



# The Longitudinal Effects of Non-injection Substance Use on Sustained HIV Viral Load Undetectability Among MSM and Heterosexual Men in Brazil and Thailand: The Role of ART Adherence and Depressive Symptoms (HPTN 063)

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## Abstract

The effect of non-injection substance use on HIV viral load (VL) is understudied in international settings. Data are from HPTN063, a longitudinal observational study of HIV-infected individuals in Brazil, Thailand, and Zambia, with focus on men with VL data (Brazil = 146; Thailand = 159). Generalized linear mixed models (GLMM) assessed whether non-injection substance use (stimulants, cannabis, alcohol, polysubstance) was associated with VL undetectability. ART adherence and depressive symptoms were examined as mediators of the association. *In Thailand*, substance use was not significantly associated with VL undetectability or ART adherence, but alcohol misuse among MSM was associated with increased odds of depression (AOR = 2.75; 95% CI 1.20, 6.32,  $p=0.02$ ). *In Brazil*, alcohol misuse by MSM was associated with decreased odds of undetectable VL (AOR = 0.34; 95% CI 0.13, 0.92,  $p=0.03$ ). Polysubstance use by heterosexual men in Brazil was associated with decreased odds of ART adherence (AOR = 0.25; 95% CI 0.08, 0.78,  $p=0.02$ ). VL suppression appears attainable among non-injection substance users. Substance use interventions among HIV-positive men should address depression, adherence, and VL undetectability.

**Keywords** Substance use · HIV · Depression · Adherence · Undetectable viral load

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## Introduction

Over time, improvements in antiretroviral therapy (ART) have lengthened lifespan and reduced HIV transmission among people living with HIV (PLH) [1–3]. Findings that ART adherence can suppress viral load (VL) and reduce HIV transmissibility during condomless sex have led to prioritizing treatment as prevention (TasP) as a key strategy to prevent HIV transmission by PLH [4]. However, high rates of substance use and depression among PLH remain key barriers to successful implementation of TasP in the U.S. and other similar settings [1]. However, empirical studies on the associations of substance use, depression, and achieving undetectable VL have not been adequately assessed in low- and middle-income settings.

Non-injection substance use is the most common form of substance use among PLH, with 40–70% reporting the use of alcohol, cannabis, non-injection stimulants (e.g., cocaine, amphetamines), and/or opioids [5, 6]. In general, PLH who use substances are less likely to access ART [7–10], are found to have lower ART adherence [11–13], are less likely to achieve viral suppression [8, 9, 11, 14], and are more likely to have faster disease progression [15–18] compared to non-substance using PLH. Moreover, this population may be the most likely to engage in condomless sex [19, 20], making it critical to understand how to improve their HIV care outcomes. Aside from behavioral risk, and the reduced ART adherence associated with substance use, emerging research indicates that substance use may have pathophysiological effects on HIV disease progression [21]. For example, stimulants have been linked to increased HIV replication—in peripheral blood mononuclear cells [22–24] and in mouse models [25].

When examining the effects of substance use on VL or other HIV outcomes, it is also important to investigate the contribution of depression as it is a highly prevalent comorbid condition. Depression is a more common comorbidity to substance use among PLH than the general population [26], and is the most common psychiatric health condition among PLH—affecting 20–33% of adults in HIV care [27, 28]. In terms of HIV clinical outcomes, depression is thought to lower ART adherence [29] and reduce the likelihood of sustained viral suppression [30]. Studies indicate that depressive symptoms may also affect HIV disease progression above and beyond sub-optimal ART adherence by reducing individuals' responsiveness to ART, decreasing CD4+ count, and increasing HIV VL [16, 31]. Depressive symptoms and substance use are prevalent among PLH and likely contribute substantially to the lack of sustained viral suppression. Despite the high prevalence of substance use and depressive symptoms among PLH,

most research examining depression, substance use, and HIV disease outcomes has been conducted in the U.S. [5].

There is little information on the type and patterns of non-injection substance use, on the prevalence of depression, and on how these common comorbidities affect viral suppression among PLH in low- and middle-income settings. There is reason to think that the association between substance use and viral load detectability may operate through decreased ART adherence and increased co-morbidity with depression among PLH. Previous research have linked substance use—including alcohol, cocaine, heroin, methamphetamines, and other stimulants—to decreased ART adherence, although these studies took place in the U.S. [32, 33]. A recent systematic review focused on ART adherence among those who engaged in substance use in low- and middle-income countries found sub-optimal adherence to treatment [34], however this review solely focused on injection drug use. In addition, a study that examined active drug use on ART adherence and viral suppression found that depression appeared to mediate the association, although the finding was only significant for HIV-infected women and not HIV-infected men [11]. Moreover, based on the *minority stress theory*—which posits that sexual minorities have adverse health outcomes as a result of heightened stress from prejudice and stigma based on their sexual minority status—it is thought that men who have sex with men (MSM) may have greater substance use and depressive symptoms than heterosexual men [35]. This greater comorbidity prevalence is hypothesized to magnify the association between substance use, depression, and viral load detectability. This is likely the case for men in low- and middle-income settings, such as Thailand and Brazil, where HIV prevalence is much greater among MSM compared to the general adult population at 9.2% (vs. 1.1% adult) and 10.5% (vs. 0.6% adult), respectively [36, 37]. Although less research has been conducted among men who identify as heterosexual in international contexts, in Brazil they comprise the largest proportion of men infected with HIV [38] and as many as 70% receive late HIV-diagnosis [39]. Furthermore, non-injection substance use often affects MSM and heterosexual men at greater rates than women [40, 41], potentially exacerbating the effects of substance use on HIV outcomes via ART adherence and depression in low- and middle-income settings.

This study aims to address this gap in research by conducting a secondary data analysis focused on MSM and heterosexual men using HPTN 063 data, a longitudinal observational study of HIV-positive individuals in HIV care in Zambia, Thailand, and Brazil [42]. First, we described the type and pattern of non-injection substance use and prevalence of depressive symptoms among men infected with HIV at baseline. Second, we examined the effect of non-injection substance use on ART adherence and HIV VL undetectability, testing ART adherence as a mediator of the association

between substance use and HIV VL undetectability. Third, we examined the effect of non-injection substance use on depressive symptoms and VL undetectability, testing depressive symptoms as a mediator of the association between substance use and HIV VL undetectability. Then, we tested whether there was evidence of effect modification due to sexual orientation, on the association between substance use, mediators (e.g., ART adherence, depressive symptoms), and HIV outcomes. For all analyses, we stratified by unique country context.

## Methods

Data were collected via HPTN 063, a multi-site, longitudinal observational cohort study of people living with HIV at high risk for sexual transmission in HIV care in Africa (Lusaka, Zambia), Asia (Chiang Mai, Thailand), and South America (Rio de Janeiro, Brazil). Recruited participants included HIV-infected heterosexual men, heterosexual women, and men who have sex with men (MSM). Structured interviews were conducted every 3 months over the course of 12 months, collecting data on socio-demographics, behavioral risk, substance use, mental health, and ARV adherence. HIV clinical variables (e.g., plasma RNA [VL], CD4+ count) were extracted from patient files. All procedures were approved at each site (Thailand—Chiang Mai University; and the Johns Hopkins Bloomberg School of Public Health, Brazil—the Evandro Chagas Clinical Research Institute; and the National Committee for Ethics in Research) and each participant provided written informed consent. The HPTN063 study design has been described in detail in previous publications [42, 43].

## Sample

This study reports findings using the data collected from HPTN 063 focused on heterosexual men and MSM in Thailand ( $n = 159$ ) and Brazil (Brazil = 146) as VL data was not available in Zambia. Men were considered MSM regardless if they reported having sex with women as well. In order to have sufficient observed data to characterize patterns of ART adherence and VL, we included participants who completed at least two of five assessments with information on substance use, depressive symptoms, ART adherence, and VL detectability. There were an average of 2.7 VL observations per individual.

## Measures

*Plasma HIV-RNA VL* was extracted from medical records at baseline and each follow-up visit and recorded if a

current VL was documented. VL was then dichotomized (0: VL  $\geq 200$  copies/ml and 1: VL  $< 200$  copies/ml).

*Non-injection substance use* was measured as the number of self-reported use days and included stimulants, cannabis, and alcohol. *Stimulant use* was measured as the number of days that non-injection cocaine (powder and crack), methamphetamine, and ecstasy use were reported in the prior 3 months. *Cannabis* was measured as the number of days that marijuana and hashish were reported in the prior three months. *Alcohol misuse* was measured using the 10-item alcohol use disorders identification test (AUDIT). Example items include how many drinks containing alcohol one has on a typical day and how often one is not able to stop drinking once started. AUDIT score was dichotomized into alcohol misuse (AUDIT score  $\geq 8$ ) versus no alcohol misuse (AUDIT score 0–7) [44].

*Polysubstance use* was measured as the total number of non-injection substances reported used in the past 3 months (yes/no), including stimulants, cannabis, and alcohol misuse, and was treated as a continuous variable (range 0–4).

*Depression symptoms* were measured using the Center for Epidemiologic Studies Depression Scale (CESD) [45]. Example items ask how often during the past week participants had a poor appetite or felt depressed. CESD score was dichotomized into severe depressive symptoms (CESD score  $\geq 16$ ) versus not severe depressive symptoms (CESD score  $< 16$ ).

*ART adherence* was measured using the self-reported question on adherence ability, “in the last 3 months, on average, how would you rate your ability to take all your antiretroviral drugs as your doctor prescribed?” [46]. Instructions provided prior to the interview normalized ART non-adherence. Participants were provided with a response card with Likert response options, ranging from very poor to excellent. This single-item, self-report adherence measure has been found as valid and reliable in prior research [47]. Due to small cell size, ART adherence ability in Thailand was recoded into two levels (very poor/poor/fair vs. good/very good/excellent). For Brazil, ART adherence ability was missing on too many participants ( $n = 43$ ; 30%) to warrant inclusion in this analysis and the dichotomized variable of taking ARTs was used in place. The self-reported measure asked, “In the last 3 months, have you taken antiretroviral drugs?”

*Socio-demographic variables* included in our analysis were age group (18–24, 25–44, and  $\geq 45$  years) and education (primary, secondary, and technical/college).

## Statistical Analysis

Data analysis began with descriptive statistics at baseline of the total sample and of heterosexual men versus men who have sex with men (MSM) on non-injection substance use,

depression, HIV outcomes, and socio-demographics. The Chi square statistic test was used for categorical variables, and t-statistic test for continuous variables, to detect statistically significant differences between groups (Table 1). Next, we described the type and number of self-reported non-injection substances used in the prior 3 months at baseline stratified by country and sub-group to understand poly-substance use in our sample (Fig. 1). Then, generalized linear mixed models (GLMMs) were applied with the logit link function for longitudinal binary outcomes to estimate the odds ratios of non-injection substance use on having an undetectable HIV VL adjusting for covariates, age and education (Tables 2, 3). The mediators, ART adherence and depression, were also estimated as an outcome of non-injection substance use using GLMM (Table 2 and 3, respectively) and mediation was controlled for when estimating the effects of non-injection substance use on undetectable HIV VL. GLMMs with the logistic link function with a random intercept and compound-symmetric covariance were used to account the correlations of observations between visits within individuals [48]. All analyses were stratified by country. For each model, an interaction term of substance use and sub-group (MSM and heterosexual men) was included to test for statistically significant differences between MSM and heterosexual men in the associations between substance use and ART adherence, depressive symptoms, and undetectable VL.

## Results

Table 1 shows the baseline characteristics of participants in the total sample stratified by study site and heterosexual men versus MSM. *In Thailand*, 43% of the total sample reported alcohol misuse. In the past 3 months, individuals, on average, reported using stimulants for zero days (range 0–30 days), cannabis one day (range 0–90 days), and used one non-injection substance (range 0–3 substances), with no significant difference by sub-group. Twenty-two percent of the total sample had severe depressive symptoms, with no significant difference by sub-group. In terms of HIV outcomes (adherence and VL), 82.4% reported good/very good/excellent adherence ability, with MSM reporting significantly better adherence ability than heterosexual men (89% vs. 77%,  $p$  value = 0.051). Seventy-seven percent of the total sample presented an undetectable VL at baseline, with no significant differences by sub-group. The median CD4+ count at baseline was significantly lower among heterosexual men compared to MSM (397.0 vs. 511.0,  $p$ -value = 0.002).

*In Brazil*, 34% of the total sample reported alcohol misuse. In the past 3 months, individuals, on average, reported using stimulants for 4 days (range 0–90 days), cannabis

for 5 days (range 0–90 days), and used one non-injection substance (range 0–4 substances), with no significant difference by sub-group. About half of the sample in Brazil had severe depressive symptoms, with no significant difference by sub-group. Seventy-one percent of the total sample reported taking ART in the past 3 months, with significantly more heterosexual men reporting taking ARTs than MSM (86% vs. 61%,  $p$ -value = 0.002). Only 53% of the total sample presented an undetectable VL at baseline, with significantly more heterosexual men presenting an undetectable VL than MSM (67% vs. 43%,  $p$ -value = 0.003). The median CD4 + count at baseline was also significantly lower among heterosexual men than MSM (462 vs. 648,  $p$ -value 0.004). In both sites, heterosexual men were significantly older and had lower education than MSM.

Figure 1 illustrates the type and pattern of non-injection substance use. *In Thailand*, of men who reported using non-injection substances in the past 3 months at baseline ( $n = 118$ ), 91% ( $n = 107$ ) used one drug and 9% ( $n = 11$ ) used two drugs simultaneously. *In Thailand*, among heterosexual men who reported using non-injection substances ( $n = 71$ ), 89% ( $n = 63$ ) reported using one substance (predominantly alcohol misuse) and 11% ( $n = 8$ ) reported using two substances (predominately alcohol misuse in combination with methamphetamines). *In Thailand*, among MSM who reported using non-injection substances ( $n = 47$ ), 94% ( $n = 44$ ) reported using one substance (predominantly alcohol misuse) and 6% ( $n = 3$ ) reported using two substances (predominately alcohol misuse in combination with methamphetamines).

*In Brazil*, of men who reported using non-injection substances ( $n = 112$ ), 67% ( $n = 75$ ) used one drug, 22% ( $n = 30$ ) used two drugs, 5% ( $n = 5$ ) used three drugs, and 2% ( $n = 2$ ) used four drugs. *In Brazil*, among heterosexual men who reported using non-injection substances ( $n = 41$ ), 63% ( $n = 26$ ) reported using one substance (predominantly alcohol misuse), 32% ( $n = 13$ ) reported using two substances (predominately alcohol misuse in combination with cocaine), and 5% ( $n = 2$ ) reported using three substances (both alcohol misuse in combination with cocaine and cannabis). *In Brazil*, among MSM who reported using non-injection substances ( $n = 71$ ), 69% ( $n = 49$ ) reported using one substance (all alcohol misuse), 24% ( $n = 17$ ) reported using two substances (predominately alcohol misuse in combination with either cocaine or cannabis), 4% ( $n = 3$ ) reported using three substances (alcohol misuse in combination with cannabis and either cocaine or ecstasy), and 6% ( $n = 4$ ) reported using four substances (alcohol misuse in combination with cannabis, cocaine, and ecstasy). For MSM in Brazil, the proportion of ecstasy users increased with the number of substances an individual reported to have taken in the past 3 months.

**Table 1** Baseline characteristics of HIV-infected men in Thailand and Brazil, HPTN 063

	Thailand				Brazil				p-value	MSM (n=88)	Heterosexual (n=58)	Total (n=146)				
	Heterosexual (n=93)		MSM (n=66)		Total (n=159)		MSM (n=88)						Heterosexual (n=58)		Total (n=146)	
	n	%	n	%	n	%	n	%					n	%	n	%
<b>Non-injection substance use<sup>a</sup></b>																
Alcohol misuse (AUDIT score 8)	44	47.3	25	37.9	0.237	69	43.4	19	32.8	31	35.2	0.671	50	34.2		
Stimulants (days used, past 3 mos.) (M, SD)	1.0	3.2	0.0	0.7	0.298	0.0	2.5	5.0	14.0	3.0	13.1	0.476	4.0	13.4		
Marijuana (days used, past 3 mos.) (M, SD)	2.0	11.2	0.0	0.9	0.245	1.0	8.6	2.0	12.5	6.0	20.1	0.209	5.0	17.5		
# of substances used (past 3 mos.) (M, SD)	1.0	0.6	1.0	0.5	0.293	1.0	0.5	1.0	0.8	1.0	0.8	0.291	1.0	0.8		
<b>Depression</b>																
Severe depressive symptoms (CESD 16)	19	20.4	17	25.8	0.429	36	22.6	27	46.6	43	48.9	0.685	70	47.9		
<b>HIV outcomes</b>																
<b>Adherence ability</b>																
Very poor, poor, or fair	21	22.6	7	10.6	0.051	28	17.6	6	10.3	9	10.2		15	10.3		
Good, very good, or excellent	72	77.4	59	89.4		131	82.4	44	75.9	44	50		88	60.3		
Took ARTs in past 3 mos.	93	100	66	100		159	100	50	86.2	54	61.4	0.002	104	71.2		
Detectable viral load	14	15.1	10	15.2	0.956	24	15.1	13	22.4	40	45.5	0.003	53	36.3		
Undetectable viral load	71	76.3	52	78.8		123	77.4	39	67.2	38	43.2	0.003	77	52.7		
Viral load missing at baseline	8	8.6	4	6.1	0.550	12	7.5	6	10.3	10	11.3	0.847	16	11.0		
CD4+ count (cells/mm <sup>3</sup> ) (M, SD)	397.0	158.0	511.0	289.1	0.002	444.0	227.8	462.0	263.2	648.0	416.6	0.004	572.0	372.4		
CD4+ count (cells/mm <sup>3</sup> ) (range)	(30, , 413)		(47, 1413)			(30, , 1413)		(126, 1508)		(13, 2614)			(13, 2614)			
<b>Socio-demographics</b>																
Age (M, SD)	41.0	7.3	38.0	8.6	0.019	40.0	8.0	40.0	9.3	35.0	8.7	0.001	37.0	9.3		
Age (range)	(21, 60)		(22, 62)			(21, 62)		(21, 60)		(19, 57)			(19, 60)			
18–24	1	1.1	4	6.1		5	3.1	2	3.4	11	12.5		13	8.9		
25–44	64	68.8	45	68.2		109	68.6	36	62.1	64	72.7		100	68.5		
45 or older		30.1	17	25.8		45	28.3	20	34.5	13	14.8		33	22.6		
<b>Education</b>																
Primary school or less	46	49.5	13	19.7	< 0.0001	59	37.1	20	34.4	23	26.2	0.033	43	29.5		
Secondary school (not/complete)	32	34.4	26	39.4		58	36.5	31	53.5	38	43.2		69	47.2		
Technical training or college (not/complete)	15	16.1	27	40.9		42	26.3	7	12.1	27	30.7		34	23.4		

<sup>a</sup>Mean and standard deviation reported for continuous variables

M mean, SD standard deviation; p-value for difference between heterosexual and MSM using  $\chi^2$  for comparisons among categorical variables and Student's t-test for comparison of means among continuous variables



**Table 2** The association of non-injection substance use with ART adherence and viral load (<200 copies/mL vs. 200 copies/mL) among men in Thailand and Brazil over 12 months, HPTN063

Outcomes	Brazil (n = 146)											
	Thailand (n = 159)					Took ART (past 3 mos.)						
	Adherence ability (good)					Undetectable viral load <sup>a</sup>						
Predictors	HM (n = 93)	MSM (n = 66)	Group diff.	HM (n = 93)	MSM (n = 66)	Group diff.	HM (n = 52)	MSM (n = 78)	Group diff.	HM (n = 50)	MSM (n = 54)	Group diff.
	AOR (95% CI)	AOR (95% CI)	p-value	AOR (95% CI)	AOR (95% CI)	p-value	AOR (95% CI)	AOR (95% CI)	AOR (95% CI)	AOR (95% CI)	AOR (95% CI)	p-value
Stimulants	0.96 (0.89, 1.03)	0.83 (0.48, 1.42)		0.94 (0.85, 1.04)	0.90 (0.34, 2.33)		0.97 (0.92, 1.02)	1.00 (0.95, 1.04)		1.00 (0.94, 1.05)	0.99 (0.96, 1.03)	
p-value	0.24	0.49	0.61	0.25	0.82	0.92	0.17	0.86	0.36	0.89	0.77	0.96
Marijuana	–	–	–	–	–	–	1.01 (0.95, 1.07)	1.00 (0.98, 1.03)		0.98 (0.94, 1.02)	1.00 (0.97, 1.03)	
p-value	–	–	–	–	–	–	0.80	0.89	0.86	0.35	0.83	0.53
Alcohol misuse	0.79 (0.37, 1.72)	0.59 (0.23, 1.51)		1.09 (0.38, 3.15)	1.28 (0.37, 4.45)		0.21 (0.04, 1.14)	0.67 (0.23, 1.99)		0.57 (0.15, 2.08)	0.34 (0.13, 0.92)	
p-value	0.56	0.27	0.63	0.87	0.70	0.85	0.07	0.48	0.25	0.39	0.03	0.54
# of substances	0.92 (0.47, 1.81)	0.85 (0.41, 1.74)		0.68 (0.26, 1.77)	1.81 (0.55, 5.99)		0.25 (0.08, 0.78)	0.70 (0.38, 1.30)		0.88 (0.37, 2.11)	1.20 (0.64, 2.25)	
p-value	0.82	0.66	0.87	0.43	0.33	0.20	0.02	0.26	0.12	0.42	0.71	0.83

All models adjusted for age and education

HM heterosexual men, MSM men who have sex with men

<sup>a</sup>Models estimating undetectable viral load in Thailand are adjusted for adherence ability

<sup>b</sup>Significance of ratio of AORs between HM and MSM

<sup>c</sup>Models estimating undetectable viral load in Brazil only includes observations where men reported taking ARTs in the last 3 months

**Table 3** The association of non-injection substance use with severe depressive symptoms and HIV viral load (<200 copies/mL vs. 200 copies/mL) among men in Thailand and Brazil over 12 months, HPTN063

Outcomes	Thailand (n = 159)						Brazil (n = 146)					
	Severe depressive symptoms			Undetectable viral load <sup>a</sup>			Severe depressive symptoms			Undetectable viral load <sup>c</sup>		
	HM (n = 93)	MSM (n = 66)	Group diff. <sup>b</sup>	HM (n = 93)	MSM (n = 66)	Group Diff.	HM (n = 52)	MSM (n = 78)	Group diff.	HM (n = 50)	MSM (n = 54)	Group diff.
AOR (95% CI)	AOR (95% CI)	p-value	AOR (95% CI)	AOR (95% CI)	p-value	AOR (95% CI)	AOR (95% CI)	AOR (95% CI)	AOR (95% CI)	AOR (95% CI)	AOR (95% CI)	p-value
Stimulants	1.03 (0.96, 1.12)	1.47 (0.82, 2.67)		0.93 (0.84, 1.03)	0.79 (0.30, 2.09)		1.00 (0.96, 1.04)	1.01 (0.98, 1.04)		1.00 (0.94, 1.05)	0.99 (0.96, 1.03)	
p-value	0.40	0.20	0.25	0.17	0.63	0.74	0.96	0.53	0.75	0.89	0.76	0.95
Marijuana	–	–	–	–	–	–	1.00 (0.97, 1.03)	1.02 (1.00, 1.04)		0.98 (0.94, 1.02)	1.00 (0.97, 1.03)	
p-value	–	–	–	–	–	–	0.99	0.12	0.43	0.33	0.82	0.51
Alcohol misuse	1.01 (0.48, 2.11)	2.75 (1.20, 6.32)		1.10 (0.38, 3.16)	1.14 (0.32, 4.09)		2.06 (0.71, 5.94)	0.73 (0.35, 1.51)		0.57 (0.15, 2.11)	0.34 (0.13, 0.92)	
p-value	0.99	0.02	0.08	0.86	0.83	0.96	0.18	0.40	0.11	0.40	0.03	0.53
#Substances	1.42 (0.73, 2.77)	1.30 (0.65, 2.58)		0.66 (0.26, 1.69)	1.68 (0.51, 5.58)		1.19 (0.64, 2.20)	1.24 (0.81, 1.88)		0.88 (0.35, 2.11)	1.20 (0.64, 2.25)	
p-value	0.30	0.46	0.85	0.38	0.40	0.22	0.59	0.32	0.91	0.77	0.57	0.57

All models adjusted for age and education

HM heterosexual men, MSM men who have sex with men

<sup>a</sup>Models estimating undetectable viral load in Thailand are adjusted for depression and adherence ability

<sup>b</sup>Ratio of ORs between HM and MSM

<sup>c</sup>Models estimating undetectable viral load in Brazil are adjusted for depression and only include observations where men reported taking ARTs in the last 3 months

non-significant interaction terms. Severe depressive symptoms did not mediate the association between substance use and undetectable VL. In Brazil, non-injection substance use was generally not significantly associated with reporting depressive symptoms. Alcohol misuse, although not significantly associated with reporting severe depressive symptoms, was significantly associated with decreased odds of having an undetectable VL over 12 months in MSM (AOR = 0.34; 95% CI 0.13, 0.92;  $p$ -value = 0.03). There were no differences between risk groups demonstrated by non-significant interaction terms. Severe depressive symptoms did not mediate the association between substance use and undetectable VL.

## Discussion

This exploratory study examined the overlap between reported non-injection substance use, severity of depressive symptoms, ART adherence, and HIV VL undetectability among men living with HIV in Rio de Janeiro (Brazil) and Chiang Mai (Thailand) over 12 months. We found varying types and patterns of non-injection substance use between countries and sub-groups. One key finding is that alcohol misuse, although not associated with reported ART adherence ability in Thailand or with taking ARTs in Brazil, was associated with significantly lower odds of achieving undetectable VL among MSM in Brazil. Another key finding is that the number of non-injection substances used was associated with lower odds of taking ARTs in the past 3 months among heterosexual men in Brazil, but not in Thailand. Lastly, alcohol misuse was associated with significantly greater odds of having depressive symptoms among MSM in Thailand, although not significantly associated with HIV VL.

Reported alcohol misuse was prevalent in this sample and was associated with significantly lower odds of achieving an undetectable VL among MSM in Brazil. Alcohol misuse was detected in 35.2% (MSM in Brazil) and 47.3% (heterosexual men in Thailand) in our of HIV-infected men. This high prevalence of alcohol misuse is consistent with one review documenting that alcohol use disorders (AUDs) can be up to two to four times more prevalent among PLH than the general population in U.S. populations [49]. Factors that might explain lack of HIV suppression in our sample could range from biological factors to the diminished cognitive function and dysfunctional behaviors caused by alcohol misuse that may lead to poor self-regulation. Alcohol misuse might directly affect HIV control by inhibiting ART metabolism [50], enhancing HIV disease progression by lowering CD4 + T-cell count [51], and/or increasing HIV replication in peripheral blood mononuclear cells (PBMCs) [52]. These biological mechanisms deserve further research in human

subjects, as the current knowledge base is largely limited to animal models. Regardless of the mechanism, our findings support the rationale for investing resources into alcohol misuse screening and prevention interventions among men with HIV/AIDS in middle-income countries, such as Brazil and Thailand.

Another key finding is that each additional substance used was associated with lower odds of taking ART among heterosexual men in Brazil. Polysubstance use among heterosexual men in Brazil involved reporting a combination of alcohol misuse, powder cocaine use, and/or cannabis use. Substance use, powder cocaine in particular, has been previously associated with poor ART adherence ability and faster HIV disease progression [8, 12, 21]. Specifically, cocaine may increase HIV disease progression by increasing HIV replication in PBMCs (in vitro) [23] and increasing circulating HIV-1 RNA (mouse models) [25]. There are fewer studies on the effect of cannabis on ART adherence ability and HIV VL with mixed findings [53]. Interestingly, non-injection substance use was associated with decreased odds of taking ART only among heterosexual men in Brazil. Previous studies that examined non-injection substance use among individuals with HIV have primarily focused on MSM [54, 55]. As there is limited research on non-injection substance use and ART adherence among HIV-infected heterosexual men, future research should examine this relationship to elucidate the contributing factors. Blips in HIV VL exams are also more frequent among people who misuse alcohol and drugs. Additional studies of ART adherence with biomarkers would enhance the understanding of how polysubstance use, ART, and HIV VL interact physiologically.

Lastly, we found that alcohol misuse was associated with significantly greater odds of having depressive symptoms among MSM in Thailand. Although our study found that depressive symptoms were not significantly associated with undetectable HIV VL, depression severity is consistently associated with inconsistent ART adherence and discontinuation [16]. Future research is needed to evaluate the efficacy of psychological and psychiatric interventions in mitigating the deleterious effects of substance use and depression on HIV disease progression. A recent critical literature review highlights some promising cognitive and behavioral and motivational interview interventions conducted among HIV-infected substance using MSM in the US [56]. Such interventions need to be adapted and evaluated in other countries and socio-cultural contexts.

The current findings should be considered in light of several limitations and strengths. First, non-injection substance use and ART adherence ability were self-reported and subject to potential biases based on recall bias or social desirability, the intentional under-reporting of sensitive or socially undesirable outcomes. There was

likely under-reporting of alcohol misuse, non-injection drug use, and ART non-adherence. Future studies should include more comprehensive measurements of substance use and ART adherence. For example, physiological biomarkers of substance use and ART adherence provide a more objective measure of chronicity and extent of substance use. Likewise, future studies would benefit from using instruments that assess substance misuse (e.g., ASI, DAST, DUDIT), as the current study assessed the number of days of non-injection drug use rather than misuse. Second, under-reporting, small sample size, and truncated variability could have decreased our statistical power to detect a significant association between key variables like stimulants, cannabis, polysubstance use, and HIV outcomes. Furthermore, it is important to highlight that significant associations were found in only one of the four sub-groups. Inconsistent findings could reflect distinct substance use and HIV care characteristics across countries and sub-groups, but could also be due to type 1 error. Third, our findings are not generalizable to populations of HIV-infected men in Thailand and in Brazil as this study focused on men engaged in care in select clinics and cities in each country.

Despite these limitations, this study contributes to evidence that achieving an undetectable VL is possible among male, non-injection substance users in low- and middle-income countries. Our results suggest TasP may be attainable among PLH who use non-injection substances. However, among MSM in Thailand and Brazil who misuse alcohol and among heterosexual men in Brazil who use multiple non-injection substances, interventions that address substance use may aim to lift mood, boost ART adherence and reduce HIV VL.

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## Compliance with Ethical Standards

**Conflict of interest** The authors declare that they have no conflict of interest.

**Ethical Approval** Ethical considerations reviewed and approved by institutional review boards (IRB) within each recruitment coun-

try. Informed consent was obtained from all individual participants included in the study prior to interview.

## References

- Gardner EM, McLees MP, Steiner JF, del Rio C, Burman WJ. The spectrum of engagement in HIV care and its relevance to test-and-treat strategies for prevention of HIV infection. *Clin Infect Dis*. 2011;52(6):793–800.
- WHO. Antiretroviral treatment as prevention (TASP) of HIV and TB. 2012.
- Cohen M, Chen Y, McCauley M, Gamble T, Hosseinipour M, Kumarasamy N, et al. Prevention of HIV-1 infection with early antiretroviral therapy. *N Engl J Med*. 2011;11(365):469–505.
- Günthard HF, Saag MS, Benson CA, Del Rio C, Eron JJ, Gallant JE, et al. Antiretroviral drugs for treatment and prevention of HIV infection in adults: 2016 recommendations of the International Antiviral Society—USA panel. *JAMA*. 2016;316(2):191–210.
- Shoptaw S, Montgomery B, Williams CT, El-Bassel N, Aramratana A, Metzger DS, et al. Not just the needle: the state of HIV prevention science among substance users and future directions. *J Acquir Immune Defic Syndr*. 1999;2013(63):S174.
- Gonzalez A, Barinas J, O' Cleirigh C. Substance use: impact on adherence and HIV medical treatment. *Curr HIV/AIDS Rep*. 2011;8(4):223.
- Bogart LM, Kelly JA, Catz SL, Sosman JM. Impact of medical and nonmedical factors on physician decision making for HIV/AIDS antiretroviral treatment. *JAIDS J Acquir Immune Defic Syndr*. 2000;23(5):396–404.
- Cofrancesco J Jr, Scherzer R, Tien PC, Gibert CL, Southwell H, Sidney S, et al. Illicit drug use and HIV treatment outcomes in a US cohort. *AIDS (London, England)*. 2008;22(3):357.
- Lucas GM, Cheever LW, Chaisson RE, Moore RD. Detrimental effects of continued illicit drug use on the treatment of HIV-1 infection. *JAIDS J Acquir Immune Defic Syndr*. 2001;27(3):251–9.
- McGowan CC, Weinstein DD, Samenow CP, Stinnette SE, Barkanic G, Rebeiro PF, et al. Drug use and receipt of highly active antiretroviral therapy among HIV-infected persons in two US clinic cohorts. *PLoS ONE*. 2011;6(4):e18462.
- Arnsten JH, Demas PA, Grant RW, Gourevitch MN, Farzadegan H, Howard AA, et al. Impact of active drug use on antiretroviral therapy adherence and viral suppression in hiv-infected drug users. *J Gen Intern Med*. 2002;17(5):377–81.
- Hinkin CH, Barclay TR, Castellon SA, Levine AJ, Durvasula RS, Marion SD, et al. Drug use and medication adherence among HIV-1 infected individuals. *AIDS Behav*. 2007;11(2):185–94.
- Mills EJ, Nachega JB, Bangsberg DR, Singh S, Rachlis B, Wu P, et al. Adherence to HAART: a systematic review of developed and developing nation patient-reported barriers and facilitators. *PLoS Med*. 2006;3(11):2039–64.
- Wilson IB, Carter AE, Berg KM. Improving the self-report of HIV antiretroviral medication adherence: is the glass half full or half empty? *Curr HIV/AIDS Rep*. 2009;6(4):177–86.
- Carrico AW. Substance use and HIV disease progression in the HAART era: implications for the primary prevention of HIV. *Life Sci*. 2011;88(21):940–7.
- Carrico AW, Riley ED, Johnson MO, Charlebois ED, Neilands TB, Remien RH, et al. Psychiatric risk factors for HIV disease progression: the role of inconsistent patterns of anti-retroviral therapy utilization. *J Acquir Immune Defic Syndr*. 2011;56(2):146.
- Ellis RJ, Childers ME, Cherner M, Lazzaretto D, Letendre S, Grant I, et al. Increased human immunodeficiency virus loads in

- active methamphetamine users are explained by reduced effectiveness of antiretroviral therapy. *J Infect Dis.* 2003;188:1820–6.
18. Cook JA, Burke-Miller JK, Cohen MH, Cook RL, Vlahov D, Wilson TE, et al. Crack cocaine, disease progression, and mortality in a multi-center cohort of HIV-1 positive women. *AIDS (London, England).* 2008;22(11):1355.
  19. Morin SF, Myers JJ, Shade SB, Koester K, Maiorana A, Rose CD. Predicting HIV transmission risk among HIV-infected patients seen in clinical settings. *AIDS Behav.* 2007;11(1):6–16.
  20. Parsons JT, Bimbi DS. Intentional unprotected anal intercourse among sex who have sex with men: barebacking—from behavior to identity. *AIDS Behav.* 2007;11(2):277–87.
  21. Rasbach DA, Desruisseau AJ, Kipp AM, Stinnette S, Kheshti A, Shepherd BE, et al. Active cocaine use is associated with lack of HIV-1 virologic suppression independent of nonadherence to antiretroviral therapy: use of a rapid screening tool during routine clinic visits. *AIDS care.* 2013;25(1):109–17.
  22. Bagasra O, Pomerantz RJ. Human immunodeficiency virus type 1 replication in peripheral blood mononuclear cells in the presence of cocaine. *J Infect Dis.* 1993;168(5):1157–64.
  23. Cabral GA. Drugs of abuse, immune modulation, and AIDS. *J Neuroimmune Pharmacol.* 2006;1(3):280–95.
  24. Nair MP, Chadha KC, Hewitt RG, Mahajan S, Sweet A, Schwartz SA. Cocaine differentially modulates chemokine production by mononuclear cells from normal donors and human immunodeficiency virus type 1-infected patients. *Clin Diagn Lab Immunol.* 2000;7(1):96–100.
  25. Tashkin DP. Evidence implicating cocaine as a possible risk factor for HIV infection. *J Neuroimmunol.* 2004;147(1):26–7.
  26. Hasin D, Kilcoyne B. Comorbidity of psychiatric and substance use disorders in the United States: current issues and findings from the NESARC. *Curr Opin Psychiatry.* 2012;25(3):165.
  27. Bing EG, Burnam MA, Longshore D, Fleishman JA, Sherbourne CD, London AS, et al. Psychiatric disorders and drug use among human immunodeficiency virus-infected adults in the United States. *Arch Gen Psychiatry.* 2001;58(8):721–8.
  28. Ciesla JA, Roberts JE. Meta-analysis of the relationship between HIV infection and risk for depressive disorders. *Am J Psychiatry.* 2001;158(5):725–30.
  29. Gonzalez JS, Batchelder AW, Psaros C, Safren SA. Depression and HIV/AIDS treatment nonadherence: a review and meta-analysis. *J Acquir Immune Defic Syndr.* 2011. <https://doi.org/10.1097/QAI.0b013e31822d490a>.
  30. Leserman J. Role of depression, stress, and trauma in HIV disease progression. *Psychosom Med.* 2008;70(5):539–45.
  31. Antoni MH, Carrico AW, Durán RE, Spitzer S, Penedo F, Ironson G, et al. Randomized clinical trial of cognitive behavioral stress management on human immunodeficiency virus viral load in gay men treated with highly active antiretroviral therapy. *Psychosom Med.* 2006;68(1):143–51.
  32. Rosen M, Black A, Arnsten J, Goggin K, Remien R, Simoni J, et al. Association between use of specific drugs and antiretroviral adherence: findings from MACH 14. *AIDS Behav.* 2013;17(1):142–7.
  33. Reback CJ, Larkins S, Shoptaw S. Methamphetamine abuse as a barrier to HIV medication adherence among gay and bisexual men. *AIDS Care.* 2003;15(6):775–85.
  34. Feelemyer J, Des Jarlais D, Arasteh K, Uusküla A. Adherence to antiretroviral medications among persons who inject drugs in transitional, low and middle income countries: an international systematic review. *AIDS Behav.* 2015;19(4):575–83.
  35. Hatzenbuehler ML, Nolen-Hoeksema S, Erickson SJ. Minority stress predictors of HIV risk behavior, substance use, and depressive symptoms: results from a prospective study of bereaved gay men. *Health Psychol.* 2008;27(4):455.
  36. Thai National AIDS Committee. Thailand ending AIDS: 2014 Thailand AIDS response progress report. 2014.
  37. UNAIDS Global AIDS Response Report. The Brazilian Response to HIV and AIDS (2015). Brasilia, D.F.
  38. Departamento de DST Aids e Hepatites Virais. AIDS in Brazil: Brazilian Policy Focus. Brasilia; 2012 July 2012.
  39. MacCarthy S, Brignol S, Reddy M, Nunn A, Dourado I. Late presentation to HIV/AIDS care in Brazil among men who self-identify as heterosexual. *Rev de Saude Publ.* 2016;50:54.
  40. Tsuyuki K, Pitpitan EV, Levi-Minzi MA, Urada LA, Kurtz SP, Stockman JK, et al. Substance use disorders, violence, mental health, and HIV: differentiating a syndemic factor by gender and sexuality. *AIDS Behav.* 2017;21:1–13.
  41. Fendrich M, Avci O, Johnson TP, Mackesy-Amity ME. Depression, substance use and HIV risk in a probability sample of men who have sex with men. *Addict Behav.* 2013;38(3):1715–8.
  42. Safren SA, Hughes JP, Mimiaga MJ, Moore AT, Friedman RK, Srithanaviboonchai K, et al. Frequency and predictors of estimated HIV transmissions and bacterial STI acquisition among HIV-positive patients in HIV care across three continents. *J Int AIDS Soc.* 2016;19(1):21096.
  43. Magidson JF, Li X, Mimiaga MJ, Moore AT, Srithanaviboonchai K, Friedman RK, et al. Antiretroviral medication adherence and amplified HIV transmission risk among sexually active HIV-infected individuals in three diverse international settings. *AIDS Behav.* 2015;20:1–11.
  44. Bush K, Kivlahan DR, McDonnell MB, Fihn SD, Bradley KA. The AUDIT alcohol consumption questions (AUDIT-C): an effective brief screening test for problem drinking. *Arch Intern Med.* 1998;158(16):1789–95.
  45. Radloff L. The CES-D scale: a self-report depression scale for researchers in the general population. *Appl Psychol Meas.* 1977;1:385–401.
  46. Lu M, Safren SA, Skolnik PR, Rogers WH, Coady W, Hardy H, et al. Optimal recall period and response task for self-reported HIV medication adherence. *AIDS Behav.* 2008;12(1):86–94.
  47. Feldman B, Fredericksen R, Crane P, Safren S, Mugavero M, Willig JH, et al. Evaluation of the single-item self-rating adherence scale for use in routine clinical care of people living with HIV. *AIDS Behav.* 2013;17(1):307–18.
  48. Hedeker D. Generalized linear mixed models. *Encyclopedia of statistics in behavioral science.* 2005. p. 1-10.
  49. Vagenas P, Azar MM, Copenhaver MM, Springer SA, Molina PE, Altice FL. The impact of alcohol use and related disorders on the HIV continuum of care: a systematic review. *Curr HIV/AIDS Rep.* 2015;12(4):421–36.
  50. Hahn JA, Samet JH. Alcohol and HIV disease progression: weighing the evidence. *Curr HIV/AIDS Rep.* 2010;7(4):226–33.
  51. Kahler CW, Liu T, Cioe PA, Bryant V, Pinkston MM, Kojic EM, et al. Direct and indirect effects of heavy alcohol use on clinical outcomes in a longitudinal study of HIV patients on ART. *AIDS Behav.* 2016;21:1–11.
  52. Baum MK, Rafie C, Lai S, Sales S, Page JB, Campa A. Alcohol use accelerates HIV disease progression. *AIDS Res Hum Retroviruses.* 2010;26(5):511–8.
  53. Lake S, Kerr T, Capler R, Shoveller J, Montaner J, Milloy M-J. High-intensity cannabis use and HIV clinical outcomes among HIV-positive people who use illicit drugs in Vancouver, Canada. *Int J Drug Policy.* 2017;42:63–70.
  54. Strathdee SA, Stockman JK. Epidemiology of HIV among injecting and non-injecting drug users: current trends and implications for interventions. *Curr HIV/AIDS Rep.* 2010;7(2):99–106.

55. Patterson TL, Semple SJ, Zians JK, Strathdee SA. Methamphetamine-using HIV-positive men who have sex with men: correlates of polydrug use. *J Urban Health*. 2005;82(1):i120–6.
56. Carrico AW, Zepf R, Meanley S, Batchelder A, Stall R. Critical review: when the party is over: A systematic review of behavioral interventions for substance-using men who have sex with men. *JAIDS J Acquir Immune Defic Syndr*. 2016;73(3):299–306.

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