



### Overview

- Hepatitis C and prisons
- Hepatitis C Incidence and Transmission Study in prisons (HITS-p)
- Nurse-led model of care
- SToP-C



THE GOOD NEWS IS THAT I HAVE FOUND A WAY FOR US TO RECEIVE FREE HEALTH CARE. THE BAD NEWS IS THAT WE'LL BE SPENDING SOME TIME IN PRISON





# Background

#### Prisons – New South Wales, Australia

- Full time custodial population ~10,000
- 30 prison centres; 800,000 Km<sup>2</sup>
- 30,000 new receptions per annum
- 150,000 movements per annum
- Short stay 24% remand; 21% sentence <6 months</li>
- Over-representation of Indigenous (20% vs 2%)
- Chronic hepatitis C prevalent 32%
- Co-morbidities prevalent
  - Mental health 49%
  - Injecting drug use (IDU) 43%

2009 NSW Inmate Health Survey

# HITS-p: incidence and predictors (n=210)

	HCV incide	ence per 100 p	person-years	Univa	riate Cox regr	ession	Multivariable Cox regression (adjusted)					
Variable	No. cases n=38	Incidence rate (%)	95% CI (%)	Hazard ratio	95% CI	$P^{a}$	Hazard ratio	95% CI	P <sup>a</sup>			
Gender												
Female	13	22.68	12.08-38.79	1								
Male	25	11.76	7.61-17.36	0.53	0.27 - 1.04	0.062						
Indigenous identity <sup>b</sup>												
No	22	10.44	6.54-15.80	1								
Yes	16	27.05	15.46-43.92	2.63	1.38 - 5.05	0.003	2.28	1.72 - 4.44	0.015			
IDU during follow-up												
No	7	4.96	1.10-10.22	1								
Yes	31	24.05	16.3-34.14	4.75	2.09 - 10.83	0.0002						
IDU daily or more often												
No	20	9.05	5.53-13.97	1								
Yes	18	36.84	21.83-58.22	4.05	2.12 - 7.76	< 0.001	2.22	1.09 - 4.52	0.028			
IDU - heroin												
No	17	7.68	4.47-12.30	1								
Yes	21	43.20	26.74-66.03	5.62	2.95 - 10.69	< 0.001	4.15	2.07 - 8.34	< 0.001			
IDU - cocaine												
No	31	12.55	8.53-17.81	1								
Yes	7	30.56	12.29-63.97	2.51	1.1 - 5.73	0.029						
IDU - methamphetamine					-112 -1112							
No	21	10.44	6.46-15.96	1								
Yes	17	24.71	14.40-39.57	2.40	1.25 - 4.59	0.008						
Break from IDU												
No	11	7.35	3.67-13.15	1								
Yes	27	22.45	14.79-32.66	2.98	1.47 - 6.04	0.024						
Sharing		-2.15	1, 52.00	,0	1							
No	21	10.52	6.51-16.09	1								
Yes	17	24.15	14.07-38.67	2.36	1.23 - 4.51	0.010						
Tattooing	-,		20.07	2.50	2001	0.010						
No	31	14.43	9.80-20.47	1								
Yes	7	12.71	5.11-26.20	0.86	0.37 - 1.98	0.731						
Bleaching- always	,	12./1	3.11 20.20	0.00	0.57 1.70	0.751						
No	28	12.27	8.15-17.73	1								
Yes	10	23.94	11.48-44.02	1.98	0.96 - 4.10	0.066						
OST	10	23.71	11.10 11.02	1.70	5.76 1.10	0.000						
No	28	12.35	8.21-17.86	1								
Yes	10	23.09	11.07-4.25	1.92	0.93 - 3.99	0.079						

Abbreviations: CI, confidence interval; OST, methadone maintenance treatment. <sup>a</sup> *P* values are 2-sided

<sup>&</sup>lt;sup>b</sup> Indigenous identity = reported Australian Aboriginal or Torres St Islander identity.





# Justice Health Hepatitis Service

- Justice Health authority separate from custodial authority
- Prison-based
- Taxpayer funded health care
- Taxpayer funded antiviral therapy (HSD S100 scheme)
- Specialist-led, nurse supported clinics
- Outcomes comparable to community standards
- <1% of those eligible, interested treated</p>

Boonwaat L et al, Medical Journal of Australia, 2010; 192:496-500



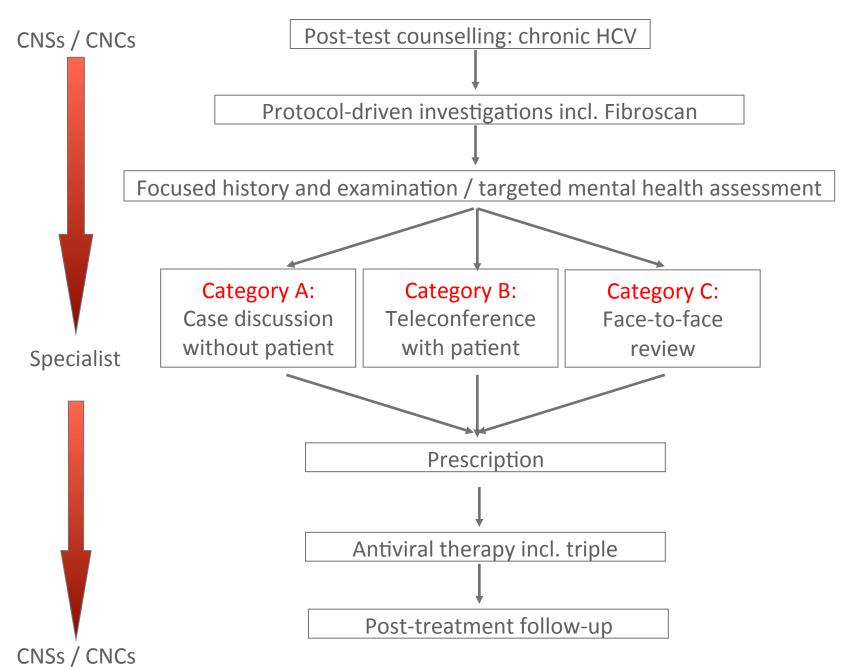


## Justice Health - Nurse-led model of care - pilot

- Lithgow, Goulburn, Long Bay
- 2009-2010
- Training of Clinical Nurse Consultants (CNCs)
- Protocol driven assessment and triage
- Qualitative and quantitative evaluation
- Safe, effective, well accepted, enhanced treatment rates

Lloyd A et al, Clinical Infectious Diseases 2013; 56(8):1078-8

### Nurse-led model of care – roll out



### Outcomes – NLMC roll out

30 months: 2011-2013

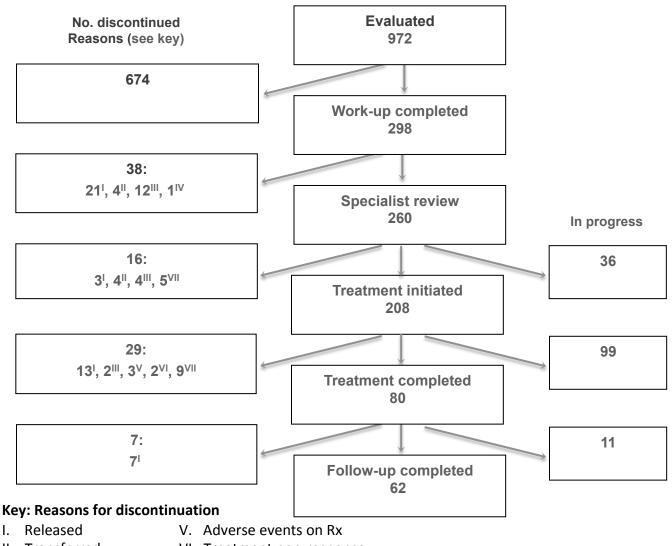
Prison sites: n=15

2 CNCs

15 CNSs (0.4FTE/site)

Portable fibroscan

Triple therapy



II. Transferred

VI. Treatment non-response

III. Not interested

VII. Other

IV. PCR negative





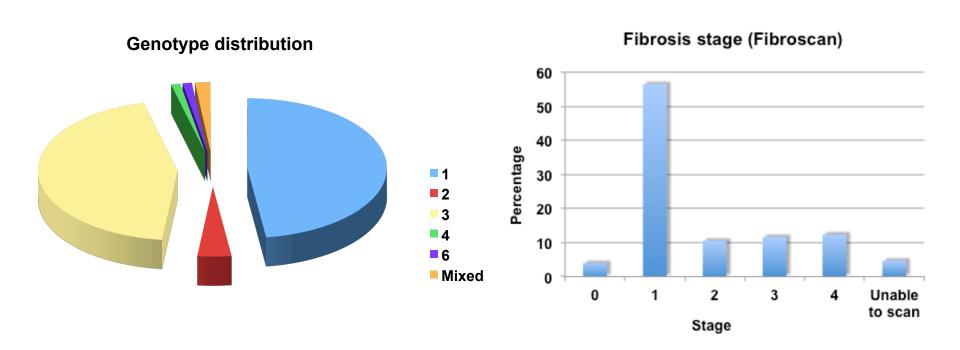
### Patient characteristics - CNC assessment (n=298)

Variable	No. (%)
Mean age; years (SD)	37 (9)
Male gender (%)	268 <sup>F</sup> (89)
Born in Australia (%)	252 (85)
Aboriginal or Torres Strait Islander (%)	80 (27)
Remand (%)	56 <sup>(</sup> (19)
Risk factors for HCV	
Lifetime - injecting drug use (IDU) (%)	283 <sup>(95)</sup>
Lifetime - tattooing (%)	250 <sup>F</sup> (84)
Recent IDU <12 mths (%)	243 <sup>(</sup> (82)
Current IDU (%)	35 <sup>₹</sup> (12)
Current methadone/buprenorphine (%)	146 <sup>F</sup> (49)
Co-morbidities	
History of execessive daily alcohol (%)	95 <sup>F</sup> (32)
History of excessive binge alcohol (%)	113 <sup><b>F</b></sup> (38)
History of major depression (%)	187 <sup><b>F</b></sup> (63)
History of anxiety disorder (%)	146 (49)
History of psychosis (%)	121 (41)
Current mood disorder (%)	70 (24)
Current psychosis (%)	10 (3)





# Patient characteristics (n=298)





### **Outcomes**

### Serious adverse events on Rx (n=28/208; 13%)

- Haematological 13; metabolic (thyroid) 8; psychiatric 4; cardiovascular
   2; gastrointestinal 1
- Hospitalisations: 2 (atrial fibrillation; myocarditis)
- Deaths: 1 (myocarditis)
- Treatment discontinuations due to serious adverse events: 3 /28 (11%)



#### **Primary objective**

 To evaluate the feasibility and impact of rapid scale-up of IFN-free HCV treatment on the incidence of HCV infection in the prison setting

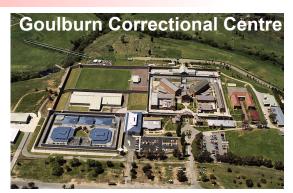
#### **Secondary objectives**

- To evaluate impact of HCV treatment on HCV prevalence
- To evaluate treatment outcomes (SVR)
- To evaluate treatment uptake and adherence
- To record HCV risk behaviors, including the impact of treatment
- To measure HCV reinfection rates following treatment
- To evaluate cost-effectiveness of scaling up HCV treatment
- To model the impact of HCV treatment in prisons on community prevalence
- To develop and framework and toolkit for roll-out across the prisons nationally



#### Phase I

- 2 maximum security prisons: active vs control
- Funding: Gilead Sciences
- Surveillance target: 80%
- Treatment target: 50%
- All infected subjects eligible
- Sofosbuvir / GS-5816
- Once daily, oral, 8-12 weeks



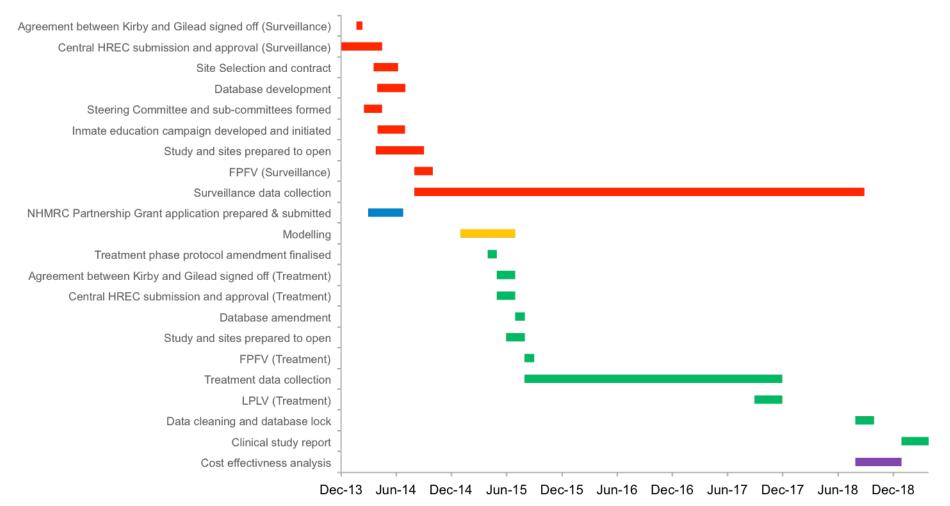


#### Implementation plan

		20	014		2015						2016			2	2017		2018			
	1 2		3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
	Start	-up																		
Lithgow			Surv	eillan	ce															
Litingow							Mod	delling							_					
							Tre	atment	scale	-up										
Goulburn			Surv	eillan	ce															
											Mo	delling							_	
								Tr	eatmen	-up										
																			Ana	lysis



#### Phase I - timelines





#### Phase II

- Rapid scale-up across network of prisons
- 3-4 medium security prison settings
- Qualitative evaluation of attitudes and barriers
- Mathematical modelling of the impact of prison-based treatment
- Cost-effectiveness and budget impact evaluation
- Development of framework and implementation toolkit for roll-out
- Funding application to NHMRC (Partnership Grant)

#### **Timelines**

		2	015				2017					2	2018			2019				
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
	Star	t-up		•	•	•	•			•	•	•	•		•	•	•	•	•	•
Roll-out			Sur	veilla	nce															
prisons 1-4							Mo	delling												
							Tre	Treatment scale-up												
							Qua	litative	eva	luatio	n				•					
Translational															Mat	hema	tical	mode	lling	
studies		Cost-effectiveness - data collection - data analysis																		
												Fra	mewo	rk an	d too	lkit				



# Future perspectives

- Taxpayer subsidised listing of direct-acting antivirals (DAAs)
- Australian burden of disease equation
  - ~250,000 currently affected
  - ~3-4,000 currently treated per annum
  - ~9-10,000 new infections per annum
- Models / venues of care key elements
  - Hepatitis- skilled nurses
  - Fibroscan
  - Triage and management protocols
  - Prisons, OST clinics, primary care
  - Liver clinics for advanced liver disease
- Treatment-as-prevention

SToP-C, TAP



# Acknowledgements

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- Fran Pekin CNC
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#### *Investigators*

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