

# Il modello HCV: tra passato e futuro - I risultati fin oggi ottenuti-

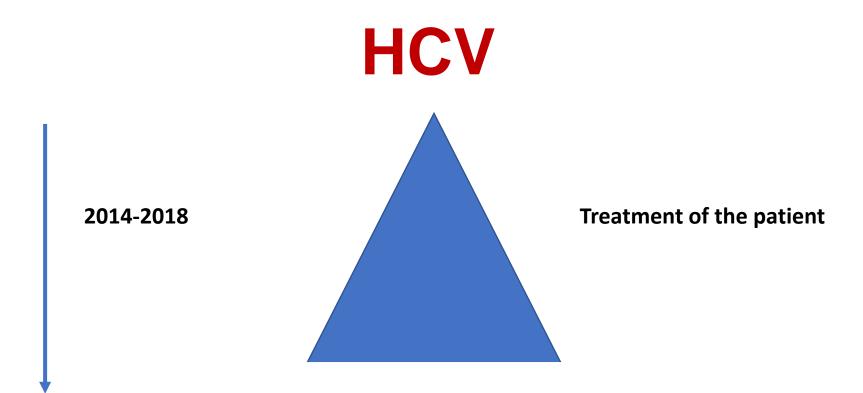
## Nicola Coppola

University of Campania, Luigi Vanvitelli, Naples, ITALY Infectious Diseases Unit, AOU Vanvitelli nicola.coppola@unicampania.it

# **Disclosures**

# NC reports:

- -grants from ViiV Healthcare, Janssen-Cilag and Gilead Science
- -personal fees from Gilead Sciences, Abbvie, Bristol-Myers Squibb, Correvio, Merk-Sharp & Dohme, Angelini

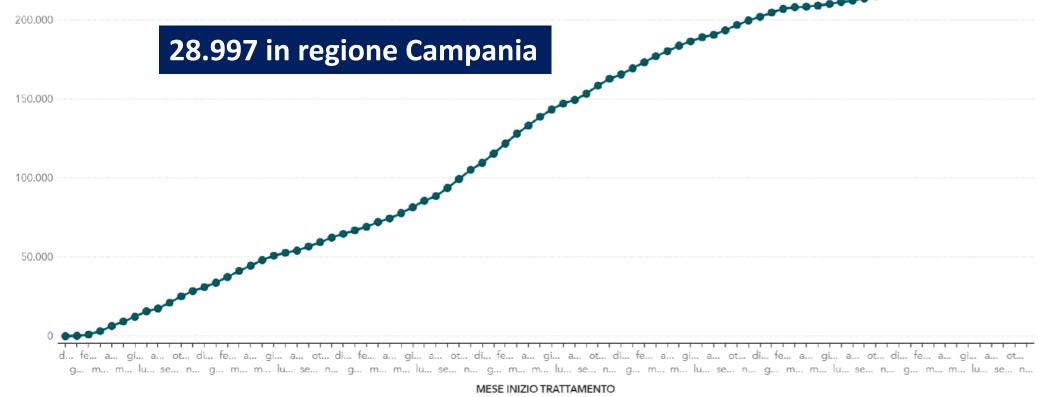


# Treated non-treated Disease control Tx of patients with liver disease

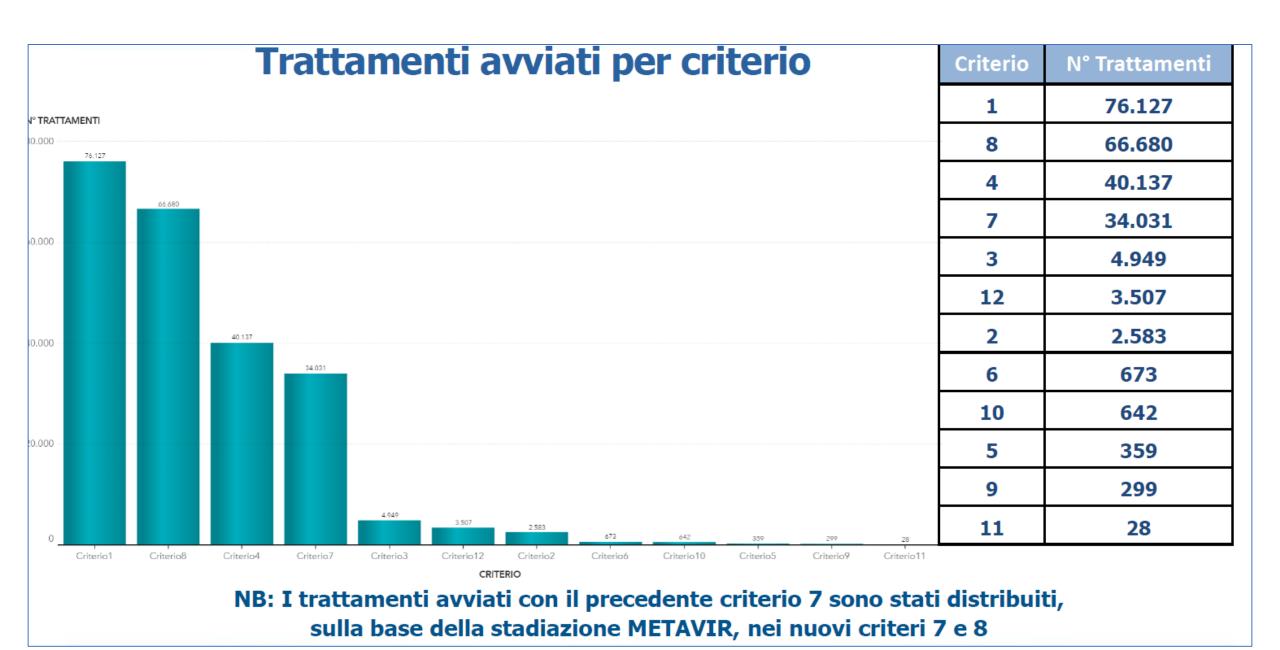


### Trend cumulativo dei trattamenti avviati

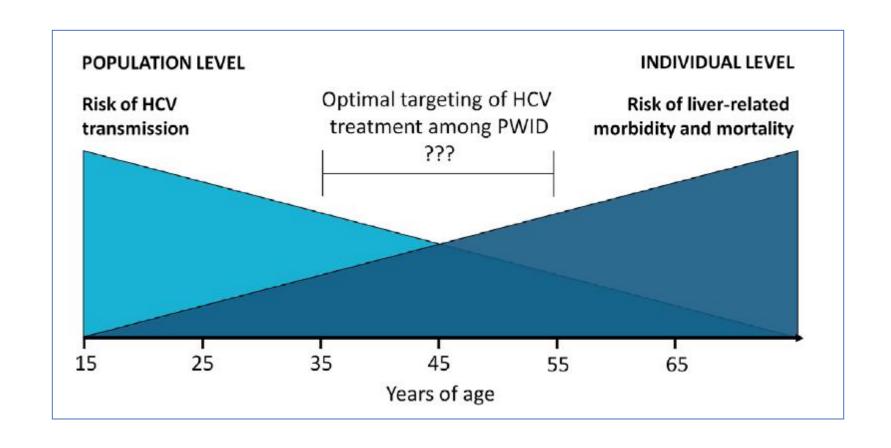


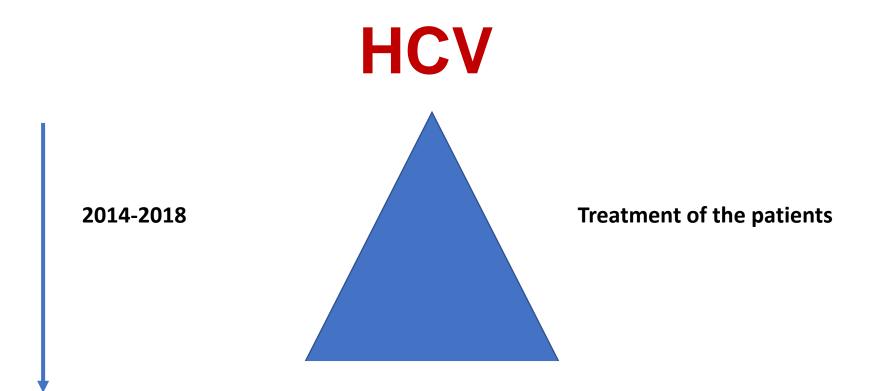


230.015 «avviati» sono i trattamenti (solo pazienti eleggibili) con almeno una scheda di dispensazione farmaco

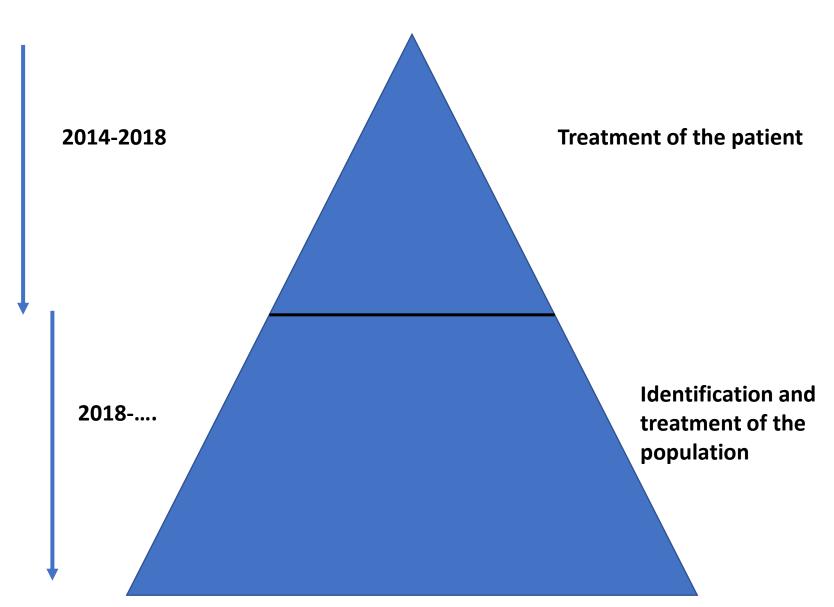


# **HCV** treatment as prevention





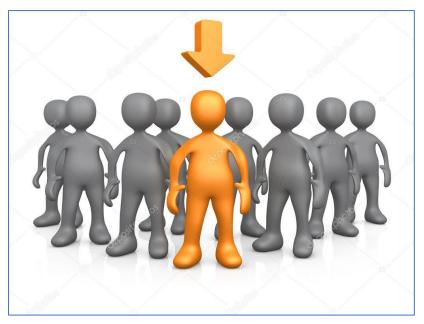




### **Individualized HCV treatment**

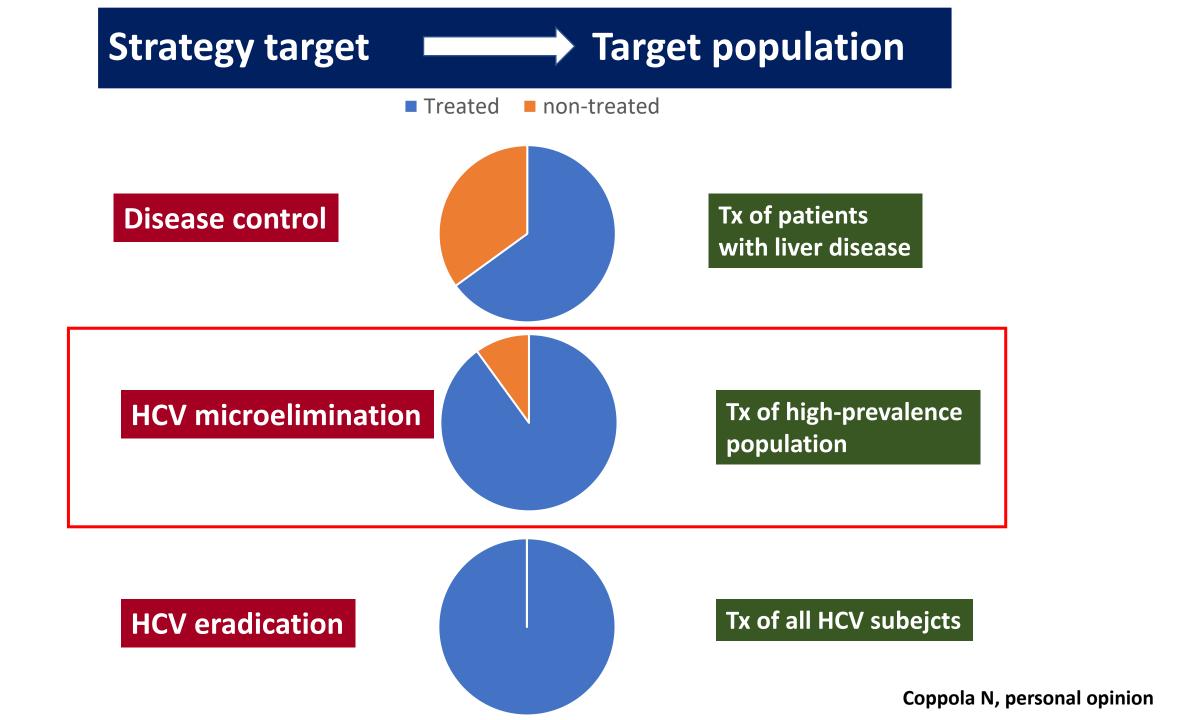


**Population-based approch** 









# **Global Call for HCV Elimination**

WHO vision: "A world where viral hepatitis transmission is stopped and everyone has access to safe, affordable, and effective treatment and care"

### 2030 Targets

90% Diagnosed

80% Treated

65% Reduced mortality

### **Feasible** by scaling up **key interventions**:

- Hepatitis B vaccination and treatment
- Safe injection practices and safe blood
- Harm reduction for PWID
- Safer sex (including condom promotion)
- Hepatitis C cure

US HBV/HCV Elimination Strategy developed by National Academies of Sciences, Engineering, and Medicine: "elimination" = 90% reduction in incidence by 2030

### Released at AASLD November 10th 2019

Call to Action for Liver Associations to Advance Progress Towards Viral Hepatitis Elimination: A Focus on Simplified Approaches to HCV Testing and Cure.











### Journal Pre-proof

EASL recommendations on treatment of hepatitis C – Final update of the series

European Association for the Study of the Liver



### Indication for DAA treatemtn

### Recommendations

- All treatment-naïve and treatment-experienced patients with recently acquired or chronic HCV infection must be offered treatment without delay (A1).
- Urgent treatment should be considered: in patients with significant fibrosis or cirrhosis (METAVIR score F2, F3 or F4), including compensated (Child-Pugh A) and decompensated (Child-Pugh B or C) cirrhosis; in patients with clinically significant extrahepatic manifestations (e.g. symptomatic vasculitis associated with HCV-related mixed cryoglobulinaemia, HCV immune complex-related nephropathy and non-Hodgkin B cell lymphoma); in patients with HCV recurrence after liver transplantation; in patients at risk of a rapid evolution of liver disease because of concurrent comorbidities (non-liver solid organ or stem cell transplant recipients, HBV and HIV coinfections, diabetes); and in individuals at risk of transmitting HCV (PWIDs, men who have sex with men with high-risk sexual practices, women of childbearing age who wish to get pregnant, patients on haemodialysis, incarcerated individuals) (A1).
- Treatment is generally not recommended in patients with limited life expectancy due to nonliver-related comorbidities (B2).

- General population
- Populations with high HCV prevalence

- General population
- Populations with high HCV prevalence

**Incarcerated populations** 

People living with HCV/HIV coinfection

Men who have sex with men

**Migrants** 

Persons who inject drugs

- General population
- Populations with high HCV prevalence

**Incarcerated populations** 

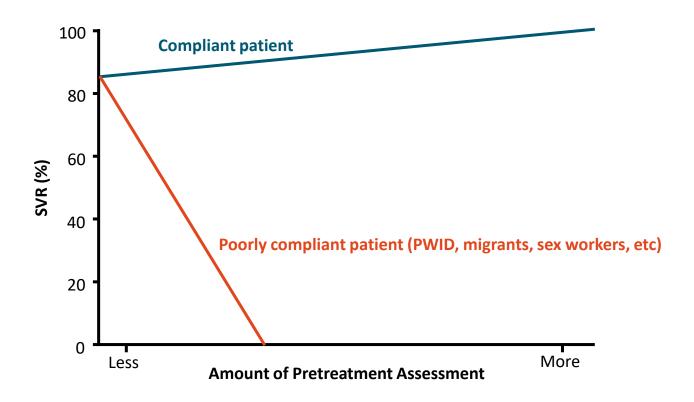
People living with HCV/HIV coinfection

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# **Illustrating the Danger of Too Many Tests**

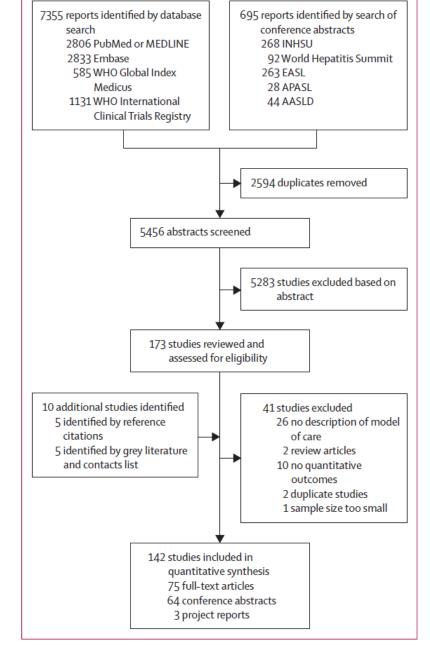


### Decentralisation, integration, and task-shifting in hepatitis C virus infection testing and treatment: a global systematic review and meta-analysis



Ena Oru, Adam Trickey, Rohan Shirali, Steve Kanters, Philippa Easterbrook





### **Lancet Glob Health 2021**

Decentralisation, integration, and task-shifting in hepatitis C virus infection testing and treatment: a global systematic review and meta-analysis



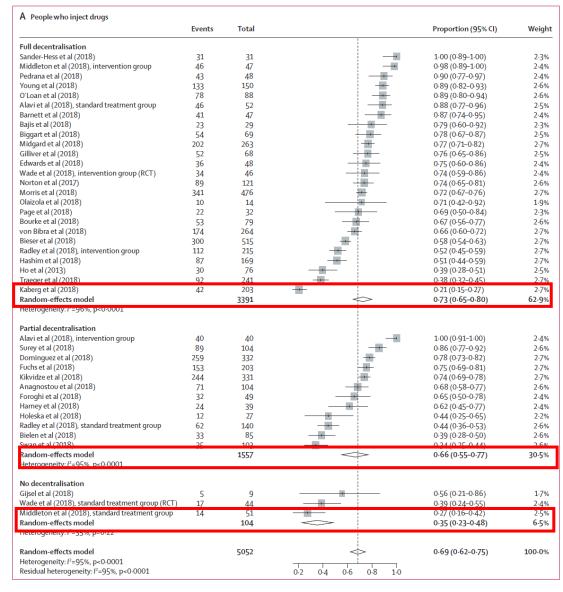
Ena Oru, Adam Trickey, Rohan Shirali, Steve Kanters, Philippa Easterbrook



### Effect of decentralisation and integration on linkage to care

	Events	Total		Proportion (95% CI)	Weight
- Full decentralisation					
Seidenburg et al (2013)	85	85	-	1.00 (0.96-1.00)	3.2%
ander-Hess et al (2018)	31	31		1.00 (0.89-1.00)	3.0%
Vade et al (2018), intervention group (RCT)	52	59		0.88 (0.77-0.95)	3.1%
lashim et al (2018)	169	211	-	0.80 (0.74-0.85)	3.2%
Plaizola et al (2018)	15	19	<del></del>	0.79 (0.54-0.94)	2.8%
'Loan et al (2018)	88	116		0.76 (0.67-0.83)	3.2%
Nidgard et al (2018)	263	348	-	0.76 (0.71-0.80)	3.2%
ack et al (2009)	86	118		0.73 (0.64-0.81)	3.2%
/ade et al (2015)	186	279		0.67 (0.61-0.72)	3.2%
edrana et al (2018)	48	76		0.63 (0.51-0.74)	3.1%
ajis et al (2018)	29	47		0.62 (0.46-0.75)	3.1%
age et al (2018)	32	71		0.45 (0.33-0.57)	3.1%
adley et al (2018), intervention group (RCT)	215	545		0.39 (0.35-0.44)	3.3%
/ilkinson et al (2008)	83	545 411	+	0.39 (0.35-0.44)	3.2%
andom-effects model	03	2416		,	3·2% 44·0%
eterogeneity: I <sup>2</sup> =98%, p<0·0001		2416		0.72 (0.57-0.85)	44.0%
eterogeneity. Jow, p to cool					
artial decentralisation					
ikvidze et al (2018)	338	350	-	0.97 (0.94-0.98)	3.2%
ielen et al (2018)	85	114		0.75 (0.66-0.82)	3.2%
wan et al (2018)	103	141	-	0.73 (0.65-0.80)	3.2%
lam et al (2012)	68	96	-	0.71 (0.61-0.80)	3.2%
/ong et al (2014)	69	98	<del>                                     </del>	0.70 (0.60-0.79)	3.2%
Masson et al (2013), intervention group (RCT)	97	149	-	0.65 (0.57-0.73)	3.2%
Martinez et al (2012)	76	125		0.61 (0.52-0.69)	3.2%
utton et al (2018)	369	710	<b>∓</b> T	0.52 (0.48-0.56)	3.3%
oroghi et al (2018)	49	97		0.51 (0.40-0.61)	3.2%
ntonini et al (2018)	2	4		0.50 (0.07-0.93)	1.9%
lagaldi et al (2018)	77	200		0.38 (0.32-0.46)	3.2%
lasson et al (2013), standard treatment group (RCT)	51	137		0.37 (0.29-0.46)	3.2%
loleska et al (2018)	34	126		,	_
adley et al (2018) standard treatment group (RCT)	34 140			0.27 (0.19–0.36)	3.2%
, , , , , , , , , , , , , , , , , , , ,		540		0.26 (0.22-0.30)	3.3%
lackburn et al (2016) orter et al (2017)	198	861	<u> </u>	0.23 (0.20-0.26)	3.3%
andom-effects model	/	44 3792		0·16 (0·0/-0·30) 0·53 (0·38-0·67)	3·1% <b>49·9</b> %
eterogeneity: /²=99%, p<0·0001		3/ 32		0.55 (0.50-0.07)	43.3%
•					
lo decentralisation					
Vade et al (2018), standard treatment group (RCT)	38	57	<del>-   •   -   -   -   -   -   -   -   -   </del>	0.67 (0.53–0.79)	3.1%
andom-effects model	9	34 <b>91</b>		0·26 (0·13-0·44) 0·47 (0·11-0·84)	3.0% 6.1%
leterogeneity: 1 = 93%, p<0.0001		3.⁴		5 47 (0·11-0·04)	0-170
andom-effects model		6299		0.61 (0.51-0.71)	100.0%
leterogeneity: I <sup>2</sup> =98%, p<0.0001					

### Effect of decentralisation and integration on DAA intake



**Lancet Glob Health 2021** 

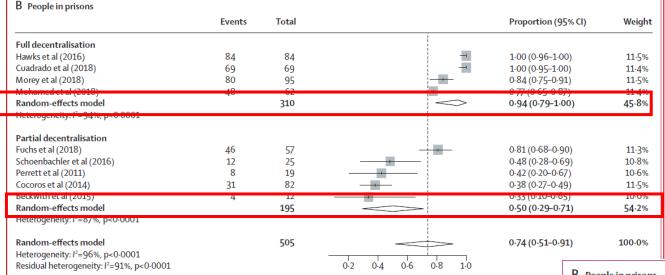
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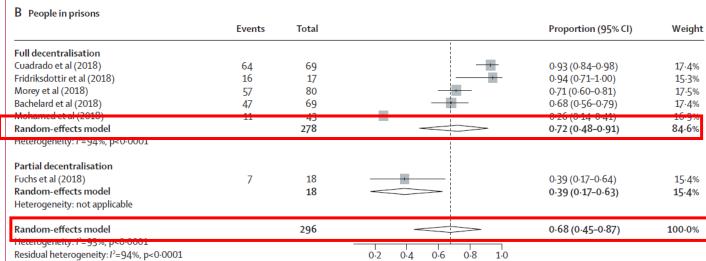
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### Effect of decentralisation and integration on linkage to care



### Effect of decentralisation and integration on DAA intake



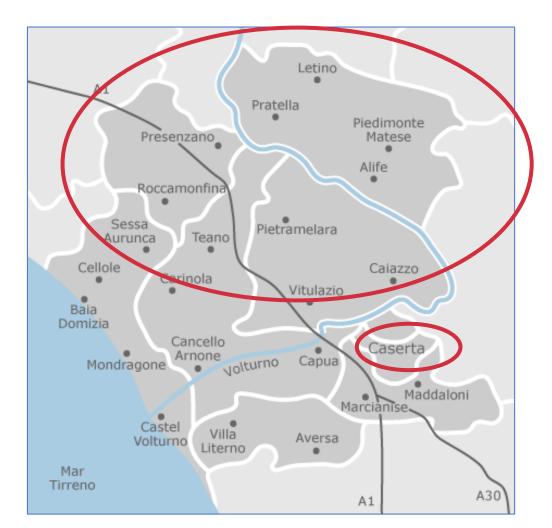
### Lancet Glob Health 2021



# Innovative procedures for micro-elimination of HCV infection in persons who use drugs

Vincenzo Messina<sup>1</sup> | Antonio Russo<sup>1,2</sup> | Enrico Parente<sup>3</sup> | Giovanni Russo<sup>3</sup> | Tiziana Raimondo<sup>3</sup> | Angela Salzillo<sup>1</sup> | Filomena Simeone<sup>1</sup> | Lorenzo Onorato<sup>1</sup> | Giovanni Di Caprio<sup>1</sup> | Mariantonietta Pisaturo<sup>1,2</sup> | Nicola Coppola<sup>1,2</sup>

A prospective, interventional, before and after study, based on audits performed by Infectious Diseases physicians in a SUD facility in Piedimonte Matese, in southern Italy, was performed. Pre-intervention period: January-December 2017 and Post-intervention period: January-December 2018.



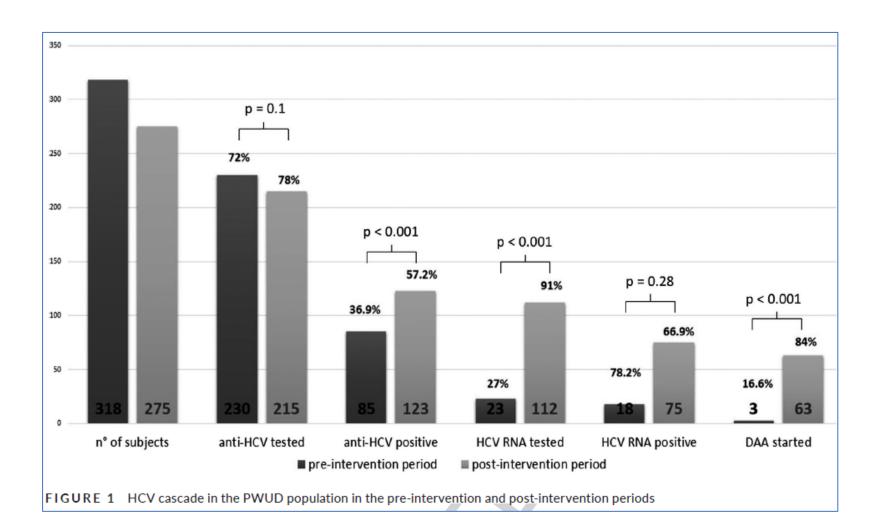
6 SerD following about 300 PWUD and acts in the northern area of Caserta with about 151 000 inhabitants



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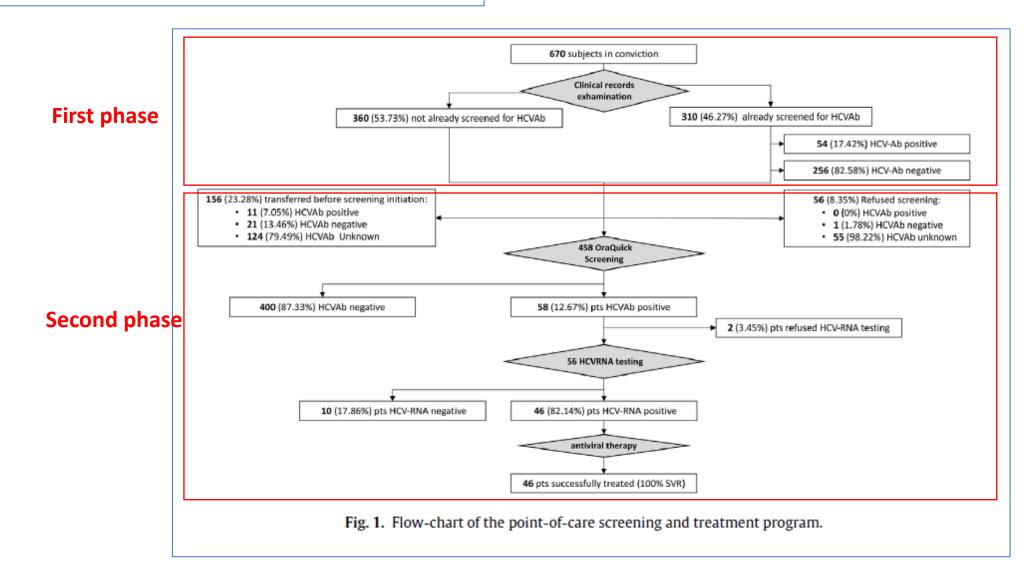
Liver, Pancreas and Biliary Tract

Hepatitis C virus infection in jail: Difficult-to-reach, not to-treat. Results of a point-of-care screening and treatment program



Mario Masarone<sup>a</sup>, Rosa Caruso<sup>a,b</sup>, Andrea Aglitti<sup>a</sup>, Carmine Izzo<sup>a,b</sup>, Giuseppe De Matteis<sup>b</sup>, Maria Rosaria Attianese<sup>b</sup>, Antonio Maria Pagano<sup>b</sup>, Marcello Persico<sup>a,\*</sup>

- A prospective observational study in two phases:
- first, all the prisoners' clinical records were reviewed, to verify HCV-Ab execution.
- Second, a universal point-of-care screening and treatment program we re performed



- General population
- Populations with high HCV prevalence

**Incarcerated populations** 

People living with HCV/HIV coinfection

Men who have sex with men

Migrants/homeless

Persons who inject drugs

# Medical points I livello (parrocchie, centri sociali, punti caritas, ARCI)

NAPOLI CASERTA POTENZA FOGGIA LAMEZIA TERME









# Medical points I livello

(parrocchie, centri sociali, punti caritas, ARCI)

**NAPOLI** 

**CASERTA** 

**POTENZA** 

**FOGGIA** 

LAMEZIA TERME

Laboratorio di malattie infettive

### Centri di secondo livello

(per approfondimenti diagnostici ed eventuale terapia)

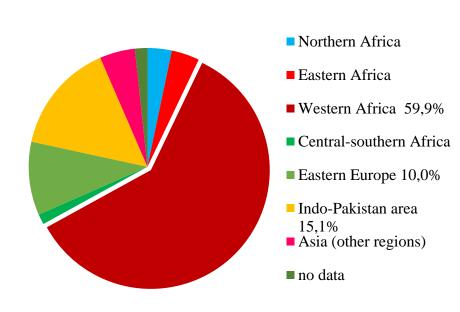
UOC Malattie Infettive, Vanvitelli, Napoli UOC Malattie Infettive, AO Caserta

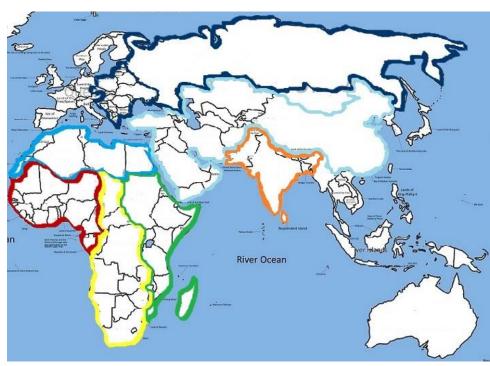
UOC Malattie Infettive, Foggia Servizio Malattie Infettive, Lamezia terme



- 4125 subjects enrolled: 3839 (93.1%) accepted the screening
- Mean age 28 years  $\pm 10$  SD
- 84.0% males, 15.9% females, 0.1% unknown

### Geographical areas of origin: 7 macro-regions

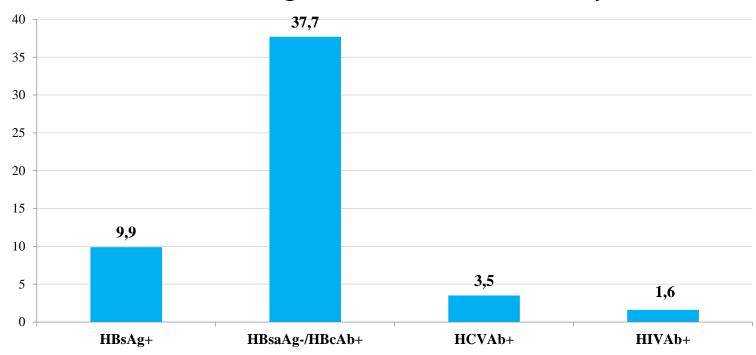




Travel Medicine and Infectious Dis 2020



### Overall serological markers in 3,839 subjects

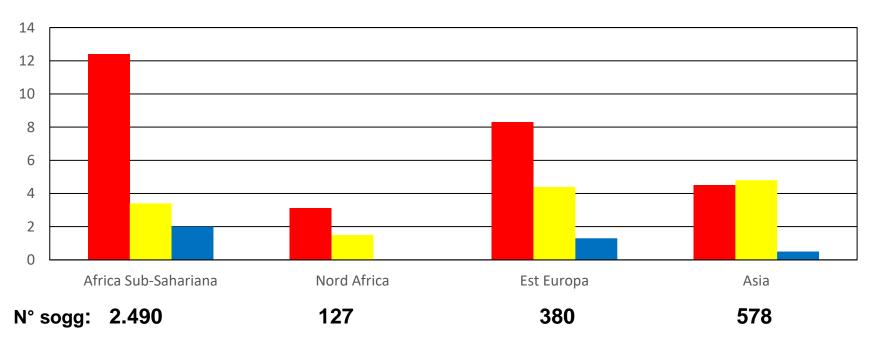




# 3.839 immigrants







# Case finding and linkage to care in migrants Interventations

Questionnaires

Meeting

**Brochures** 

Burocratic tutor





### Vieni a trovarci presso i nostri Centri Clinici!!

### Il test del sangue è completamente gratuito

A Caserta presso:

"ex Centro Sociale Liberato", viale Ellittico 9

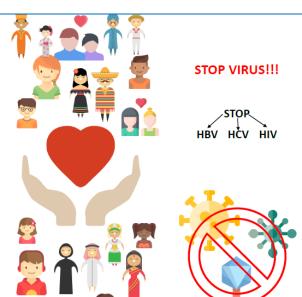
A Napoli presso:

"Tutela della Salute degli Immigrati", via Egiziaca a Forcella 34

"Centro di accoglienza vicolo Verticoeli", via Tribunali

"Centro di accoglienza femminile", via Tommasoni, villa Tommasoni,

Saviano (NA)



### Cosa sono i virus HBV e HCV?

L'HBV e l'HCV sono virus che colpiscono il fegato causando l'Epatite virale, possono progredire nel tempo provocando la cirrosi epatica (alterazione di forma e funzione del fegato) e/o il tumore del

### Cos'è l'HIV?



L'HIV è il virus responsabile dell'AIDS (Sindrome da Immunodeficienza Acquisita) e consiste in una progressiva diminuzione delle difese del corpo. Nel tempo compaiono altre malattie infettive e non, anche molto gravi.



Spesso queste malattie sono asintomatiche Altre volte si possono manifestare con malessere generale, nausea, vomito e diarrea.

### Come si trasmettono i virus HBV, HCV e HIV?

- Rapporti sessuali non protetti (vaginali e anali)
- Sangue: scambio di siringhe infette, strumenti per tatuaggi e piercing contaminati con sangue infetto, uso in comune di oggetti personali (rasoi, tagliaunghie, spazzolino da denti, etc) contaminati da sangue
- madre infetta al bambino (durante gravidanza, parto e allattamento al



Con il test specifico (un prelievo di sangue) per la ricerca degli anticorpi nel sangue è possibile sapere se si è stati contagiati da questi virus.





Nei primi mesi dopo il contagio il test può essere negativo. perciò è bene ripeterlo, fino a 3 mesi dopo il contagio.

### Come si può prevenire il contagio dai virus

### HBV, HCV e HIV?

- · Usare sempre il profilattico durante i rapporti sessuali genitali, anali e orali
- Usare soltanto siringhe sterili
- · Assicurarsi che gli strumenti utilizzati per piercing e tatuaggi siano utilizzati una volta sola (monouso)
- Evitare il contatto diretto con il sangue
- Molte infezioni possono trasmettersi durante gravidanza, parto e allattamento bambino: madre sottoporsi al test del sangue e ai controlli per le infezioni è la migliore protezione per i







Esiste un vaccino per proteggersi dall'HBV, e farmaci contro HCV e HIV.





### Uje kwenye zahanati yetu!

### Kipimo hakina

gharama vovote

Caserta:

"ex Centro Sociale Liberato". viale Ellittico 9

Napoli:

"Tutela della Salute degli Immigrati", via Egiziaca a Forcella 34

"Centro di accoglienza vicolo Verticoeli", via Tribunali

"Centro di accoglienza femminile", via Tommasoni, villa Tommasoni, Saviano (NA)

### STOP VIRUSI!!!









Mara nyingi magonjwa haya hayana dalili. Magonjwa haya huweza kuanza kama kichefuchefu, kutapika na kuhara.

HBV na HCV ni virusi vinavyoshambulia ini na kusababisha ugonjwa wa homa ya ini (Hepatitis B), virusi hivi vinaweza pia kusababisha sirosisi (Cirrhosis yaani mabadiliko ya umbo na kazi za ini) na/au saratani ya ini.

### HIV ni nini?

HBV na HCV ni nini?

HIV, VVU ni virusi vinavyosababisha ugonjwa wa UKIMWI (Upungufu wa Kinga Mwilini) na kuharibu mfumo wa kingamwili. Magonjwa mengine ya kuambukiza na yasiyo ya kuambukiza huweza kushambulia mwili kwa urahisi kutokana na maambukizi ya VVU.





### HBC, HCV na HIV huambukizwaje? Kwa njia ya ngono zembe (iwe kwa njia ya uke au kwa njia ya haja kubwa)

- Damu: kuchangia sindano ambayo haijachemshwa baada ya mtu mwingine kuitumia, kutumia vifaa vya kuchorea bombwe na kupenyeza piercing mwilini vilivyochafuliwa na damu yenye maambukizi, kubadilishana vitu binafsi (kama wembe, mashine ya kukatia kucha, mswaki n.k)
- Kutoka kwa mama aliye na maambukizi ya virusi kwenda kwa mtoto wakati wa ujauzito, kujifungua au kunyonyesha





Uchunguzi wa kimaabara (kipimo cha damu) wa kingamwili kwenye damu huweza kuashiria kama mtu ameambukizwa virusi hivi.



Miezi ya mwanzo baada ya kuambukizwa kipimo kinaweza kisionyeshe uwepo wa maambukizi wa virusi, hivyo ni vema kupimwa baada ya miezi mitatu ya maambukizi.

### Kuzuia maambukizi ya virusi vya HBC, HCV na HIV

- Tumia kondomu wakati wa kujamiiana (ngono kwa njia ya uke, njia ya haja kubwa, na mdomoni)
- Tumia sindano ilivochemshwa
- Hakikisha kwamba vyombo vya kuchorea bombwe na kupenyeza piercing mwilini vinatumiwa mara moja tu
- Epuka kugusa au kuloanishwa na damu
- Maradhi mengi husambaa kutoka kwa mama aliye na virusi kwenda kwa mtoto wakati wa ujauzito, kujifungua na kunyonyesha: kupima ni njia bora ya kuepusha maambukizi kwa mtoto wako.





Chanjo ya kuzuia VVU ipo, na vile vile madawa dhidi va HCV na HIV.



# Case finding and linkage to care in migrants Interventations

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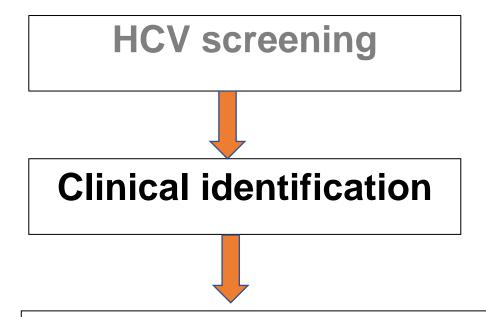
Burocratic tutor



Milan, 17 February 2017

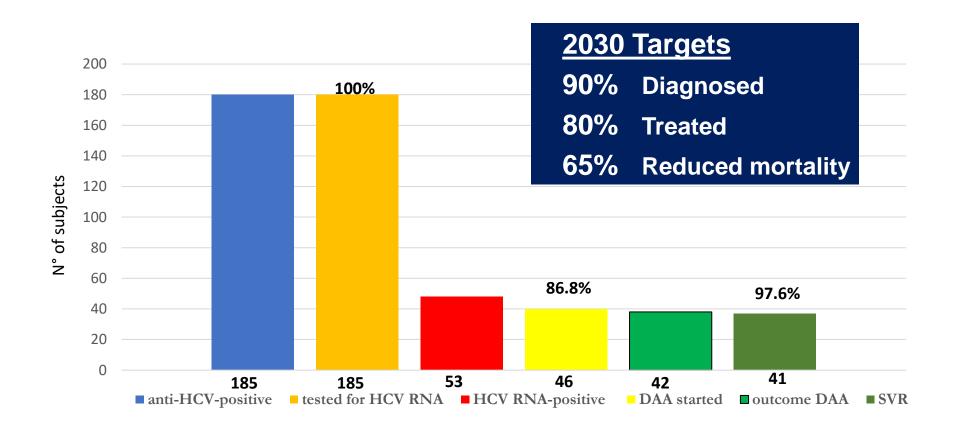
Department of mental health and public medicine Second University of Naples Largo Madonna delle Grazie 80133 Napoli - Italy

RE: EVALUATION OF AN INNOVATIVE MODEL TO ELIMINATE HCV INFECTION IN A HIGH-RISK POPULATION OF UNDOCUMENTED MIGRANTS AND LOW-INCOME REFUGEES



Sofosbuvir/velpatasvir for all HCV RNA positive subjects identified

## HCV care cascade in immigrants





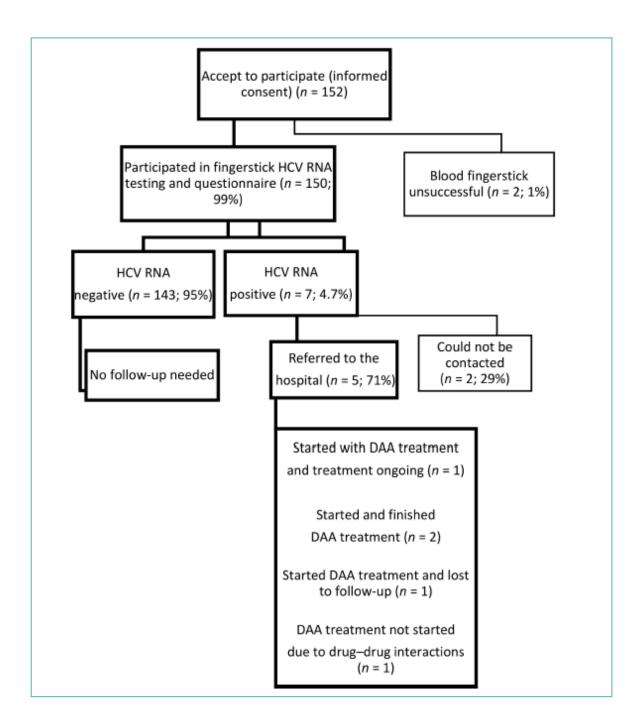


Articl

A Feasibility Study to Increase Chronic Hepatitis C Virus RNA Testing and Linkage to Care among Clients Attending Homeless Services in Amsterdam, The Netherlands

Ellen Generaal  $^{1,*}$ , Hilje Logtenberg van der Grient  $^2$ , Eberhard Schatz  $^2$ , Daniela K. van Santen  $^{1,3}$ , Anders Boyd  $^{1,4}$ , Sara K. Woods  $^5$ , Bert L. C. Baak  $^6$  and Maria Prins  $^{1,7}$ 

To improve HCV case finding, the feasibility of rapid HCV RNA testing in homeless services in Amsterdam was evaluated.



# Challenges in HCV Case Finding and Linkage to Care

- General population
- Populations with high HCV prevalence

**Incarcerated populations** 

People living with HCV/HIV coinfection

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Migrants

Persons who inject drugs

# Treatment as Prevention to Eliminate HCV in HIV-infected MSM

Phase A (10/2015-06/2016): a population-based and systematic screening for HCV-RNA among MSM

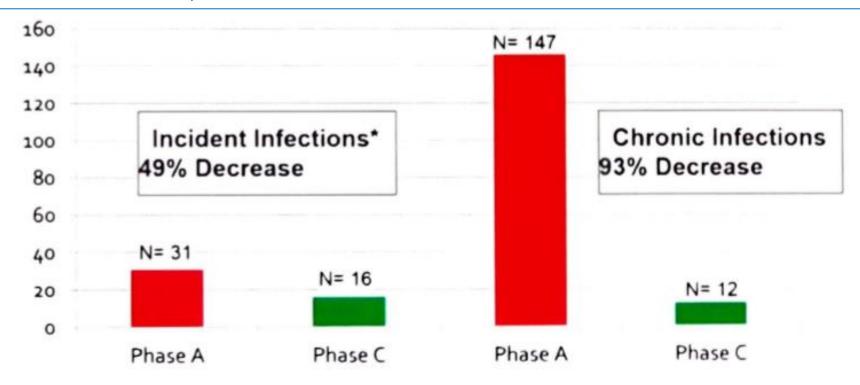
Phase B (06/2016-02/2017): DAAs treatment for MSM identified with a replicating HCV infection.

Phase C (03/2017-11/2017): re-screen to all MSM for HCV-RNA

# Treatment as Prevention to Eliminate HCV in HIV-infected MSM

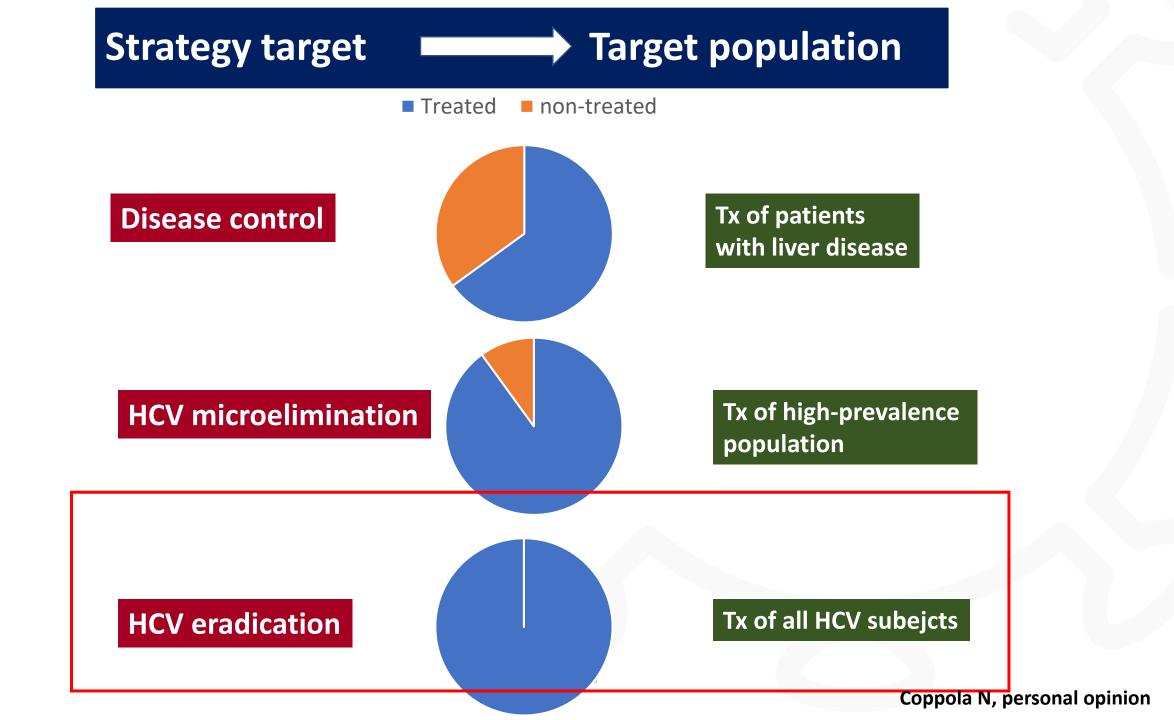
**Phase A** (10/2015-06/2016): a population-based and systematic screening for HCV-RNA among MSM **Phase B** (06/2016-02/2017): DAAs treatment for MSM identified with a replicating HCV infection.

Phase C (03/2017-11/2017): re-screen to all MSM for HCV-RNA



Negative serologic assay before 2015

Braun DL, CID 2020



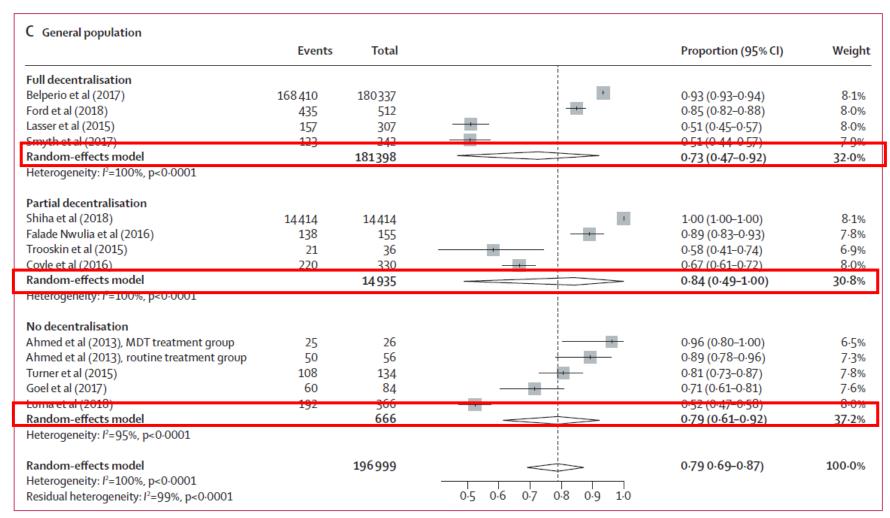
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oa OPEN ACCESS

#### Effect of decentralisation and integration on linkage to care



#### **Lancet Glob Health 2021**

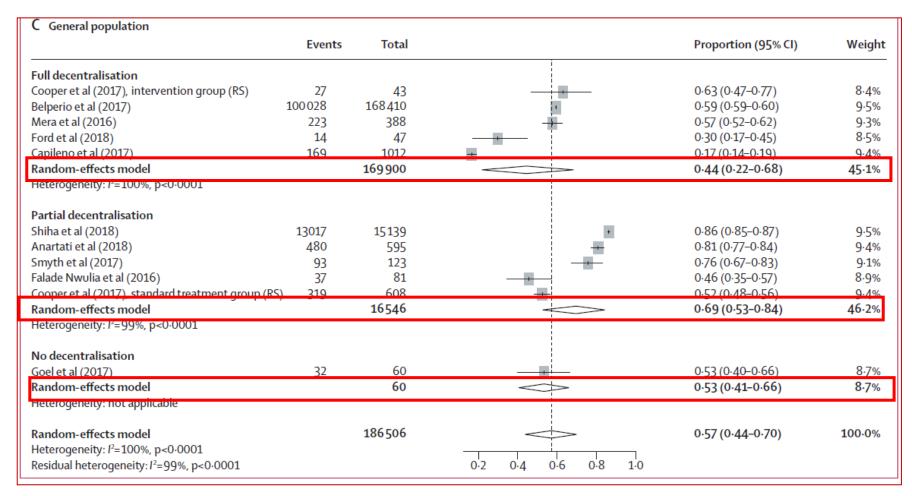
Decentralisation, integration, and task-shifting in hepatitis C virus infection testing and treatment: a global systematic review and meta-analysis



Ena Oru, Adam Trickey, Rohan Shirali, Steve Kanters, Philippa Easterbrook



#### Effect of decentralisation and integration on DAA intake



#### **Lancet Glob Health 2021**

# Challenges in HCV Case Finding and Linkage to Care

- General population
- Populations with high HCV prevalence
  - Opportunistic case-finding
  - Case finding in specific setting
  - Case finding in general population

## HCV screening during Flu-vaccination, season 2019/2020

Case finding dei soggetti HCV-positivi non noti tra la popolazione che accede al medico di medicina generale per la vaccinazione anti-influenzale, stagione 2019-2020

10 studi di medicina generale del distretto 35, ASL Caserta

Soggetti anti-HCV non noti che eseguono

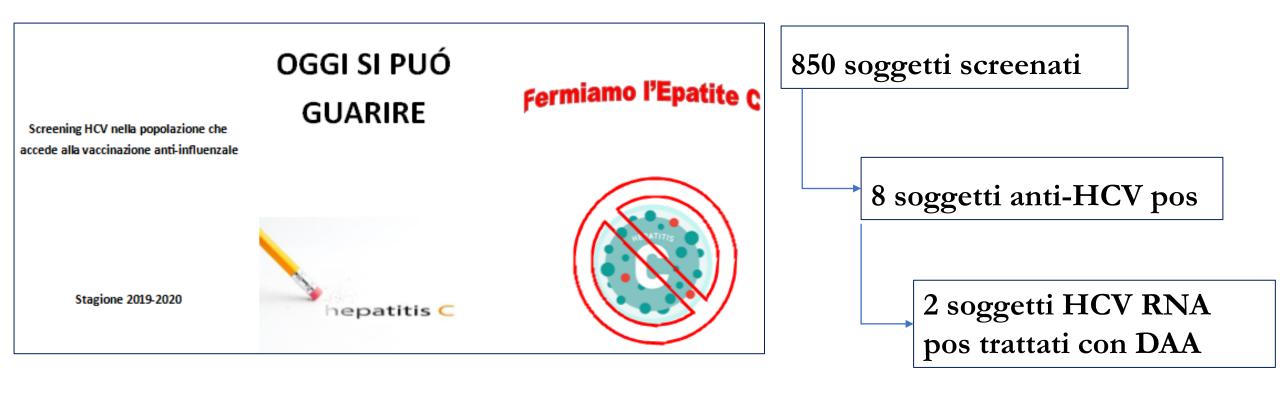
vaccinazione anti-influenzale



Test rapido per anti-HCV

## HCV screening during Flu-vaccination, season 2019/2020

Case finding dei soggetti HCV-positivi non noti tra la popolazione che accede al medico di medicina generale per la vaccinazione anti-influenzale, stagione 2019-2020



## Associated screening for HCV and SARS-Cov2 infection in an urban area of Southern Italy: A cohort study

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Carmine Coppola<sup>1</sup> | Mario Masarone<sup>2</sup> | Marco Bartoli<sup>3</sup> | Laura Staiano<sup>1</sup> | Roberta Coppola<sup>2</sup> | Pietro Torre<sup>2</sup> | Massimiliano Conforti<sup>3</sup> | Daniela Amoruso<sup>1</sup> | Ivan Gardini<sup>3</sup> | Marcello Persico<sup>2</sup> |
```

A prospective observational cohort study was set up with the aim of testing for both SARS-CoV2 and anti-HCV antibodies using rapid blood tests in all the available populations of Casola Di Napoli, a small town in the southern province of Naples

Variable	Overall (% on overall)	quick HCV-Ab negative (%)	quick HCV-Ab positive (%)	p
N (%)	2738 (100%)	2684 (98.03%)	54 (1.97%)	
Age mean (SD)	45.57 (19.45)	45.19 (19.34)	64.46 (14.73)	<0.0001
Sex % (M/F)	45.5/54.5	45.8/54.2	33.3/66.7	0.074
Not Italian (%)	0.6% (n16)	0.59%	0	0.56
HCV rapid test positive	54 (1.97%)	-	54	-
HCVAb confirmation positive	41 (1.49%)	-	41	-
HCV Already Known	36/41(87.8%)	-	36/54 (66.67%)	
HCVRNA positive	5 (0.18%)		5/54 (9.26%)	-
HCVRNA positivity not already known	3 (0.11%)	-	3/54 (5.55%)	
SARS-COV2 Ab positive	39 (1.4%)	38 (1.41%)	1/54 (1.9%)	0.54
IgM+IgG	17 (0.6%)	16 (0.59%)	1/54 (1.9%)	0.75
IgM	20 (0.7%)	20 (0.74%)	0	0.53
IgG	2 (0.1%)	2 (0.07%)	0	0.17
SARS-Cov2 NS swabs (PCR)	0	0	0	-
Age classes	Overall population	SARS-Cov2 IgM/IgG rapid blood test positive	HCV-Ab quick blood test positive	
<20 years	348 (12.7%)	0 (0%)	1 (0.3%)	
21-40 years	721 (26.3%)	2 (0.3%)	2 (0.3%)	
41-60 years	996 (36.4%)	14 (1.4%)	14 (1.4%)	
61-80 years	604 (22.1%)	20 (3.3%)	33 (5.5%)	
>81 years	69 (2.5%)	3 (1.4%)	4 (5.8%)	

# Challenges in HCV Case Finding and Linkage to Care

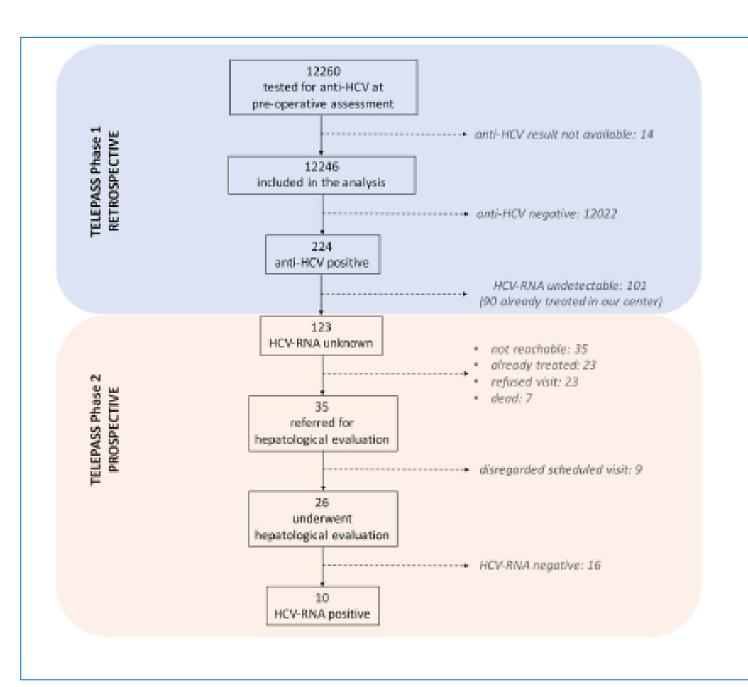
- General population
- Populations with high HCV prevalence
  - Opportunistic case-finding
- Case finding in specific setting
  - Hospital setting
  - Emergency department
- Case finding in general population

Missed linkage to care for patients who screened positive for Hepatitis C in a tertiary care centre: Results of the Telepass project

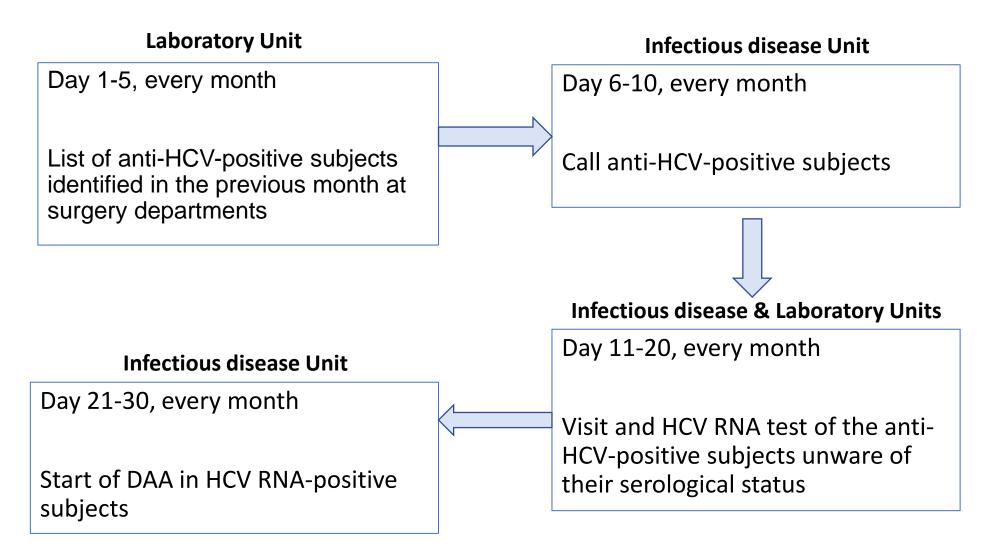
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Francesca Romana Ponziani<sup>1</sup> | Francesco Santopaolo<sup>2</sup> | Massimo Siciliano<sup>1</sup> | Antonio Giulio De Belvis<sup>3</sup> | Annalisa Tortora<sup>2</sup> | Vincenzina Mora<sup>1</sup> | Caterina Fanali<sup>1</sup> Alisha Morsella<sup>4</sup> | Fulvio Balducci<sup>5</sup> | Giuseppe Vetrugno<sup>6</sup> | Maria Elena D'Alfonso<sup>7</sup> | Andrea Cambieri<sup>8</sup> | Roberto Cauda<sup>9</sup> | Rocco Bellantone<sup>10</sup> | Maurizio Sanguinetti<sup>11</sup> | Maurizio Pompili<sup>1</sup> | Antonio Gasbarrini<sup>1</sup> |
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The 'Telepass' project was structured in two phases:

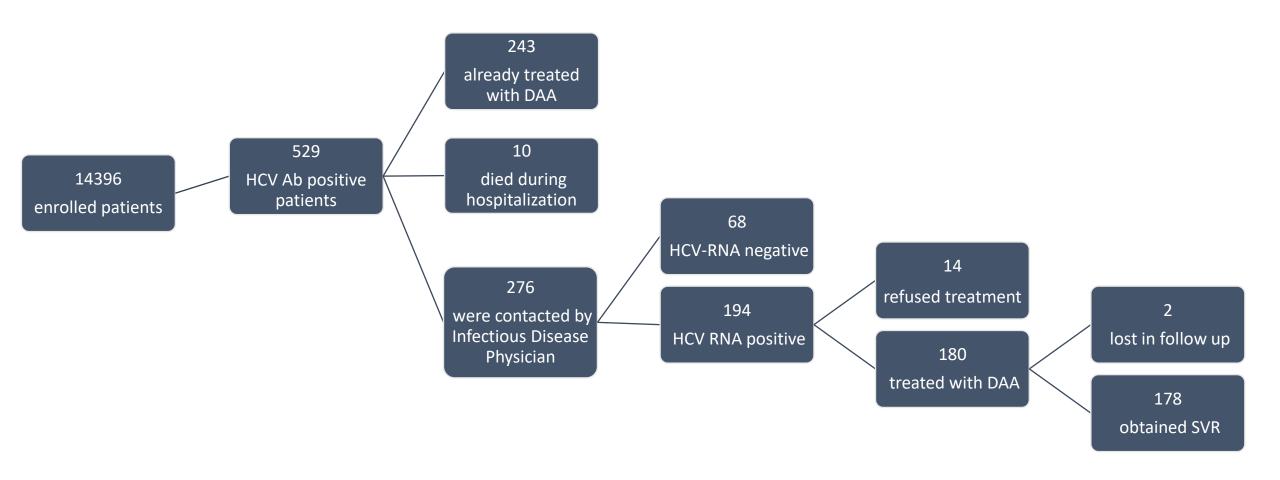
- a retrospective analysis first identified all anti—HCVpositive subjects among patients who underwent pre-operative assessment in the facility in the course of one year;
- a following prospective phase, aimed to recall patients in need either of further diagnostic tests (ie HCV-RNA) or treatment



# HCV-free Caserta-Hospital project, January-December 2019



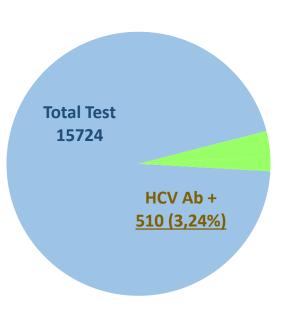
# HCV-free Caserta-Hospital project, January-December 2019







## Periodo Gennaio 2020 - giugno 2021

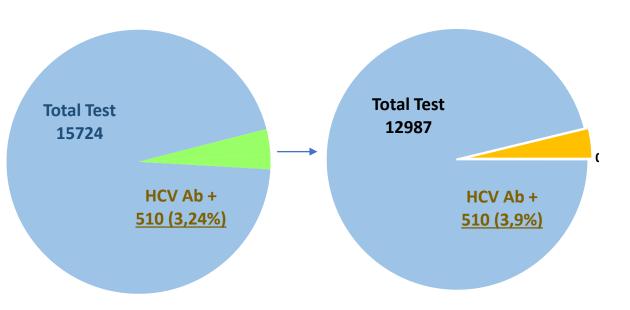


Analisi sul totale dei pazienti





### Periodo Gennaio 2020 - giugno 2021



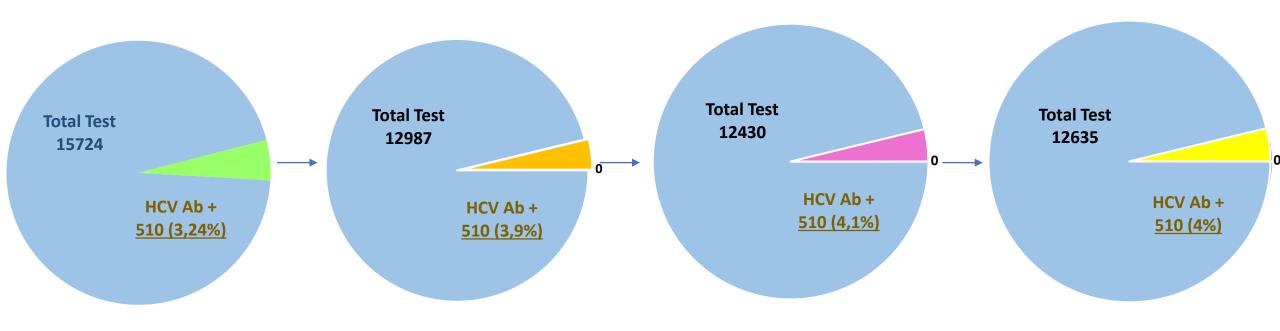
Analisi sul totale dei pazienti

Esclusi i pazienti neonati (Ostetricia + TIN)





#### Periodo Gennaio 2020 - giugno 2021



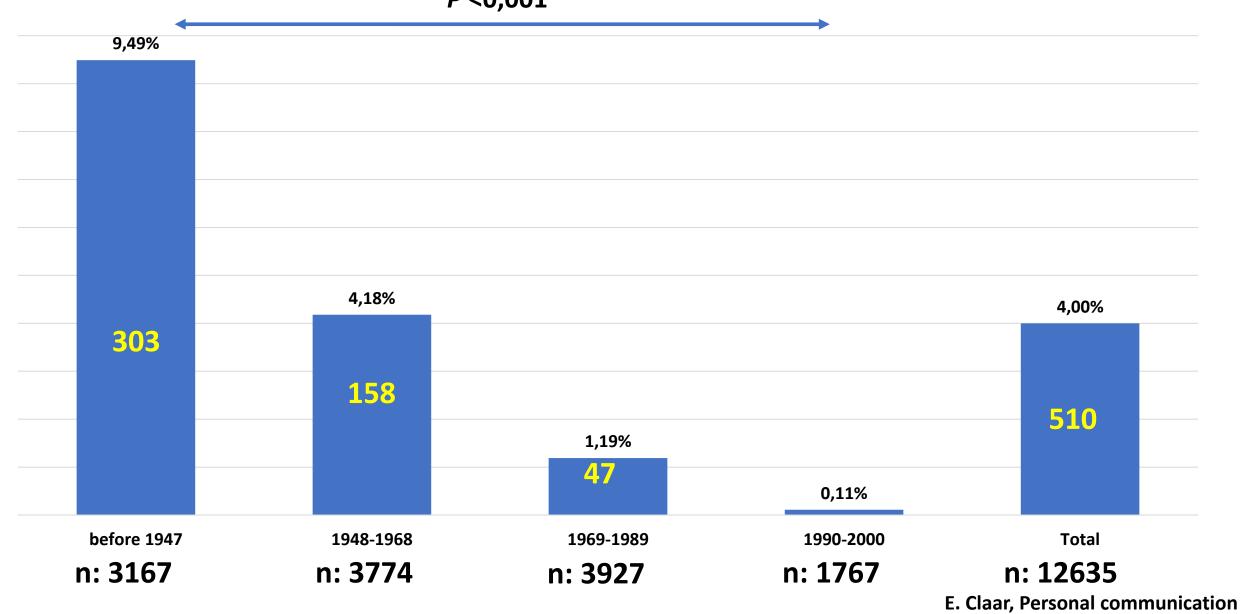
Analisi sul Totale dei pazienti Esclusi i pazienti neonati (Ostetricia + TIN) Esclusi i pazienti 0 – 23 anni Esclusi i pazienti nati dopo 2000



#### **HCV** ab prevalence matched by age of birth



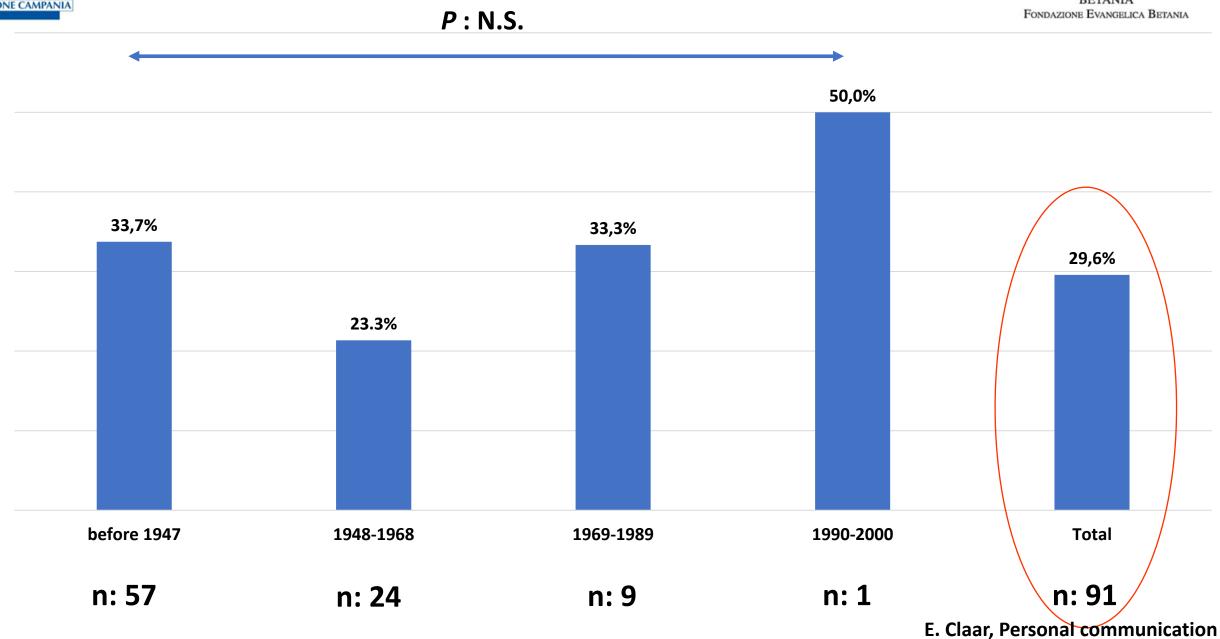






#### **HCV RNA** prevalence on HCV ab matched by age of birth

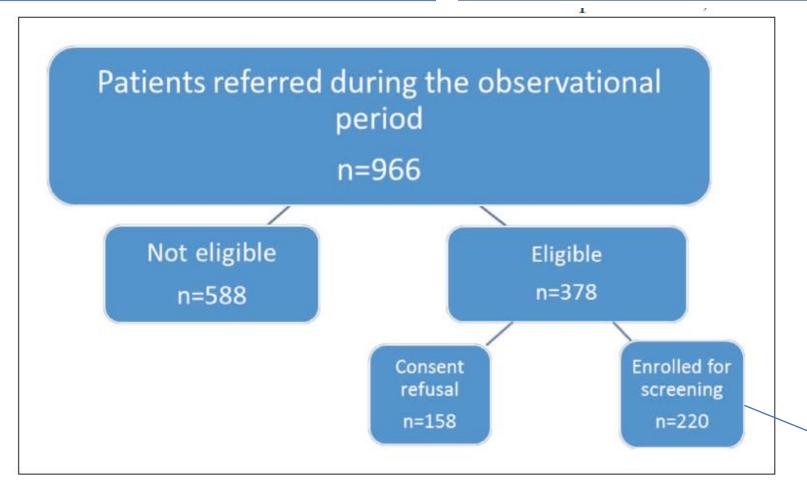




# Hepatitis C screening in the Emergency Department of a large hospital in Southern Italy: results of a pilot study

Ivan Gentile¹, Biagio Pinchera¹, Giulio Viceconte¹, Manuel Crispo¹, Davide Simeone¹, Riccardo Scotto¹, Emanuela Zappulo¹, Alberto Enrico Maraolo¹, Fiorella Paladino², Raffaella Tortora², Giovanni Giuseppe Di Costanzo², Antonio Riccardo Buonomo², Guglielmo Borgia¹

To evaluate the feasibility of a rapid salivary, point-of-care assay for anti-HCV, performed in patients aged between 45 and 80 years old who were referred to the emergency department from May to August 2017 and were all unaware of their HCV serostatus



2 anti-HCV-positive

0 linkaged-to-care

## Emergency department testing is feasible but ineffective to eliminate Hepatitis C in Denmark

Jessica Jennifer Wentworth<sup>1,2</sup>, Anne Lindebo Holm Øvrehus<sup>1,2</sup>, Janne Fulgsang Hansen<sup>1</sup>, Peter Biesenbach<sup>3</sup>, Peer Brehm Christensen<sup>1,2</sup>

During a three-month period (May-August 2020), emergency department patients at Odense University Hospital were screened for risk factors and offered point-of-care HCV-Antibody testing

#### Figure 1: Flow of Participants **ED** presentations during study time Excluded (n= 1.457, 44.3%) (n=3,288)Multiple reasons possible -Age <18 or >80 (n=957) -Isolated (n=269) -Altered mental status (n=130) -Assessed ineligible by nurse (n=121) SCREENED -Critically ill (n=60) -Other (n=104) Eligible (n= 1,831, 55.7%) Left ED before inclusion could be offered. (n= 1,158, 63.2%) Approached (n= 673, 36.8%) Declined participation (n= 159, 23.6%) Enrolled, Risk factor INCLUDED survey performed (n= 514, 76.4%) Risk factor negative, test declined. (n=25)Participation concluded. **HCV**- antibody quicktest performed (n= 489), of which: 339 risk factor positive 150 risk factor negative Quicktest negative (n= 485) Participation concluded. HCV-Ab positive (n= 4) TESTED Offered RNA testing Declined venous blood test (n= 2) Participation concluded. Venous blood test for HCV-RNA (n= 2) RNA negative (n= 2) Participation concluded. HCV-RNA positive (n= 0) Linked to care (n= 0)

# Challenges in HCV Case Finding and Linkage to Care

- General population
- Populations with high HCV prevalence
  - Opportunistic case-finding
  - Case finding in specific setting
- Case finding in general population





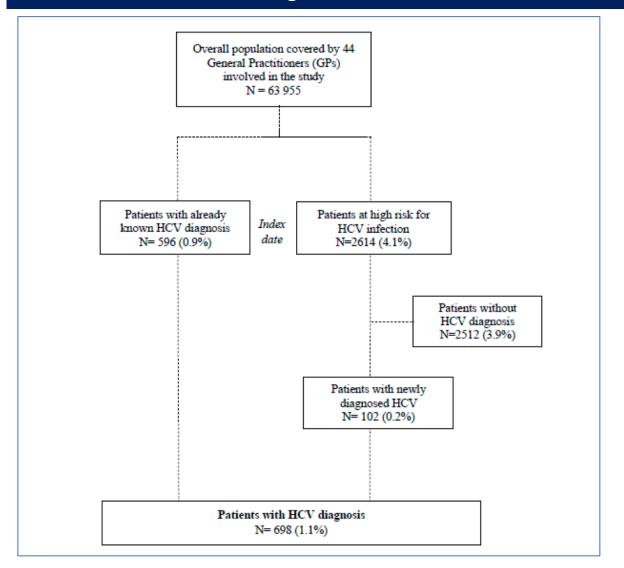
Artic

Screening, Linkage to Care and Treatment of Hepatitis C Infection in Primary Care Setting in the South of Italy

Anna Citarella <sup>1,\*,†</sup>, Simona Cammarota <sup>1,†</sup>, Francesca F. Bernardi <sup>2</sup>, Carmine Coppola <sup>3</sup>, Maria D'Antò <sup>4</sup>, Marianna Fogliasecca <sup>1</sup>, Elio Giusto <sup>5</sup>, Mario Masarone <sup>6</sup>0, Angelo Salomone Megna <sup>7</sup>, Carmine Sellitto <sup>8</sup>0, Rosa Servodio <sup>9</sup>, Massimo Smaldone <sup>10</sup>, Laura Staiano <sup>3</sup>, Ugo Trama <sup>2</sup>, Valeria Conti <sup>8,‡</sup> and Marcello Persico <sup>6,‡</sup>

A retrospective cohort study of 44 general practitioners (GPs) who managed 63,955 inhabitants in the Campania region.

Adults with already known HCV diagnosis or those with HCV high-risk profile at June 2019 were identified and reviewed by GPs to identify newly diagnosed of HCV and to assess the linkage to care and treatment for the HCV patients.



Life 2020





Articl

Screening, Linkage to Care and Treatment of Hepatitis C Infection in Primary Care Setting in the South of Italy

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Table 1. Demographic and clinical characteristics of patients with hepatitis C virus (HCV) diagnosis.

	Overall $(N = 698)\%$	Patients with Already Known HCV Diagnosis (N = 596)%	Patients with Newly Diagnosed HCV (N = 102)%	p Value
Age Groups				
<40	5.2	5.4	3.5	0.13
40-49	9.7	10.0	8.0	
50-59	13.7	12.4	21.8	
60-69	21.5	22.4	16.1	
≥70	49.9	49.8	50.6	
Gender				
Male	48.9	47.4	57.6	0.06
Female	51.1	52.6	42.4	
Comorbidities				
Diabetes	14.7	14.3	16.7	0.54
CKD	4.9	4.2	8.8	0.05
Obesity	4.3	3.7	7.8	0.06



40-49

50-59

Age group

■ Not treated with DAA and not referred to the specialist centre

■ Referred to the specialist centre, treated and responded to DAA

■ Referred to the specialist centre for DAA treatment

<40

41.5

16.2

60 - 69

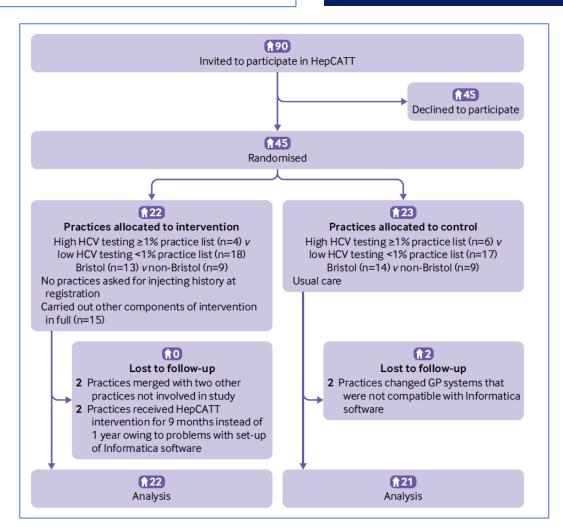
43.5

>70

#### Cost effectiveness of an intervention to increase uptake of hepatitis C virus testing and treatment (HepCATT): cluster randomised controlled trial in primary care

Kirsty Roberts, <sup>1</sup> John Macleod, <sup>1,2</sup> Chris Metcalfe, <sup>1,3</sup> Will Hollingworth, <sup>1</sup> Jack Williams, <sup>4,5</sup> Peter Muir, <sup>6</sup> Peter Vickerman, <sup>1,2</sup> Clare Clement, <sup>2</sup> Fiona Gordon, <sup>7</sup> Will Irving, <sup>8</sup> Cherry-Ann Waldron, <sup>9</sup> Paul North, <sup>6</sup> Philippa Moore, <sup>6</sup> Ruth Simmons, <sup>5,10</sup> Alec Miners, <sup>4,5</sup> Jeremy Horwood <sup>1,2</sup> Matthew Hickman <sup>1,2</sup>

Electronic algorithm and flag on practice systems identifying patients with HCV risk markers (such as history of opioid dependence or HCV tests with no evidence of referral to hepatology), staff educational training in HCV, and practice posters/leaflets to increase patients' awareness. Flagged patients were invited by letter for an HCV test (with one follow-up) and had on-screen pop-ups to encourage opportunistic testing.



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Table 3 | Hepatitis C virus (HCV) antibody testing, HCV positive test yield, polymerase chain reaction (PCR) tests for chronic infection, and referral to secondary care in intervention and control practices, with intervention effect estimated as rate ratio from random effects Poisson regression model that accommodates any variations in testing between practices

Number (%)			
Intervention (n=13097)	Control (n=11376)	Rate ratio (95% CI)	P value
2071 (15.8)	1163 (10.2)	1.57 (1.18 to 2.09)	0.002
		1.59 (1.21 to 2.08)	< 0.001
189/2930 (6.5)	80/3315 (2.4)	2.73 (1.95 to 3.82)	-
1882/10167 (18.5)	1083/8061 (13.4)	1.45 (1.08 to 1.95)	-
-	-	1.91 (1.45 to 2.52)	< 0.001
129 (1.0)	51 (0.4)	2.30 (1.41 to 3.75)	0.001
		2.24 (1.47 to 3.42)	< 0.001
43 (0.3)	13 (0.1)	3.17 (1.38 to 7.31)	0.007
		2.96 (1.34 to 6.58)	0.008
20 (0.2)	3 (<0.1)	6.25 (1.67 to 23.38)	0.007
		5.78 (1.55 to 21.61)	0.009
ınalysis)			
27 (0.2)	7 (<0.1)	3.43 (1.36 to 8.65)	0.009
		3.40 (1.35 to 8.52)	0.009
	Intervention (n=13 097)  2071 (15.8)  189/2930 (6.5) 1882/10 167 (18.5)  129 (1.0)  43 (0.3)  20 (0.2)  Inalysis)	Intervention (n=13097) Control (n=11376)  2071 (15.8) 1163 (10.2)  189/2930 (6.5) 80/3315 (2.4) 1882/10167 (18.5) 1083/8061 (13.4)	Intervention (n=13097)   Control (n=11376)   Rate ratio (95% CI)

<sup>\*</sup>Adjusted for practice location (Bristol versus elsewhere) and historical HCV testing rate (low versus high, as indicated by Public Health England). †Subgroups defined by history of opioid/injecting drug use.

<sup>‡</sup>Estimated ratio of rate ratios in two subgroups (opioid/injecting drug use and no opioid/injecting drug use, and control practices as reference within each), with interaction test P value estimated from model with covariates as in above\*.

#### Cost effectiveness of an intervention to increase uptake of hepatitis C virus testing and treatment (HepCATT): cluster randomised controlled trial in primary care

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Task	Intervention (n=12922)	Control (n=10 974)	Difference (95% CI)
Training cost	£1.22	£0	-
Screening cost	£2.06	£0	-
Mean HCV antibody test cost per patient	£3.54	£2.33	£1.21 (£1.02 to £1.40)
Mean HCV PCR test cost per patient	£0.89	£0.41	£0.48 (£0.28 to £0.68)
No (%) HCV related consultations: no; yes	12 187 (94); 735 (6)	10 467 (95); 507 (5)	
Mean HCV related consultation cost per patient	£2.27	£2.10	£0.17 (-£0.09 to £0.44
Mean hepatology referral cost per patient	£0.44	£0.12	£0.32 (£0.12 to £0.52)
Total mean case finding cost per patient	£10.42	£4.96	£7.10 (£4.75 to £9.45)
No (%) patients referred to hepatology for treatment	20 (0.15)	3 (0.03)	-
Cost per additional patient referred to hepatology for treatment	-	-	£5569

PCR=polymerase chain reaction.

#### CONCLUSION

HepCATT had a modest impact but is a low cost intervention that merits optimisation and implementation as part of an NHS strategy to increase HCV testing and treatment.

<sup>\*</sup>Adjusted mean difference from mixed effects linear regression, clustered by practice, adjusted for previous HCV testing, Bristol practice, and length of follow-up.

### **Conclusions**

- HCV treatment is safe and efficacious
- Today case finding and linkage to care are the milestones in the HCV management
- Delocalization approach may be useful in special population
- In general population case new finding and linkage-tocare approaches are needed

