



POPPY

Pharmacokinetic and clinical
observations in people over 50

Health and wellbeing outcomes in PLWH with and without depression

Daide De Francesco, Caroline A. Sabin and Alan Winston
on behalf of the POPPY study group

Department of Infection & Population Health, UCL, London, UK



UK-CAB Meeting

14th July 2017

Background

- Depression is the most common psychiatric condition in people living with HIV (PLWH)
- Depression is consistently reported with high but varying prevalences among PLWH (as high as 80% but 40-42% on average)¹
- Reported prevalence in PLWH can be up to three-times higher than in the general population²

¹Ciesla, JA et al. 2001 ²Kessler, RC et al. 2008

Why is depression more prevalent in PLWH?

- Evidence support the hypothesis that HIV infection and depression have a bidirectional association
- Link is likely to be a complex mix of:

Psychosocial factors

Burden of having a life-threatening, chronic, difficult to manage disease

Stigma

Social isolation and discrimination

Occurrence of other comorbidities, complications and ART side effects

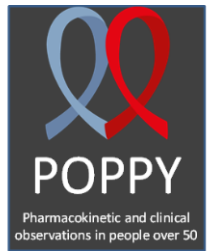
Need for a strict adherence to ART

Biological factors

Neurological changes caused by the penetration of the virus into the CNS

- Increase concentrations of inflammatory markers linked to depression

How does depression impact lives of PLWH?



- Depression has been found to negatively affect:
 - ART treatment prescription and adherence
 - CD4⁺ T cell count and HIV viral load
 - risk behaviours (substance/alcohol abuse)
- This may ultimately lead to disease progression and poorer health

Aims

- To compare prevalence of diagnosed depression in PLWH and matched HIV-negative controls
- To explore lifestyle factors associated with the occurrence of depression in PLWH
- To investigate the association between depression, quality of life and general health outcomes

The POPPY Study

- Multicentre, prospective, observational study to examine the effects of ageing on the clinical outcomes of people living with HIV in UK and Ireland
- Cohorts of HIV-positive people aged ≥ 50 years and demographically matched < 50 HIV-positive and ≥ 50 HIV-negative individuals

≥ 50 years PLWH	< 50 years PLWH	≥ 50 years HIV-
N=698	N=374	N=304
White/black African ethnicity	Frequency matched on gender, ethnicity, sexual orientation and location (in/out London)	Frequency matched on age, gender, ethnicity, sexual orientation and location (in/out London)
Acquired HIV via sexual routes		

Methods

- Participants underwent a structured interview with trained staff to report:
 - medically diagnosed depression (treated by a doctor)
 - occurrence of comorbidities
 - concomitant medication (other than ART)
 - healthcare utilisation in the previous year (hospital admissions)
- Quality of life was assessed by the SF-36 questionnaire to measure physical health

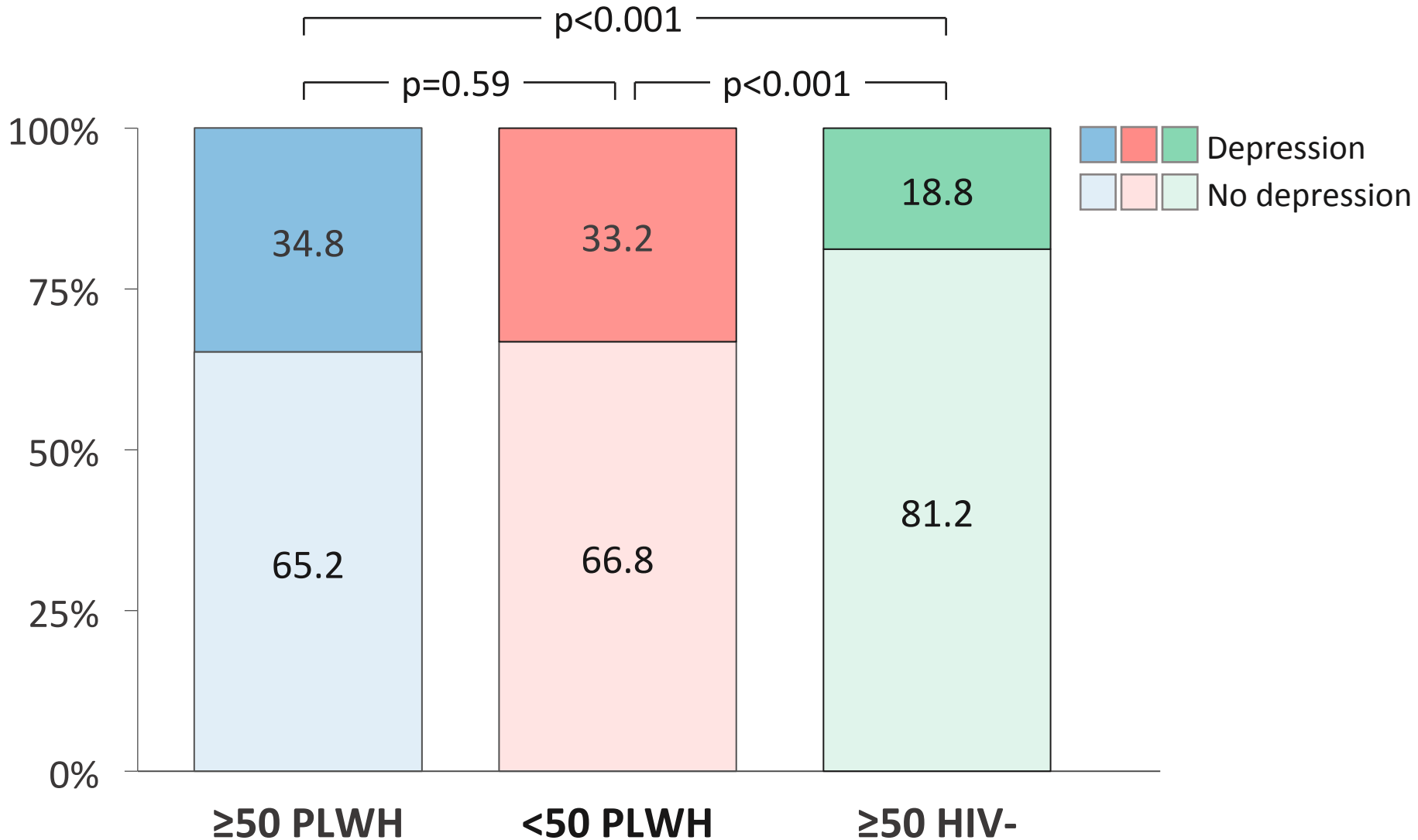
Statistical methods

- Group differences in the prevalence of depression were evaluated using the chi-squared test
- Associations between depression and lifestyle in PLWH were assessed using logistic regression
- Logistic, ordinal or linear regression to evaluate associations between health outcomes and both study group and depression

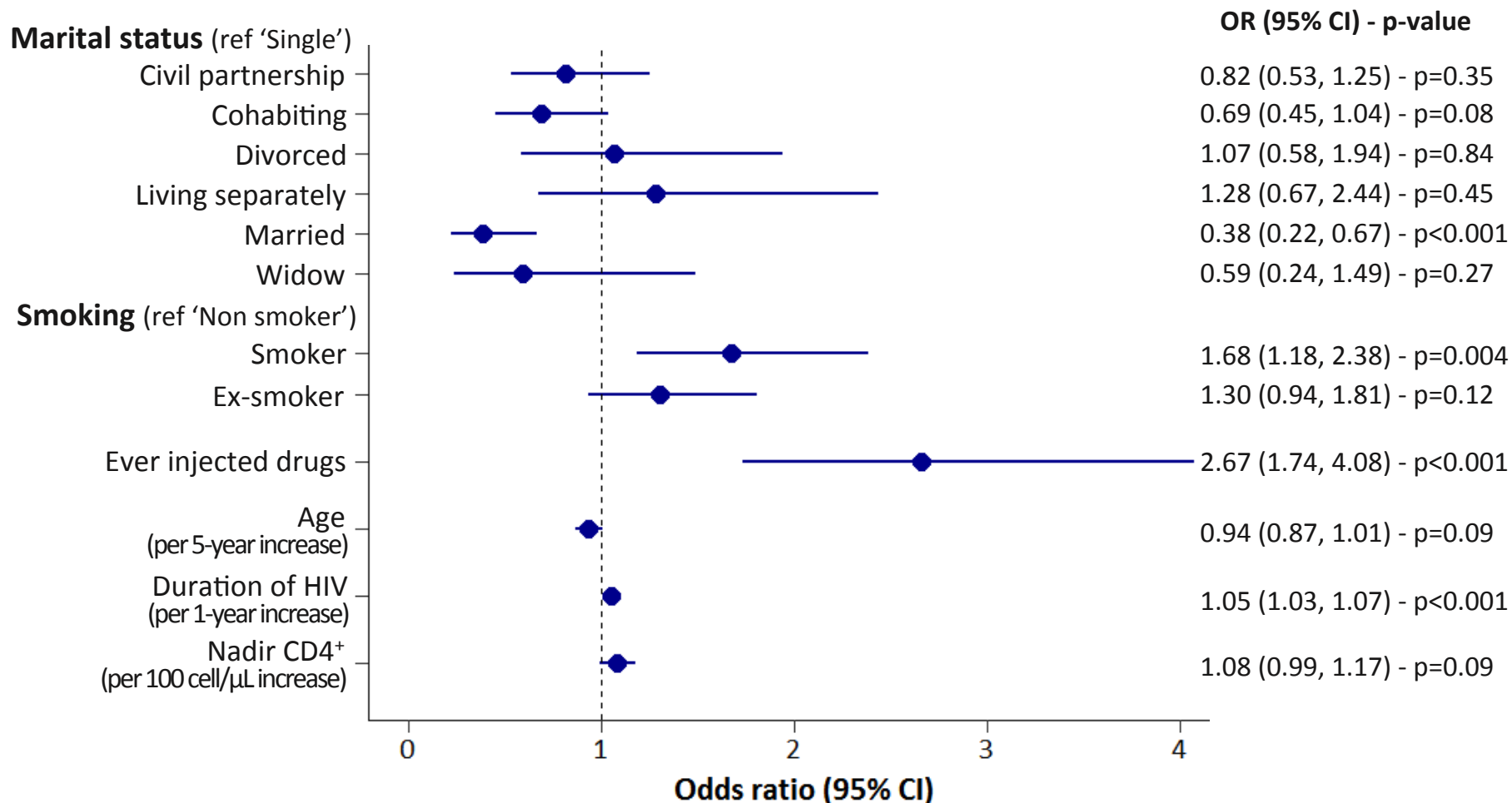
Baseline characteristics

		≥50 PLWH (N=698)	<50 PLWH (N=374)	≥50 HIV- (N=304)
Gender, n (%)	Female	86 (12.3%)	72 (19.3%)	109 (35.9%)
	Male	612 (87.7%)	302 (80.8%)	195 (64.1%)
Ethnicity, n (%)	Black-African	95 (13.6%)	75 (20.1%)	31 (10.2%)
	White	603 (86.4%)	299 (80.0%)	273 (89.8%)
Age [years], median (IQR)		57 (53, 62)	43 (37, 47)	58 (53, 63)
Sexual orientation	MSM	550 (78.8%)	269 (71.9%)	144 (47.4%)
	Heterosexual	148 (21.2%)	105 (28.1%)	160 (52.6%)
Viral load <50 copies/mL, n (%)		641 (92.1%)	323 (86.8%)	N/A
CD4⁺ count [cells/μL], median (IQR)		610 (466, 792)	661 (499, 847)	N/A
Nadir CD4⁺ count [cells/μL], median (IQR)		155 (85, 273)	178 (151, 376)	N/A

Depression

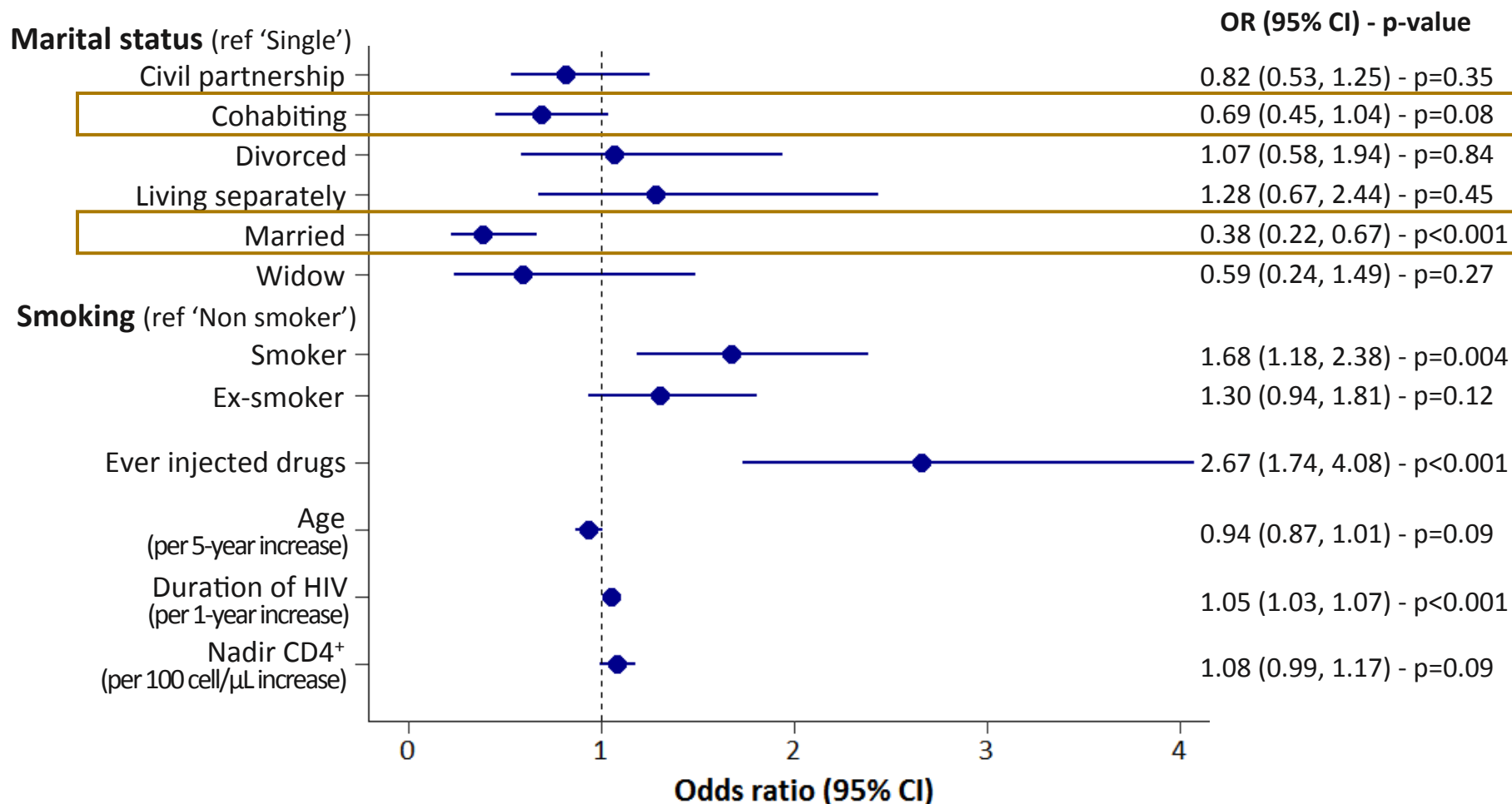


Who is more likely to report depression?



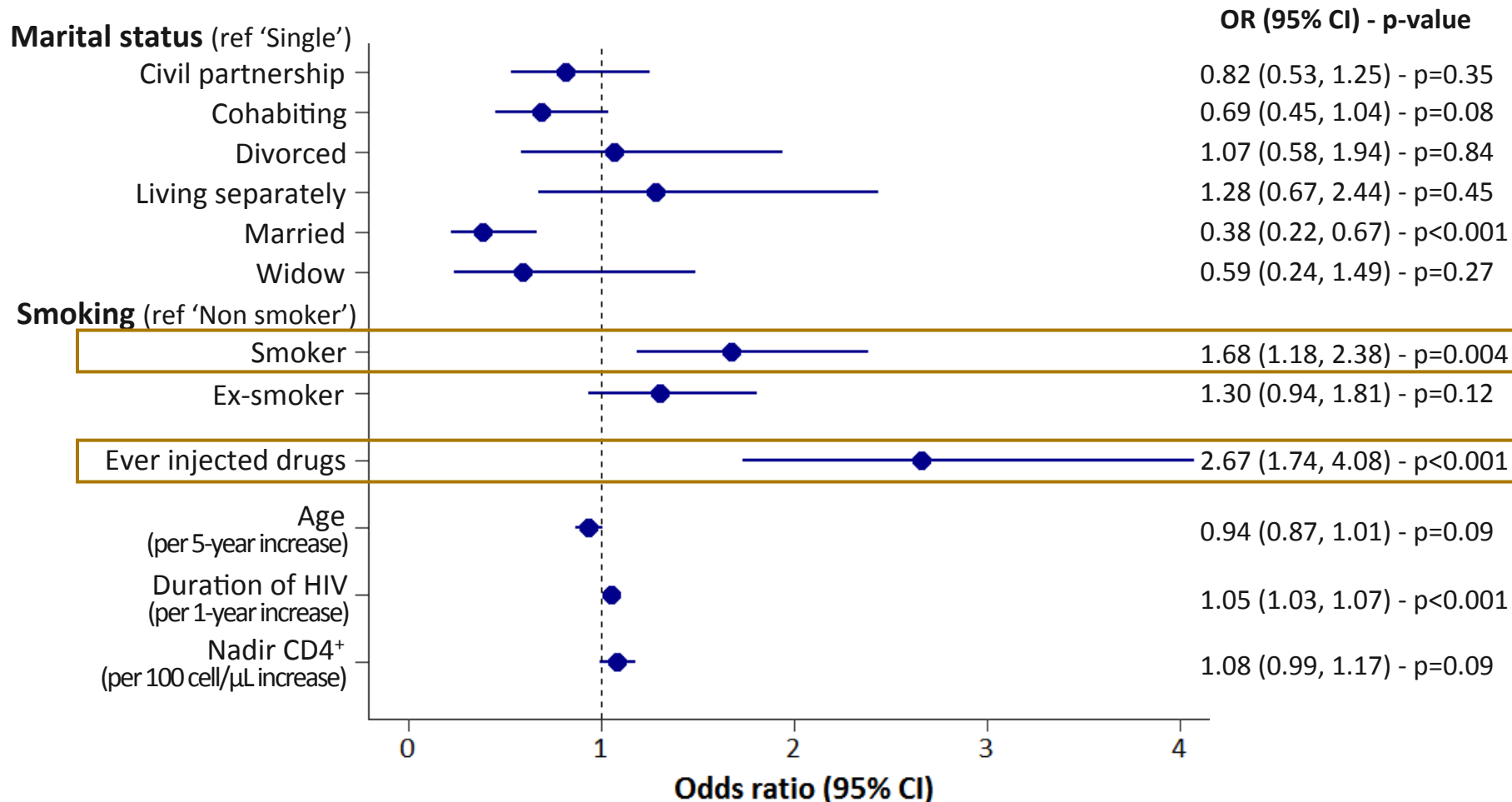
Gender, sexuality, race, alcohol consumption, rec. drugs use and VL<50 copies/mL were not significantly associated

Who is more likely to report depression?



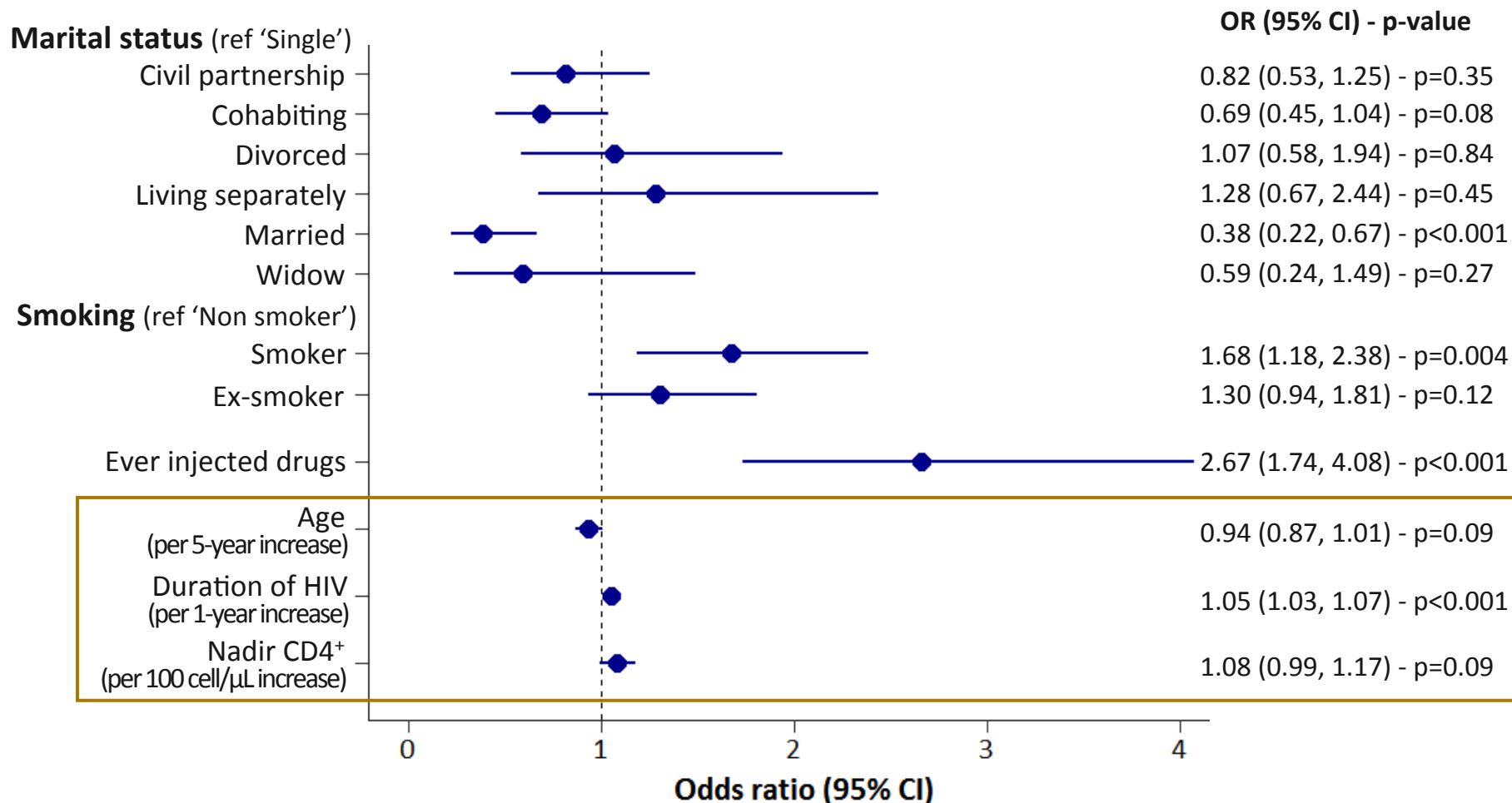
Gender, sexuality, race, alcohol consumption, rec. drugs use and VL<50 copies/mL were not significantly associated

Who is more likely to report depression?



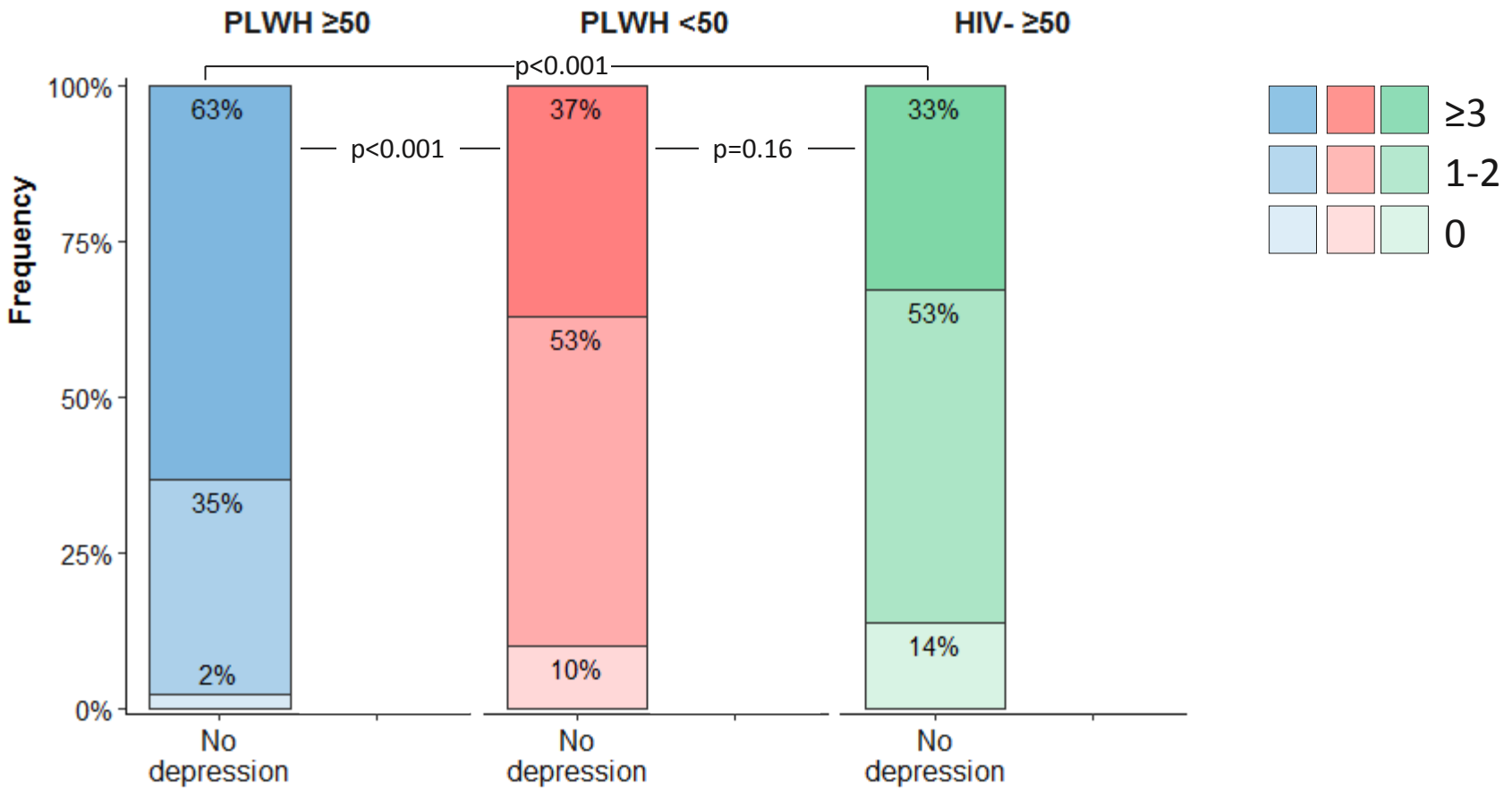
Gender, sexuality, race, alcohol consumption, rec. drugs use and VL<50 copies/mL were not significantly associated

Who is more likely to report depression?



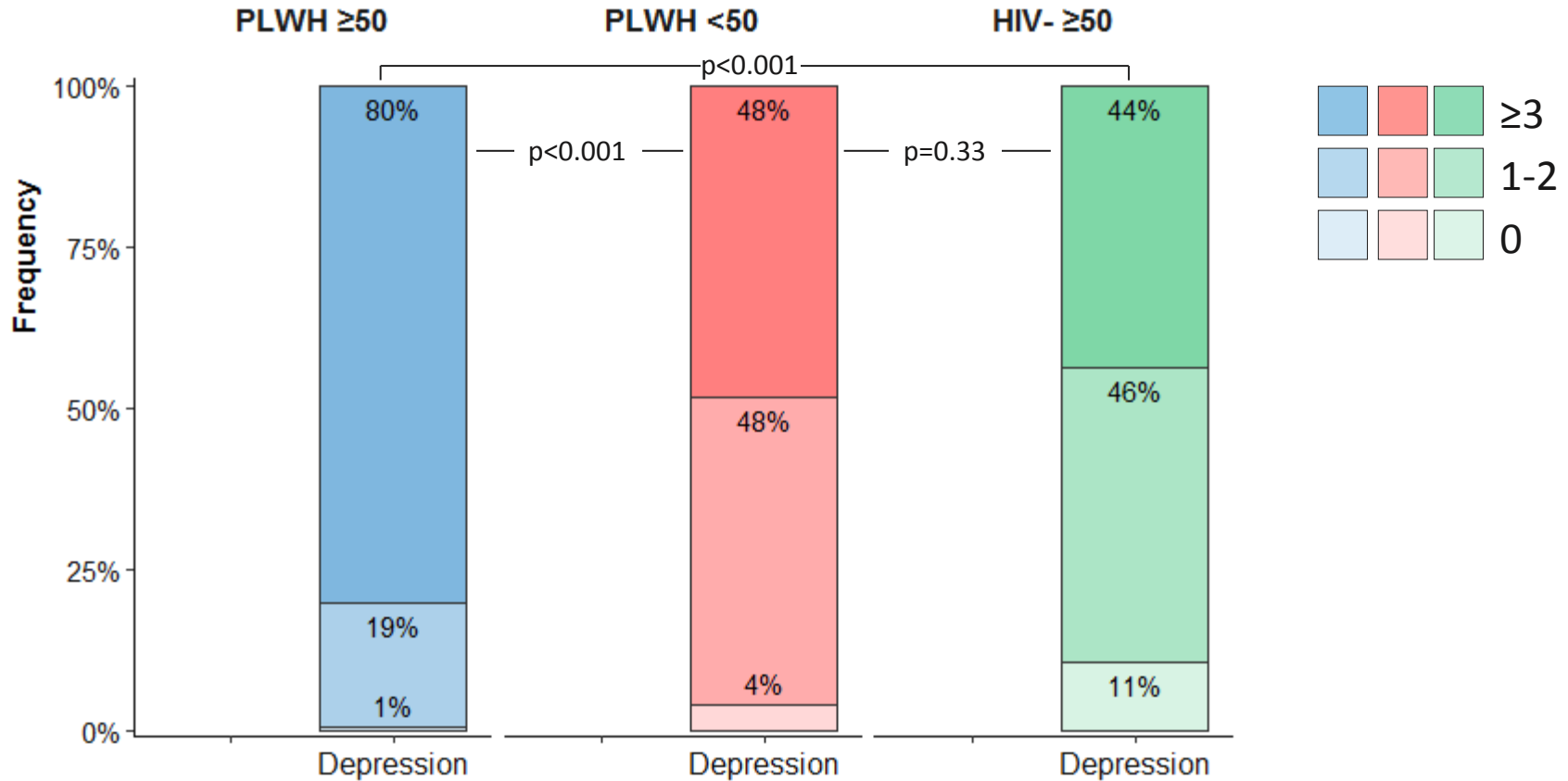
Gender, sexuality, race, alcohol consumption, rec. drugs use and VL<50 copies/mL were not significantly associated

Number of comorbidities*



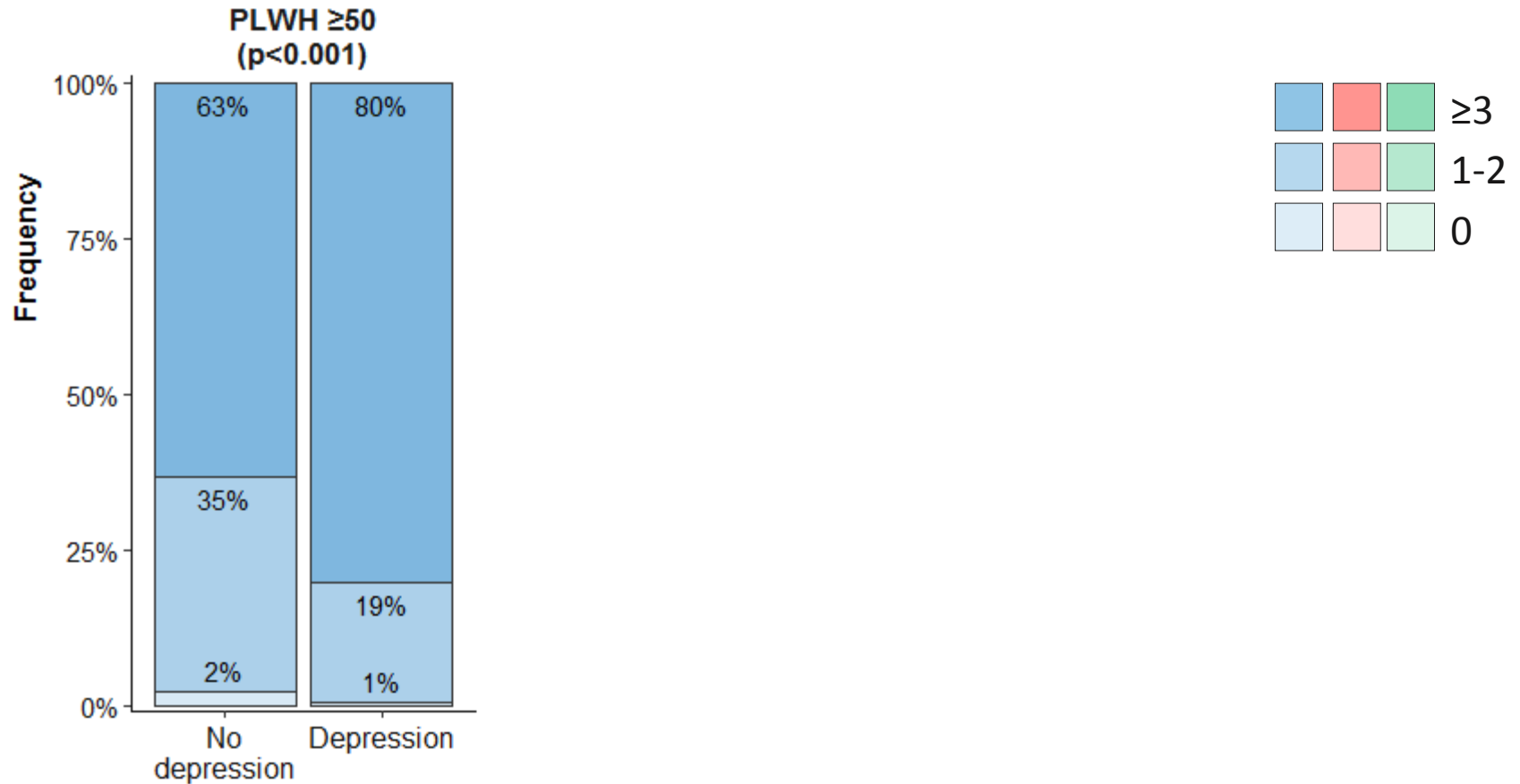
*between diabetes mellitus, CVDs, chest problems, Hepatitis B and C, AIDS events neurological problems, genitourinary problems, cancer, STDs, joint problems

Number of comorbidities*



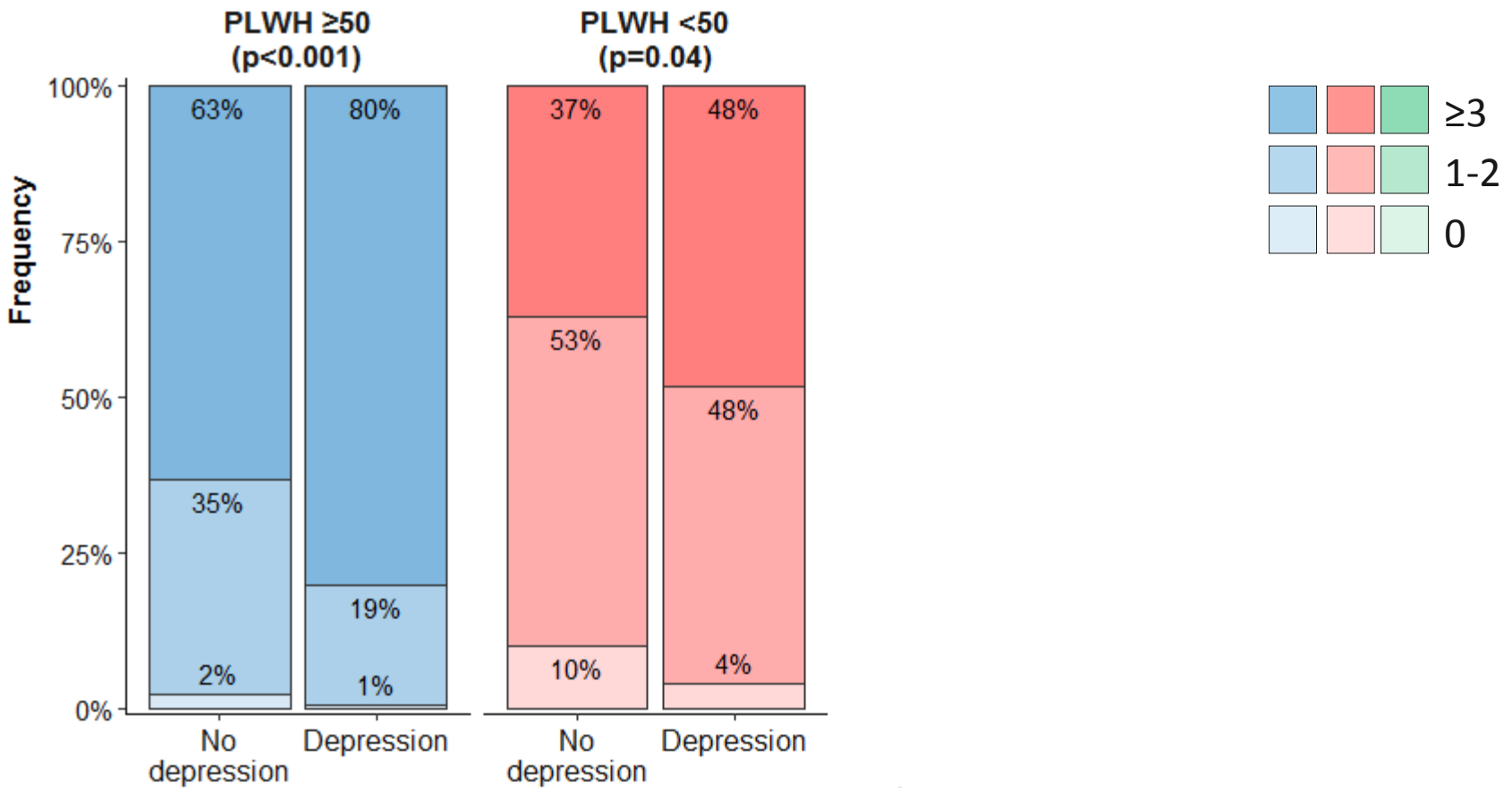
*between diabetes mellitus, CVDs, chest problems, Hepatitis B and C, AIDS events neurological problems, genitourinary problems, cancer, STDs, joint problems

Number of comorbidities*



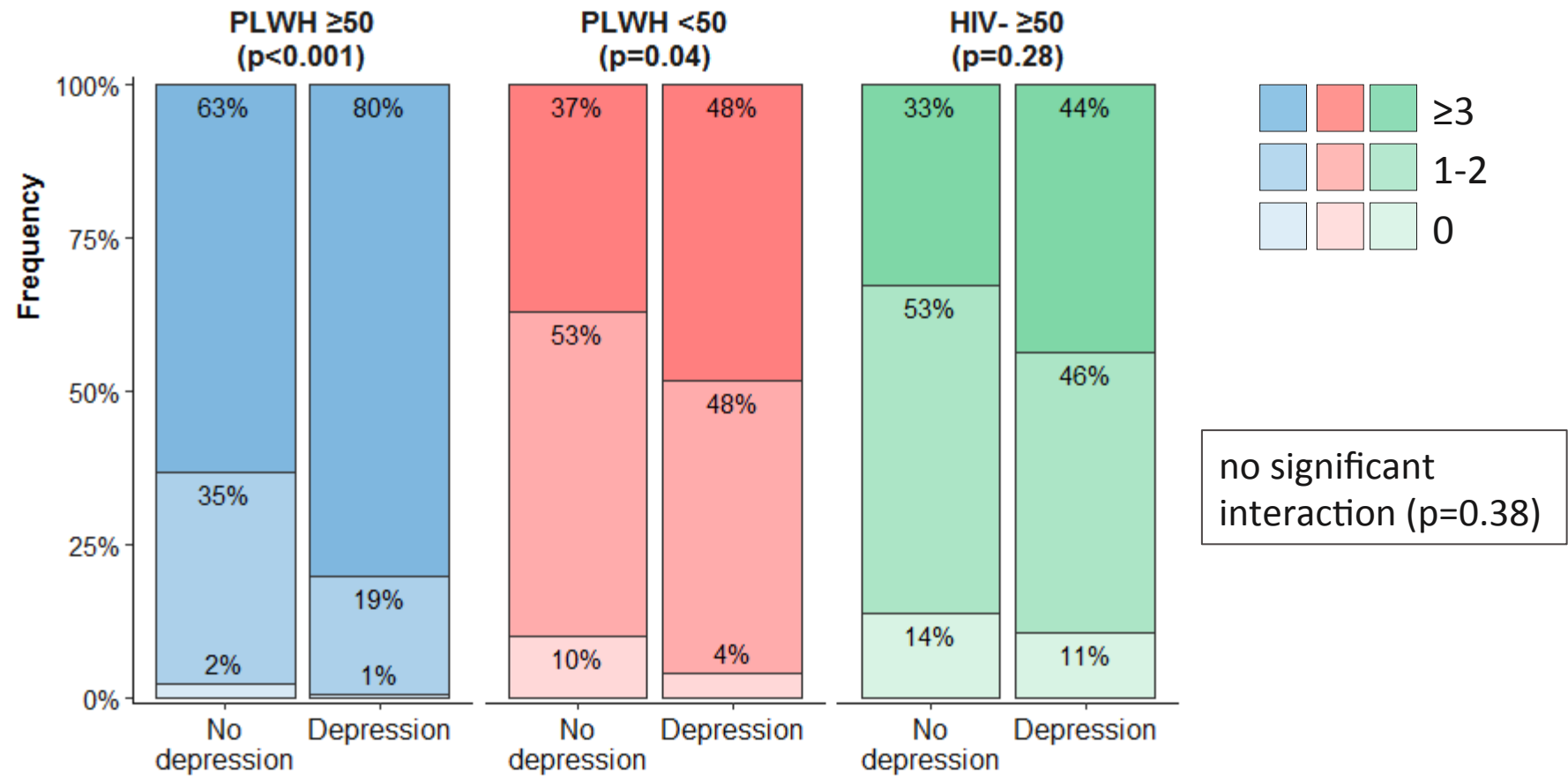
*between diabetes mellitus, CVDs, chest problems, Hepatitis B and C, AIDS events neurological problems, genitourinary problems, cancer, STDs, joint problems

Number of comorbidities*



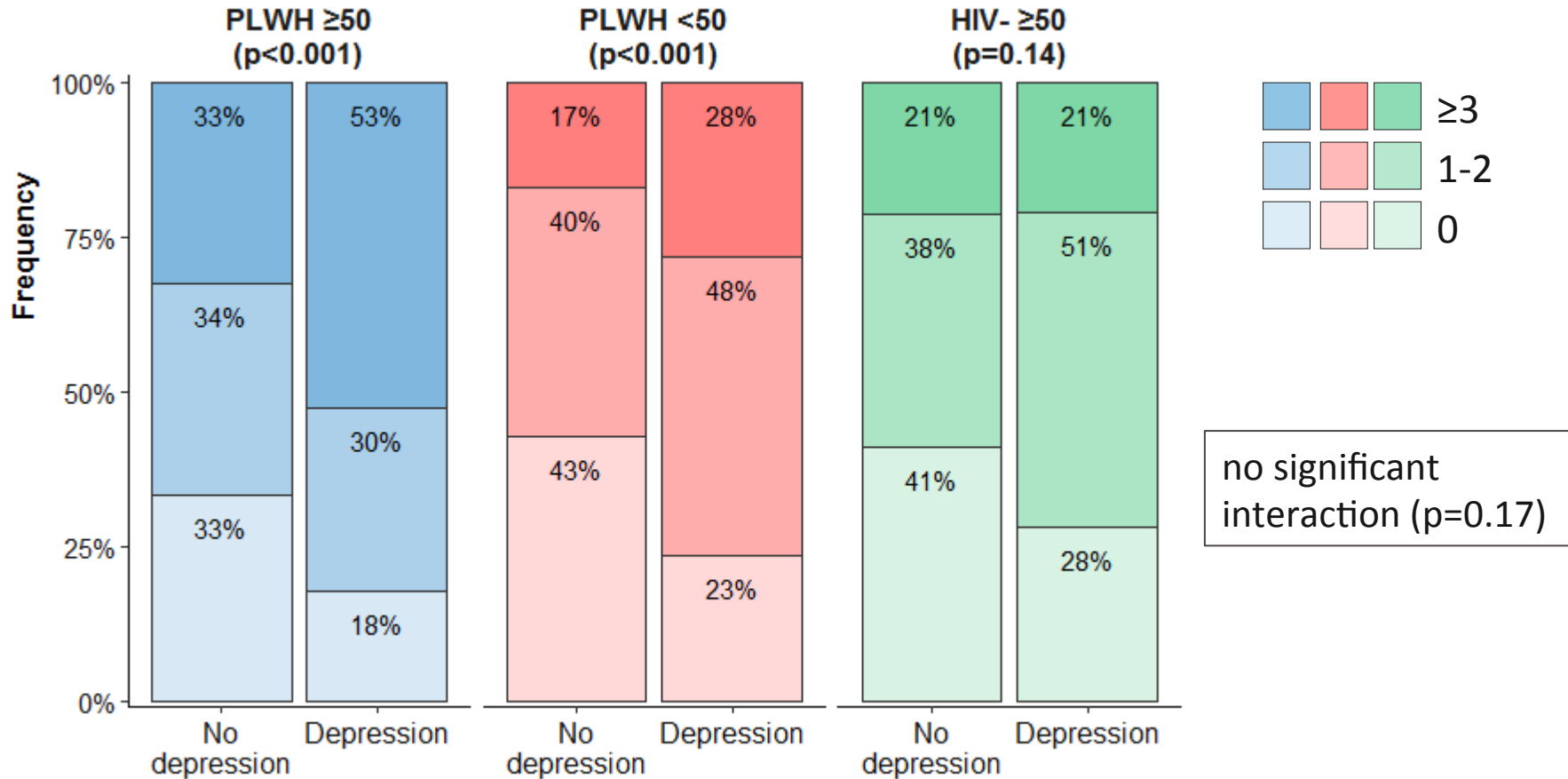
*between diabetes mellitus, CVDs, chest problems, Hepatitis B and C, AIDS events neurological problems, genitourinary problems, cancer, STDs, joint problems

Number of comorbidities*



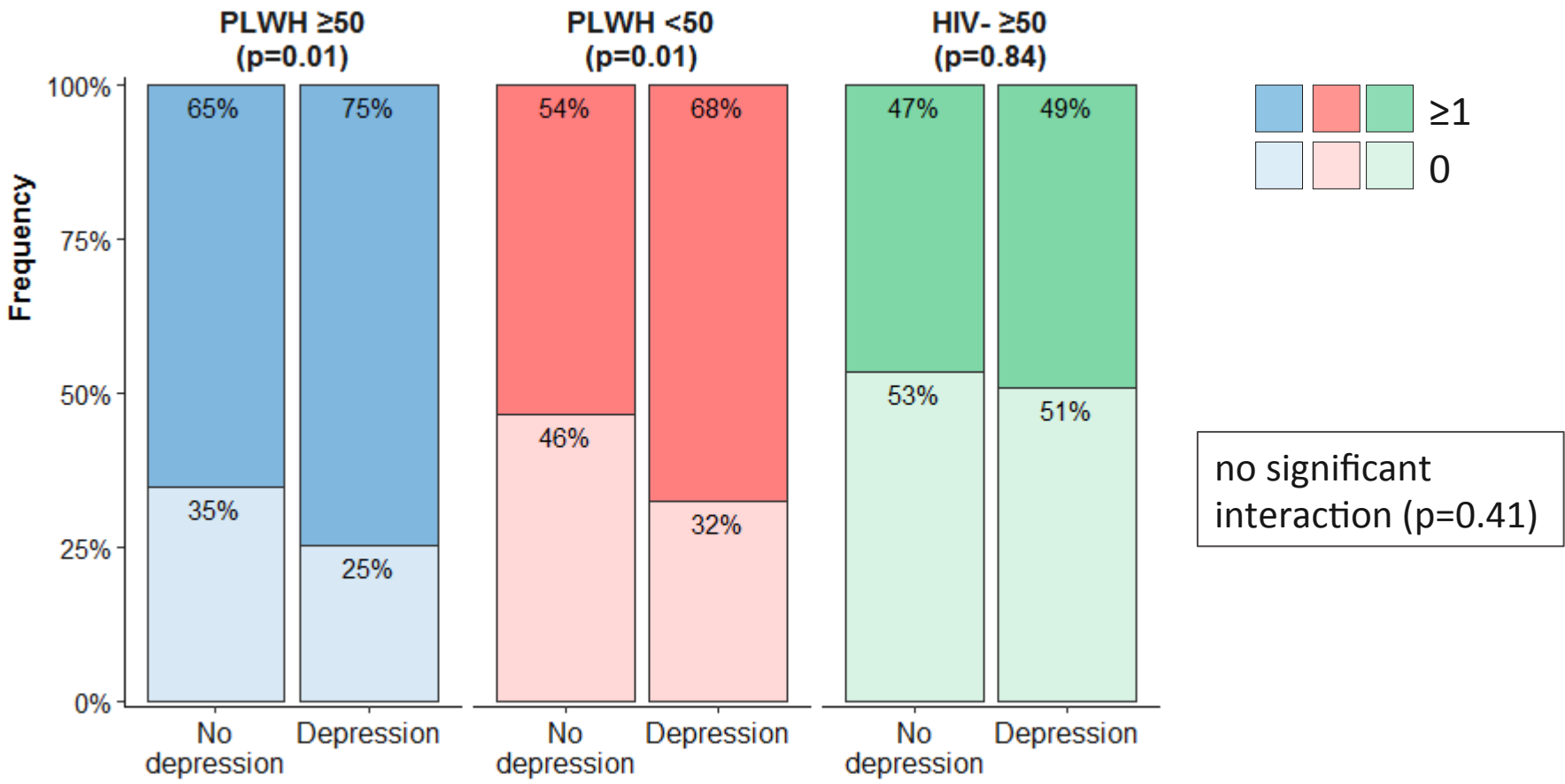
*between diabetes mellitus, CVDs, chest problems, Hepatitis B and C, AIDS events neurological problems, genitourinary problems, cancer, STDs, joint problems

Concomitant medications*



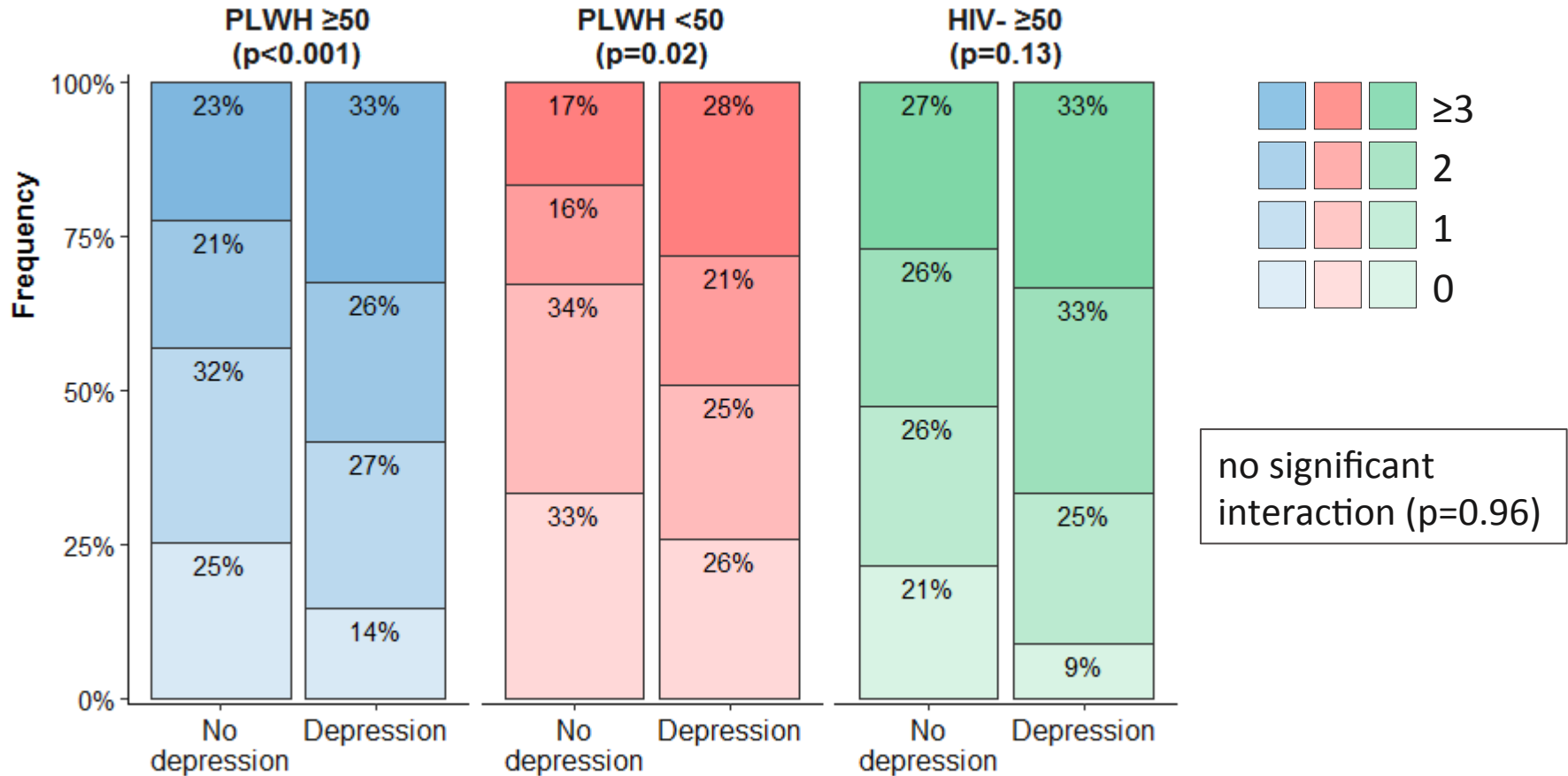
*including analgesic, antihistamines, antimicrobial, antiplatelet, antiviral, chest, dermatology, endocrine, gastrointestinal, hypertension, lipids lowering, mental health, sleeping pills, neurological, anti-rheumatic/steroids, supplements, urogenital/STD

Hospital admission*



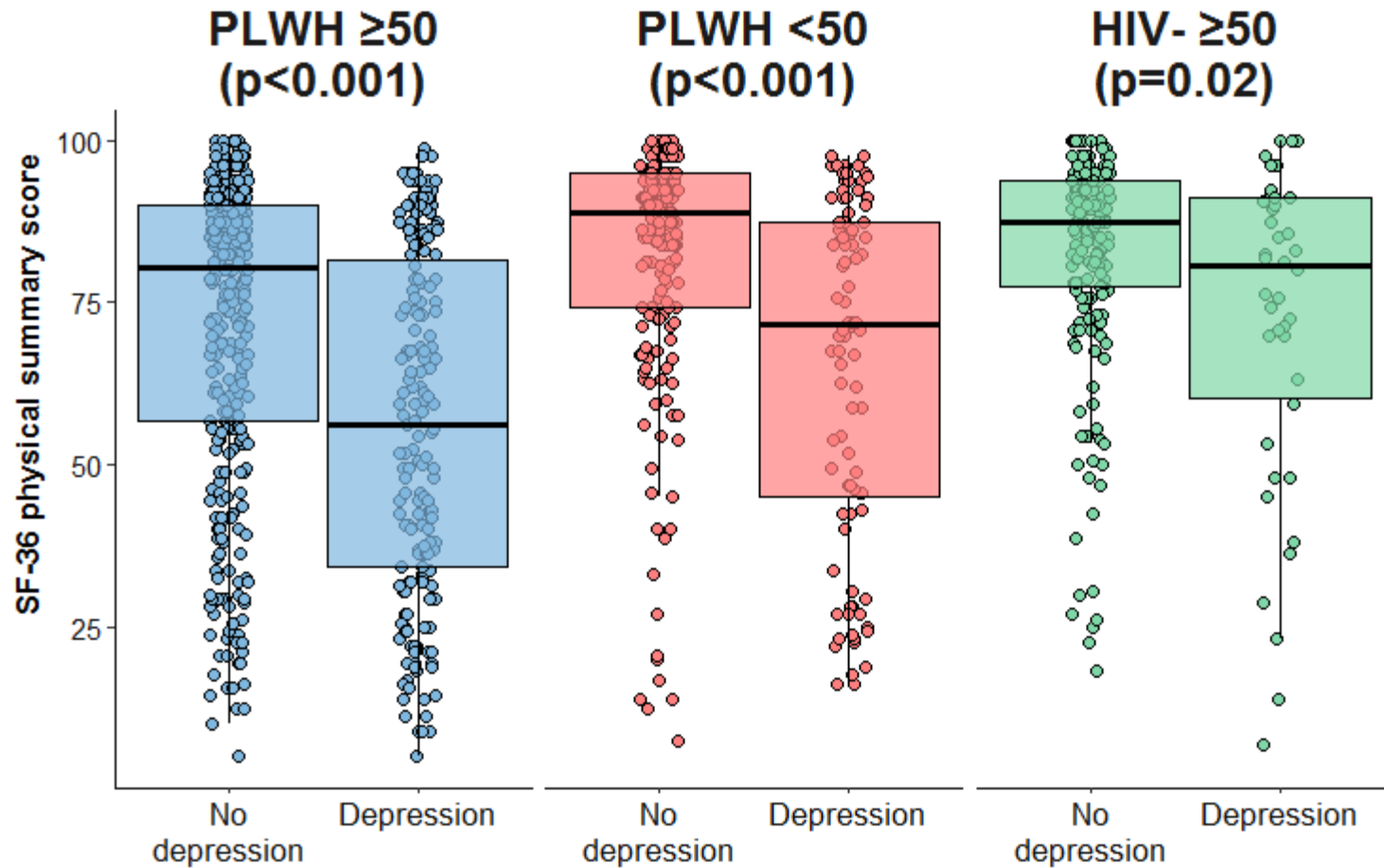
*including A&E visit, hospital investigation and hospital procedure

GP visits (non HIV-related)



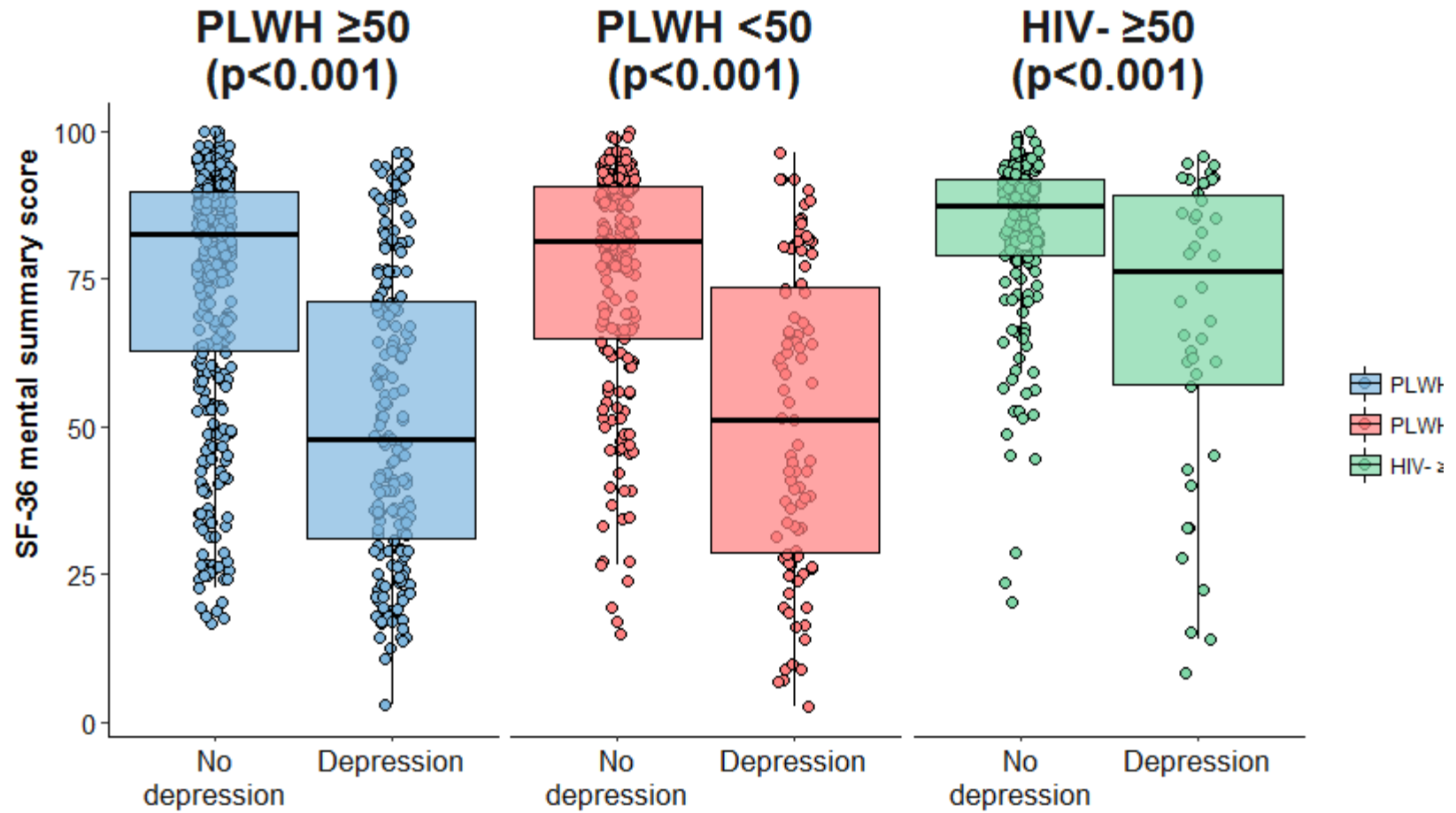
Physical health

no significant
interaction ($p=0.36$)



Mental health

no significant
interaction ($p=0.12$)



Conclusions

- PLWH are more likely to have medically-diagnosed depression even when compared to HIV-negative people with similar lifestyles
- Depression and HIV are each independently associated with several health outcomes
- Given the cross-sectional nature of the analysis, we are unable to demonstrate causality or the direction of the associations



Acknowledgments

POPPY Management Team: Marta Boffito, Paddy Mallon, Frank Post, Caroline Sabin, Memory Sachikonye, Alan Winston

POPPY Scientific Steering Committee: Jane Anderson, David Asboe, Marta Boffito, Lucy Garvey, Paddy Mallon, Frank Post, Anton Pozniak, Caroline Sabin, Memory Sachikonye, Jaime Vera, Ian Williams, Alan Winston

POPPY Sites and Trials Unit:

- Elton John Centre, Brighton and Sussex University Hospital (Martin Fisher, Amanda Clarke, Jaime Vera, Andrew Bexley, Celia Richardson)
- St Stephen's Centre, Chelsea and Westminster Hospital (Marta Boffito, David Asboe, Anton Pozniak, Chris Higgs, Elisha Seah, Stephen Fletcher, Michelle Anthonipillai, Ashley Moyes, Katie Deats, Irtiza Syed, Clive Matthews, Peter Fernando)
- Homerton Sexual Health Services, Homerton University Hospital (Jane Anderson, Sifiso Mguni, Rebecca Clark, Rhiannon Nevin-Dolan, Sambasivarao Pelluri)
- Caldecot Centre, King's College Hospital (Frank Post, Lucy Campbell, Selin Yurdakul, Sara Okumu, Louise Pollard)
- HIV Molecular Research Group, School of Medicine, University College Dublin (Paddy Mallon, Alan Macken, Bijan Ghavani-Kia, Joanne Maher, Maria Byrne, Ailbhe Flaherty, Sumesh Babu)
- Department of Infection and Population Health, University College London (Ian Williams, Damilola Otiko, Laura Phillips, Rosanna Laverick, Michelle Beynon, Anna-Lena Salz)
- St. Mary's Hospital London, Imperial College Healthcare NHS Trust (Alan Winston, Lucy Garvey, Jonathan Underwood, Matthew Stott, Linda McDonald)
- Imperial Clinical Trials Unit, Imperial College London (Andrew Whitehouse, Laura Burgess, Daphne Babalis)
- Ian Charleson Day Centre, Royal Free Hospital (Margaret Johnson, Nnenna Ngwu, Nargis Hemat, Martin Jones, Anne Carroll)

POPPY methodology/statistics/analysis: Caroline Sabin, Davide De Francesco, Emmanouil Bagkeris

Funders: The POPPY study is funded from investigator initiated grants from BMS, Gilead Sciences, Janssen, MSD and ViiV Healthcare.