

Acknowledgements

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- NSP attendees, staff and managers
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Disclosures

- GD is a consultant/advisor and has received research grants from Merck, Gilead, Bristol-Myers Squibb, and AbbVie.
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Monitoring exposure, treatment uptake and viraemic prevalence

Australian Needle Syringe Program Survey

 Bio-behavioural sentinel surveillance system conducted annually since 1995



Hepatitis C infection among people who inject drugs in Australia

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- DBS testing: HIV/HCV antibody



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- Self-administered questionnaire & provision of dried blood spot (DBS)
- DBS testing: HIV/HCV antibody
- Conducted at ~50 NSPs nationally
- 2000-2500 respondents per annum
- 75% metropolitan NSPs, 25% regional/remote

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 Representative of NSP attendees at sentinel sites¹



Source: 1. Topp et al, JAIDS, 2011











HCV RNA testing using dried blood spots

- Feasibility study conducted in 2012
- · Laboratory feasibility conducted by St Vincent's HIV Reference Lab/AMR
- Financial support from jurisdictions (NSW, NT, QLD, VIC & WA)
- Commenced annual ANSPS HCV RNA testing in 2015 (baseline)
- Combined HCV antibody/RNA test results and self-reported treatment uptake to determine:
 - Non-exposed
 - Spontaneous clearance
 - Treatment induced clearance (cured)
 - Active infection
- Monitor elimination efforts among people who inject drugs: from baseline (2015 pre DAAs) into the future

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^ Treatment eligible respondents: Ever exposed excluding those with spontaneous clearance

___# Post stratification weightings adjusted for previous and recent HCV treatment and gender among RNA tested sample 12



[^] Treatment eligible respondents: Ever exposed excluding those with spontaneous clearance



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Post stratification weightings adjusted for previous and recent HCV treatment and gender among RNA tested sample 14

[#] Post stratification weightings adjusted for previous and recent HCV treatment and gender among RNA tested sample



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___ # Post stratification weightings adjusted for previous and recent HCV treatment and gender among RNA tested sample 15



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 Unable to assess SVR12 → provides cross sectional serial point prevalence to enable monitoring trends over time

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Limitations

- Unable to assess SVR12 → provides cross sectional serial point prevalence to enable monitoring trends over time
- Not all respondents have sufficient DBS for HCV RNA testing → unable to report data by jurisdiction due to small sample size

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 Captured people engaged with NSP services → generalisability wider population is uncertain 	of results	to

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Conclusions

 Highlights the value and importance of conducting HCV RNA testing in surveillance projects

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- Rapid increase in treatment uptake \rightarrow decline in viraemic prevalence
- Some emerging concerns re inequity of treatment access/uptake

Conclusions

- Highlights the value and importance of conducting HCV RNA testing in surveillance projects
- Rapid increase in treatment uptake \rightarrow decline in viraemic prevalence
- Some emerging concerns re inequity of treatment access/uptake
- · Contribute to monitoring of HCV elimination efforts