or no MAT
- ANCHOR colocated care model may provide a critical opportunity to cure HCV while simultaneously treating OUD in PWID in order to prevent reinfection, HIV acquisition and overdose-related death

## Acknowledgements

### DC PFAP

- Elana Rosenthal, MD
- Poonam Mathur, DO, MPH
- Rachel Silk, RN, MPH
- Chloe Gross, RN
- Elizabeth Akoth, RN, MS
- Kristi Hill, BA
- Nadeera Sidique, BA
- Laura Nussdorf, BA
- Ben Emmanuel, MPH
- Henry Masur, MD
- Shyam Kottlilil, MD, PhD

### HIPS Staff

- David Sternberg
- Phyllis Bijole
- Miriam Jones
- Dana McCullough
- Randy Kier

### Gilead Sciences

**The patients of Washington DC**