



# Liver Disease Burden and Clinical Follow-Up During a Liver Health Promotion Intervention Integrating Non-Invasive Liver Disease Screening in Drug and Alcohol Settings: The LiveRLife Study

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## Partners



# Developing the campaign

## **PHASE I: Message Development**

- Targeted focus groups with community peers
- Assess knowledge, attitudes & beliefs about liver disease, testing and treatment

## **PHASE II: Message Testing**

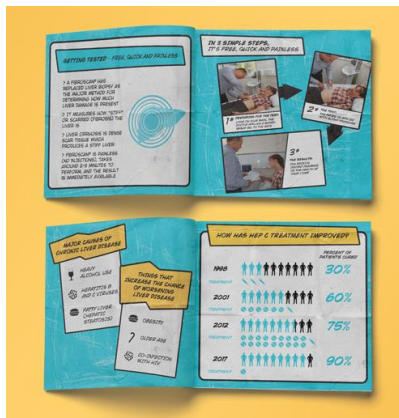
- Focus test the messaging and resources with community peers

## **OUTCOME**

- Target one achievable behavior

# LiveRLife resources

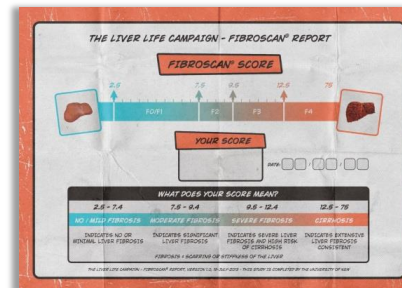
## ✓ PRINTED RESOURCE



## ✓ POSTER CAMPAIGN



## ✓ FIBROSCAN REPORT

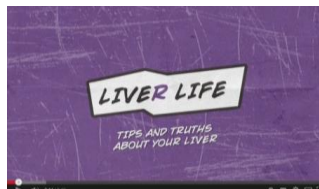


## ✓ STUDY WEBSITE

LIVERLIFE.ORG.AU



## ✓ SHORT FILM



# Developing the campaign

## PHASE III: Campaign Implementation

To evaluate the impact of a healthy liver campaign on liver disease knowledge, assessment and treatment among people attending drug & alcohol services

### Inclusion

- ☒ ≥18yrs of age
- ☒ History of injecting drug use

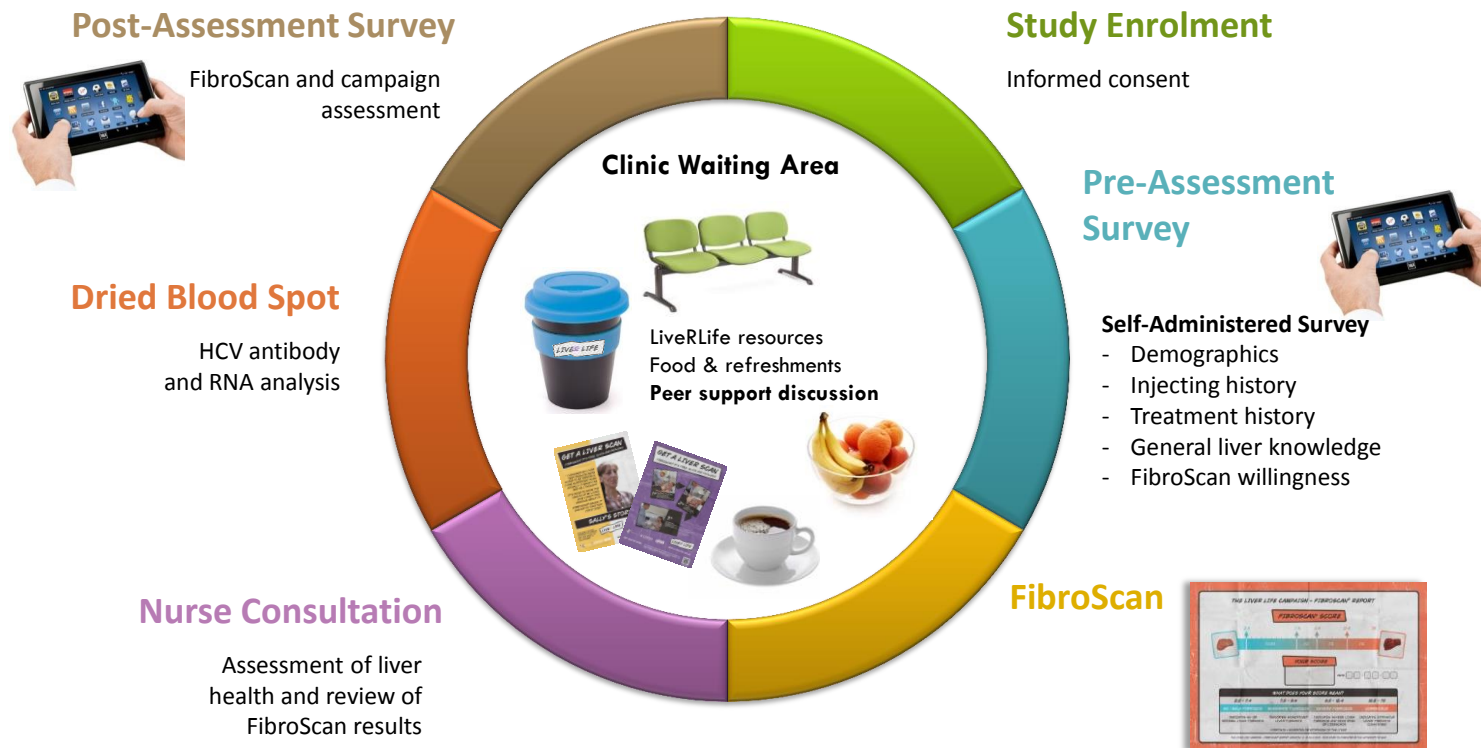
### Exclusion

- ☒ Pregnant women

### Recruitment

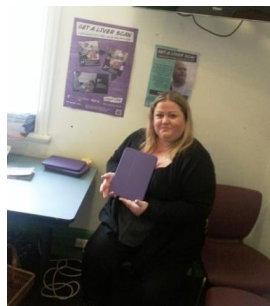
- Through one community-based primary health care clinic, two opioid substitution treatment clinics, and one medically supervised injecting centre in New South Wales, Australia

# Enhanced liver disease assessment – FibroScan®



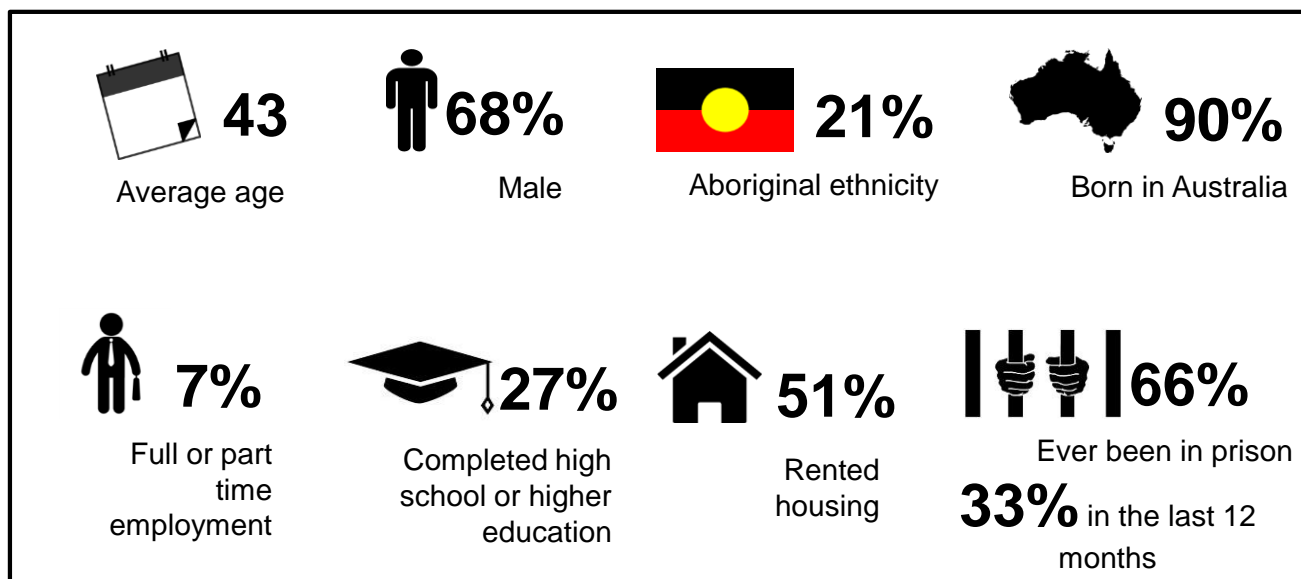
## Campaign days

- A team of staff attended each campaign day
- Support from the service was key to building interest and participation
- Clients were keen to participate and have their liver health assessed
- An opportunity to engage individuals with significant disease staging



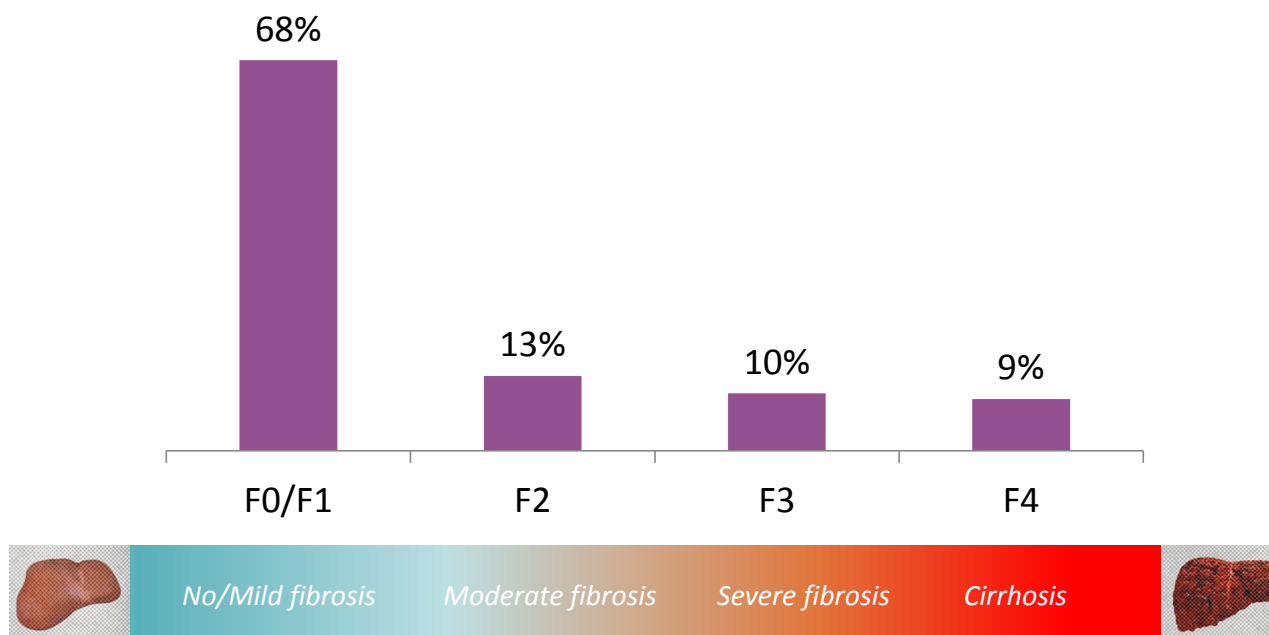
## Participant characteristics

- LiveRLife has been run at 4 clinics (n=253)





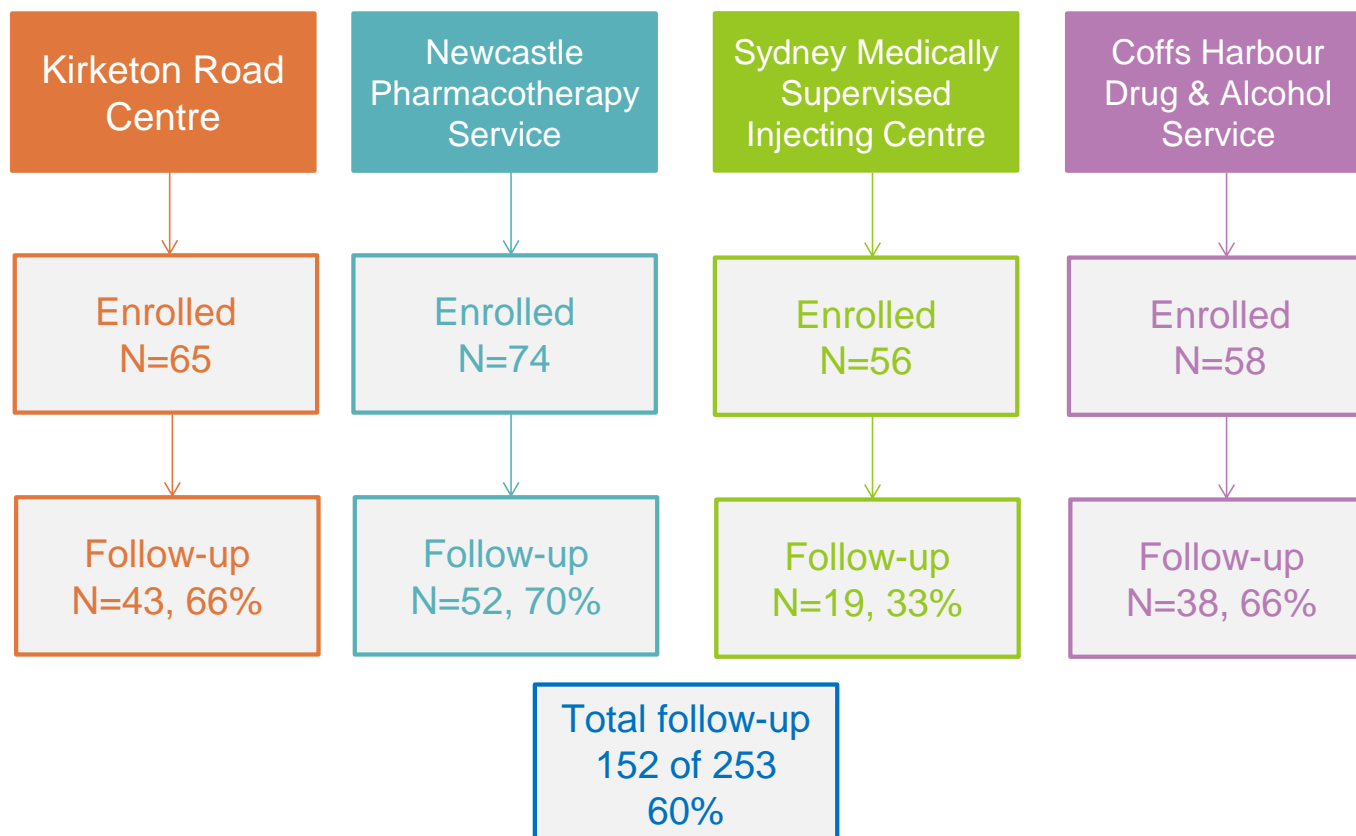
# Disease staging



## Factors associated with F3/4 disease staging

	Number with F3/4 (%)	Unadjusted OR (95% CI)	<i>P</i>
<b>Age</b>			
<35 years	3 (5%)	1.00	-
>=35-<45 years	13 (16%)	3.69 (1.00, 13.58)	0.050
≥45 years	29 (31%)	8.48 (2.45, 29.31)	0.001
<b>Gender</b>			
Female	9 (13%)	1.00	-
Male	34 (21%)	1.82 (0.82, 4.02)	0.142
<b>HCV RNA</b>			
Undetectable	9 (13%)	1.00	-
Detectable	35 (23%)	2.09 (0.95, 4.63)	0.068

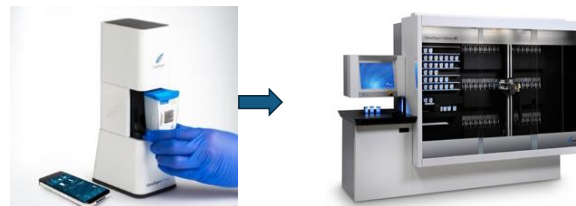
## Clinical follow-up



# Xpert® HCV Viral Load point-of-care assay



- ✓ • Automated, self-contained, single use, random access
- ✓ • European CE-IVD approved (plasma)
- ✓ • Single platform for integration (HIV, HPV, TB)
- ✓ • Minimal training, rapid (60-105min), capillary blood (alpha testing)
- ✓ • Multiple configurations



## Method: Venous blood and finger-prick samples

### Venous whole blood



- 1 Collect venous whole blood by venepuncture



- 2 Centrifuge



- 3 Load plasma into Xpert® HCV Viral load (RUO) cartridge



- 4 Result < 100min

### Finger-prick capillary blood (interim)



- 1 Collect 100µL capillary blood by finger-prick into a Minivette



- 2 Load blood into Xpert® HCV Viral load (RUO) cartridge



- 3 Add 1mL dilution buffer



- 4 Result < 100min

**Gold standard comparator: Abbott RealTime HCV assay, v7, m2000**

## Results: Sensitivity and specificity (detectable)

Abbott plasma	Xpert® HCV VL plasma		Total
	Undetected -	Detected +	
Undetected -	114	1	115
Detected +	0	51	51
Total	114	52	166
Sensitivity	<b>100%</b> (95%CI, 93-100%)		
Specificity	<b>99.1%</b> (95%CI, 95.3-100%)		

### One discrepant result:

1201-61410-018 Abbott = 0 Xpert = 3,380,000



Abbott plasma	Xpert® HCV VL finger-prick		Total
	Undetected -	Detected +	
Undetected -	111	2	113
Detected +	2	47	49
Total	113	49	162
Sensitivity	<b>95.9%</b> (95%CI, 86-99.5%)		
Specificity	<b>98.2%</b> (95%CI, 93.8-99.8%)		

### Four discrepant results:

1201-61410-018 Abbott = 0 Xpert = 7,686,000

1201-61249-030 Abbott = 38 Xpert = 0

1201-61249-104 Abbott = 0 Xpert = 5 (<110)

1201-61223-002 Abbott = <12 Xpert = 0

## Results: Sensitivity and specificity (quantifiable)

	Xpert® HCV VL plasma		
Abbott plasma	Unquantifiable	Quantifiable	Total
Unquantifiable	114	0	114
Quantifiable	1	48	49
Total	115	48	163
Sensitivity	98% (95%CI, 89.1-99.9%)		
Specificity	100% (95%CI, 96.8-100%)		

**Note: Outlier excluded**

**One discrepant result:**

1201-61249-030 Abbott = 38, Xpert = <10

	Xpert® HCV VL finger-prick		
Abbott plasma	Unquantifiable	Quantifiable	Total
Unquantifiable	112	0	112
Quantifiable	1	46	47
Total	113	46	159
Sensitivity	97.9% (95%CI, 88.7-99.9%)		
Specificity	100% (95%CI, 96.8-100%)		

**One discrepant result:**

1201-61249-030 Abbott = 38, Xpert = 0

**Excluding those on treatment, n=10 (69, 4.8%)**

Sensitivity	97% (95%CI, 88-99.9%)
Specificity	100% (95%CI, 96.7-100%)

Sensitivity	97.7% (95%CI, 87.7-99.9%)
Specificity	100% (95%CI, 96.7-100%)

## Conclusions

- Demonstrated considerable liver disease burden in this population
- A high proportion attended post-LiveRLife clinical follow-up
- Provided an opportunity to address other health issues (e.g. HAV/HBV vaccinations)
- Developed key partnerships between services, clinical providers, and researchers
- Demonstrated the feasibility of interventions to enhance health outcomes among people in drug and alcohol settings

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*It's good to know the health of my liver, now I don't feel anxious about it. FibroScan makes it easier to take that first step!*

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### SALLY'S STORY





## Future directions

- Additional 250 participants have been recruited from homelessness settings, drug and alcohol clinics and NSPs (including POC HCV RNA testing) in Australia
- Planned project to evaluate LiveRLife in Bangkok, Thailand in collaboration with HIV-NAT
- Simplified LiveRLife intervention planned to increase testing, linkage to care and DAA therapy for ETHOS-II study (to begin in March 2017)

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MERCK  
*Be well*