

REINFECTION FOLLOWING SUCCESSFUL DIRECT-ACTING ANTIVIRAL THERAPY FOR HEPATITIS C VIRUS INFECTION AMONG PEOPLE WHO INJECT DRUGS: THE SHARP-C STUDY

Authors:

Joshua Ang¹, Behzad Hajarizadeh¹, Shane Tillakeratne¹, Carla Treloar², Janaki Amin³, Jodi van Dyk¹, Louisa Degenhardt⁴, Tanya Applegate¹, Adrian Dunlop⁵, Chris Fraser⁶, Brian Conway⁷, Alexander Wong⁸, Dennaye Fuchs⁹, Jeff Powis¹⁰, Kate Mason¹¹, Edward J Gane¹², Gregory J Dore¹, Martin Weltman^{13,14}, Phillip Read^{1,15}, Marianne Martinello^{1,20}, Emily Rowe^{16,17}, David A Baker¹⁷, Alexandra Wade^{18,19}, Gail Matthews¹, Lise Lafferty¹, Andrew Lloyd¹, Jason Grebely¹, and Evan B Cunningham¹

¹Kirby Institute, UNSW, Sydney, NSW, Australia, ²Centre for Social Research in Health, UNSW, Australia, ³Macquarie University, Sydney, NSW, Australia, ⁴National Drug and Alcohol Research Centre, UNSW Sydney, Sydney, Australia, ⁵Drug & Alcohol Clinical Services, Hunter New England Health, & University of Newcastle, Newcastle, NSW, Australia, ⁶Cool Aid Community Health Center, Victoria, BC, Canada, ⁷Vancouver Infectious Diseases Centre, Vancouver, BC, Canada, ⁸Department of Medicine, University of Saskatchewan, Regina, SK, Canada, ⁹ID Clinic, Saskatchewan Health Authority, Regina, SK, Canada, ¹⁰Michael Garron Hospital, Toronto, ON, Canada, ¹¹South Riverdale Community Health Centre, Toronto, ON, Canada, ¹²University of Auckland, Auckland, New Zealand, ¹³Department of Gastroenterology and Hepatology, Nepean Hospital, Kingswood, Australia, ¹⁴Nepean Clinical School, University of Sydney, Kingswood, Australia, ¹⁵Kirketon Road Centre, South Eastern Local Health District, Sydney, Australia, ¹⁶Royal Adelaide Hospital, Central Adelaide Local Health Network, Adelaide, SA, Australia, ¹⁷The University of Adelaide, Adelaide, SA, Australia, ¹⁸Mid North Coast Liver Clinic, Mid North Coast Local Health District, Australia, ¹⁹Drug and Alcohol Services, Mid North Coast Local Health District, Australia, ²⁰Blacktown and Mt Druitt Hospital, Western Sydney Local Health District, Blacktown, NSW, Australia

Background:

Reinfection following successful Hepatitis C Virus (HCV) treatment can reduce the individual and population benefits of HCV treatment. This study aimed to evaluate the incidence of HCV reinfection and associated factors following successful direct acting antiviral (DAA) therapy among people with recent injecting drug use.

Methods:

The SHARP-C study was a prospective, observational cohort study, which recruited participants from hospital-based HCV clinics, community-based drug treatment clinics, and community healthcare clinics in Australia, New Zealand, and Canada between 2018 and 2020. Participants with recent injecting drug use (prior 6 months) with either: 1) chronic HCV infection who were commencing DAA therapy; or 2) had a documented sustained virological response (SVR) were recruited. Participants were evaluated every three months, with reinfection confirmed via viral sequencing. Follow-up was calculated from end of treatment (those commencing treatment during study) or SVR (those previously treated). Person-time of observation and Cox proportional hazard models were used to calculate reinfection incidence and associated factors.

Results:

A total of 111 participants were enrolled (median age 43 years; 66% male, 85% reported injecting drugs in the last month). Twelve reinfection cases were confirmed over 113 person-years, for an incidence of 10.6/100 person-years (95% confidence interval [CI], 6.0-18.7). Median time to reinfection was 22 weeks. Reinfections exclusively occurred in people with recent injecting drug use (last month) during study follow-up, for an incidence of 12.2/100 person-years in this population (95% CI, 5.8-20.0). Among people who reported sharing needles/syringes, incidence was 32.3/100 person-years (95% CI, 8.0-76.8).

Conclusion:

These findings underscore the importance of monitoring for HCV reinfection following treatment among people who continue to inject drugs. They also highlight the need for research into holistic models of HCV care which are integrated into harm reduction services in order to reduce the risk of reinfection.

Disclosure of interest:

EBC, and BH are funded by an NHMRC Investigator Grant. GJD is funded by an NHMRC Investigator Grant and has received research income from Gilead Sciences and AbbVie. AW has received honoraria and consulting fees from Gilead and AbbVie. CF has received research grants from AbbVie and Gilead. MW has acted as an education provider to Gilead Sciences.