# CD4 cell count changes following acute HCV infection in chronically HIV infected patients

Luuk Gras<sup>1</sup>, Colette Smit<sup>1</sup>, Maria Prins<sup>2,3</sup>, Frank de Wolf<sup>1,4</sup>, Janke Schinkel<sup>3</sup>, Femke Lambers<sup>3</sup>, Joost van Hommeriq<sup>2</sup>, Jan van der Meer<sup>3</sup>, Ronald Geskus<sup>2,3</sup> and the ATHENA national observational HIV cohort <sup>1</sup>Stichting HIV Monitoring, Amsterdam, the Netherlands; <sup>2</sup>Municipal Health Service, Amsterdam, the Netherlands, <sup>3</sup>Academical Medical Centre of the University of Amsterdam, Amsterdam, the Netherlands 4Imperial College School of Medicine, London, U.K.

# Background

- Outbreaks of acute HCV among HIV-positive men who have sex with men (MSM) have been reported.
- HCV co-infection contributes to morbidity and mortality and complicates patient management.
- Limited data is available on the impact of acute HCV co-infection on CD4 cell counts during chronic HIV infection.
- Objective: To study changes in CD4 cell counts before and after acute HCV infection but before the start of anti-HCV therapy, among both cART treated and untreated chronically HIV-infected natients.

### Methods

### HIV-1 infected MSM selected from the national observational ATHENA cohort with:

- Acute HCV co-infection: a negative HCV RNA test result followed within 1 year by a positive HCV RNA test result (midpoint = estimated date of HCV infection).
- At least 6 months between HIV diagnosis and HCV infection.

#### Outcome:

- CD4 cell counts 2 years prior and 1 year after the date of HCV infection.
- CD4 cell counts sampled after the start of anti-HCV treatment or after a change in the HIV treatment status (change from HIV untreated to treated with any HIV-antiretroviral or vice-versa) were censored.

#### Statistical analysis:

- CD4 cell counts were longitudinally modelled usina:
  - mixed effects models.
  - pattern-mixture models (to account for possible bias arising from censoring CD4 cell counts after start of anti-HCV treatment). This model includes an interaction term with CD4 slope after HCV infection and timing of anti-HCV treatment initiation (< and ≥6 months from HCV infection).
- Slopes were allowed to change at date of HCV seroconversion.
- A random intercept (representing CD4 cell counts at HCV infection) and 2 random slopes for each patient were included.
- HIV treatment status was included in all models.

#### N(%) Total 39 (100) **HCV** genotype 22 (56) 2 2(5)7 (18) not determined 8 (21) 29 (74%) On cART Median (IQR) Known duration of HIV (years) 5.9 (2.1-10.1) Last CD4 cell count prior to HCV 480 (390-620) infection (cells/mm<sup>3</sup>) Year of HCV infection 08 (06-09) 4.5 (3.3-6.4) Months between last negative first positive HCV RNA test

Table 1. Characteristics at estimated HCV infection.

 Anti-HCV treatment within 6 months from HCV diagnosis in 17 patients (44%).

	model	ture model
CD4 cell count at HCV infection (cells/mm³)		
On cART HIV-untreated	514 (458-570) 445 (348-541)	515 (461-568) 448 (357-540)
Slope CD4 cell count prior to HCV infection (cells/mm³/yr)		
On cART HIV-untreated	18 (-5, 42) -70 (-118, -21)	19 (-5, 43) -65 (-114, -16)
Slope CD4 cell count after HCV infection		

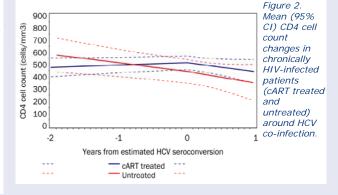
(cells/mm<sup>3</sup>/yr) On cART HIV-untreated

-72 (-163,20) -82 (-198, 35) -95 (-228, 38) -134 (-200, 46)

Mixed effects Pattern-mix-

Table 2. Estimates of mean (95% CI) CD4 cell count at HCV co-infection and changes in CD4 cell counts around HCV co-infection.

· In cART treated patients differences in slope before and after HCV infection were 90 cells/mm<sup>3</sup> (-179, 0; p=0.04) for the mixed effects model and  $-101 \text{ cells/mm}^3$  (-219,17; p=0.09) for the pattern-mixture model.



#### Results

- Characteristics at HCV co-infection of the 39 included patients are shown in Table 1.
- · Analyses included 360 CD4 cell counts.

## Conclusions

- Acute HCV infection in cART-treated chronically HIV-infected patients may be associated with a decrease in CD4 cell counts.
- The small number of patients limited subgroup analyses.

Luuk Gras Stichting HIV Monitoring E:I.a.gras@amc.uva.nl www.hiv-monitoring.nl