A COST-EFFECTIVENESS ANALYSIS OF PRIMARY VERSUS HOSPITAL-BASED SPECIALIST CARE FOR DIRECT ACTING ANTIVIRAL HEPATITIS C TREATMENT

<u>Palmer A</u>¹, Wade A^{1,2}, Draper B^{1,3}, Howell J^{1,3,4,5}, Doyle J^{1,6}, Petrie D⁷, Thompson A^{4,5}, Wilson D¹, Hellard M^{1,3,6,8,9}, Scott N^{1,3}

¹Disease Elimination Program, Burnet Institute, Melbourne, Victoria, Australia ²Department of Infectious Diseases, Barwon Health, Geelong, Victoria, Australia ³Department of Epidemiology and Preventive Medicine, Monash University, Clayton, Victoria, Australia

⁴Department of Medicine, The University of Melbourne, Parkville, Australia ⁵Department of Gastroenterology, St Vincent's Hospital Melbourne, Fitzroy, Victoria, Australia

⁶Department of Infectious Diseases, The Alfred and Monash University, Melbourne, Victoria, Australia

⁷Centre for Health Economics, Monash University, Melbourne, Victoria, Australia ⁸Peter Doherty Institute for Infection and Immunity, Parkville, Victoria, Australia ⁹School of Population and Global Health, University of Melbourne, Parkville, Victoria, Australia

Background: Hepatitis C virus elimination may be possible by scaling up direct-acting antiviral (DAA) treatment. Due to the safety and simplicity of DAA treatment, primary-based models of care are now feasible, efficacious and may be cheaper than hospital-based specialist care. The Prime Study was a randomised controlled trial comparing the uptake of DAA treatment and cure outcomes between primary and hospital-based care settings. In this paper, we use Prime Study data to estimate the cost of initiating treatment for people diagnosed with hepatitis C in primary care compared to hospital-based care.

Methods: The total economic costs associated with delivering DAA treatment (post hepatitis C diagnosis) within the Prime study – including health provider time/training, medical tests, equipment, logistics and pharmacy costs – were collected. Appointment data were used to estimate the number/type of appointments required to initiate treatment in each case, or the stage at which loss to follow up occurred.

Results: Among the hepatitis C patients randomised to be treated within primary care, 43/57 (75%) commenced treatment at a mean cost of A\$1,007 (range: A\$934-1,099) per patient initiating treatment. In hospital-based care, 18/53 hepatitis C patients (34%) commenced treatment at a mean cost of A\$2,197 (range: A\$2,127-2,469) per patient initiating treatment – more than twice as high as primary care.

Conclusion: Compared to hospital-based care, providing hepatitis C services in primary care can improve treatment uptake and reduce the costs of treatment initiation. To improve treatment uptake and cure, countries should consider primary-based care as the main model for hepatitis C treatment scale-up.

Disclosure of Interest Statement: MH, JD and the Burnet Institute receive investigator-initiated research funding from Gilead Sciences, Merck, AbbVie and Bristol-Myers Squibb (BMS). AT is an advisory board member for Gilead Sciences,

AbbVie, BMS, Merck and Roche Diagnostics, and a speaker for Gilead, Merck, BMS, AbbVie, Roche Diagnostics. AW has received investigator-initiated research funding from AbbVie. JD has received honoraria for advisory boards or speaking from Gilead Sciences, AbbVie and Merck. NS has received investigator-initiated research funding from Gilead Sciences. JH has received an Australia Fellowship (investigator-initiated project) from Gilead Sciences.