

AN UNMET NEED FOR HEPATITIS C TESTING AT NEEDLE AND SYRINGE SERVICES – LESSONS LEARNED FROM THE RAPID-EC FEASIBILITY STUDY.

Authors:

Williams B¹, Pedrana A^{1,2}, Howell J^{1,2,3,4}, Doyle J^{1,5}, Thompson A^{3,4}, Bramwell F⁶, Membrey D⁶, Mcpherson M⁷, Layton C⁶, Draper B¹, Roney J⁸, Latham N⁹, Hellard ME^{1,2,5}.

¹Disease Elimination Program, Burnet Institute, Melbourne, Australia, ² School of Public Health and Preventive Medicine, Monash University, Melbourne, Australia, ³ Department of Gastroenterology, St Vincent's Hospital, Melbourne, Australia, ⁴Department of Medicine, University of Melbourne, Melbourne, Australia, ⁵Department of Infectious Diseases, The Alfred and Monash University, Melbourne, Australia, ⁶Cohealth, General Practice, Melbourne, Australia, ⁷North Richmond Community Health, General Practice, Melbourne, Australia, ⁸ Department of Infectious Diseases, The Alfred, Melbourne, Australia, ⁹ Department of Infectious Diseases, Monash University, Melbourne, Australia

Introduction:

Point-of-care (POC) tests for hepatitis C (HCV) can facilitate testing in novel settings, overcoming some of the barriers to care facing people who inject drugs, the population key to Australia's HCV elimination efforts. We tested the feasibility and acceptability of POC testing for HCV from the needle and syringe program (NSP) desk.

Methods:

NSP workers, community health workers (CHWs), and nurses offered POC tests to clients accessing the NSP. The POC tests used were the OraQuick HCV antibody test using saliva (20 minutes to result) and the Xpert HCV viral load using serum (105 minutes to result). Participants were offered same-day results, and booked for follow-up review for treatment work-up and linkage to care if required. Participants completed behavioural, demographic and acceptability surveys. Participants were reimbursed AUD 30 for study involvement.

Results:

A total of 174 participants completed POC testing for HCV antibodies; 150 (86%) had a reactive result and of these, 140 (93%) underwent a POC HCV RNA test, with 76 (54%) positive. 167 (96%) had been tested for HCV previously, of which only 44 (26%) reported testing within 12 months, and 73 (44%) reported their last test as RNA positive. 154 (89%) reported injecting drug use and 82 (47%) receptive sharing of any injecting equipment in the preceding six months. Of the 140 participants who underwent POC RNA testing, only 7 (5%) waited on-site to receive the result, but 104 (74%) attended follow up by study completion. Surveys showed strong support for CHW and NSP worker involvement (104/106, 98%) and preference for POC tests (97/116, 84%).

Conclusion:

Provision of POC testing through NSPs was feasible and highly acceptable to clients at high risk of HCV. Despite few participants waiting on site for RNA results, we observed a high follow-up rate, indicating successful linkage to care.

Disclosure of Interest Statement:

The authors acknowledge funding support from the Shepherd Foundation, St Vincent's Hospital Foundation, in-kind support from Cepheid Ltd., and an investigator initiated research grant from Gilead Sciences for this project. The Burnet also receives funding support from the National Health and Medical Research Council, Abbvie, GSK and Merck for investigator initiated research. The authors also acknowledge the contribution to this work of the Victorian Operational Infrastructure Support Program received by the Burnet Institute. The authors gratefully acknowledge the contribution of Rapid-EC participants, implementing sites and staff.