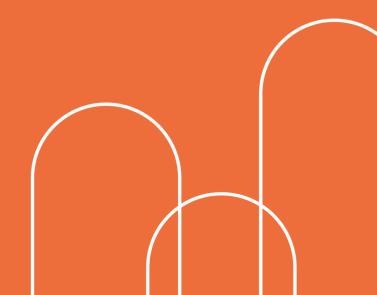
Supervised Injecting Room Cohort Study

Ash Stewart on behalf of the SIRX study team





At Burnet Institute, we proudly acknowledge the Boon Wurrung people of the Kulin Nations as the Traditional Custodians of the land on which our office is located. We pay our respect to Elders past and present, and extend that respect to all First Nations people.





Overview

- Early evidence & the Insite evaluation
- The Melbourne MSIR
- Findings from the MSIR Reviews
- The SIRX Cohort Study



Early evidence & the Insite evaluation





History of SIFs

- Established in the mid-1980s
- In response to epidemics of public injecting, overdose, and HIV transmission
- At the end of 2024, SIFs were legally operating in 18 countries, globally





SIFs aim to

REDUCE

- Fatal and non-fatal overdose
- Blood borne virus transmission
- Ambulance attendance and hospital admissions for overdose
- Public injecting and unsafe disposal of syringes

IMPROVE

- Linkage to health and social services
- Public amenity (poorly defined and rarely applied to other health services¹)



Early SIF studies

- SIF were operating in many European cities and in Sydney (Aus) by the early 2000s
- Various reports credited SIFs with several public health and community benefits



Findings from early SIF studies



OPIOID OVERDOSE

Observed reductions in overdose^{1,2}



BLOOD BORNE VIRUSES

Reductions in BBV risk behaviours (needle sharing & condomless sex)^{4–7}



SERVICE ACCESS

Improved health of SIF clients was observed.^{7,8} Ceasing SIF use associated with obtaining stable housing.⁹



PUBLIC AMENITY

Observed reductions in public injecting.^{8,10}
Fewer negative encounters with residents or police in the area surrounding the SIF⁹



CLIENT SATISFACTION

Acceptability and uptake by people who inject drugs¹⁰ Reasons for not using SIFs were opening hours, wait times, & distance⁸



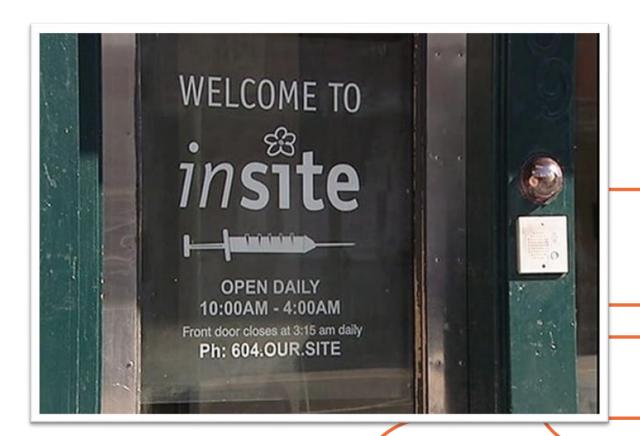
Findings from early SIF studies

- No funding allocated to comprehensive evaluations
- Unavailability of cohort data to examine the impact of SIFs
- Ecological data limited by discerning between SIFs and concurrent changes
 - Policy changes (drug law reform)
 - Harm reduction initiatives
 - Drug markets



Insite, Vancouver

- In September 2003, Vancouver opened North America's first government sanctioned SIF pilot study¹
- This was conditional on the implementation of a rigorous scientific evaluation of the health and social impacts of the SIF

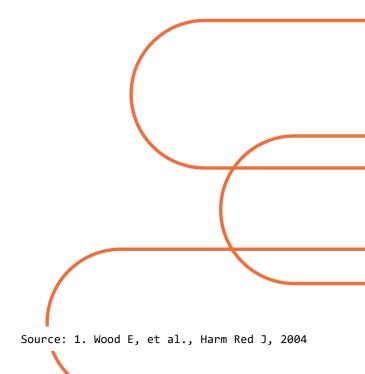




Evaluating Insite, Vancouver

STUDY DESIGN

- Prospective cohort study design
- Longitudinal measurement of outcomes
 - Blood-borne virus infections
 - Overdose incidence
 - Risk behaviours
 - Drug use practices (public drug use)
 - Health service use





Evaluating Insite – The cohorts





Community Health and Safety Evaluation

Data linkage cohort of Downtown Eastside residents



VIDUS

The Vancouver Injection Drug Users Study

Semi-annual blood testing and survey, and data linkage



SEOSI

The Scientific Evaluation of Supervised Injecting

Insite clients - service use survey and data linkage



Findings from Insite

- A reduction in overdose deaths and risk of death
- Increases in the uptake of detox services
- Increased entry into drug treatment
- SIF clients had a reduction in syringe sharing
- Public injecting and discarded syringes declined
- Increases in treatment for injecting-related skin infections
- Wide acceptance of the service and success in attracting a particularly high-risk population



Summary of findings from the evaluation of a pilot medically supervised safer injecting facility

Evan Wood, Mark W. Tyndall, Julio S. Montaner, Thomas Kerr

∞ See related article page 1395

ABSTRACT

In many cities, infectious disease and overdose epidemics are occurring among illicit injection drug users (IDUs). To reduce these concerns, Vancouver opened a supervised safer injecting facility in September 2003. Within the facility, people inject pre-obtained illicit drugs under the supervision of medical staff. The program was granted a legal exemption by the Canadian government on the condition that a 3-year scientific evaluation of its impacts be conducted. In this review, we summarize the findings from evaluations in those 3 years, including characteristics of IDUs at the facility, public injection drug use and publicly discarded syringes, HIV risk behaviour, use of addiction treatment services and other community resources, and drug-related crime rates. Vancouver's safer injecting facility has been associated with an array of community and public health benefits without evidence of adverse impacts. These findings should be useful to other cities considering supervised injecting facilities and to governments considering regulating their use.

CMAJ 2006;175(11):1399-404

any cities are experiencing infectious disease and overdose epidemics as a result of illicit injection drug use, ¹⁻⁴ an activity that is also associated with a number of negative community impacts, including public drug use. ⁵⁻⁶ Despite these harms, innovative public health programs for reducing health and community concerns remain highly controversial in North America and other settings where HIV infection is spreading rapidly among injection drug users (IDUs). ⁷⁻⁹

In Canada, Vancouver has been an epicentre of drug-

federal government that allowed operation of the facility was limited to 3 years and was granted on the condition that an external 3-year scientific evaluation of its impacts be conducted. Given the controversial nature of the program, 14 stakeholders agreed that all findings from the evaluation, including this report, should be externally peer-reviewed and published in the medical literature before dissemination. In this review we report on the 3 years' findings.

Program and evaluation methods

As described previously, 13 the Vancouver safer injecting facility has 12 injection stalls where IDUs inject pre-obtained illicit drugs under the supervision of nurses. Nurses respond to overdoses and address other health needs (e.g., treating injection-site abscesses), and the facility has an addiction counsellor and support staff who seek to meet the needs of IDUs or refer them to appropriate community resources (e.g., housing services, addiction treatment). 13

Although the best strategy for evaluating the safer injecting facility would be to randomly assign IDUs to either full access or no access to the program, interventional study designs for the evaluation of such facilities have been deemed unethical; thus, the evaluation of the Vancouver facility was structured primarily around prospective cohort studies involving IDUs who used the facility and those who did not. In accordance with the Transparent Reporting of Evaluations with Nonrandomized Designs (TREND) criteria for observational research, to a detailed description of the evaluation









Findings from the evaluation of

Vancouver's Pilot Medically Supervised Safer Injecting Facility – Insite

Revised June 2009

American Journal of Preventive Medicine



REVIEW ARTICLE

Supervised Injection Facilities as Harm Reduction: A Systematic Review

Timothy W. Levengood, MPH, Grace H. Yoon, MS, Melissa J. Davoust, MSc, 1 Shannon N. Ogden, MPH, Brandon D.L. Marshall, PhD, Sean R. Cahill, PhD, 1,3 Angela R. Bazzi, Phn4,5

Other studies have come from

Context: Supervised injection facilities are harm reduction inject drugs to use previously obtained substances under Australia (SYD), Spain, & Norway Although currently considered illegal under U.S. federal 14W, SEVE implementing supervised injection facilities anyway as a response to 16 OF THE 22 STUDIES

The objective of this review is to determine the effectiveness of sup pared with that of control conditions, for harm reduction and comm

WERE FROM CANADA **Evidence Acquisition:** Studies were identified from 2 sources:

examining supervised injection facility-induced benefits and harn January 2014) and an updated search using the same search strategy (January 2014-September 2019). Systematic review methods developed by the Guide to Community Preventive Services were used (screening and analysis, September 2019-December 2020).

Evidence Synthesis: A total of 22 studies were included in this review: 16 focused on 1 supervised



Melbourne Medically Supervised Injecting Rooms





What happened?

- The Labour government first proposed a SIF trial in 1999
- Drug Policy Expert Committee Report: community support for a SIF trial injecting was high
- Opposing resident groups (e.g. Footscray Matters) claimed the government was ignoring community views
- Major CBD Traders Myer & David Jones threatened to leave if city didn't become a drug free area
- By December 2000, the Victorian Government abandoned the trial and increased funding for drug treatment services – the 'Saving Lives' strategy
- In 2010 the drug market moved to North Richmond
- Between 2009 2017 over 1,300 people died from fatal overdoses with heroin (most with other drugs)
- In 2017, the Victorian coroner recommends a SIF trial

Timeline



Source: Canberra Times, 2017

The North Richmond
MSIR opens on a twoyear trial basis

2018 – June



MSIR service review released
'Hamilton review'
Trial is extended for three
most years to gather further
evidence of its effectiveness

2020 – June



2017

Victorian government announces plans for a trial of the MSIR in North Richmond Source: RACGP, 2019



2019 – July

Purpose built facility opens



2023 - Feb

MSIR service review released 'Ryan review'

Timeline

North Richmond MSIR announced as an ongoing service and commitment for a CBD SIF trial

2023 – March



Source: Canberra Times, 2017

Report assessing the feasibility of a CBD site was released.

2024 - April



Source: ABC News, 2024

NRCH successful in recommissioning process and announced as operators of the MSIR

2024 - April

Source: VIC Gov, 2023

Strengthening MSIR To Keep **Saving Lives**

Monday 6 March 2023

BREAKING Victoria's Medically Supervised Injecting Boom (MSIR) in North Riv will become an ongoing service, with critical support services ext. **VEWS** often an independent review Found the trial has soved 31 lives successfully managing almost 6,000 overdoses, taking a ospitals and reducing ambulance call outs.

The Victorian Government will today introduce legislation to establish he health service as an angoing service – with immediate measures to be taken to further boost safety and amenity in the North Richmond

2023 - Aug

Victorian government announces Ken Lay to undertake an independent review for the CBD SIF site



2024 - April

The Victorian Government withdraws plans to proceed with the CBD trial.





Resident reveals reality near Richmond injecting after Victorian governm decision to scrap CBD

Victorian government scraps plans for a sec supervised injecting room in Melbourne

By state political reporters Richard Wi

Melbourne's safe inj upsetting locals

A second safe injecting room for Melbourne has been rejected. Here's how we got here

The Victorian government announced a new trial facility will not proceed after concluding there was no site that could balance the needs

drug users with the broader of **SW3NE:**

Sydney Today

22 °/ 29 °



Residents 'at a loss' over Melbourne's saf injecting room being made permanent

THE **INJECTING**

Entertainment Technology

Serena Seyfort • Afternoon Editor 7:29am Mar 8, 2023

Lifestyle > Health > Health Problems

Melbourne injecting room to allow ice users to s up

DRUG users will be permitted to shoot up ice in a new injecting room despite the government previous

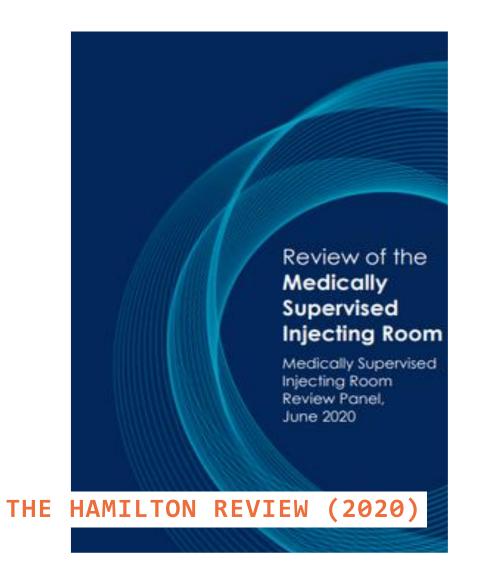


Findings from the MSIR Reviews





MSIR reviews



Review of the Medically Supervised Injecting Room FINAL REPORT: Key findings and recommendations February 2023 THE RYAN REVIEW (2023)



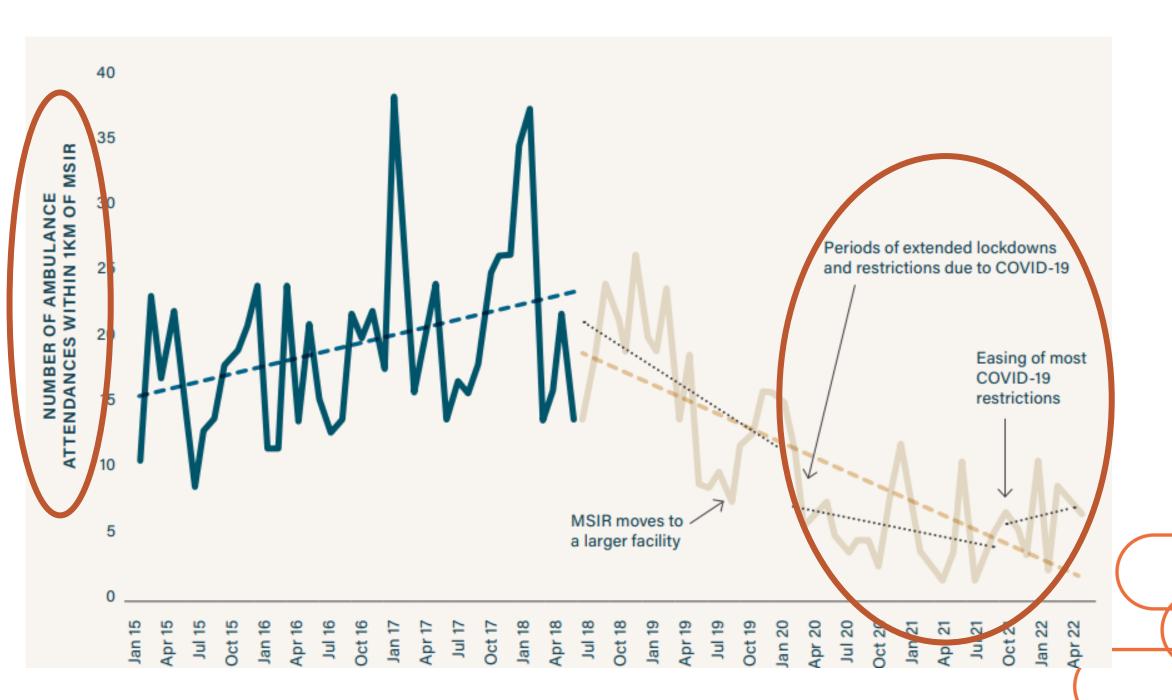
Methodology

ECOLOGICAL STUDIES

- Coroners court data
- Emergency department and hospital admissions
- Ambulance data
- Needle collection data
- BBV notifications (Ryan Review)

OBSERVATIONAL STUDIES

- MSIR service data
- Staff/service user consultations/surveys
- Surveys with residents/business owners for 'public amenity'
- SuperMIX cohort data (Hamilton Review)







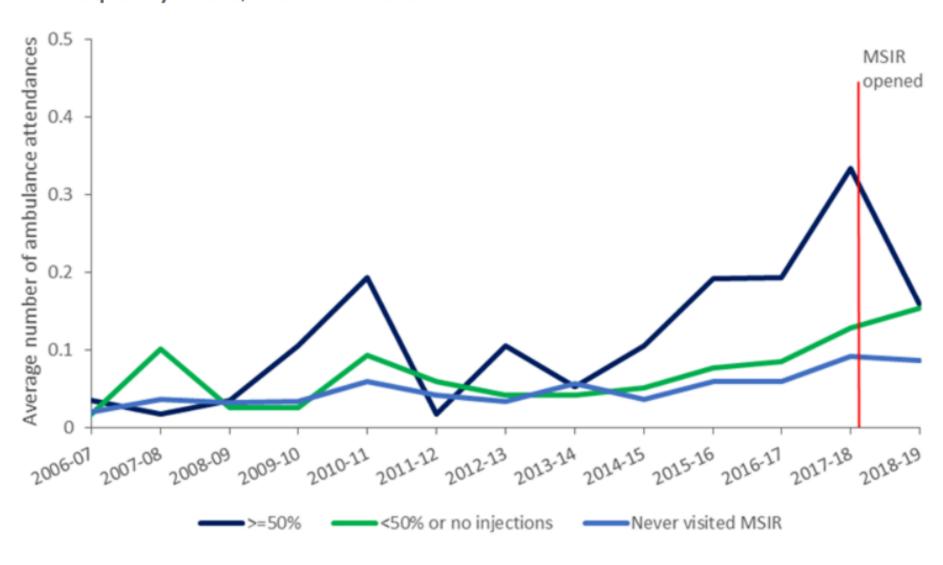
SuperMIX study

- The Melbourne Injecting Drug User Cohort Study
- Community recruited cohort of people who inject drugs in Melbourne
- Established in 2008 ongoing
- Participant surveys at baseline and annual follow-up
- Sociodemographics, drug use and behaviours, drug market characteristics, health and health service use
- Blood specimens (HIV, HCV, HBV)
- Administrative data linkage to health and social databases



Figure 31: Average number of ambulance attendances with naloxone administration per year by MSIR frequency of use, 2006–07 to 2018–19





Source: Burnet Institute 2019, p. 38



American Journal of Preventive Medicine

RESEARCH ARTICLE

The Melbourne Safe Injecting Room Attracted People Most in Need of Its Service



Wijnand Van Den Boom, PhD,¹ Maria del Mar Ouiroga. PhD.^{1,2} Dagnachew Muluye Fetene. PhD.¹ Paul A. Agius, MSc,^{1,3} Peter G. Higo Mark A. S

Introduction: In 2018, the fit officially opened. This study a were socially vulnerable, and who bidity and mortality risk.

People who used the MSIR were more likely to report homelessness, unemployment, daily injecting, injecting in public, and past year imprisonment.

Methods: This was a cross-sectional analysis of the frequency of Medically Supervised Injecting Room use during the first 18 months after opening (July 2018—December 2019) among 658 people who inject drugs participating in the Melbourne Injecting Drug User Cohort Study (SuperMIX). To examine the differences between no Medically Supervised Injecting Room use, infrequent use (<50% injections within the facility), and frequent use (≥50% of injections within the facility), RRRs were estimated using bivariate multinomial logistic regression analyses and postestimation Wald tests. Analyses were conducted in 2020.

Results: A total of 451 participants (68%) reported no Medically Supervised Injecting Room use, 142 (22%) reported infrequent use, and 65 (10%) reported frequent use. Participants who reported either





MSIR Reviews

- Review findings were largely drawn from ecological data sources
 - Difficult to discern whether changes were due to pandemic or the MSIR
- Main outcomes from SuperMIX analyses were assessed to 2019, limiting the timeframe to evaluate and observe effects
- Methodology used to generate findings for the Ryan review was not clearly presented
 - SuperMIX was not asked to contribute to this review



The Supervised Injecting Room Cohort Study (SIRX)



Study design

- SIRX draws on the methodology used to evaluate Insite SIF in Vancouver
- Cohort and quasi-experimental design to measure varying levels of exposure to the Melbourne MSIRs and their on-site services across a range of outcomes
- SIRX is made up of two cohort studies



The cohorts

THE MELBOURNE INJECTING DRUG USER COHORT STUDY (SUPERMIX)

- Community-recruited people who inject drugs
- Annual surveys
- Blood tests (HIV, HCV, HBV)
- Administrative data linkage to MSIR database, health and social databases

THE SIRX-REGISTRATION COHORT STUDY (SIRX-R)

- Registered clients of the MSIR
- Once-off survey (nofollow-up)
- Administrative data linkage to MSIR database, health and social databases



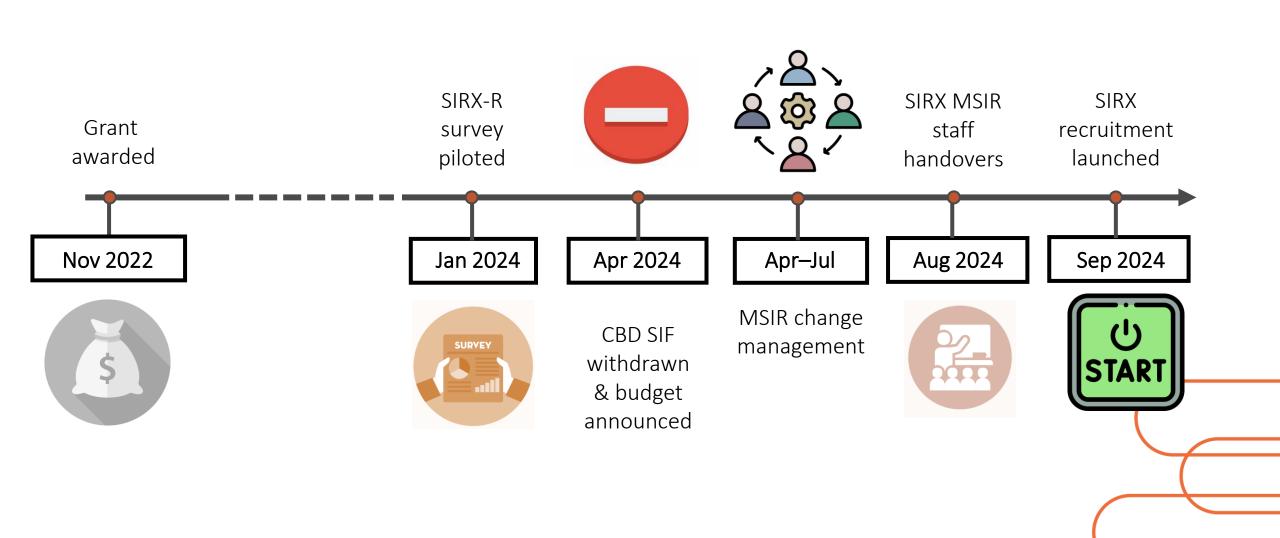
The study design

SIRX-R N=1,800 1,200 North Richmond 600 CBD

Ongoing record linkage (all participants), 2022/23 - 2027 440 North Richmond SIRX-R SIRX-R participants who consent to 400 CBD (Registration record linkage only Cohort) SuperMIX/ SIRX ongoing cohort SuperMIX participants who are also MSIR clients **SuperMIX** Cohort SuperMIX participants who don't use the MSIR Ongoing questionnaires and interviews, 2022/23 - 2027



Study timeline





Participant recruitment



694

Participants in SuperMIX that report ever using the MSIR 71% men



RECENT INTERVIEW

586

Reported MSIR use in most recent interview.

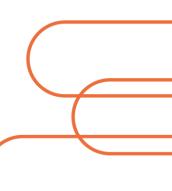
67% were interviews conducted in 2023/24



29

Participants enrolled into SIRX-R

*polit period





Critical evidence

PRIMARY OUTCOMES

- Drug-related and allcause mortality
- Ambulance attendance for opioid overdose

SECONDARY OUTCOMES

- Hospitalisations for skin and soft-tissue infections
- Opiate agonist treatment uptake
- Use of non-acute health services





Critical evidence

ECONOMIC EVALUATION

- Total annual cost savings
 - E.g., MSIR running costs vs. averted healthcare costs (e.g., ambulance attendances for overdose and for blood borne viruses and skin infections)
- The cost per life saved
- The cost per quality adjusted life year gained
 - OAT uptake, treatment of co-morbidities (HCV), changes to employment, and averted deaths

BMJ Open Supervised Injecting Room Cohort Study (SIRX): study protocol

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ABSTRACT

Background Supervised injecting facilities (SIFs) are designed to reduce the harms associated with injecting drug use and improve access to health and support services for people who need them. The Supervised Injecting Room Cohort Study (SIRX) aims to provide evidence of the effects, including cost-effectiveness, of a SIF embedded within a community health service, the Melbourne Medically Supervised Injecting Room (MSIR), which has a range of integrated harm reduction, health and social support services on-site.

Methods and analysis The SIRX study design involves two prospective cohort studies that collect behavioural data and retrospectively and prospectively linked administrative data for primary and tertiary health services, criminal justice records, and mortality. The two cohorts are: (1) participants drawn from the existing Melbourne Injecting Drug User Cohort Study (SuperMIX; established in 2008–ongoing) through which participants consent to annual behavioural surveys (including serological testing for MIV and hepatitis R and C viruses) and linkage to

STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ The Supervised Injecting Room Cohort Study (SIRX) uses a cohort and quasi-experimental design to measure varying levels of exposure to the Melbourne Medically Supervised Injecting Room (MSIR) and its on-site services and the effect of these exposure levels across a range of outcomes.
- ⇒ Comprehensive longitudinal behavioural data and linkage to MSIR visits and routinely collected administrative health and social databases.
- ⇒ The SuperMIX Cohort may be subject to attrition bias through lost-to-follow-up; however, data linkage for primary outcomes minuses this risk.
- Consistent with all observational studies, confounding may impact the observed associations, but casual inference methods are being applied to minimise this.

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(en short infographic summaries) for participants will be



Acknowledgments: SIRX Team

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