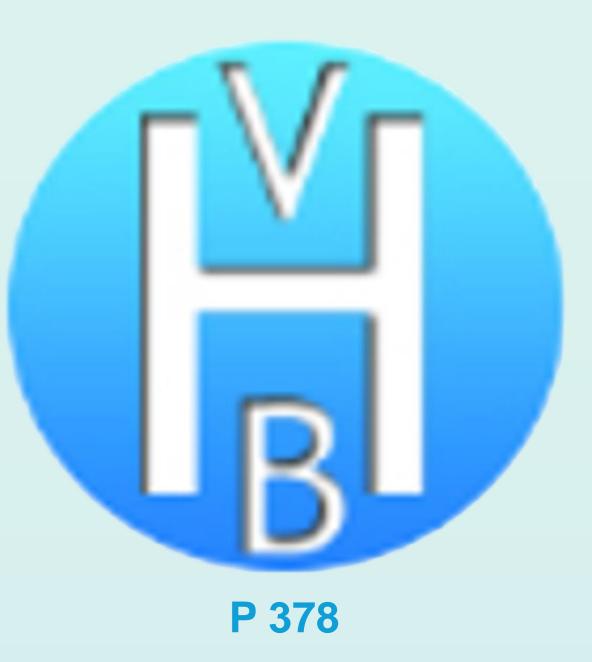


# HCV CURE WITH DIRECT-ACTING ANTIVIRALS IN HIV/HCV COINFECTED PATIENTS BELONGING TO KEY POPULATIONS (HepCURE study)

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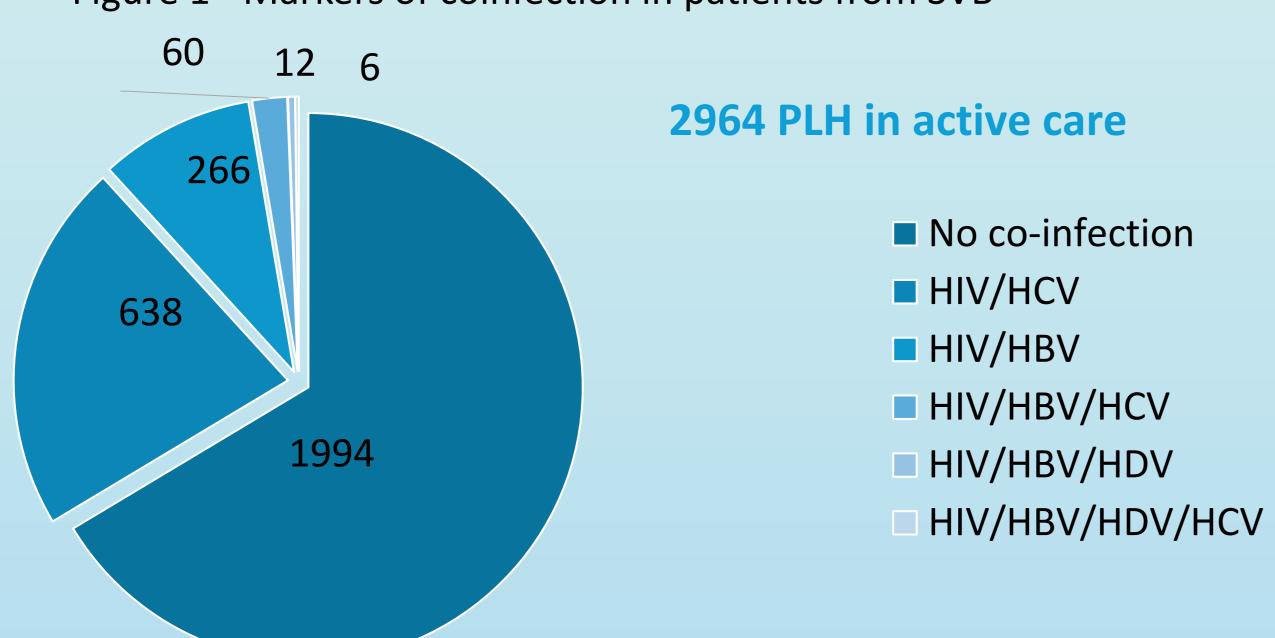
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### Background

Direct-acting antiviral (DAA) treatment is a priority as HIV/HCV co-infection accelerates liver disease progression. This study aimed to assess the response to DAA treatment in HIV/HCV coinfected patients belonging to key populations.

Figure 1 - Markers of coinfection in patients from SVB



#### Results

In total, 122 HIV/HCV coinfected patients started treatment with DAAs. Results on socio-demographic and clinical characteristics are presented in <u>Table 1</u> and <u>Figure 3</u>. Socio-economic barriers and the lack of pan-genotypic DAAs limited outcomes in the first years of the study. HCV reinfection was diagnosed in one case. There was a marked increase in the use of DAAs during the years, from 4.5% in 2017 to 100% of the newly diagnosed cases in 2023.

Figure 2 – Modes of HCV acquisition in patients from key populations 3 (2%)

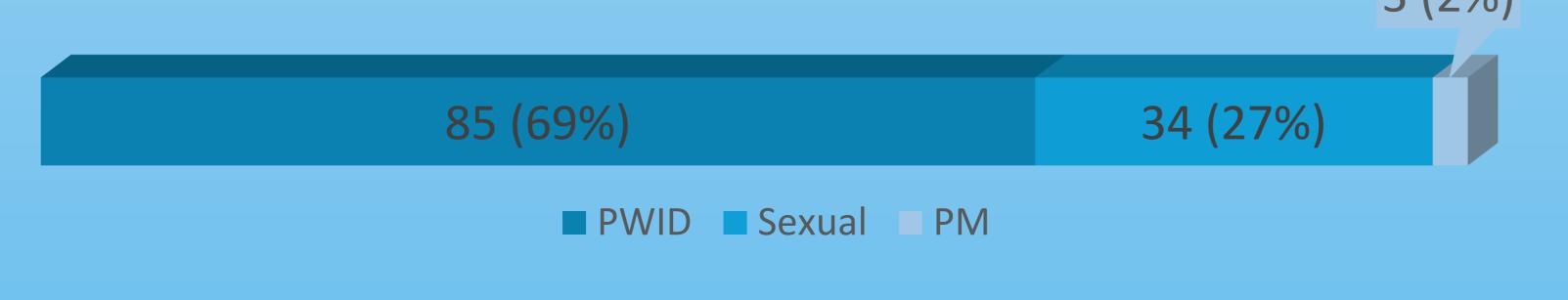
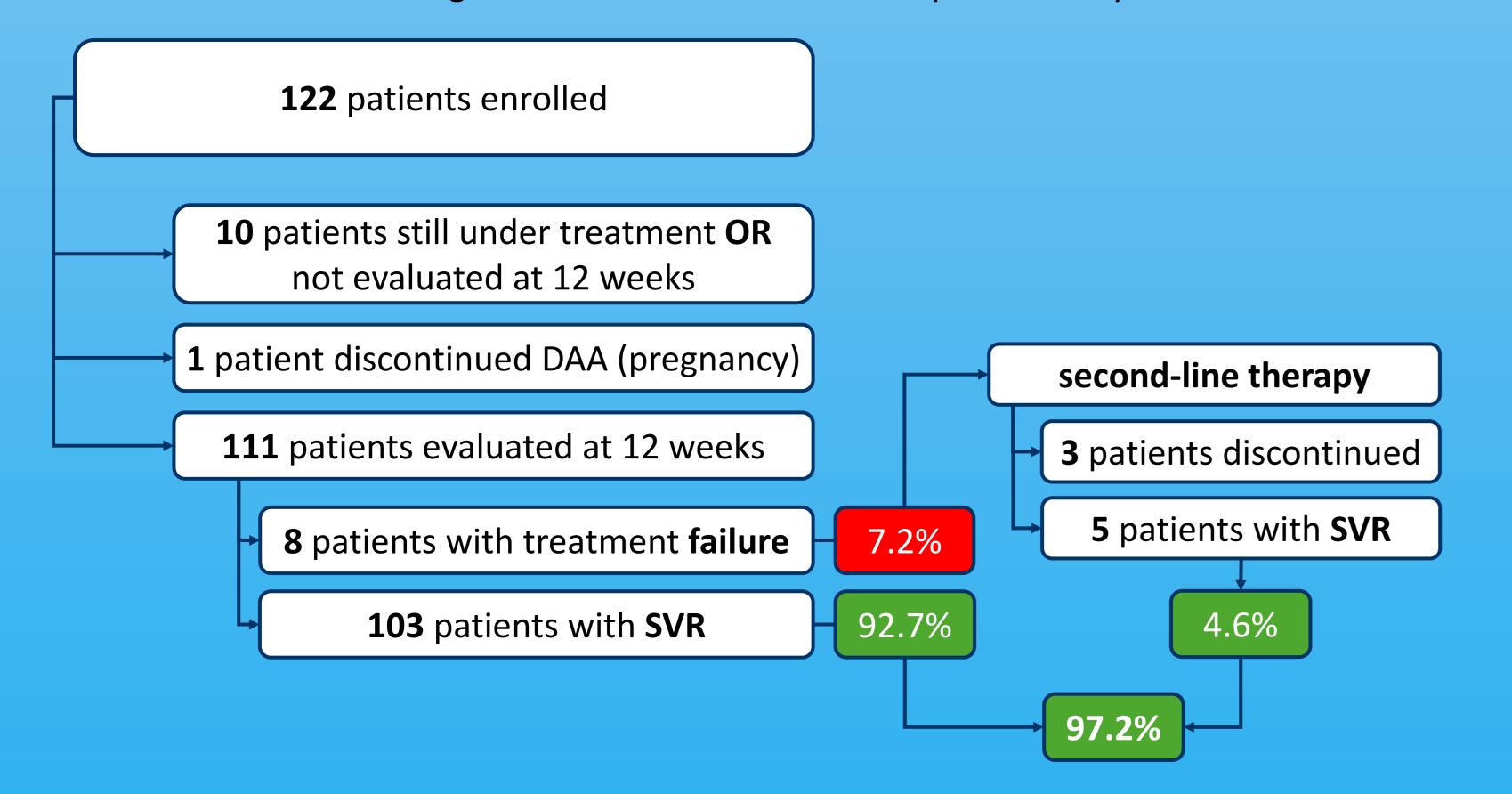


Figure 3 – Patients enrolled in HepCURE study



## Methods

Prospective study performed on PLH, with confirmed HCV infection, in active care at "Victor Babes" Hospital for Infectious and Tropical Diseases (SVB), Bucharest, between 1 January 2017 and 31 October 2023. Patients were stratified by modes of HCV acquisition in: injecting drug use (PWIDs), sexual transmission, and parenteral mode, during early childhood (Figure 2).

Table 1 - Demographic and clinical characteristics - comparison by modes of HCV acquisition

Characteristics		Total n=122	PWIDs n=85	Non-PWIDs n=37	P value			
Gender (male)	n (%)	88 (72.7)	67 (79.7) 21 (56.7)		0.007			
Λαο (νιοονο)	median	41	40	42	0.005			
Age (years)	(IQR)	(35, 48)	(35, 43)	(34, 52)				
CD4 cell count/μL	median	663	660	665	0.764 5)			
	(IQR)	(385, 882)	(423, 867)	(336, 955)				
HIV-RNA < 50 c/mL	n (%)	95 <b>(80.5</b> )*	65 (76.4)**	31 (88.5)***	0.044			
Prior IFN α	n (%)	8 (6.5)	3 (3.5)	5 <b>(13.5)</b>	0.039			
Fibrosis stage								
F0-F1	n (0/)	55 (46.6)	38 (41.1)	17 (45.9)				
F1-F2	n (%)	43 (36.4)	29 (34.1)	15 (40.5)	0.886			
F3-F4		20 (16.9)	14 (16.4)	5 (13.5)				
*out of 118 available; **out of 85 available; *** out of 35 available								

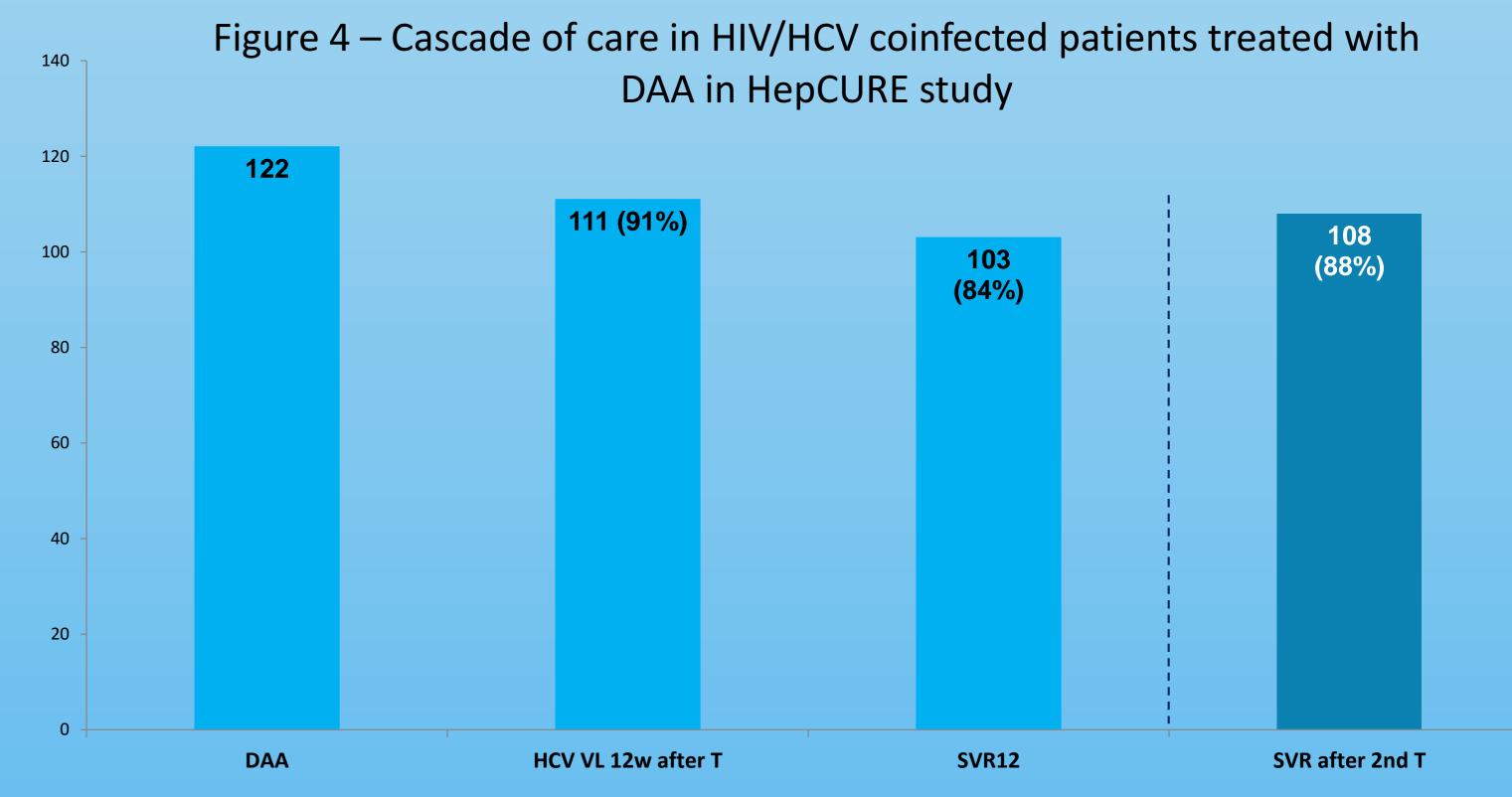


Table 2 Availability of DAA during the study years

Period	Drug		Fibrosis stage	
2017	Ombitasvir/paritaprevir /ritonavir	1%	F2, F3, F4	4.1%
2018 - 2019	Grazoprevir/elbasvir Sofosbuvir/ledipasvir		F1, F2, F3, F4	
2020	Sofosbuvir/velpatasvir Sofosbuvir/ledipasvir		F0, F1, F2, F3, F4	47.1%
2022 - 2024	Sofosbuvir/velpatasvir/ Sofosbuvir/velpatasvir/ voxilaprevir		F0, F1, F2, F3, F4 for those with treatment failure	1%

# Conclusions

DAA treatment success rate in HIV/HCV coinfected patients from key population was high and comparable to those monoinfected. The SVR rates were similar in PLH infected by sexual mode or in PWIDs, irrespective of the CD4 cell count or HIV-VL. Elimination of HCV requires a targeted scale-up of DAA treatment and behavioral interventions in particular among high-risk populations. This study underlines the need for strong interventions meant to improve linkage and care retention in HIV/HCV coinfected patients.