

Change over 12-months of an Integrated Treatment Program for Opiate Addiction and HIV in Ho Chi Minh City, Vietnam:

highlighting the role of counseling





T. Nguyen^{1,2}, C.Denis^{3,4}, A. Voisin⁵, T. Doan², H. Tran², L. Nguyen², V. Trias², L. Huang², M. Auriacombe^{4,3}, université G. Raguin², S. Mai Thi Hoai⁶, G. Le Truong⁶, J-P. Daulouède^{5,4,2}, C. O'Brien³, D. Metzger^{3,7}

¹Ho Chi Minh City University of Medicine and Pharmacy; ² Expertise France, Paris, France; ³ University of Pennsylvania, Philadelphia, PA, USA; ⁴ CNRS USR 3413 SANPsy, University of Bordeaux, Bordeaux, France; ⁵ Bizia Addiction Clinic, Bayonne, France; ⁶ HCMC Provincial AIDS Committee, Ho Chi Minh City, Vietnam; ⁷ Treatment Research Institute, Philadelphia, PA, USA

Abstract	Methods	Results (cont')	
Background: As part of the integrated drug treatment program implemented in an HIV treatment setting at Go Vap clinic, Ho Chi Minh City, Vietnam, we have developed structured counseling sessions. Methods: Participants were assessed at baseline, weekly for 12 weeks and then	 Efficacy of treatment at 12 months was assessed by Treatment retention Heroin use 	 Change in craving, mood, family relationships, satisfaction with income/work, satisfaction with recovery 	
monthly. Here we describe the retention, the treatment outcomes over the 12- month follow-up. Results: 448 heroin injectors (97.2% males, 32.3 y.o.) were enrolled (268 receiving methadone – MET, and 180 BUP/NX -Suboxone [®]). The retention in	Self-reports of drug use: collected on a weekly basis for 12 weeks and monthly thereafter Urine drug screens: randomly once a week for the first 12 weeks of treatment and twice a month thereafter Agreement between self-report use and urine drug screen: high (89%)	- Significant decrease of craving regardless of the treatment (F(12,232)=20.2, p<0.0001) BUP/NX > MET	

methadone treatment at 12-month was 89.9% for MET, 57.0% for BUP/NX. Treatment adherence was high. The three treatment approaches were equally effective with a significant decrease in the days of heroin use (F(12,277)= 21.8, p<.0001) along with a significant decrease of reported craving (F(12,232)= 20.2, p<.0001), a significant improvement of mood (F(12,232)= 5.88, p<.0001), satisfaction with income/work (F(12,232)= 4.64, p<.0001), and satisfaction with recovery (F(11,249)= 3.78, p<.0001). Change in heroin use was found highly correlated with all these previously listed outcomes (correlations range: 0.20-0.72). The baseline characteristics of the participants who dropped out treatment did not differ significantly from those who completed the 12-month program. Lower self-rated mood was associated with dropping out of treatment (aOR=1.3, 95% CI=1.1-1.5), highly correlated with family relationships (p= 0.32, p< 0.0001) and satisfaction with income/work (p= 0.48, p<0.0001).

Conclusion: As part of an integrated treatment program, structured counseling sessions using both relapse prevention and cognitive-behavioral techniques that address treatment observance, substance outcomes, client's psychosocial needs and their family relationships enhance the program retention and effectiveness.

Background

- Vietnam:
- 271,000 people using drugs (UNAIDS 2015)
- 14,000 new HIV infections in 2015, and most of them are driven by drug users
- 2006: Scale up of HIV prevention programs targeting people who inject drugs (PWID) (Giang et al., 2013)

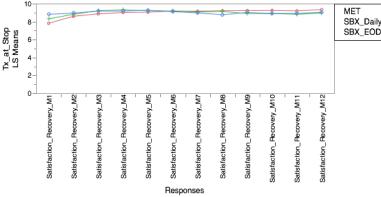
- Self-evaluation during counseling sessions 12 weekly sessions then 10 monthly sessions thereafter - Craving
 - Mood
 - Family relationships
 - Satisfaction with income/work
 - Satisfaction with recovery
- Comparative assessments for each treatment strategies have been performed using Chi², t-tests, and analyses of variance. Multivariable models were performed to evaluate the factors associated with retention and treatment outcomes. Spearman's correlation were used to evaluate the correlations between variables.

Results

Participant characteristics at baseline

	All	MET	BUP/NX	Test value, p-value
	n=448	n=268	n= 180	71
Age Mean (SD)	32.6 (5.9)	32.5 (6.1)	32.7 (5.6)	t=0.52, p= 0.59
Gender – Males n (%)	434 (96.9)	260 (97.0)	174 (96.7)	$\chi^2 = 0.04, p = 0.83$
Education – High school or higher n (%)	69 (35.9)	51 (40.8)	51 (40.8)	$\chi^2=0.70, p=0.40$
Living condition	368 (82.5)	215 (80.5)	154 (86.0)	χ ² =5.37, p=0.25
With parents/ family n (%)				
Currently have a job n (%)	235 (52.7)	153 (57.3)	82 (45.8)	$\chi^2 = 0.68, p = 0.02$
Serology n (%)				
HIV-positive	152 (34.2)	96 (36.0)	56 (31.5)	$\chi^2 = 0.96, p = 0.33$
Hepatitis C-positive	323 (72.4)	184 (69.0)	139 (77.7)	χ^2 =4.17, p=0.05
Opiate Use				
Years of use - Mean (SD)	7.8 (6.2)	7.9 (6.9)	7.6 (4.9)	t=0.68, p=0.49
Days of use past 30 days - Mean (SD)	29.8 (1.4)	29.8 (1.2)	29.6 (1.8)	t=1.31, p=0.19
No. previous drug treatment Mean (SD)	5.4 (4.7)	5.9 (5.2)	4.6 (3.7)	t=3.14, p=0.002
Other substance use $-n$ (%) of users past 30 days				
Alcohol	86 (19.3)	48 (18.0)	38 (21.2)	$\chi^2 = 0.73$, p=0.39
Amphet./ Methamphetamines	91 (20.4)	31 (11.6)	60 (33.5)	$\chi^2 = 31.67, p < 0.0001$
Benzodiazepines	67 (15.0)	27 (10.1)	40 (22.3)	$\chi^2 = 12.56, p = 0.0004$
Tobacco	440 (98.6)	264 (98.9)	176 (98.3)	$\chi^2 = 0.25 \text{ p} = 0.67$

- First 3 months of treatment Significant improvement of mood regardless of the treatment (F(12,232)=5.88, p<0.0001) BUP/NX > MET First 3 months of treatment - Significant improvement of satisfaction with income/work regardless of the treatment (F(12,232)=4.64, p<0.0001) - No change in family support that stayed high over the 12-month follow-up period (F(12, 232)=1.21, p=0.28) - Significant improvement of satisfaction with recovery
 - regardless of the treatment (F(11,249)=3.78, p<0.0001) BUP/NX > MET

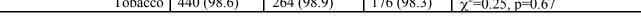


Factors associated with dropping out treatment program Lower self-rated mood (aOR=1.3, 95% CI=1.1-1.5)

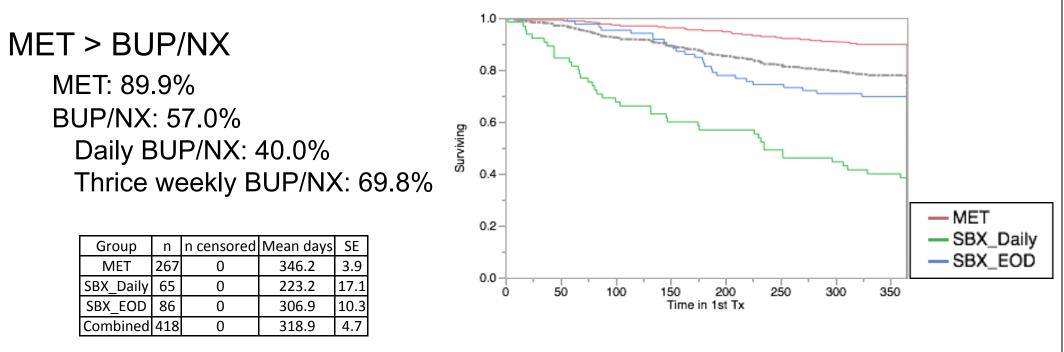
Sterile injection equipment, the introduction and expansion of methadone treatment, and increased access to antiretroviral therapy (Government of Vietnam, 2014)

- Although the proportion of HIV infections among PWID has declined from 30% in 2005 to 22% in 2013, PWID remain the largest risk group living with HIV infection (Vietnam Administration of HIV/AIDS Control, 2013)
- Since November 2013: Implementation of an integrated drug treatment program within an HIV treatment setting at Go Vap clinic, Ho Chi Minh City, Vietnam.
- Integrated Treatment provided
 - A pharmacological opiate maintenance treatment with methadone (MET) or buprenorphine/naloxone (BUP/NX)
 - Introduction of BUP/NX (Suboxone[®]) in January 2015
 - First time in Vietnam
 - Counseling: 12 weekly sessions and 10 monthly sessions thereafter
 - HIV screening and HIV treatment if needed
 - HCV screening
- Structured and manual-based standardized counseling sessions
 - Rooted in cognitive behavioral therapy
 - Sessions assess the need for intervention in six areas of functioning: 1) Adherence to SUD, HIV, TB; 2) continued drug use and related drug and sex risk; 3) cravings for drug use; 4) psychological status (depression, anxiety, symptoms of psychiatric disorder); 5) confidence in and satisfaction with SUD treatment; and, 6) strategies for the next month
 - Data recorded and available for review with the participant as a clinical tool to show progress and promote self-monitoring.

Objectives



High retention in treatment at 12 months (78.0%)



Wilcoxon chi²= 110.9, p<0.0001

6

.9)

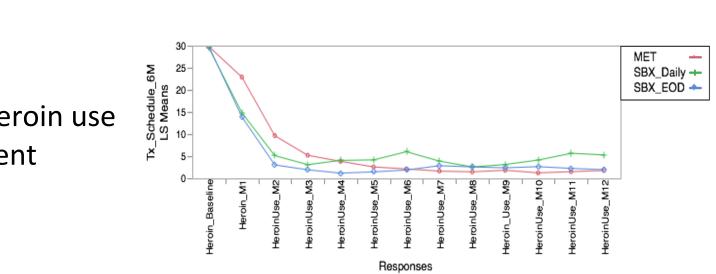
Average daily dose (mg/day (SD)) **HIV-positive HIV-negative** All 118.5 (61.8) 163.5 (64.7) MET 101.6 (52.0)

BUP/NX daily	18.5 (5.6)	19.0 (7.4)	18.2 (4.
BUP/NX 3x/wk	14.8 (1.8)	15.1 (1.7)	14.7 (1.



Significant decrease of heroin use regardless of the treatment (F(12,277)=21.8, p<0.0001)

BUP/NX > MET First 3 months of treatment



Highly correlated with

First 3 months of treatment

Family relationships (ρ = 0.32, p<0.0001) Satisfaction with income/work (p=0.48, p<0.0001)

Conclusion

- The three approaches to treatment appear to be as efficient to treat opiate use disorder
 - As part of a comprehensive and integrated treatment program i.e. OMT and structured counseling sessions

BUP/NX quit heroin earlier (within the first 3 months of Tx)

- In a context where no take-home dose is allowed, BUP/NX thrice weekly
 - Enhance retention and adherence in treatment
 - Reduce heroin use sooner
 - Less time spend traveling to and from the clinic (reduces the cost for the client)
 - Less disturbance that prevents from working (less missed time from work)
- Importance of counseling sessions that address: substance use, craving, psychological functioning, family and social relationships Enhance retention in treatment program

To describe the retention and treatment outcomes over the 12month follow-up period

Dosing schedule: All participants were required to receive their medications in directly observed dosing as Vietnamese law does not allow take-home medication.

Methadone: Daily dosing at the clinic

Buprenorphine/naloxone: Daily dosing at the clinic at the induction of treatment

After four weeks of stable dose, switch to thrice weekly dosing (if clinically possible)

- Thus, we measured the efficacy of three opiate maintenance treatment strategies: daily methadone (MET), daily buprenorphine/naloxone (BUP/NX) and thrice weekly BUP/NX

To identify the factors associated with treatment retention

Funding: NIDA R01- DA033671

Conflict of Interest: Suboxone[®] tablets were donated by Reckitt-Benckiser/Indivior PLC. The pharmaceutical company has no role in study design, data collection, nor data analyses

(F(12,702)=0.69, p<0.0001)

No change in other substance use

Change in heroin use correlated with

	MET	BUP/NX	
Craving	0.72	0.64	
Mood	0.54	0.56	
Family	0.27	0.20	
Income Satisfaction	0.30	0.31	
w/ recovery	0.56	0.52	All p<0.0001

• Not able to predict who will respond better on one medication versus another

More treatment options available, more likely one will find a more suitable treatment option

Long-acting medication within a comprehensive addiction treatment could be valuable to enhance treatment engagement, retention and adherence

Contact

Cécile Denis, PhD	cdenis@upenn.edu
David Metzger, PhD	dsm@upenn.edu
Charles O'Brien, MD, PhD	obrien@upenn.edu