

Hepatitis C Single Attendance Test Assess and Treat  
 (“**Hep STAT**”) minimizes care cascade attrition and achieves  
 high rates of treatment initiation, completion and cure  
 in people who inject drugs



B. Stone, S. Emerson, K. Jenkins, J. Havercroft, I. Shaukat, B. McClean, G. Dinsey & G. Hill

South Yorkshire, Bassetlaw and North Derbyshire Hepatitis C Operational Delivery Network, UK

# Disclosures

- Cepheid® GeneXpert® and Xpert® HCV VL Fingerstick assays were provided by Cepheid UK Ltd.
- Peer Support Lead salary provided by AbbVie Inc.
- No other pharmaceutical grants were received in the development of this study.

# Objectives

- To design and pilot the simplest and shortest possible pathway from hepatitis C testing to diagnosis, assessment and treatment initiation
- To minimize number of investigations and attendances and reduce care cascade attrition
- To engage individuals at high risk of hepatitis C with pre-existing potential barriers limiting access to traditional models of care

# “Hep STAT”

- **Hepatitis C Single Attendance Test, Assess and Treat**

## Resources:

- NHSE pan-genotypic access (Epclusa®)
- Portable FibroScan™
- Cepheid® GeneXpert® II
- Cepheid® Xpert® HCV Viral Load cartridges
- Cepheid® Xpert® HIV Viral Load cartridges
- Hep C Trust Peer Support Lead (AbbVie Inc.)
- Peer volunteers



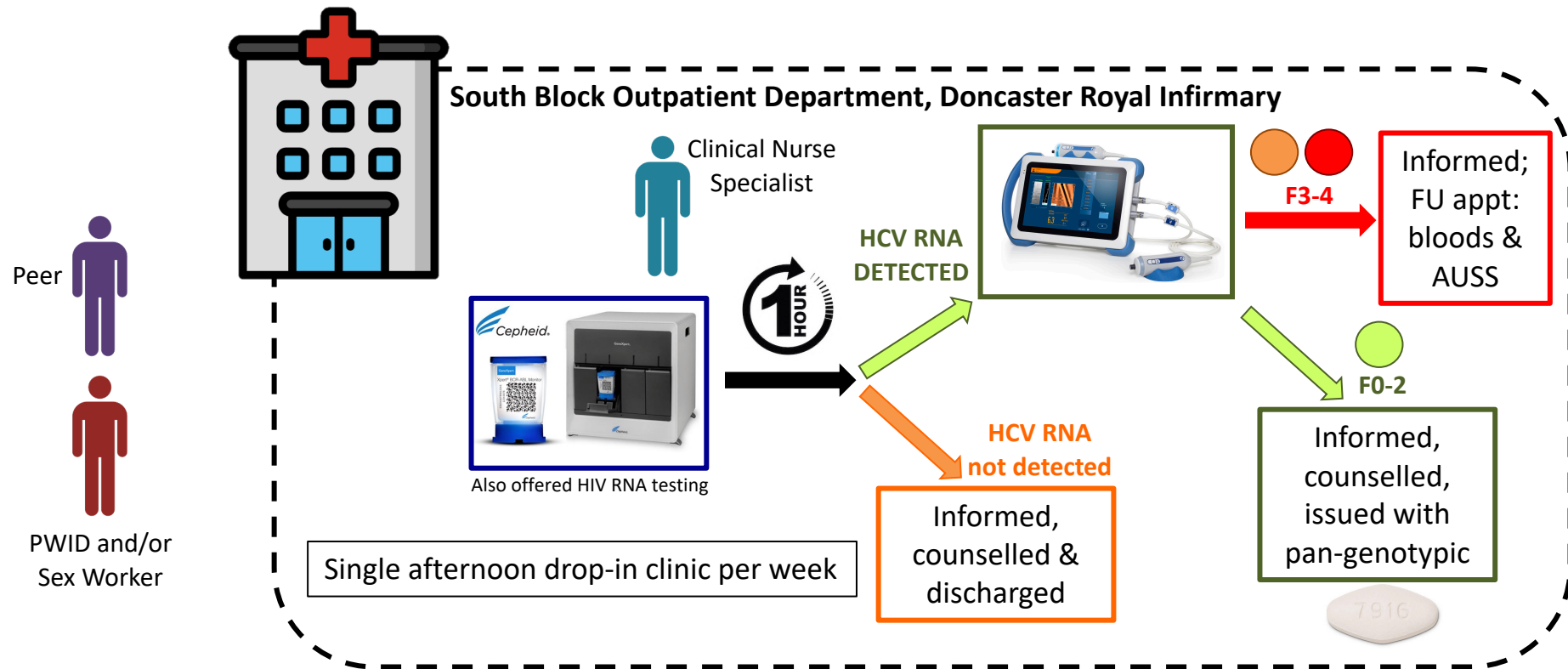
## Target population:

- Adults aged  $\geq 18$  years old
- History of current or recent past injecting drug use and/or sex work
- No known current or past hepatitis C infection diagnosis or treatment

AUSS = abdominal ultrasound scan

PWID = person who injects drugs

# “Hep STAT” Patient Pathway



At risk patients identified by Clinical Nurse Specialists and Peers from Homeless Hostels, Drug Services & other community services for PWIDs in Doncaster and Bassetlaw (Workshop) areas

# “Hep STAT” Results

- **27 high-risk individuals identified and attended with peer-support** over 16-week period (12 clinics held March to end July 2019; average 2.3 patients per half-day clinic)
  - **25/27 (92.6%) current IDU** of whom **3/25 (12.0%) with sex work history**; 1/27 (3.7%) past IDU
  - All consented to HCV and HIV testing
- HIV RNA detected: 0/25 (test failed in 2 individuals)
- **HCV RNA detected: 9/27 (33.3%)**
  - Median TE  $\leq 9.5$  kPa (METAVIR F0-2) = **6/9 (66.7%)**
  - Median TE  $> 9.5$  kPa (METAVIR F3-4) = **3/9 (33.3%)** (median TE range 15.6 – 26.6 kPa)
- **Median time arrival to departure: 113 minutes (range 75 to 195)\***
  - For HCV RNA detected: shortest time from arrival to departure = 90 minutes

\*Excluding 2 patients unwilling to wait for HCV RNA result (both HCV RNA not detected)

# Outcomes for patients with detectable HCV RNA

Patient	METAVIR	Attended follow up	1 <sup>st</sup> 4 weeks issued	2 <sup>nd</sup> 4 weeks issued	3 <sup>rd</sup> 4 weeks issued	Reached PTW12	Attended PTW12 bloods	SVR12 achieved
1	F0-2	Not applicable	✓	✓	✗	✓	✓	✓
2	F0-2		✓	✓	✓	✓	✓	✓
3	F0-2		✓	✓	✓	✓	✓	✓
4	F0-2		✓	✓	✓	✓	✓	✓
5	F0-2		✓	✓	✓	✓	✓	✓
6	F0-2		✓	✓	✓	✓	✗	?
7	F4	✓*	✓	✓	✓	✓	✓	✓
8	F4	✓*	✓	✓	✓	✓	✓	✓
9	F4	✓*	✓	✓‡	✓‡	✓	✓	✓

\*All Child Pugh A(5); 1/3 attended for AUSS (no focal lesions); 2 x G3a, 1 x unable to genotype; all commenced Eplusa

\*Treatment re-initiated after interruption after week 4

PTW12 = post-treatment week 12

# Conclusions & Future Application

- “Hep STAT” minimizes care cascade attrition and achieves high rates of hepatitis C treatment initiation, completion and cure in a high prevalence setting
- Embedding into drug services or prison reception screening may improve hepatitis C micro-elimination potential
- Roll out dependent on availability of: peer support, dedicated health professional time, Cepheid® GeneXpert®, portable FibroScan® and on-site stock of pan-genotypic treatment
- Upscale requires sustained investment and careful consideration of most appropriate highest yield locations and settings



# Acknowledgements

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- Community participants

