

Good practices of integrated POC testing for HCV, HIV other STIs and TB with harm reduction strategies for people who use drugs

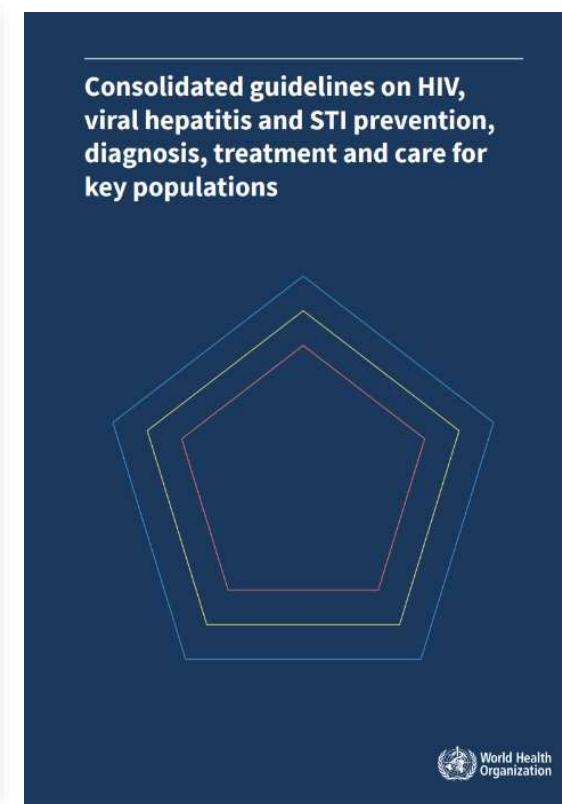
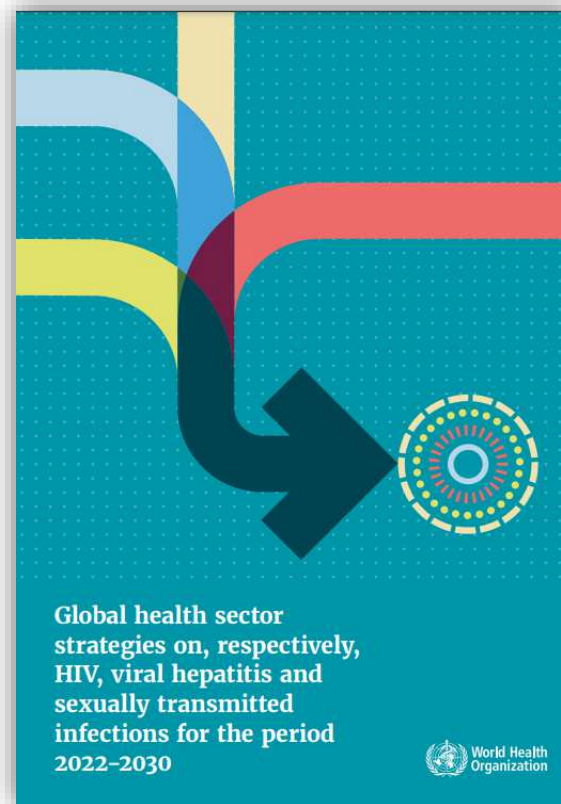
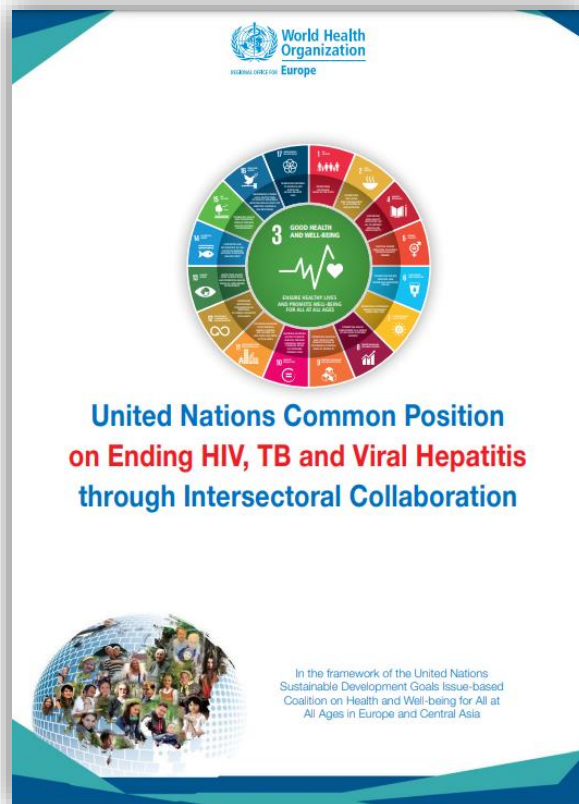
Elisa Martró

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Microbiology Department, Germans Trias i Pujol University Hospital
CIBER in Epidemiology and Public Health (CIBERESP)



Integration of strategies

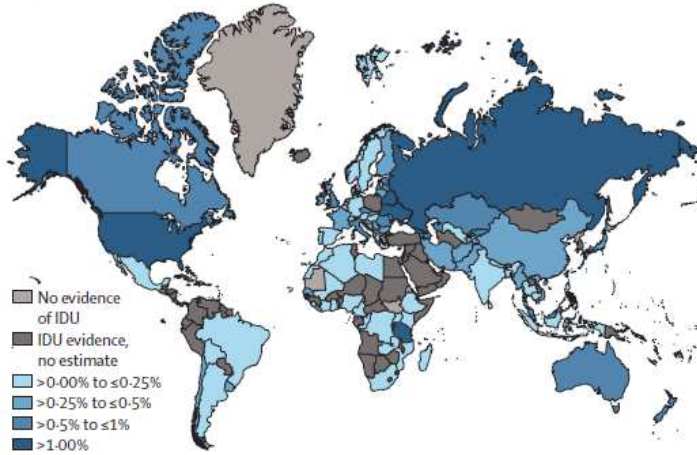
- The global objective under target 3.3 of the United Nations Sustainable Development Goals (SDGs) seeks to **end by 2030 the epidemics of AIDS, tuberculosis and combat hepatitis**, among others
- In **key populations**, such as people who use drugs, **point-of-care (PoC) tests** and **simplified diagnostic algorithms** are required, with **decentralized care and treatment**.



Burden of blood-borne viruses in people who inject drugs (systematic review)

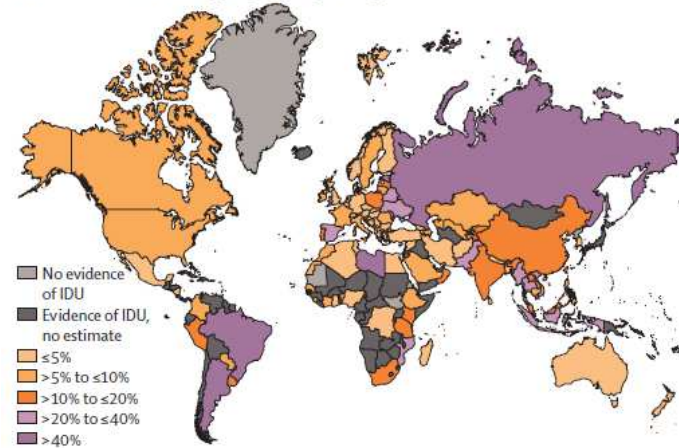
IDU
14.8 M

A Prevalence of IDU



HIV
15.2%
2.3 M

B Prevalence of HIV among people who inject drugs



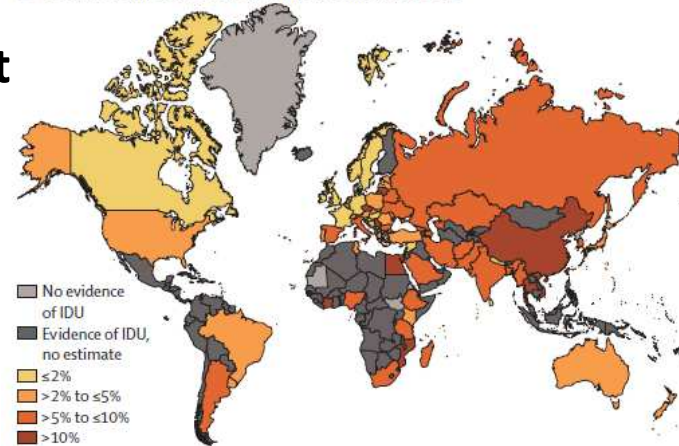
Current HCV
38.8%
5.8 M

C Prevalence of current HCV infection among people who inject drugs



Current HBV
8.4%
1.2 M

D Prevalence of HBV infection among people who inject drugs



- **24.8%** had experienced recent **homelessness or unstable housing**
- **58.4%** had a lifetime history of **incarceration**
- **14.9%** (95% CI, 8.1–24.3) had recently engaged in **sex work**

HIV and HCV prevalence remain high globally requiring improvements in **harm reduction coverage** and prevention of drug-related harm (including **provision of antiviral treatment and care**)

Recommended package for people who inject drugs

Essential for impact: health interventions

Prevention of HIV, viral hepatitis and STIs

Harm reduction (NSPs, OAMT and naloxone for overdose management)
Condoms and lubricant
Pre-exposure prophylaxis for HIV²⁴
Post-exposure prophylaxis for HIV and STIs
Prevention of vertical transmission of HIV, syphilis and HBV
Hepatitis B vaccination
Addressing chemsex

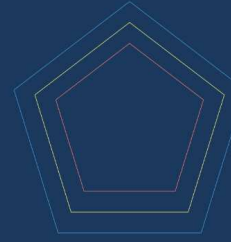
Diagnosis

HIV testing
STI testing
Hepatitis B and C testing

Treatment

HIV treatment
Screening, diagnosis, treatment and prevention of HIV associated TB
STI treatment
HBV and HCV treatment

Consolidated guidelines on HIV, viral hepatitis and STI prevention, diagnosis, treatment and care for key populations



Jul 2022



Essential for impact: enabling interventions

Removing punitive laws, policies and practices
Reducing stigma and discrimination
Community empowerment
Addressing violence

Essential for broader health: health interventions

Conception and pregnancy care
Contraception
Mental health
Prevention, assessment and treatment of cervical cancer
Safe abortion
Screening and treatment for hazardous and harmful alcohol and other substance use
TB prevention, screening, diagnosis and treatment

- **High prevalence of blood-borne viruses**
- **Increased risk of TB**, irrespective of their HIV status, a leading cause of HIV-related mortality
- **May be at increased risk of STIs** (particularly those engaging in chemsex or those using stimulants); global estimates not known

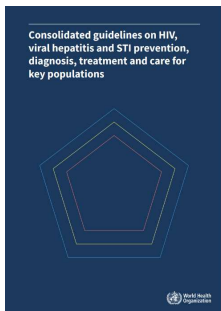
Service delivery

- To improve **access**, **acceptability** and **availability** of services for key populations (from **prevention** and **diagnosis** to **treatment** and **care**)

**Community-led
services**

Task sharing: peers,
nurses, outreach workers

Integration:
“one-top-shop”



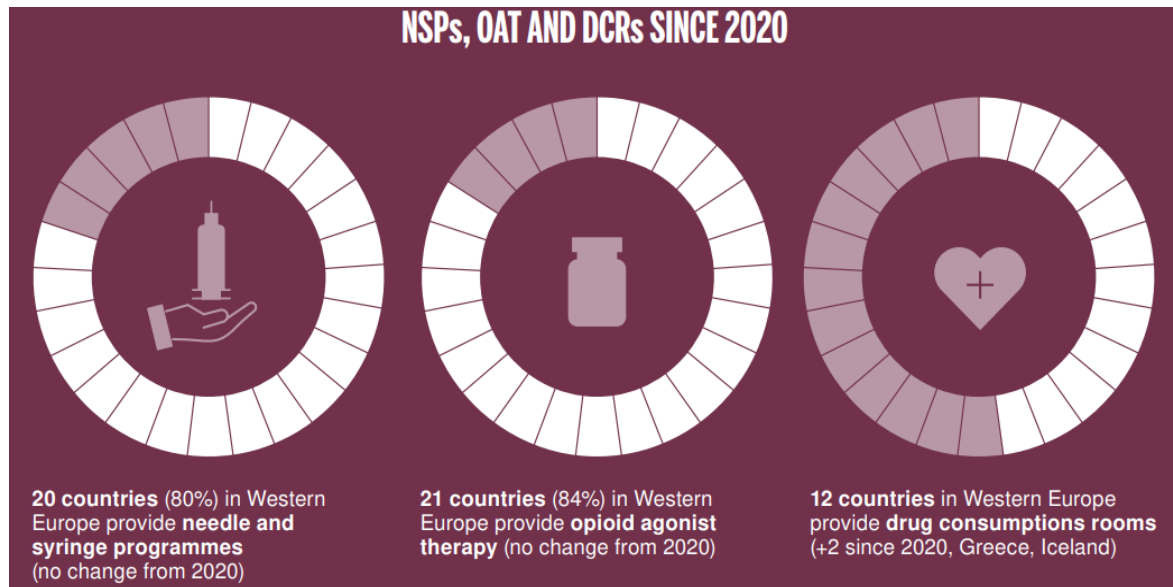
Harm reduction: the opportunity of integrated screening and care

Preventing major public and individual health harms (HIV, viral hepatitis, overdose) without necessarily stopping drug use:

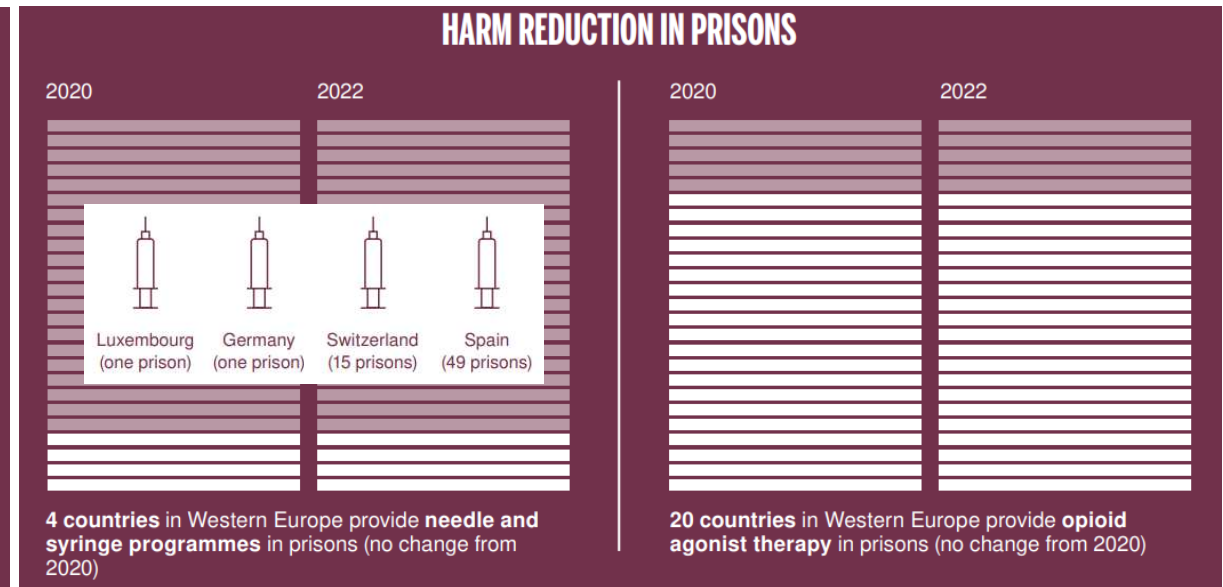
- **needle and syringe programmes (NSPs)**
- **drug consumption rooms (DCRs)**
- **opioid agonist therapy (OAT)**
- **naloxone programmes** for overdose management

Community centers, prisons, pharmacies, outreach settings (**mobile units**)

Missing in many countries
Western Europe: long tradition of harm reduction but **uneven distribution of services within countries** (rural areas underserved)



Only Spain, Luxembourg and Norway meet the WHO targets (200 syringes/p/y and 40% of people who use opioids on OAT)



Predominantly only available to people who started OAT before going to prison

Available PoC tests for HCV, HIV other STIs and TB



Rapid diagnostic tests (RDTs) for point-of-care (PoC) testing

Blood-borne viruses, single RDTs:

- **HIV:** Ab/Ag (4th gen.) in whole blood, oral fluid
- **HCV:** HCV-Ab in whole blood, oral fluid
- **HBV:** HBsAg in whole blood



Molecular confirmatory tests (viral load)

*Regional approval,
WHO prequalification*
https://extranet.who.int/prequal/sites/default/files/document_files/231020_prequalified_IVD_products_list.pdf

Multiplex RDTs:

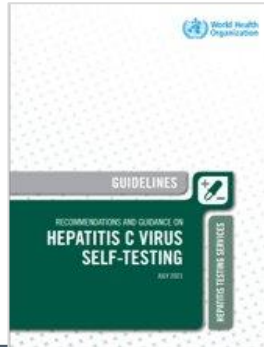
Test	Manufacturer
Detect 3 HIV/HCV/HBV combo kit	Artron Laboratories (Canada)
Triplex HIV, HCV, HBsAg	Biosynex (France)
Hep B, Hep C, HIV Combination Rapid Test	Maternova (US)
Multiplo HBc/HIV/HCV	MedMira (Canada)
HBsAg/HCV Ab Rapid Test	Spectrum Diagnostics (Egypt)
Rapid HBsAg/HCV/HIV/Syphilis Combo	Euro Genomas (Lithuania)
OnSite HBsAg/HCV Ab Rapid Test	CTK Biotech (US)
COMBIQUIC HIV/HCV	Qualpro Diagnostics (India)
TriQuick HIV/HCV/HCV	Genlantis Diagnostics

Field validation is needed to assess diagnostic accuracy of these tests





Source: E. Ivanova, FINIS

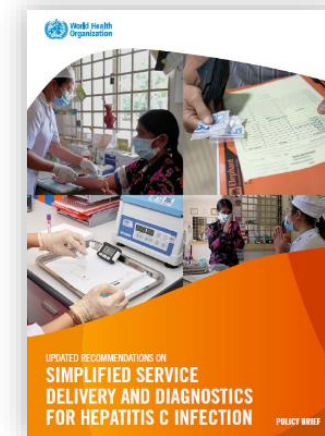
Self-testing:

Company	HCV self-test status	Specimen type	HCV RDT for prof use
OraSure Inc, US	WHO PQ submission HIV	Oral fluids	CE, FDA, WHO PQ
Premier Medical Corporation, US/India	Clinical studies completed, WHO PQ submission in prep	Blood	CE + submitted to WHO PQ (under review)
bioLytical (Canada)	Studies ongoing HIV	Blood	WHO PQ under review
Wondfo	In development HIV	Blood	WHO PQ in prep
Abbott Rapid Diagnostics Korea; SD Biosensor	Development plans	Blood	WHO PQ



Molecular near-PoC tests

PLATFORM	Xpert HCV VL assay 	Xpert HCV Fingerstick VL assay 	GeneDrive HCV ID assay Discontinued 	Truenat™ HCV assay 
SAMPLE TYPE	Plasma	Capillary blood	Plasma	Plasma, serum, capillary blood
SENSITIVITY	100%	98%	99%	95%
SPECIFICITY	97%	99%	100%	99%
SAMPLE PREP	Integrated	Integrated	Off-board (several pipetting steps)	Separate kit and instrument for sample prep
TIME TO RESULT	110 min	60 min	90 min	20 min (sample prep), 40 min (analysis)
REGULATORY STATUS	CE-IVD, WHO PQ	CE-IVD, WHO PQ	CE-IVD, WHO PQ	Approved in India. Submitted to CE-IVD
POWER SUPPLY	Need electricity supply		Need electricity supply	Battery integrated
DATA ANALYSIS	PC		Integrated	Integrated
TEST MENU	TB, HIV, HBV, CV19 and many others		CV19, MTB/RIF in development	TB, HIV, CV19 and many others
TEST COST	US\$ 14.95 ex works (HBDC price)		\$25-30	Not disclosed
INSTRUMENT COST	US\$ 17,500 (4 module)		US\$ 5,000*	US\$ 18,000 (4 tests at a time)**



HCV DIAGNOSTICS: USE OF POINT-OF-CARE (POC) HCV RNA ASSAYS FOR DETECTION OF HCV VIRAEMIC INFECTION TO GUIDE TREATMENT, AND AS TEST OF CURE

<https://www.who.int/publications/i/item/9789240052697>

→ **“Test & Treat” strategies**

Source: E. Ivanova, FIND

HCV: Intermediate-high RNA prevalence (50-74%) → RNA screening recommended (one-step)

Scott N, et al. *J Viral Hepat* 2018; 25(12):1472-1480

Importance of PoC testing for HCV-RNA

Studies in people who inject drugs and/or were homeless (28); people incarcerated in prison (4); general or mixed populations (4); and in people living with HIV (4)



Lower turnaround time between HCV-Ab screening and treatment initiation:

Onsite PoC: **19 d** [95% CI 14–53]

vs.

Laboratory-based PoC: **64 d** [64–64]

Laboratory-based SoC: **67 d** [50–67]

Higher treatment uptake:

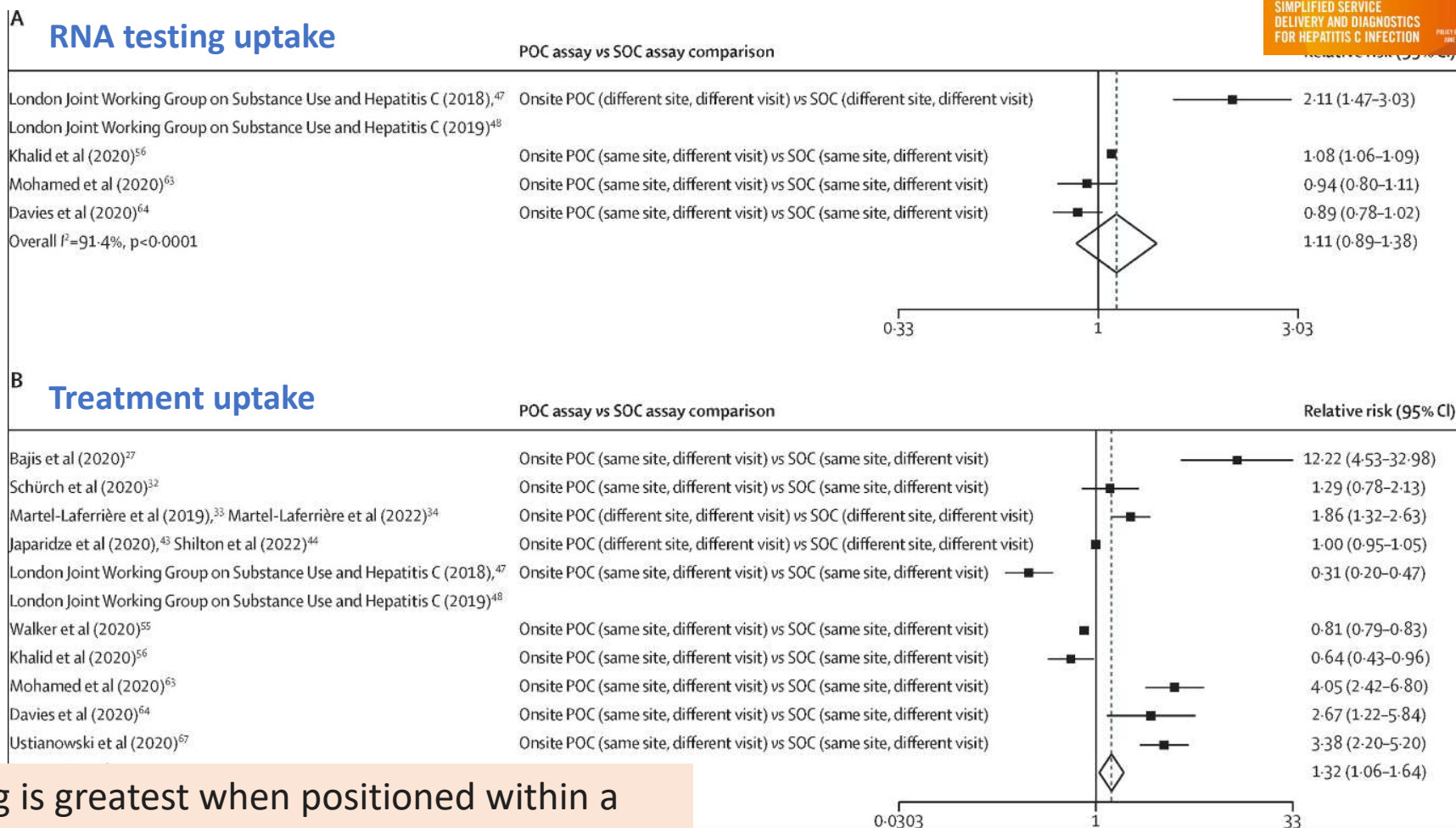
Onsite PoC: **77%** [95% CI 72–83]

Mobile PoC: **81%** [60–97]

vs.

SoC assays: **53%** [31–75]

The effect of POC viral load testing is greatest when positioned within a simplified care model in which **testing and treatment are provided at the same site** and, where possible, on the **same day**.



Molecular near-PoC tests

SD Biosensor	Bioneer
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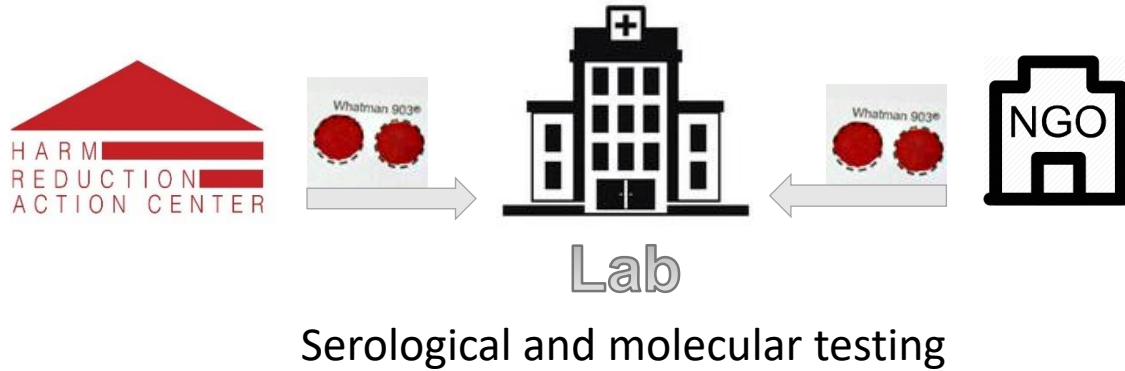
Value to Market	Established company with large manufacturing capacity and strong global health focus with a number of assays in the pipeline. Can run isothermal and PCR assays	Several assays in late development stage, high multiplexing capacity, strong global health focus
	COVID/Flu/RSV, TB, Zika/Dengue/ChikV, HIV, HCV, HBV VL, HPV,	COVID/Flu/RSV, TB, HCV,
Timeline	HCV in development	HCV expected in
Data analysis	Integrated	Integrated

Molecular true-PoC tests in the pipeline

	Minute Molecular, DASH	Mirai Genomics, GenPad	PlusLife, Mini Dock	ThermoFisher Scientific, Accula™
Sample preparation	Chemical lysis, RNA filtering†	Chemical lysis, RNA filtering‡	Thermal and chemical lysis	Thermal and chemical lysis
Amplification method	RT-qPCR	Smart Amp, proprietary isothermal technology	RHAM (proprietary isothermal technology)	RT-PCR
Turnaround time	15 min	40 min	15-35min	30 min
Tests menu (commercially available)	SARS-COV2	SARS-COV2 SARS-COV2/Flu A/B	SARS-COV2 SARS COV2/Flu A/B Monkeypox (RUO)	SARS-COV2 Flu A/B
Tests in development	HCV, HIV, STDs, Flu	Strep A, STDs	HPV, HCV, M. tuberculosis, Strep A, STDs	N/A

During the **COVID-19** pandemic, countries expanded molecular testing capacity
This infrastructure should now be repurposed to diagnose VH/HIV/STIs/TB

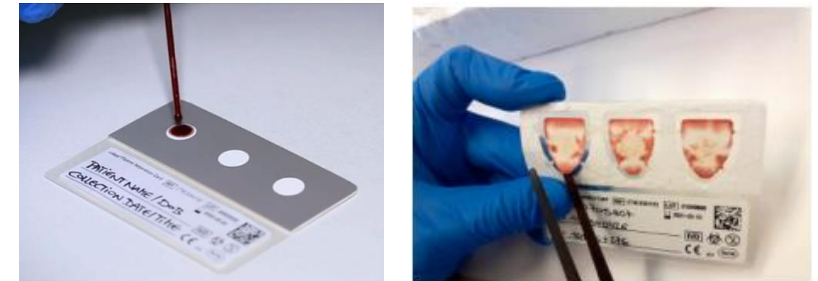
Dried blood spots (DBS) testing: decentralised sample collection



Conventional dried blood spots:
CE-IVD for **HCV** and **HIV** viral load
(Abbott Molecular Inc)



Plasma separation cards:
CE-IVD for **HIV** viral load (Roche Diagnostics)



Velásquez-Orozco F, et al. *Diagnostics* 2021

The Author(s) *BMC Infectious Diseases* 2017, **17**(Suppl 1):700
DOI 10.1186/s12879-017-2777-y

BMC Infectious Diseases

RESEARCH Open Access

Diagnostic accuracy of serological diagnosis of hepatitis C and B using dried blood spot samples (DBS): two systematic reviews and meta-analyses

Berit Lange^{1,2*}, Jennifer Cohn³, Teri Roberts⁴, Johannes Camp¹, Jeanne Chauffour⁵, Nina Gummad⁶, Azumi Ishizaki⁷, Anupriya Nagarathnam⁸, Edouard Tuailon^{9,10}, Philippe van de Perre^{9,10}, Christine Pichler¹¹, Philippa Easterbrook² and Claudia M. Denlinger⁴

The Author(s) *BMC Infectious Diseases* 2017, **17**(Suppl 1):693
DOI 10.1186/s12879-017-2776-z

BMC Infectious Diseases

RESEARCH Open Access

Diagnostic accuracy of detection and quantification of HBV-DNA and HCV-RNA using dried blood spot (DBS) samples – a systematic review and meta-analysis

Berit Lange^{1,2*}, Teri Roberts³, Jennifer Cohn⁴, Jamie Greenman⁴, Johannes Camp¹, Azumi Ishizaki⁵, Luke Messac⁴, Edouard Tuailon^{6,7}, Philippe van de Perre^{6,7}, Christine Pichler¹, Claudia M. Denlinger³ and Philippa Easterbrook²

PLOS MEDICINE

RESEARCH ARTICLE

The performance of using dried blood spot specimens for HIV-1 viral load testing: A systematic review and meta-analysis

Lara Vojnov^{1*}, Sergio Carmona², Clement Zeh³, Jessica Markby⁴, Debrah Boeras⁵, Marta R. Prescott⁶, Anthony L. H. Mayne⁷, Souleymane Sawadogo⁸, Christiane Adje-Toure⁹, Guoging Zhang⁹, Mercedes Perez Gonzalez⁴, Wendy S. Stevens^{2,8}, Meg Doherty¹⁰, Chunfu Yang¹¹, Heather Alexander¹², Trevor F. Peter¹, John Nkengasong¹³, the DBS for VL Diagnostics Investigation Consortium⁴

The Journal of Infectious Diseases

MAJOR ARTICLE

Diagnostic Accuracy of Assays Using Point-of-Care Testing or Dried Blood Spot Samples for the Determination of Hepatitis C Virus RNA: A Systematic Review

Beth Catlett^{1,2*}, Behzad Hajarizadeh¹, Evan Cunningham¹, Brett Wolfson-Stofko³, Alice Wheeler⁴, Benazir Khandaker-Hussain⁴, Jordan J. Feld⁵, Elisa Martz^{6,7}, Stéphane Chevaliez⁷, Jean-Michel Pawlotsky⁷, Chrianna Bharat⁷, Philip H. Cunningham^{1,2}, Gregory J. Dore⁸, Tanya Applegate⁹, and Jason Grebely¹⁰

PoC tests for STIs

- **Syphilis RDTs:**

Treponemal tests: Ab to *Treponema pallidum* (10-30 min), cannot distinguish active/past treated infections.

First dual **treponemal and non-treponemal** RDT:

DPP Syphilis Screen and Confirm Assay (Chembio Diagnostic Systems), 15-20 min, whole blood

- **Combined HIV/syphilis RDTs**

- **Chlamydia and gonorrhea RDTs:** suboptimal sensitivity, improved assays needed

- **Molecular near-PoC tests for *Chlamydia trachomatis*, *Neisseria gonorrhoeae*, *Trichomonas vaginalis*,** require equipment (35-90 min), urine/swab samples

Platform	Technology	<i>C. trachomatis</i>	<i>N. gonorrhoeae</i>	<i>C. trachomatis/N. gonorrhoeae</i>	<i>T. vaginalis</i>
Available platforms/assays					
GeneXpert Cepheid	PCR-based NAAT	N/A	N/A	√ CE-IVD FDA	√ CE-IVD FDA
Solana QuidelOrtho	iNAAT-HDA	N/A	N/A	N/A	√ FDA CE-IVD
careHPV System QIAGEN	Nucleic acid hybridization	N/A	N/A	N/A	N/A
Truelab RT micro PCR Molbio	Real-time PCR	√ CE-IVD	√ CE-IVD	√ CE-IVD	√ CE-IVD
io Diagnostic System binx health	PCR-based NAAT; electrochemical detection	N/A	N/A	√ FDA CE-IVD	
EasyNAT Ustar	iNAAT – CPA	N/A	√ CE-IVD	√	√ CE-IVD
Visby Medical	PCR-based NAAT	N/A	N/A	√ CT/NG/TV FDA	√
HG Swift HiberGene Diagnostics	Isothermal LAMP; Fluorometric detection	N/A	N/A	√ CE-IVD	N/A
Genie II & Genie III eazyplex Amplex/OptiGene	iNAAT; fluorescence	√ CE-IVD		√ Plus <i>C. trachomatis</i> , <i>N. gonorrhoeae</i> , <i>U. urealyticum</i> , <i>M. hominis</i> , <i>M. genitalium</i> and <i>T. pallidum</i> combo test CE-IVD	√
Vivalytic Randox/Bosch	PCR-based NAAT	N/A	N/A	√ <i>C. trachomatis/N. gonorrhoeae/T. vaginalis/M. genitalium/HSV-1 and HSV-2</i> , plus CE-IVD	√

Tuberculosis

Screening for LTBI

Confined to those at high risk of progressing to disease and who will benefit from chemoprophylaxis (e.g.: PLWH)

Tuberculin skin test (TST)

- high proportion of **FN** and **FP** results (BCG vaccination, NTM)
- subjective interpretation
- a need for a second visit (48-96 h)



Interferon- γ release assays (IGRAs):

- higher sensitivity and specificity
- only one visit (lab testing)



Limitations (both tests): **cannot accurately differentiate between LTBI / active TB**, nor reactivation / reinfection

In high-incidence settings, the focus of prevention and control is on **identifying and treating active TB cases**.



Screening for active TB

In populations in which TB screening is recommended, **systematic screening for TB disease** may be conducted using (alone or in combination):

WHO consolidated guidelines on tuberculosis
Module 2: Screening
Systematic screening for tuberculosis disease

2021



Symptom screen



Chest X-ray



Molecular WHO-recommended rapid diagnostic tests (mWRDs)

→ Community-wide systematic screening using an accurate screening and diagnostic algorithm may be used in settings with a TB prevalence of 0.5% and higher, based on new evidence of public health benefit.

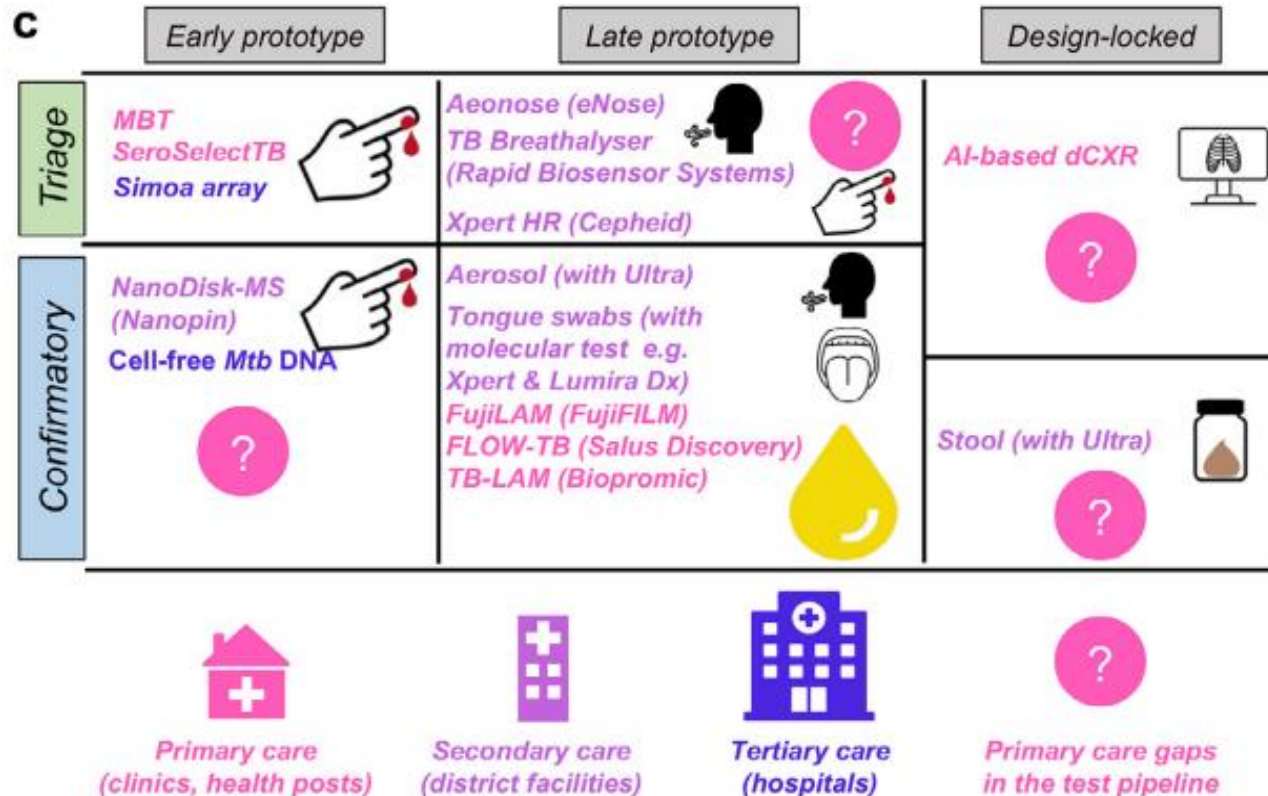
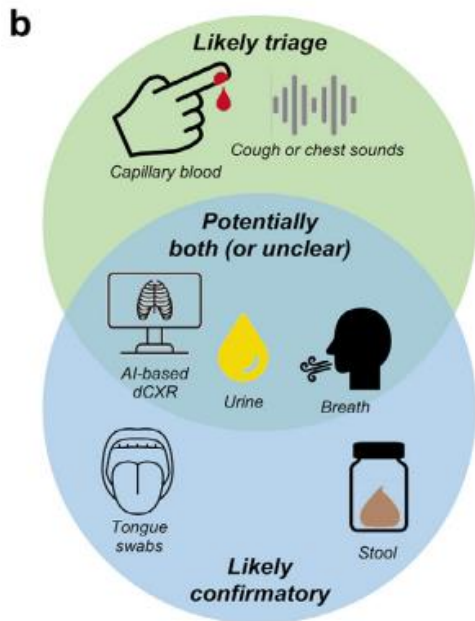
Screening and testing in **communities by mobile teams**. Needs **confirmation by a diagnostic test and clinical evaluation**.

**Sputum collection generates infectious aerosols, should only be performed at a distance from other people, preferably in open spaces, or in rooms with negative pressure and adequate air exchange.*

Tuberculosis: Need for sputum-free tests for decentralised testing

Moving from **passive case-finding** to **active case-finding (systematic screening)** for TB disease

- Sputum: difficult to obtain in high-risk groups, such as PLWH, and early disease patients;
- High proportion in surveys of **a- or pre-symptomatic cases (subclinical TB)** missed by current symptom screening approaches, may be responsible for more than half of TB transmission.



Insufficient late-stage tests useful for:

- **facility-based PoC testing**
- **community-based testing** (diagnostically and operationally more challenging)

Examples of good practices

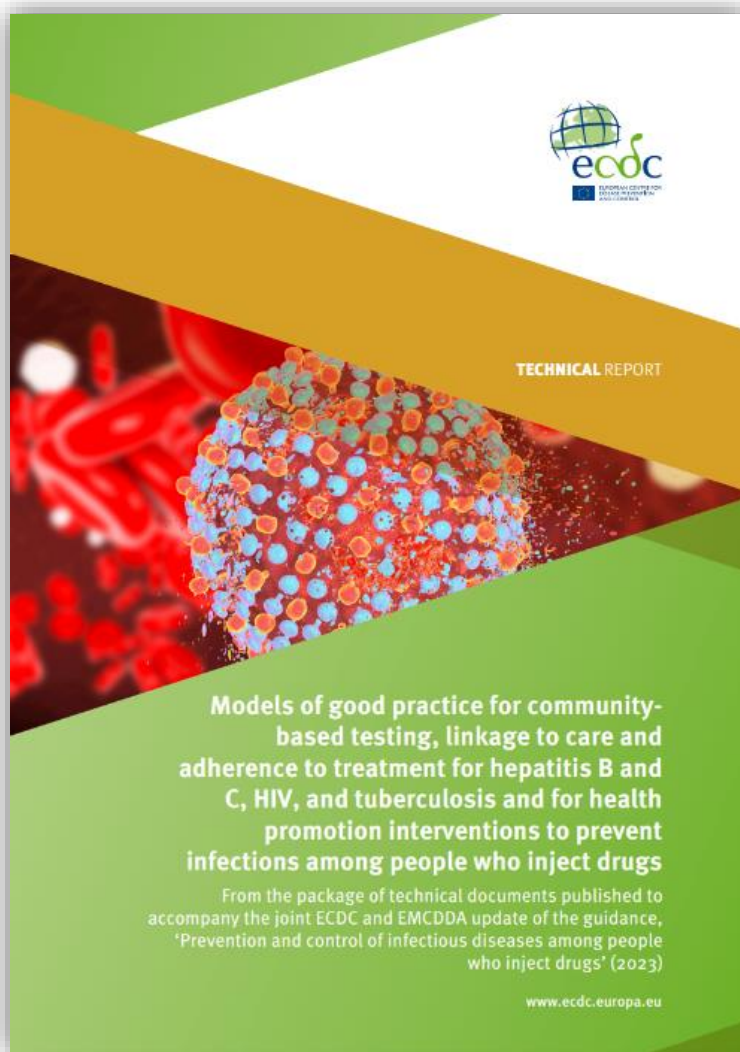


Community-based VH/HIV/TB models of good practice in people who inject drugs emerged among the 12 projects/programmes:

- **Peer involvement (7/12):**
 - ✓ to enhance community-based testing (e.g. outreach peer support, peer-to-peer recruiting, involvement in PoC testing)
 - ✓ to increase linkage to care (e.g. peer navigators supporting referrals, outreach tracing of those not in care)
 - ✓ to increase adherence to treatment (e.g. keep regular contact during treatment).
- **Integration of nurses** in the treatment cascade (5/12)
- **Multidisciplinary approach**, cooperation between drug services and specialised healthcare services often using a **low-threshold approach** (in particular to increase linkage to care).



<http://www.peerinvolvement.eu/>



<https://www.ecdc.europa.eu/en/publications-data/models-good-practice-community-based-testing-linkage-care-and-adherence-treatment>

Find & Treat (London, UK)



1. Active case finding of people with active tuberculosis (homeless people, drug or alcohol users, vulnerable migrants and people who have been in prison)

- **Multidisciplinary team:** former TB patients as peer advocates, TB nurse specialists, social and outreach workers, radiographers and expert technicians.
- Screen 8,000 high risk people every year using a mobile **digital X-ray unit** (12.5% onward referral)
- Supports Public Health England to manage TB outbreaks nationally



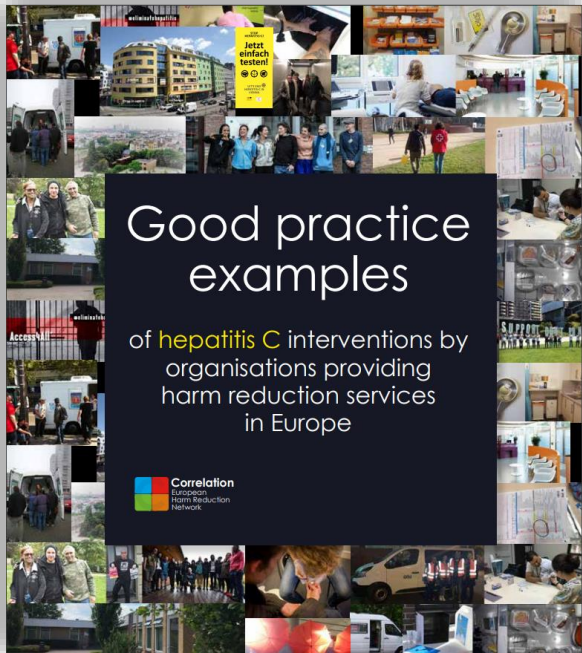
Digital X-ray
GeneXpert and RDTs
Fibroscan
Screening & treatment

2. Extended to BBVs: trained peer support workers (PSWs) + clinical team (nurses, medical staff, social worker)

- **HCV confirmatory testing and treatment initiation on the same day** and DAAs delivery and post-treatment tests are performed on-site. Complex cases are accompanied to specialist treatment services.
- **Video Supported Care** via smart phone with a secure app that is used to monitor treatment adherence and other healthcare support interventions.

3. Expansion to outreach testing activities for **HBV** and **HIV**.

Partners: **harm reduction providers** and **NGOs** (homeless and drug services) in developing screening interventions, **peers** in developing the models



https://www.correlation-net.org/wp-content/uploads/2019/09/good-practice_-example_web.pdf

Portugal APDES

Where - Services provided:

low-threshold drug service and counselling; opioid substitution therapy (OST); outreach/street-work; mobile unit.

Keywords: Treatment on site | mobile unit | paid peer workers,



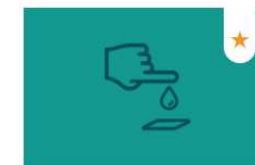
- **On-site HIV, HCV, HBV and syphilis testing** and counseling, including **rapid diagnostic tests, DBS, venipuncture**, and external HCV core antigen assay as well as fibroscan-elastography.
- Two GIRUGaia staff, a nurse and a harm reduction worker, conduct the tests.
- Clients can receive daily treatment in the Combined Therapy Programme on-site, enabling **HIV and tuberculosis treatment** and **psychiatric medicines** to be dispensed at the same time at GiruGaia.



Welcome to our **Intervention Toolkit**, developed to inform and inspire global hepatitis C testing, diagnosis, linkage to care and treatment for people who use drugs.

How-to guides

Below you will discover practical guides on how to implement evidence-based HCV interventions in your service, with more How-To Guides coming soon.



Dried Blood Spot (DBS) Testing for Hepatitis C



Hepatitis C Care Navigation



Medical Record Auditing for Hepatitis C



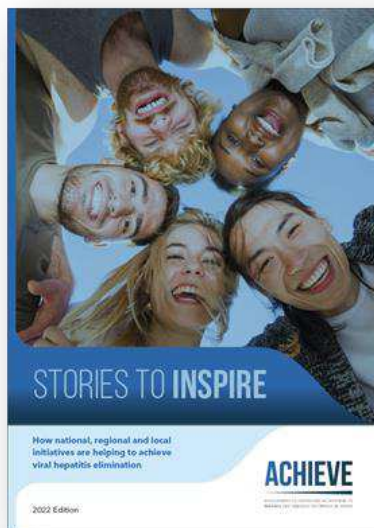
Peer Support for Hepatitis C



Point-of-Care Antibody Testing for Hepatitis C



Point-of-Care RNA Testing for Hepatitis C



A PEER-LED APPROACH TO HEPATITIS C



Access to naloxone



Access to a primary care provider



Access to HIV testing and treatment



Hepatitis B testing & vaccination



Tuberculosis screening & treatment





Access to OST/OAT











Access to needles, syringes and other equipment

WHAT HCV SERVICES DO WE PROVIDE?

-  HCV education and information
-  Peer support
-  HCV testing
-  HCV diagnosis
-  HCV treatment
-  Liver disease assessment
-  Genotyping for HCV

WHAT INTERVENTIONS DO WE USE?

-  Peer support
-  Opt-out screening
-  On-site testing
-  Point-of-care HCV antibody testing
-  Point-of-care HCV RNA testing
-  Dried blood spot testing
-  Pre-test counselling and education
-  Patient navigation

This Médecins du Monde programme uses a **peer-led approach** to **refer people** at risk for HCV to **existing HIV / harm reduction clinics/mobile unit**, where testing and treatment is provided

NSP AND HEPATITIS C TESTING AND TREATMENT SERVICE | QUEENSLAND, AUSTRALIA



Access to naloxone



Access to a primary care provider



Access to HIV testing and treatment



Access to needles, syringes and other equipment



Sexual Health Screening



Links to social support

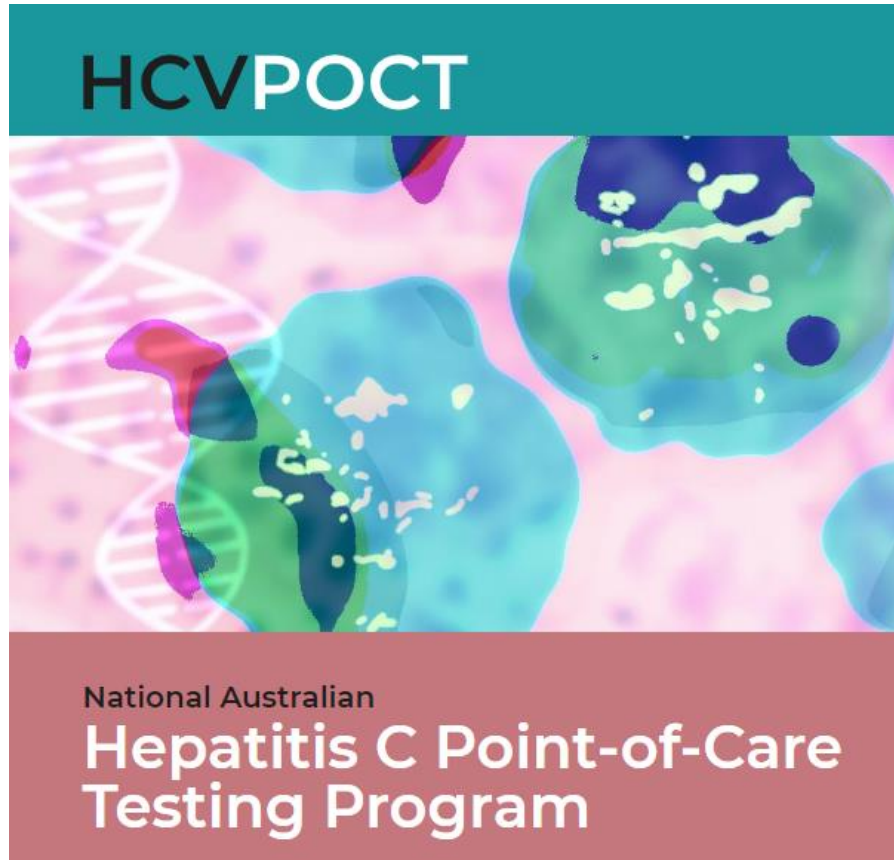


Point-of-care testing for hepatitis C, HIV and Syphilis



Access to alcohol & other drug treatment

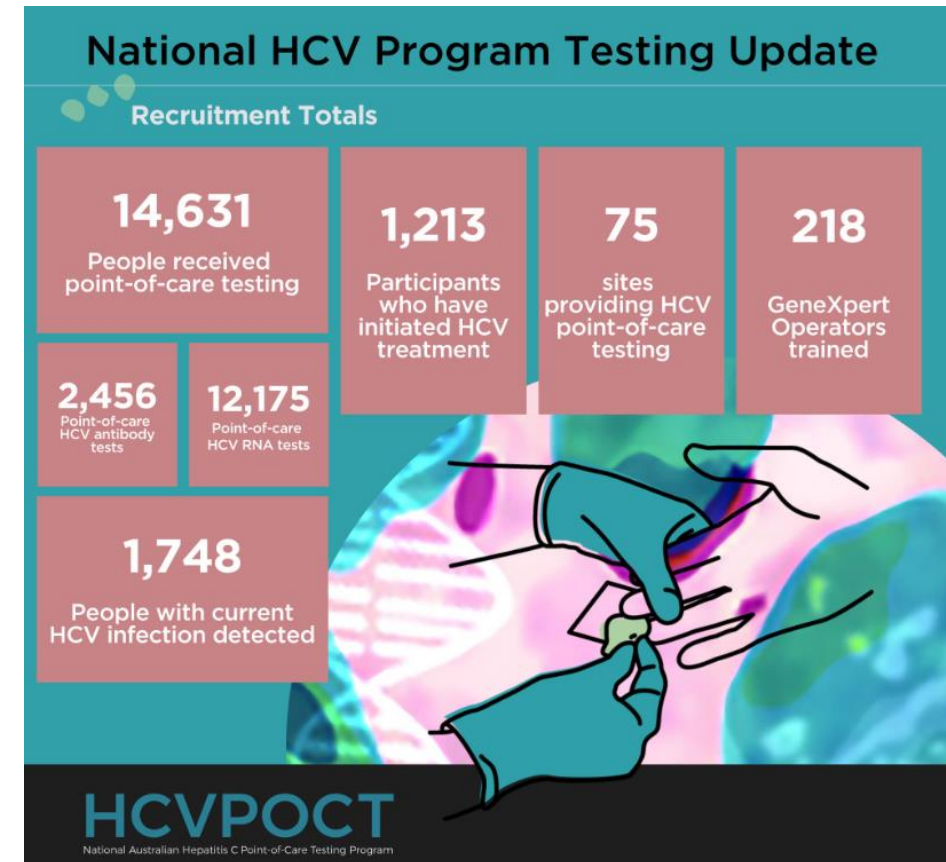
The Queensland Injectors Health Network (QuIHN), extended its existing harm reduction services to provide a '**one-stop-shop**' where people who use drugs could access **testing and treatment** in a safe and familiar environment.



HCVPOCT

National Australian
**Hepatitis C Point-of-Care
Testing Program**

- HCV Ab
- HCV RNA
- HIV Ab/Ag
- HBsAg



Drug treatment clinics, NSPs, prisons, mental health, mobile outreach models, homelessness services, Aboriginal Community Controlled Health Organisations

<https://x.com/hcvpoc/status/1713696575120437650?s=20>

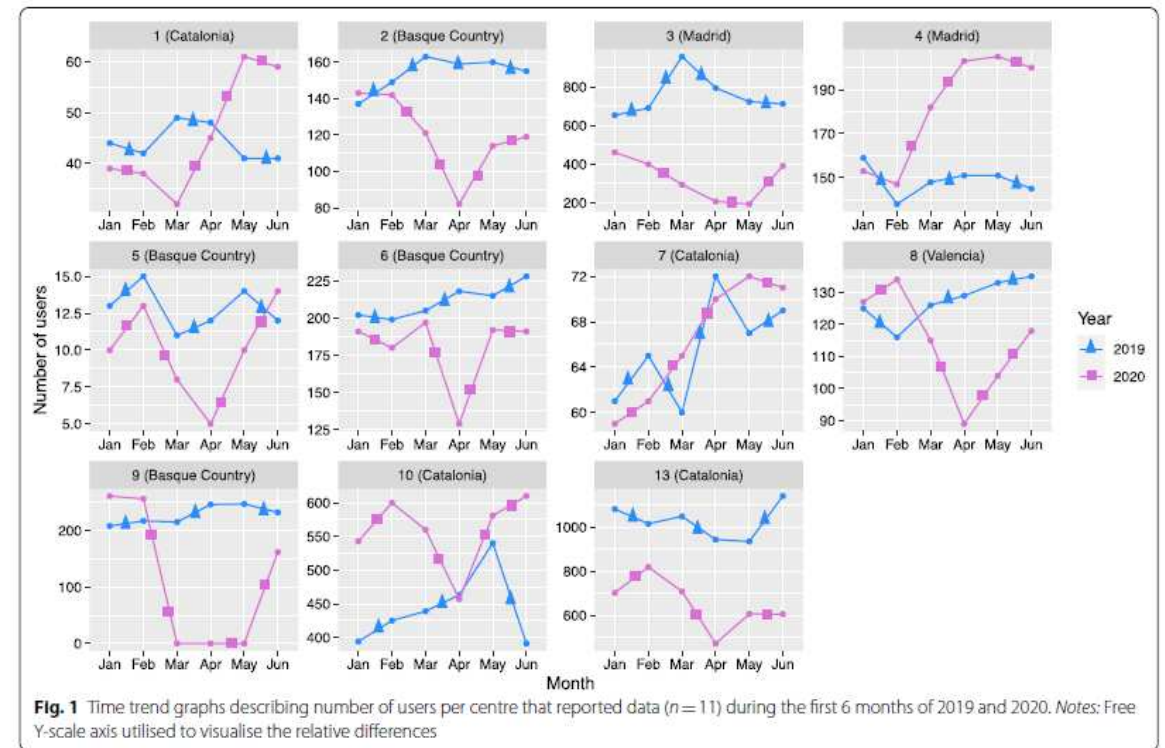
16/10/23

Integration of services in harm reduction centres in Spain

Centre (city name)	HCV DAA	HIV ART	HBV testing	HCV testing	HIV testing	TB testing *
Centre 1 (Badalona)						
Centre 2 (Bilbao)				✓		✓
Centre 3 (Madrid)	✓	✓	✓	✓	✓	✓
Centre 4 (Madrid)	✓	✓	✓	✓	✓	✓
Centre 5 (Vitoria-Gasteiz)					✓	
Centre 6 (Bilbao)	✓	✓	✓	✓	✓	
Centre 7 (Reus)						
Centre 8 (Valencia)	✓	✓	✓	✓	✓	✓
Centre 9 (Bilbao)						
Centre 10 (El Prat de Llobregat)		✓		✓	✓	
Centre 11 (Barcelona)	✓		✓	✓	✓	✓
Centre 12 (Barcelona)	✓		✓	✓	✓	✓
Centre 13 (Sant Adrià de Besos)	✓			✓	✓	
Total	7	5	6	9	9	6

*LTBI testing with the tuberculin skin test.

COVID-19-related disruptions to harm reduction services: reduced outreach activities and low threshold harm reduction service capacities in general, leading to reduced HIV and hepatitis C testing availability

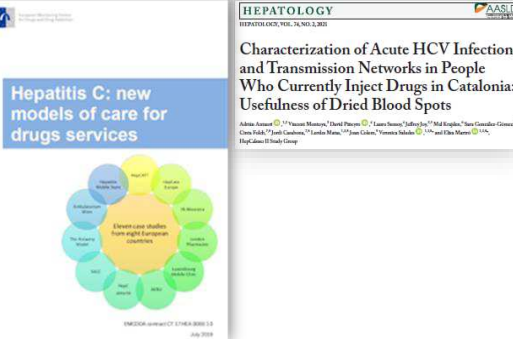


Integrated screening of HCV/HBV/VIH in people who inject drugs in harm reduction services in Catalonia



HepCdetect II
2016-17, N=410

1. Decentralised DBS collection for one-step HCV-RNA testing



HepTestFarma: community pharmacies (harm reduction)
2023-24, N=400



HepCdetect II
2018-19, N=100

2. Decentralised HCV-RNA testing



CHIME
2019-to date, N=750
Dx & test of cure



3. Decentralised Test & Treat pilot

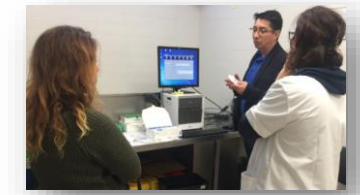
M. Gálvez. HepHIV - PO3/03



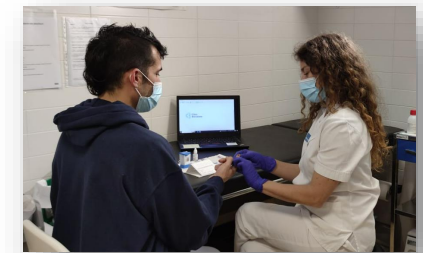
Drug centres
2021-to date



Training and remote validation of GeneXpert results

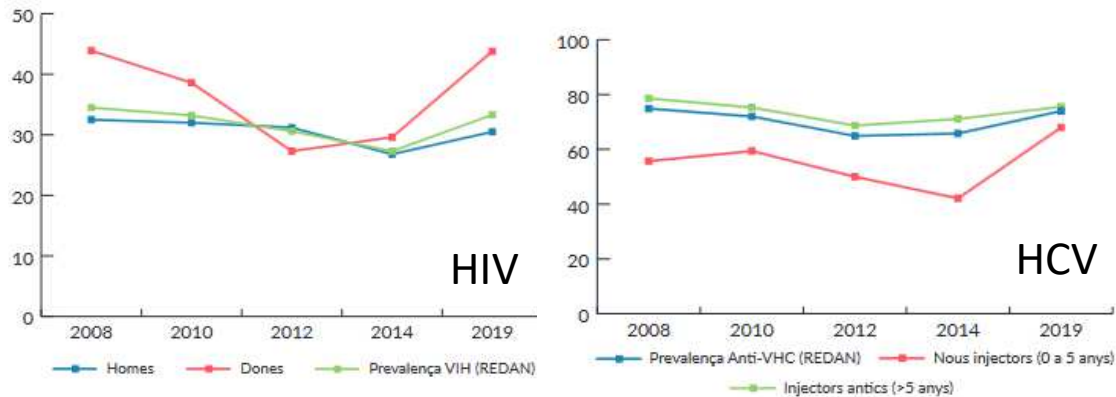


DBS:
- Baseline vs. post-tx HCV sequencing (differentiation between reinfection and tx failure)
- Validation as 'test of cure'



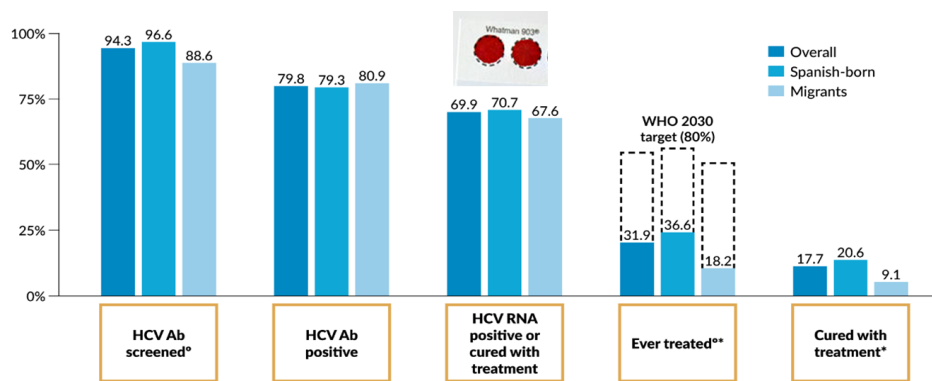
Biobehavioural monitoring in people who inject drugs in harm reduction centers in Catalonia

HIV and HCV screening in oral fluid (Ab)



SIVES 2020. CEEISCAT, 2021.

HCV screening with RDTs (Ab) and DBS (RNA)



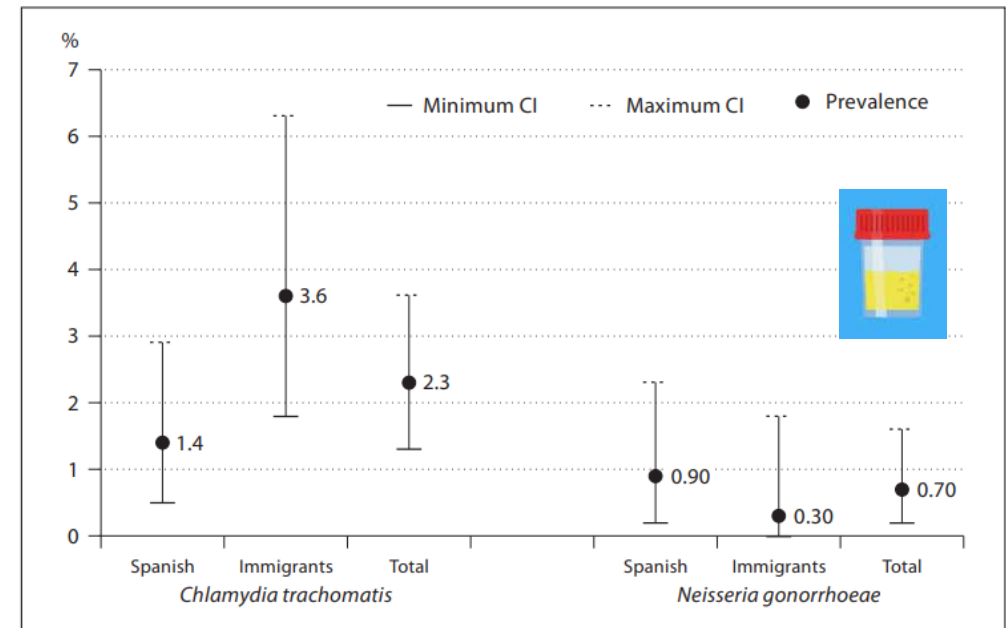
* significant differences between proportions (p<0.05) ** among PWID HCV RNA positive or cured with treatment

HCV care cascade in people who actively inject drugs attending harm reduction services in Catalonia (HepCdetect II Study)

Folch C, et al. Int J Drug Policy 2021;90:103057

STI screening in urine (laboratory testing)

Respondents aged ≤25 years had a higher risk of STIs (OR 3.39), as did women (OR 3.08).



Folch C, et al. Eur Addict Res 2011;17(5):271-8.

Integrated screening of HCV/HBV/VIH in drug centres and homelessness services in Catalonia



20 min



50 min



15 min

Mobile unit



- Nurse and social educator
- RDTs for **HCV/HBV/VIH**
- HCV-RNA confirmatory testing and 'test of cure' (GeneXpert)
- Transient elastography for liver fibrosis assessment
- Access to **decentralized HCV treatment** through visited centres

- **2023: Screening of STIs** (*C. trachomatis*, *N. gonorrhoeae*, *T. vaginalis*) by decentralized sample collection



CEEIS
Cat

- **2024: Expanding to active TB testing**



Conclusions

What will it take to achieve elimination of these infections in people who use drugs?

- ✓ **Radical simplification of diagnosis and care pathways**
- ✓ **Integrated services**

- **Training** of testing teams and **quality assurance**
- **Careful choice of specific tests and diagnostic algorithms** (balance suitability for decentralized settings, multiplexing capacity, low cost, high diagnostic accuracy)
- Involvement of all necessary partners to **ensure access to confirmatory diagnosis and treatment** if not provided on-site
- **Connectivity** and **reporting** systems, **monitoring** of the elimination progress

Thank you for your attention



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