

# HCV late presentation

## (severe fibrosis/cirrhosis at first HCV therapy or first HCV diagnosis) in the Swiss Association for the Medical Management in Substance Users (SAMMSU) cohort since 2017

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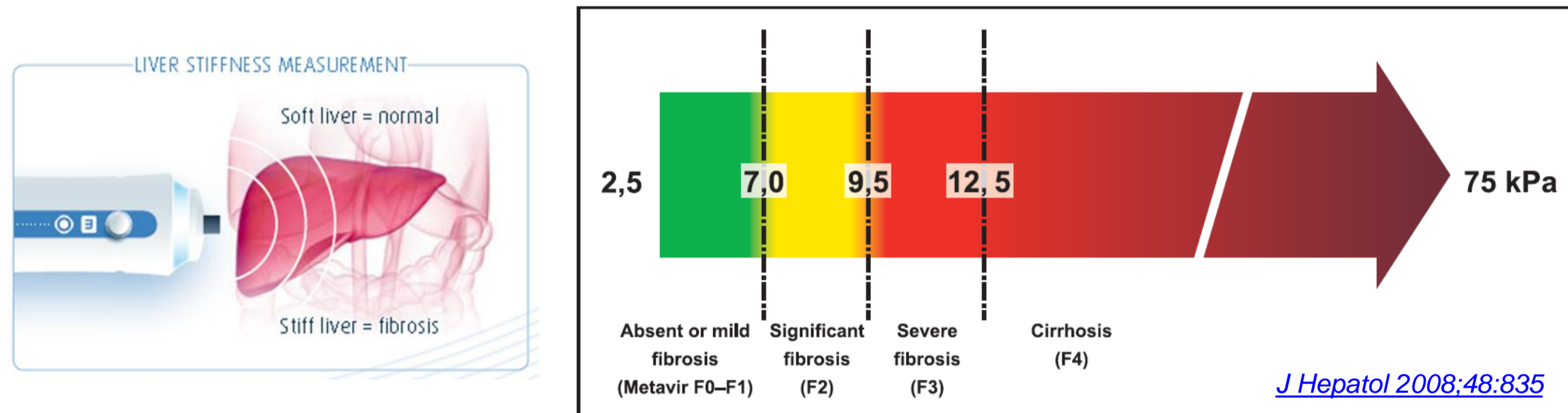
### BACKGROUND

- In Switzerland, direct-acting antivirals (DAAs) are reimbursed without liver fibrosis restriction since 2017.
- We determined the prevalence of HCV late presentation in times of unrestricted DAA access.

### METHODS

- The SAMMSU-cohort is an open cohort with yearly follow-up, enrolling >18-year-old patients with current/previous opioid agonist therapy (OAT) in eight different centers throughout Switzerland since 2014.
- By 02/01/2024, 1,390 patients have been enrolled (76% male, median age: 44y, 75% ever intravenous drug use, 61% HCV- and 10% HIV-antibody-positive).
- HCV late presentation (LP) was defined as F3/F4-fibrosis (Fibroscan® >9.5kPa) at first HCV therapy and first HCV diagnosis, respectively.

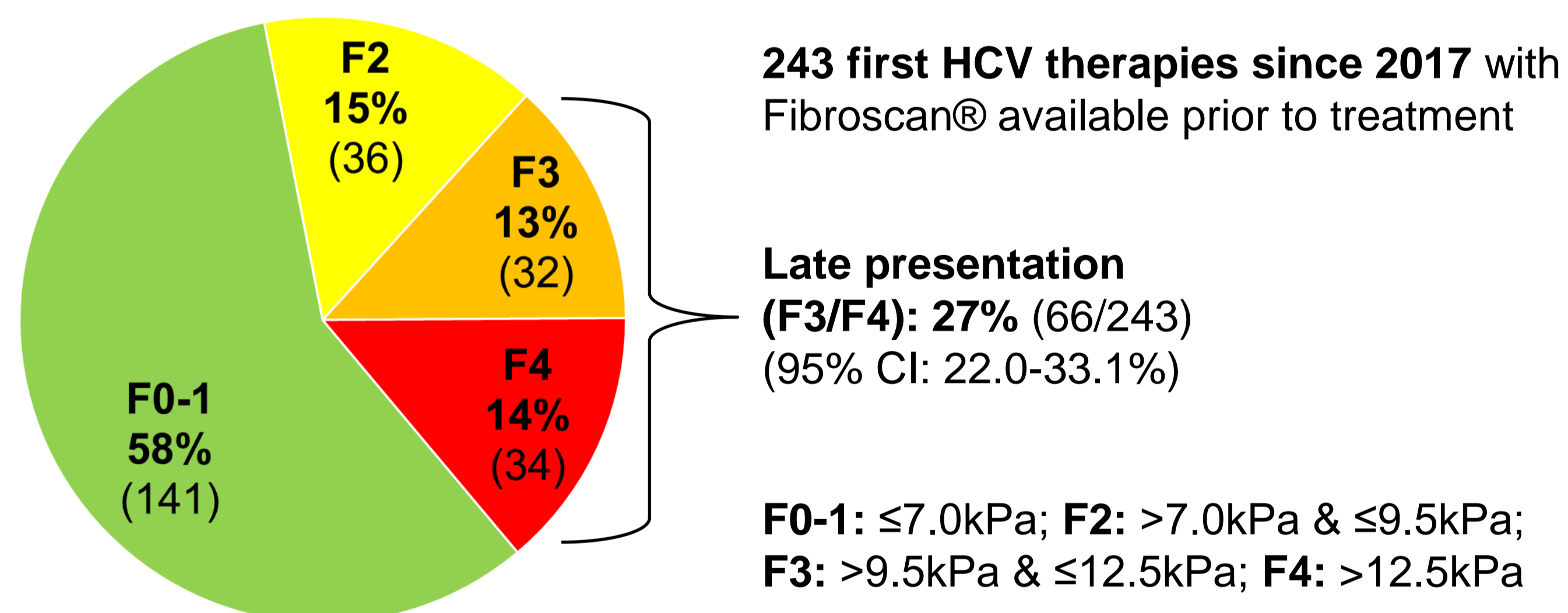
#### Liver fibrosis assessment with Fibroscan® (transient elastography)



### RESULTS – Late presentation at first HCV therapy

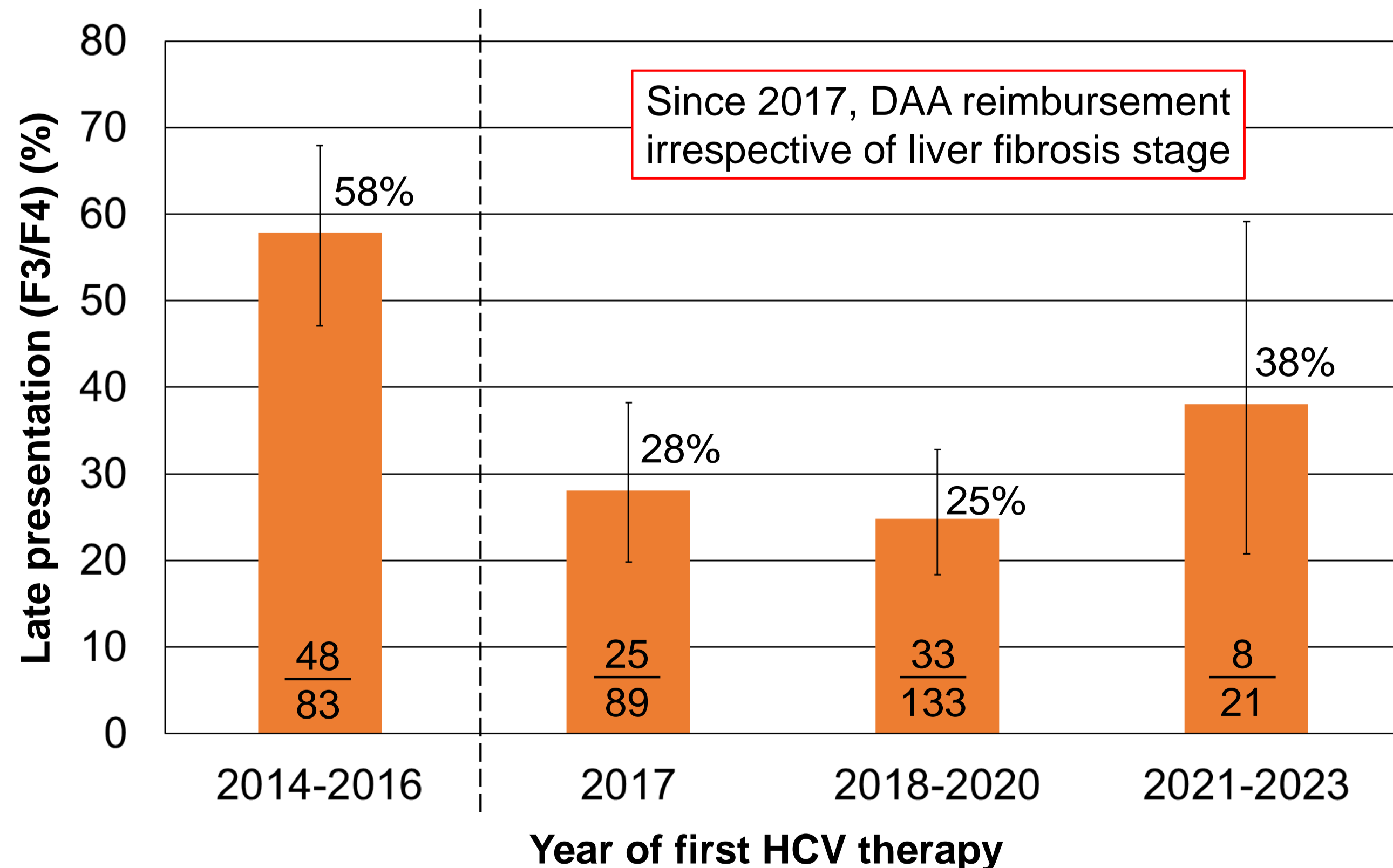
- For 90.3% (243/269) first HCV therapies since 2017, Fibroscan® was available prior to treatment: 58.0% had no/mild fibrosis (F0/F1), 14.8% significant fibrosis (F2), 13.2% severe fibrosis (F3) and 14.0% cirrhosis (F4) [LP: 27.2% (66/243)].

#### Liver fibrosis stage (Fibroscan®) at first HCV therapy (2017 onwards)



- LP prevalence was 57.8% (48/83) in 2014-2016, 28.1% (25/89) in 2017, 24.8% (33/133) in 2018-2020 and 38.1% (8/21) in 2021-2023.

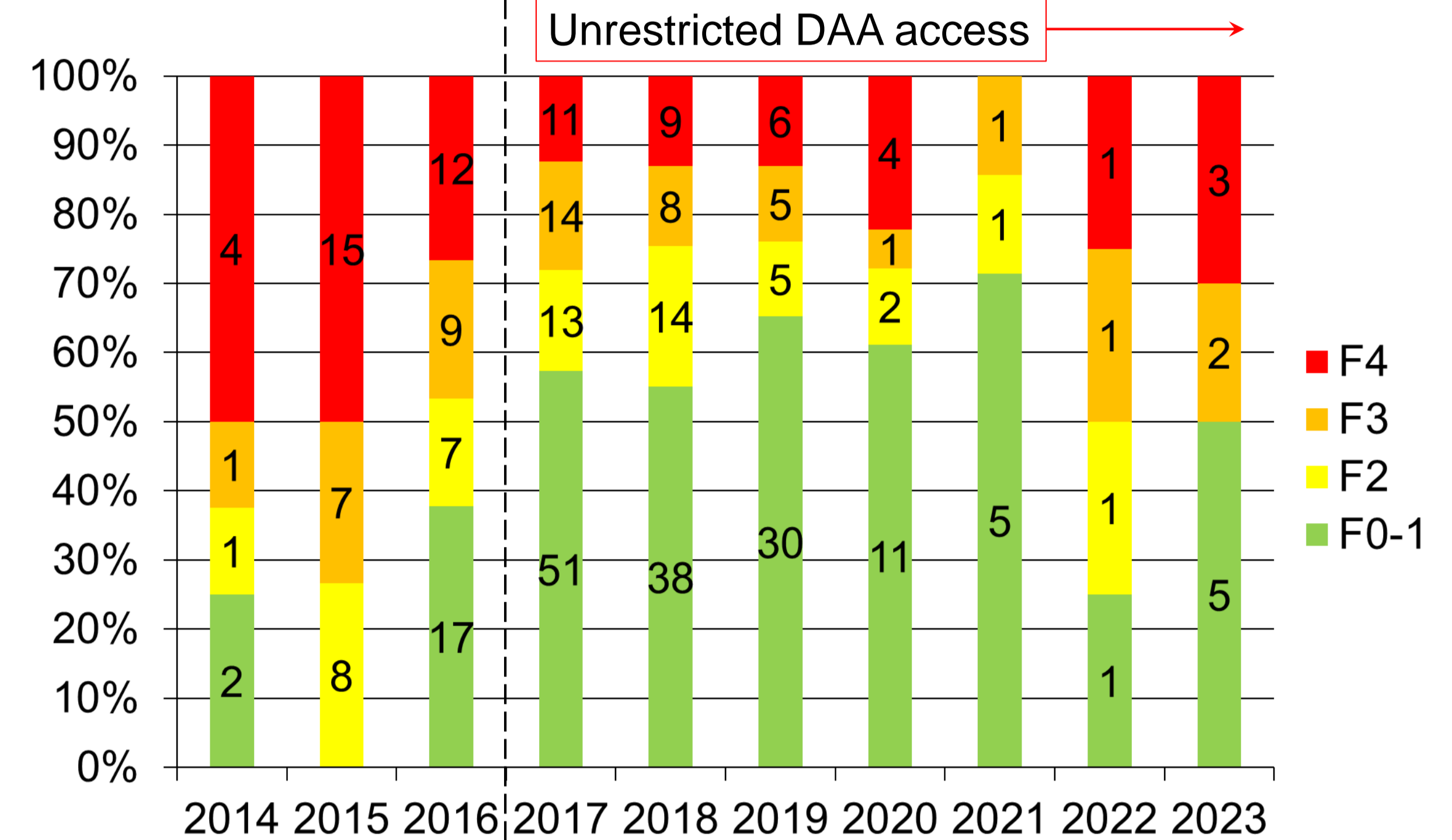
#### Proportion with late presentation at first HCV therapy (before and after 2017)



### ACKNOWLEDGEMENT/SPONSORING

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#### Liver fibrosis stage at first HCV therapy according to calendar year



F0-1: ≤7.0kPa; F2: >7.0kPa & ≤9.5kPa; F3: >9.5kPa & ≤12.5kPa; F4: >12.5kPa

- Risk factors** for LP were **male sex** (OR 1.9, p=0.078) and **alcohol** (≥49g/d) (OR 3.2, p<0.001), whereas diagnosis of **schizophrenia** was **negatively associated** (OR 0.4, p=0.015).
- Comparing LP versus non-LP at first HCV therapy, median age was 47.1 versus 45.9 years (p=0.250), median time since HCV diagnosis 12.5 versus 11 years (p=0.472) and median time since first intravenous drug use (IDU) 28 versus 25 years (p=0.357).

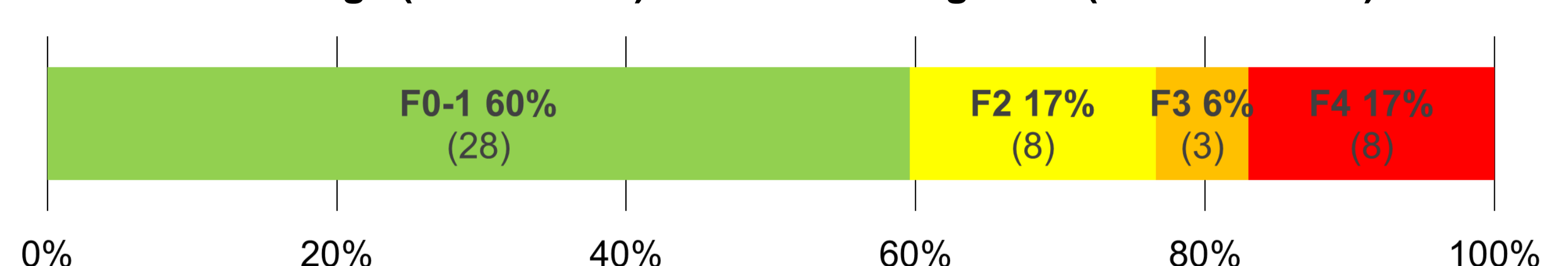
#### Patient characteristics (Late Presentation (LP) versus non-LP) (n=243)

	LP (n=66)	Non-LP (n=177)	p	OR (95% CI)
<b>Male (%)</b>	81.8% (54/66)	70.6% (125/177)	0.078	1.9 (0.9-3.8)
<b>Median age (y) at HCV therapy (IQR)</b>	47.1 (39.7-52)	45.9 (37.8-50.5)	0.250	
<b>HIV-antibody-positive</b>	16.7% (11)	14.1% (25)	0.620	1.2 (0.6-2.6)
<b>Alcohol ≥49 g/d</b>	28.8% (19)	11.3% (20)	0.001	3.2 (1.6-6.4)
<b>Obese (BMI&gt;30kg/m<sup>2</sup>)</b>	24.2% (16)	18.6% (33)	0.333	1.4 (0.7-2.8)
<b>Schizophrenia</b>	10.6% (7)	24.9% (44)	0.015	0.4 (0.2-0.8)
<b>Affective disorder</b>	51.5% (34)	45.8% (81)	0.424	1.3 (0.7-2.2)
<b>Median time (y) since HCV diagnosis (IQR)</b>	12.5 (5-19) (n=66)	11 (4-18) (n=172)	0.472	
<b>Median time (y) since first IDU</b>	28 (16-34) (n=54)	25 (15-30) (n=156)	0.357	

### RESULTS – Late presentation at first HCV diagnosis

- There were **78 HCV first diagnoses since 2017**: 69.2% (54) male; median age at HCV diagnosis: 39 years (IQR: 33-47); median time since first IDU: 16 years (IQR: 5-26; n=63); 88.5% diagnosed at/before enrolment; for 70.5%, no pretest recorded; 37.9% (22/58) diagnosed with a rapid test.
- 27.4% (20/73) cleared spontaneously, while 72.6% (53) developed chronic infection (5 unclear/unknown).
- Treatment-uptake was 86.8% (46/53), mostly within one year.
- Of the 88.7% (47/53) **chronically infected individuals with Fibroscan® available, 23.4% (11/47) already had F3/F4-fibrosis.**

#### Liver fibrosis stage (Fibroscan®) at first HCV diagnosis (2017 onwards)



For 47 chronically infected individuals with first HCV diagnosis since 2017, Fibroscan® was available.

### CONCLUSIONS

- In times of unrestricted DAA access, late diagnosis and late treatment of chronic hepatitis C are still common (one in four), suggesting insufficient care.
- But, once diagnosed or relinked to care, patients are promptly treated.

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