



A point-of-care HCV testing intervention to improve hepatitis C diagnosis and treatment uptake among people attending Aboriginal Community Controlled Health Services: the SCALE-C study

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Introduction

Hepatitis C Virus (HCV) poses a significant global health challenge, disproportionately affecting marginalized communities with limited healthcare access. In Australia, First Nations Peoples are a priority population for HCV elimination, with 16% of all hepatitis C notifications in 2021 reported among them.

"Strategies for hepatitis C testing and treatment in Aboriginal communities that Lead to Elimination" (SCALE-C) was a community-based "test and treat" intervention integrating point-of-care HCV testing, non-invasive liver fibrosis assessment, and linkage to care in Aboriginal Community Controlled Health Services in regional Australia.

Method

Participants were enrolled between 28 May 2019, and 21 July 2022, from four sites (2

Results



- in South Australia and 2 in New South Wales).
- At enrollment, a standardized risk assessment questionnaire guided choice of hepatitis C point-of-care test:
- Participants with no history of HCV infection, injecting drug use, incarceration, or opioid substitution therapy were considered at **no or low risk** and received an HCV antibody test, followed by an HCV RNA test if the antibody test was positive.
- Participants with a history of HCV infection, injecting drug use, incarceration, or opioid substitution therapy were considered **at risk** and received an HCV RNA test.







Figure 1. HCV infection status among SCALE-C participants, overall and stratified by key demographic and risk characteristics.

Participants who had a detectable HCV RNA or reported risk in the last 12 months were enrolled in the SCALE-C Cohort, with follow-up visits every 6 months for up to three years.

Participants with detectable HCV RNA were offered DAA therapy (sofosbuvir-velpatasvir 12 weeks or glecaprevir-pibrentasvir 8 weeks) per standard of care, with

Infection status was categorised as: never infected (anti-HCV Ab negative [n=421] OR anti-HCV Ab unknown + HCV RNA negative [n=9]), past infection (anti-HCV Ab positive, HCV RNA negative [n=50] – spontaneous clearance, n=2; HCV treatment, n=12; unknown treatment history, n=36), current infection (HCV RNA positive [n=47]), and unknown (no or inadequate testing to determine status [n=9])

a) 1 transgender person was not included; b) among those with available data

Abbreviations: Ab, antibody; F, female; HCV, hepatitis C virus; M, male; m, month(s); OAT, opioid agonist therapy; RNA, ribonucleic acid.

Table 1. Factors associated with current HCV infection among people with reported lifetime risk

acteristics					Unadjusted Analysis	
					OR (95% CI) P	
Never					-refref	-
ing drug use		Ever but not in the last month			8.89 (1.14, 69.02) 0.03	\$7
		Last 1 month			13.76 (1.79, 105.72) 0.01	2
		Never			-refref	
d agonist ther	ару	Ever but not currently				<i>9</i>
		Currently			2.03 (0.73, 5.65) 0.17	<u>'</u> 6
	62%					
		57%	100%	100%		
	n-20	r -10	n - 4 0	n-10	Abbreviations: RNA, ribonucleic acid; SVR, su	staine
	agonist thera		Ever but no Currently 62% 57%	I agonist therapy Ever but not currently Currently 62% 57% 100%	Ever but not currently Currently 62% 57% 100% 100%	Ever but not currently 2.25 (0.97, 5.24) 0.05 Currently 2.03 (0.73, 5.65) 0.17 62% 57% 100% 100% Figure 2. HCV treatment cas Abbreviations: RNA, ribonucleic acid; SVR, sus

on- and post-treatment assessments.

This analysis evaluated:

- HCV infection status among the SCALE-C population, overall and stratified by socio-demographic characteristics and risk behaviour.
- Factors associated with current HCV infection among those with known HCV status and reported lifetime risk.
- Factors associated with current HCV infection were analyzed using logistic regression.



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Risk-based screening using point-of-care HCV testing facilitated diagnosis at regional Aboriginal Community Controlled Health Services. HCV infection was associated with a history of injecting drug use.

Additional interventions are required to improve treatment uptake and reduce HCV burden among atrisk Aboriginal and Torres Strait Islander Australians, particularly people who inject drugs.

Disclosure of Interest: The study was supported by a National Health and Medical Research Council grant (1148093) and Cepheid (GeneXpert platforms). The Kirby Institute is funded by the Australian Government Department of Health and Ageing. The views expressed in this publication do not necessarily represent the position of the Australian Government.