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Correlates of cocaine use during methadone treatment: implications for screening and clinical management (ANRS Methaville study)

Perrine Roux^{1,2,3*}, Caroline Lions^{1,2,3}, Antoine Vilotitch^{1,2,3}, Laurent Michel^{4,5,6}, Marion Mora^{1,2,3}, Gwenaëlle Maradan^{1,2,3}, Fabienne Marcellin^{1,2,3}, Bruno Spire^{1,2,3}, Alain Morel⁷, Patrizia M. Carrieri^{1,2,3} and The ANRS Methaville study group

Abstract

Background: Cocaine use is frequent in patients receiving methadone maintenance treatment (MMT) and can jeopardize their treatment response. Identifying clinical predictors of cocaine use during methadone treatment can potentially improve clinical management. We used longitudinal data from the ANRS Methaville trial both to describe self-reported occasional and regular cocaine use during MMT and to identify clinical predictors.

Methods: We selected 183 patients who had data on cocaine (or crack) use at months 0 (M0), M6, and/or M12, accounting for 483 visits. The outcome was “cocaine use” in three categories: “no,” “occasional,” and “regular” use. To identify factors associated with the outcome over time, we performed a mixed multinomial logistic regression.

Results: Time on methadone was significantly associated with a decrease in occasional but not in regular cocaine use from 14.7 % at M0 to 7.1 % at M12, and from 10.7 % at baseline to 6.5 % at M12, respectively. After multiple adjustments, opiate injection, individuals screening positive for attention deficit hyperactivity disorder (ADHD) symptoms, and those presenting depressive symptoms were more likely to regularly use cocaine.

Conclusions: Although time on MMT had a positive impact on occasional cocaine use, it had no impact on regular cocaine use. Moreover, regular cocaine users were more likely to report opiate injection and to present ADHD and depressive symptoms. Early screening of these disorders and prompt tailored pharmacological and behavioral interventions can potentially reduce cocaine use and improve response to MMT.

Trial registration: The trial is registered with the French Agency of Pharmaceutical Products (AFSSAPS) under the number 2008-A0277-48, the European Union Drug Regulating Authorities Clinical Trials, number Eudract 2008-001338-28, the ClinicalTrials.gov Identifier: NCT00657397, and the International Standard Randomised Controlled Trial Number Register ISRCTN31125511.

Keywords: Methadone, Opioid dependence, Cocaine, ADHD, Maintenance, Depression, Injection

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Background

Opioid maintenance treatment (OMT), especially methadone maintenance treatment (MMT), is widely recognized as a gold standard for managing opioid dependence [1]. However, many socio-economic, behavioral, and clinical conditions and determinants may impair response to MMT and consequently negatively impact the primary goal of this treatment which is abstinence from street-opioid use. Studies have pointed out that many opioid-dependent individuals are also either cocaine-dependent or are cocaine users [2–4] and that the effectiveness of methadone treatment is lower in this population [5]. Moreover, no pharmacological treatment currently exists for harmful cocaine use or dependence. Treatment is based on psychological intervention using cognitive behavioral therapy (CBT) or contingency management [6, 7].

Some studies have shown that retention in OMT has a positive impact on cocaine use, especially in less severe cocaine users [4, 8, 9]. However, other studies highlight that people who continue to use cocaine while on OMT are more likely to have a poorer response to treatment, in terms of retention and abstinence from street-opioid use [10, 11], and may also have a lower level of pharmacokinetic exposure to methadone, the consequence being a decreased effect of methadone [12]. Moreover, studies on methadone-related mortality [13, 14] have shown that cocaine use while on OMT is associated with a higher risk of overdose. A recent study conducted in San Francisco showed that more than one third of all fatal opioid overdoses involved also cocaine [15]. As it is known that cocaine use during methadone treatment may be associated with premature discharge [16], it seems important to identify predictors of cocaine use, in order to improve clinical management of these patients. Some hypotheses have been put forward regarding the possible association between psychiatric comorbidities—such as depression [17], attention deficit/hyperactivity disorder (ADHD) [18], alcohol dependence [19], and severe opioid dependence [20]—and more severe patterns of cocaine use [21, 22]. However, it would seem more relevant to investigate correlates of cocaine use more thoroughly and to provide clinicians with indications as to how they can better manage MMT patients whose cocaine use is deemed “at-risk.” We used longitudinal data from the ANRS Methaville trial, which enrolled opioid-dependent individuals starting methadone maintenance, to describe the pattern of cocaine use during MMT and to identify clinical correlates.

Methods

Study design

From January 2009 to January 2010, the ANRS Methaville study—a multi-site, open-label, randomized, controlled,

non-inferiority trial—recruited 195 men and women in 10 sites in France. The study aimed to compare methadone initiation in France in specialized centers (standard care) with initiation in primary care. This study was approved by the ethics committee for the protection of patients in Paris, France. All individuals provided written, informed consent before participating in the study. The full protocol is described elsewhere [23]. Each participant was followed up for 12 months. Medical visits, completion of self-administered questionnaires, and phone interviews occurred at enrolment (M0), 3, 6, and 12 months (M3, M6, and M12, respectively).

Variables and instruments

Computerized assisted phone interviews (CATI) were used to collect the following information at enrolment and during follow-up visits: (1) socio-demographic characteristics: age, gender, employment status, housing status (renter or owner); (2) history of drug use: age at first drug use, history of overdoses, history of drug injection; (3) current drug, alcohol, and tobacco use; (4) perception of the adequacy of the prescribed methadone dose, categorized into 3 possible answers: too high, adequate, too low. This was done to assess each interviewee’s satisfaction with methadone treatment.

Patients reporting that they had used cocaine or crack use once during the previous month were considered “occasional cocaine users,” while those reporting cocaine (or crack) use at least twice in the previous month were considered “regular cocaine users.”

Alcohol consumption assessment was based on the alcohol use disorders test (AUDIT) with a threshold of 13 indicating alcohol dependence [24], while depressive symptoms were assessed using the Center for Epidemiologic Studies-Depression Scale (CES-D) with a threshold of 17 indicating depression for males and 23 for females [25]. The Fagerstrom Scale, comprising 6 items, evaluated tobacco dependence, a threshold of 3 (on a 0 to 10 scale) defining dependence [26].

Drug use was assessed using the Opiate Treatment Index (OTI) questionnaire, which documented the previous 3 days when drugs were used (last 3 days when drugs were taken and amounts consumed) during the previous month [27].

The self-administered questionnaire at M0, M6, and M12 included two screening tools. The first was the Adult ADHD Self-Report Scale-Version 1.1 (ASRS-V1.1) which evaluates attention deficit hyperactivity disorder (ADHD) in patients. The scoring algorithm used was the sum score obtained adding up the scores (0–4) of the first 6 items [28]. Then, we defined the diagnosis of ADHD using 14 as a cut-off score [29]. The second tool was the Beck Hopelessness Scale (BHS), a 20-item self-reported inventory, where a score of 9 or more indicates suicide risk [30].

During medical visits, physicians collected data on withdrawal symptoms using the Objective Opioid Withdrawal Scale (OOWS) which comprises a list of 13 withdrawal symptoms [31].

During the phone interviews, patients were asked about their prescribed methadone dose. At each follow-up visit, physicians noted the methadone dose prescribed to the patient in the medical questionnaire.

Statistical analyses

From the 195 patients included in the trial, we used data of 183 patients (177 at baseline) who answered the OTI section about cocaine use at least once during follow-up, accounting for 486 visits. For these analyses, we used only the M0, M6 and M12 visits where data for most of the variables were available. First, we compared the individual characteristics of the study sample between those who consumed cocaine or crack at enrolment (i.e. before methadone initiation) and those who did not. These comparisons were performed using a chi-squared test (for categorical variables) and a Wilcoxon test (for continuous variables). To identify the factors associated with cocaine use during treatment, we used a multinomial mixed model with a 3-category outcome: no cocaine use (reference), occasional cocaine use, and regular cocaine use.

We tested the following factors as possible explanatory variables for the consumption of cocaine: (1) socio-demographic characteristics (sex, age, employment, housing (home owner or renter)); (2) drug use-related factors (history of drug injection, previous drug overdose, opioid injection); (3) clinical factors: methadone-associated characteristics (methadone induction arm, methadone dose: ≥ 60 mg vs. < 60 mg, perceiving methadone dose to be too low vs. adequate or too high); clinical factors (depressive symptoms, risk of suicide, ADHD symptoms, at least 1 withdrawal symptom); tobacco dependence and alcohol dependence. We first performed a univariate mixed multinomial logistic regression adjusted on methadone treatment duration. A liberal p value of < 0.20 in the univariate analyses was chosen to select eligible variables for the multivariate model. A stepwise procedure was used to identify the best model by removing variables one at a time based on a p value of > 0.05 . All analyses were performed using the SPSS v15.0 (SPSS, Inc, Chicago, IL) and Intercooled Stata 12 (StataCorp LP, College Station, TX) software packages using the GLLMM procedure for multinomial analysis.

Results

Sample description

Among the 177 patients with available data at enrollment, 29 (16.4 %) were female and median [IQR] age was 32 [27–38] years (Table 1). At the baseline visit, i.e.

before starting methadone treatment, half of the patients (50.9 %) were employed and two thirds (62.1 %) rented or owned their home. With respect to psychiatric comorbidities, at the baseline visit, 39.4 % of the patients had depressive symptoms, 30.9 % had risk of suicide, and 32.2 % had ADHD symptoms. Sixty-two percent of the patients had at least one withdrawal symptom at the baseline visit. At the baseline visit, 13.9 % reported alcohol dependence, half reported a history of drug injection, and 12.4 % reported overdosing at some point in their life. Only 1.3 % of patients received more than 60 mg of methadone a day. Finally, 25.4 % reported cocaine use at baseline. At baseline, patients who reported using cocaine were significantly more likely to be unemployed, to have depressive symptoms, a risk of suicide, a history of drug injection and ADHD symptoms, than those who did not use cocaine (Table 1).

Cocaine use during the first 12 months of methadone maintenance treatment

Figure 1 shows the percentage of patients reporting cocaine use at each visit: baseline (M0), M6, and M12. At the baseline visit, of those patients who had data on cocaine use ($n = 177$), 25.4 % reported using cocaine during the previous month, 14.7 % occasionally and 10.7 % regularly. The M6 visit showed a decrease in cocaine use with only 15.6 % patients reporting they used it (8.4 % occasionally and 7.1 % regularly) and this decrease remained stable at M12 with 13.6 % (7.1 % occasionally and 6.5 % regularly). Compared with enrolment therefore, the proportion of patients reporting cocaine use decreased significantly at the M6 and M12 visits (Table 2). Among the 45 cocaine users at baseline, 8 were lost to follow-up, 10 continued to use cocaine, and 27 stopped using it.

Factors associated with occasional cocaine use

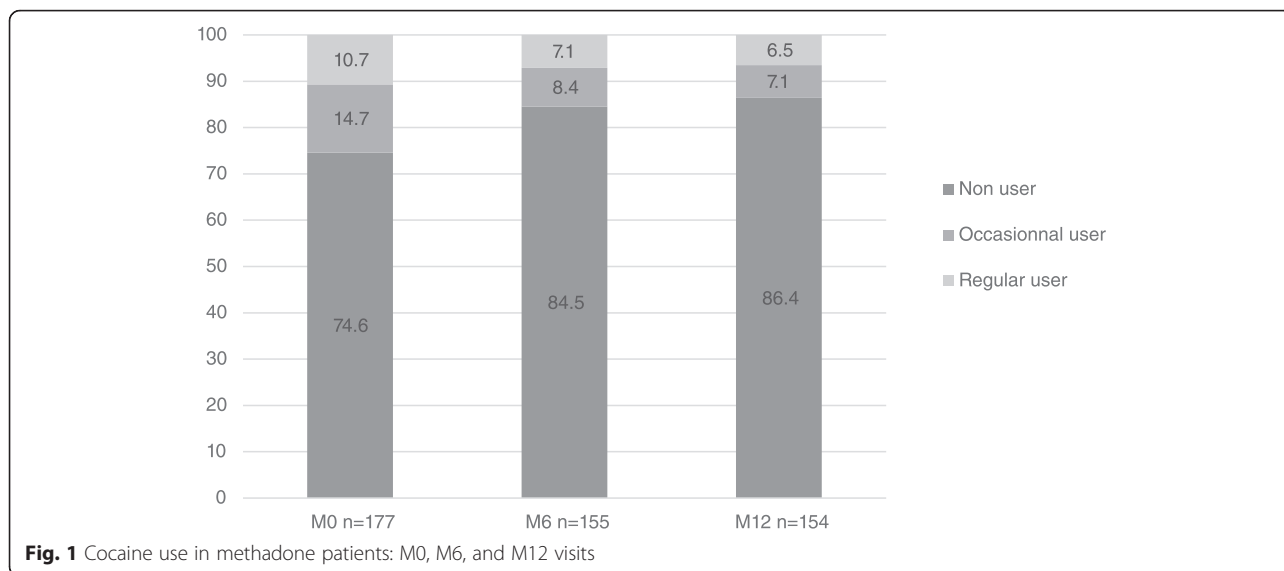
Univariate analysis (Table 2) highlighted the eligibility of several variables for the multivariate model ($p < 0.20$). First, no socio-demographic variable was associated with occasional cocaine use. A history of drug injection, current opioid injection, depressive symptoms, and alcohol dependence were considered eligible variables to explain occasional cocaine use. A significant decrease in occasional cocaine use was observed as the duration of methadone treatment increased.

After multivariate analysis (Table 3), two variables remained positively associated with the outcome: a longer duration on methadone treatment (at the M12 visit only) was associated with less occasional cocaine use and current opioid injection was associated with higher risk of being occasional cocaine user. Those who had depressive symptoms were slightly more likely to be occasional cocaine users.

Table 1 Description of the sample: socio-demographic and behavioral characteristics at baseline

	No cocaine use (n = 132)	Occasional cocaine use (n = 26)	Regular cocaine use (n = 19)	Total (n = 177)	p value ^a
Gender, n (%)					
Male	110 (83.33)	23 (88.46)	15 (78.95)	148 (83.62)	0.70
Female	22 (16.67)	3 (17.54)	4 (21.05)	29 (16.38)	
Methadone induction arm					
Specialized centers	31 (23.5)	7 (26.9)	4 (21.1)	42 (23.7)	0.89
Primary care	101 (76.5)	19 (73.1)	15 (78.9)	135 (76.3)	
Age ^b , median (IQR)	32 (27–38)	30 (27–37)	32 (27–40)	32 (27–38)	0.77
Employed, n (%)					
No	55 (44.35)	15 (57.69)	13 (68.42)	83 (49.11)	0.09
Yes	69 (55.65)	11 (42.31)	6 (31.58)	86 (50.89)	
Home owner or renter, n (%)					
No	43 (34.68)	10 (38.46)	11 (57.89)	64 (37.87)	0.15
Yes	81 (65.32)	16 (61.54)	8 (42.11)	105 (62.13)	
Depressive symptoms, n (%) ^c					
No	80 (66.67)	12 (46.15)	8 (42.11)	100 (60.61)	0.03
Yes	40 (33.33)	14 (53.85)	11 (57.89)	65 (39.39)	
Risk of suicide, n (%) ^d					
No	79 (73.15)	16 (69.57)	8 (44.44)	103 (69.13)	0.05
Yes	29 (26.85)	7 (30.43)	10 (55.56)	46 (30.87)	
ADHD symptoms, n (%)					
No	76 (70.37)	17 (73.91)	8 (44.44)	101 (67.79)	0.07
Yes	32 (29.63)	6 (26.09)	10 (55.56)	48 (32.21)	
Previous drug overdose, n (%)					
No	117 (88.64)	22 (84.62)	16 (84.21)	155 (87.57)	0.64
Yes	15 (11.36)	4 (15.38)	3 (15.79)	22 (12.43)	
History of drug injection, n (%)					
No	66 (53.66)	12 (46.15)	5 (26.32)	83 (49.40)	0.08
Yes	57 (46.34)	14 (53.85)	14 (73.68)	85 (50.60)	
Age at first drug use, median (IQR) ^b	18 (17–22)	18 (16–21)	18 (17–19)	18 (17–21)	0.23
At least one withdrawal symptom, n (%)					
No	49 (42.24)	6 (25.00)	5 (26.32)	60 (37.74)	0.16
Yes	67 (57.76)	18 (75.00)	14 (73.68)	99 (62.26)	
Methadone dose >60 mg, n (%)					
No	110 (99.10)	20 (100.00)	17 (94.44)	147 (98.66)	0.25
Yes	1 (0.90)	0	1 (5.6)	2 (1.34)	
Alcohol dependence, n (%) ^{a,e}					
No	108 (88.52)	21 (84.00)	14 (73.68)	143 (86.14)	0.21
Yes	14 (11.48)	4 (16.00)	5 (26.32)	23 (13.86)	

^aChi-squared or exact Fisher test/Kruskal–Wallis^bIn years^cCES-D score >17 for males and >23 for females^dBeck ≥9^eAUDIT score ≥13



Factors associated with regular cocaine use

For regular cocaine use, univariate analysis highlighted the same variables as those for occasional cocaine use. However, additional variables were associated with regular cocaine use. With respect to socio-demographic variables, regular cocaine users were less likely to be owners or renters of their house and to be employed. In addition, perceiving methadone dose as too low, suicidal risk and ADHD symptoms were also eligible to enter the multivariate model. After the latter was analyzed, three variables remained associated with regular cocaine use: depressive symptoms, ADHD symptoms and current opioid injection. It is worth noting that time on methadone was not associated with a reduction in regular cocaine use.

The methadone induction arm (specialized center versus primary care) was not associated with either occasional or regular cocaine use.

Discussion

Our results showed that cocaine use was highly prevalent among opioid-dependent individuals initiating methadone maintenance treatment, with almost one third of the sample reporting it at baseline. Furthermore, depressive symptoms, ADHD disorders, and current opioid injection were the main predictors of regular cocaine use. These results may have important repercussions on clinical management of these patients.

Our results about cocaine use prevalence are in line with other studies where 30 to 50 % of opioid-dependent individuals seeking care for opioid dependence used cocaine concomitantly [32–34]. More specifically, in our study, at baseline, 14.7 and 10.7 % of the study sample were occasional and regular

cocaine users, respectively. This is in line with other findings [35].

Cocaine use decreased during the 12-month methadone maintenance treatment (MMT) follow-up, with 27 (15 %) patients stopping use altogether. This result corroborates findings in other studies [9, 36] and suggests that despite treatment, certain cocaine users do not reduce their cocaine consumption. In turn, this may act as a barrier to MMT optimization. Accordingly, these vulnerable patients deserve special investigation, particularly those who continue to use cocaine regularly.

To better understand patterns of cocaine use in MMT patients, we separated occasional users from regular users at all visits and studied the factors associated with occasional and regular cocaine use over 12 months of follow-up. Univariate analyses revealed that patients with the most severe characteristics, that is to say, users with a longer history of drug use, those with a history of injecting practices and polydrug users, were more likely to use cocaine during MMT.

One of the main results from the present study is that ADHD was associated with regular cocaine use. It is known that ADHD is a common psychiatric comorbidity among cocaine-dependent patients [37], as cocaine may be used as a means of self-medication for the disease [38]. Furthermore, it is well known that substance use and ADHD are closely correlated [39]. Consequently, screening for ADHD at methadone initiation and appropriate clinical management during methadone treatment, together with the monitoring of cocaine use, can potentially help patients to improve their response and reduce overdose risks [13, 14]. Few clinical responses currently exist for cocaine dependence [40] and cocaine use during MMT [41–43]. Standard treatment includes

Table 2 Factors associated with cocaine use: logistic mixed model adjusted for follow-up visit

	Number of visits (%) or median (IQR)	Number of patients	Occasional cocaine use vs no use		Regular cocaine use vs no use	
			OR [95 % CI]	p	OR [95 % CI]	p
Follow-up						
M0	177 (36.4)	177 (96.7)	1		1	
M6	155 (31.9)	155 (84.7)	0.35 (0.15–0.82)	0.02	0.41 (0.16–1.01)	0.05
M12	154 (31.7)	154 (84.2)	0.28 (0.12–0.69)	0.01	0.35 (0.14–0.90)	0.03
Methadone induction arm						
Specialized centers	130 (20.5)	43 (23.4)	1		1	
Primary care	503 (79.5)	141 (76.6)	0.87 (0.28–2.69)	0.80	1.28 (0.37–4.40)	0.70
Gender						
Male	411 (84.6)	154 (84.2)	1		1	
Female	75 (15.4)	29 (15.8)	0.49 (0.12–2.10)	0.33	1.10 (0.29–4.23)	0.89
Age ^a	32 (27–39)		0.89 (0.46–1.73)	0.74	1.03 (0.52–2.02)	0.94
Employed						
No	131 (41.7)	99 (56.9)	1		1	
Yes	183 (58.3)	122 (70.1)	1.02 (0.43–2.39)	0.97	0.52 (0.21–1.30)	0.16
Home owner or renter						
No	107 (34.1)	76 (43.7)	1		1	
Yes	207 (65.9)	128 (73.6)	0.77 (0.31–1.92)	0.57	0.33 (0.13–0.86)	0.02
Previous drug overdose						
No	429 (88.3)	161 (88.0)	1		1	
Yes	57 (11.7)	22 (12.0)	2.07 (0.51–8.33)	0.31	2.30 (0.55–9.57)	0.25
History of drug injection						
No	244 (51.6)	88 (50.6)	1		1	
Yes	229 (48.4)	86 (49.4)	1.99 (0.75–5.30)	0.17	3.98 (1.39–11.37)	0.01
Age at first drug use ^a	18 (17–21)		0.92 (0.83–1.02)	0.12	0.86 (0.75–0.98)	0.03
Methadone dose >60 mg						
No	305 (66.7)	164 (92.7)	1		1	
Yes	152 (33.3)	84 (47.5)	1.37 (0.51–3.68)	0.54	0.82 (0.29–2.32)	0.71
Perceiving methadone dose to be too low						
No	265 (94.3)	151 (96.8)	1		1	
Yes	16 (5.7)	13 (8.3)	1.23 (0.08–18.09)	0.88	6.06 (0.68–53.78)	0.11
Depressive symptoms ^b						
No	310 (68.3)	142 (81.6)	1		1	
Yes	144 (31.7)	86 (49.4)	2.12 (0.88–5.11)	0.09	4.32 (1.71–10.92)	0.002
Risk of suicide ^c						
No	279 (72.7)	133 (81.1)	1		1	
Yes	105 (27.3)	69 (42.1)	1.62 (0.68–3.86)	0.28	3.93 (1.59–9.76)	0.003
ADHD symptoms						
No	304 (79.0)	148 (89.7)	1		1	
Yes	81 (21.0)	59 (35.8)	1.48 (0.55–4.01)	0.44	6.17 (2.34–16.22)	<0.001
At least 1 withdrawal symptom						
No	296 (72.0)	148 (87.1)	1		1	
Yes	115 (28.0)	103 (60.6)	1.42 (0.52–3.89)	0.50	1.51 (0.52–4.38)	0.45
Alcohol dependence ^d						

Table 2 Factors associated with cocaine use: logistic mixed model adjusted for follow-up visit (*Continued*)

No	274 (88.1)	157 (91.3)	1		1	
Yes	37 (11.9)	27 (15.7)	3.08 (0.88–10.80)	0.08	4.10 (1.15–14.61)	0.03
Opiate injection ^e						
No	414 (91.4)	164 (95.3)	1		1	
Yes	39 (8.6)	24 (13.8)	6.06 (1.29–28.44)	0.02	12.42 (2.72–56.61)	0.001

Univariate analyses: *N* = 486 visits; 183 persons^aIn years^bCES-D score >17 for males and >23 for females^cBeck ≥ 9 ^dAUDIT score ≥ 13 ^eDuring the previous 4 weeks

contingency management and cognitive behavioral therapy [6, 44, 45]. However, findings regarding the efficacy of psychotherapy for cocaine abuse in MMT patients are mixed. For example, in one article, counseling therapy for cocaine users in a population of methadone-maintained patients was not effective [46], while another article showed that CBT may indeed have a positive impact [6]. More generally, long-term studies highlight the weakness in providing only CBT [47].

We also found that depressive symptoms were associated with cocaine use. The relationship between cocaine use and depression is complex, and the causality direction is unclear. It is known that cocaine-dependent individuals are more likely to have depressive symptoms, depression perhaps being a preliminary condition for cocaine dependence [48]. However, one recent study has also highlighted that depression may also be a consequence of cocaine use [49]. In any case, depression should be diagnosed and treated in a timely fashion in

order to reduce its negative impact on the effectiveness of OMT. Although it has been shown that certain antidepressants are not effective in reducing cocaine use in cocaine and opioid co-dependent patients [50, 51], some studies have demonstrated that access to adequate care for depression in cocaine-dependent patients may lead to decreased cocaine use [52, 53].

In addition, persistent opiate injection during methadone maintenance (i.e., non-response to opioid dependence treatment) was associated with persistent (both occasional and regular) cocaine use. This suggests that methadone treatment (at the prescribed dosages) may be an inadequate opioid dependence treatment; and that for such patients, methadone treatment has a negligible impact on cocaine use. In addition, two associations with regular cocaine use found in the univariate analyses may help us better understand the relationship between heroin injection and regular cocaine use: reporting withdrawal symptoms and a tendency to perceive methadone

Table 3 Factors independently associated with occasional and regular cocaine use: logistic mixed model

	Occasional cocaine use vs no use		Regular cocaine use vs no use	
	ORa [95 % CI]	<i>p</i>	ORa [95 % CI]	<i>p</i>
Follow-up				
M0	1		1	
M6	0.48 [0.18–1.27]	0.14	0.77 [0.26–2.26]	0.63
M12	0.33 [0.12–0.95]	0.04	0.38 [0.11–1.32]	0.13
Depressive symptoms ^b				
No	1			
Yes	2.41 [0.92–6.31]	0.07	3.49 [1.21–10.04]	0.02
ADHD symptoms				
No	1			
Yes	1.40 [0.47–4.18]	0.55	5.23 [1.74–15.71]	0.003
Opiate injection ^a				
No	1			
Yes	5.73 [1.26–25.99]	0.02	8.98 [1.87–43.22]	0.01

Multivariate analyses: *N* = 353 visits; 156 persons^aDuring the previous 4 weeks^bCES-D score >17 for males and >23 for females

dose as too low were both associated with regular cocaine use (but not occasional cocaine use). This may also suggest that cocaine has a pharmacokinetic impact on methadone which decreases the effect of the latter [12]. To conclude, persistent heroin injection during MMT may be explained by underdosing of methadone [54] and/or inappropriate treatment for opioid dependence [55].

Finally, it is interesting to note that a longer time on methadone (measured in terms of each follow-up visit) was associated with reduced occasional cocaine use but no such association was seen for regular cocaine use. This suggests that higher levels of cocaine use in methadone-maintained patients may not be influenced by medical follow-up for opioid dependence.

Some limitations of this study have to be acknowledged. First, we did not identify cocaine dependence in our sample. This data would have been interesting to analyze as cocaine-dependent patients may have been those who did not respond to MMT. The role of cocaine use in ADHD should be investigated more thoroughly, as the ASRS-v1.1 recently reported low specificity in the detection of ADHD among populations with substance use disorders [56]. The second limitation is that the validity of self-reported behaviors is often questioned due to the risk of underreporting linked to social desirability bias. However, this effect modifies OR estimates in a conservative manner. Furthermore, the reliability of self-reports based on using the Opiate Treatment Index (OTI) treatment questionnaire in drug-using populations has already been demonstrated [57].

Conclusions

As suggested by previous findings, in our study, time on MMT had a positive impact on occasional cocaine users. However, regular cocaine use was not influenced by MMT duration but was associated with psychiatric comorbidities (ADHD and depression) and more severe addictive profiles (opioid injection). Early screening of these disorders and prompt tailored pharmacological and behavioral interventions have the potential to reduce cocaine use and to improve response to MMT.

Abbreviations

ADHD: attention deficit/hyperactivity disorder; ASRS: Adult ADHD Self-Report Scale-Version; AUDIT: alcohol use disorders test; BHS: Beck Hopelessness Scale; CATI: computerized assisted phone interviews; CBT: cognitive behavioral therapy; CES-D: Center for Epidemiologic Studies-Depression; IQR: interquartile range; MMT: methadone maintenance treatment; OMT: opioid maintenance treatment; OOWS: Objective Opioid Withdrawal Scale; OTI: Opiate Treatment Index.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

RP designed the study, supervised the analyses, and wrote the first draft. LC coordinated and conducted the statistical analysis, interpreted the results, and contributed to the draft of the manuscript. ML contributed to the design of the study and revised the final version of the manuscript. MF contributed to the statistical analysis in particular for methodological issues. MM, MG, and SB contributed to the design of the study and to the socio-behavioral data collection. MA designed the study and contributed to the data collection. CMP designed the study, supervised the analyses, and contributed to the different versions of the manuscript. All authors contributed to and approved the current version of the manuscript.

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References

- Connock M, Juarez-Garcia A, Jowett S, Frew E, Liu Z, Taylor RJ, et al. Methadone and buprenorphine for the management of opioid dependence: a systematic review and economic evaluation. *Health Technol Assess.* 2007;11(9):1–171. iii-iv.
- Magura S, Kang SY, Nwakeze PC, Demsky S. Temporal patterns of heroin and cocaine use among methadone patients. *Subst Use Misuse.* 1998; 33(12):2441–67.

3. Savant JD, Barry DT, Cutter CJ, Joy MT, Dinh A, Schottenfeld RS et al. Prevalence of mood and substance use disorders among patients seeking primary care office-based buprenorphine/naloxone treatment. *Drug Alcohol Depend.* 2012. doi:10.1016/j.drugalcdep.2012.06.020.
4. Hubbard RL, Craddock SG, Anderson J. Overview of 5-year followup outcomes in the drug abuse treatment outcome studies (DATOS). *J Subst Abuse Treat.* 2003;25(3):125–34.
5. Senbanjo R, Wolff K, Marshall EJ, Strang J. Persistence of heroin use despite methadone treatment: poor coping self-efficacy predicts continued heroin use. *Drug Alcohol Rev.* 2009;28(6):608–15. doi:10.1111/j.1465-3362.2009.00064.x.
6. Tzilos GK, Rhodes GL, Ledgerwood DM, Greenwald MK. Predicting cocaine group treatment outcome in cocaine-abusing methadone patients. *Exp Clin Psychopharmacol.* 2009;17(5):320–5. doi:10.1037/a0016835.
7. Dutra L, Stathopoulou G, Basden SL, Leyro TM, Powers MB, Otto MW. A meta-analytic review of psychosocial interventions for substance use disorders. *Am J Psychiatry.* 2008;165(2):179–87. doi:10.1176/appi.ajp.2007.06111851.
8. Peles E, Schreiber S, Adelson M. Factors predicting retention in treatment: 10-year experience of a methadone maintenance treatment (MMT) clinic in Israel. *Drug Alcohol Depend.* 2006;82(3):211–7.
9. Bravo MJ, Llorens N, Barrio G, Brugal MT, Santos S, Sordo L, et al. Methadone maintenance treatment: a protective factor for cocaine injection in a street-recruited cohort of heroin users. *Drug Alcohol Depend.* 2010; 112(1-2):62–8. doi:10.1016/j.drugalcdep.2010.05.014.
10. Salamina G, Diecidue R, Vigna-Taglianti F, Jarre P, Schifano P, Bargagli AM, et al. Effectiveness of therapies for heroin addiction in retaining patients in treatment: results from the VEdETE study. *Subst Use Misuse.* 2010;45(12): 2076–92. doi:10.3109/10826081003791932.
11. Sullivan LE, Moore BA, O'Connor PG, Barry DT, Chawarski MC, Schottenfeld RS, et al. The association between cocaine use and treatment outcomes in patients receiving office-based buprenorphine/naloxone for the treatment of opioid dependence. *Am J Addict.* 2010;19(1):53–8. doi:10.1111/j.1521-0391.2009.00003.x.
12. McCance-Katz EF, Jatlow P, Rainey PM. Effect of cocaine use on methadone pharmacokinetics in humans. *Am J Addict.* 2010;19(1):47–52. doi:10.1111/j.1521-0391.2009.00009.x.
13. Albion C, Shkrum M, Cairns J. Contributing factors to methadone-related deaths in Ontario. *Am J Forensic Med Pathol.* 2010;31(4):313–9. doi:10.1097/PAF.0b013e3181ca4b1e.
14. Laberke PJ, Bartsch C. Trends in methadone-related deaths in Zurich. *Int J Legal Med.* 2010;124(5):381–5. doi:10.1007/s00414-010-0442-8.
15. Visconti AJ, Santos GM, Lemos NP, Burke C, Coffin PO. Opioid overdose deaths in the city and county of San Francisco: prevalence, distribution, and disparities. *J Urban Health.* 2015;92(4):758–72. doi:10.1007/s11524-015-9967-y.
16. Proctor SL, Copeland AL, Kopak AM, Hoffmann NG, Herschman PL, Polukhina N. Predictors of patient retention in methadone maintenance treatment. *Psychol Addict Behav.* 2015. doi:10.1037/adb0000090.
17. Leventhal AM, Mooney ME, DeLaune KA, Schmitz JM. Using addiction severity profiles to differentiate cocaine-dependent patients with and without comorbid major depression. *Am J Addict.* 2006;15(5):362–9. doi:10.1080/10550490600860148.
18. Dunne EM, Hearn LE, Rose JJ, Latimer WW. ADHD as a risk factor for early onset and heightened adult problem severity of illicit substance use: an accelerated gateway model. *Addict Behav.* 2014;39(12):1755–8. doi:10.1016/j.addbeh.2014.07.009.
19. Rubio G, Manzanares J, Jimenez M, Rodriguez-Jimenez R, Martinez I, Iribarren MM, et al. Use of cocaine by heavy drinkers increases vulnerability to developing alcohol dependence: a 4-year follow-up study. *J Clin Psychiatry.* 2008;69(4):563–70.
20. Marsch LA, Stephens MA, Mudric T, Strain EC, Bigelow GE, Johnson RE. Predictors of outcome in LAAM, buprenorphine, and methadone treatment for opioid dependence. *Exp Clin Psychopharmacol.* 2005;13(4):293–302.
21. Delavenne H, Ballon N, Charles-Nicolas A, Garcia FD, Thibaut F, Lacoste J. Attention deficit hyperactivity disorder is associated with a more severe pattern of cocaine consumption in cocaine users from French West Indies. *J Addict Med.* 2011;5(4):284–8. doi:10.1097/ADM.0b013e31821b4038.
22. Herbeck DM, Hser YI, Lu AT, Stark ME, Paredes A. A 12-year follow-up study of psychiatric symptomatology among cocaine-dependent men. *Addict Behav.* 2006;31(11):1974–87. doi:10.1016/j.addbeh.2006.01.013.
23. Roux P, Michel L, Cohen J, Mora M, Morel A, Aubertin JF, et al. Initiation of Methadone in primary care (ANRS-Methaville): a phase III randomized intervention trial. *BMC Public Health.* 2012;12(1):488. doi:10.1186/1471-2458-12-488.
24. Gache P, Michaud P, Landry U, Accietto C, Arfaoui S, Wenger O, et al. The Alcohol Use Disorders Identification Test (AUDIT) as a screening tool for excessive drinking in primary care: reliability and validity of a French version. *Alcohol Clin Exp Res.* 2005;29(11):2001–7.
25. Fuhrer R, Rouillon F. La version française de l'échelle CES-D. Description and translation of the autoevaluation scale (in French). *Psychiatrie et Psychobiologie.* 1989;4:163–6.
26. Heatherton TF, Kozlowski LT, Frecker RC, Fagerstrom KO. The Fagerstrom Test for Nicotine Dependence: a revision of the Fagerstrom Tolerance Questionnaire. *Br J Addict.* 1991;86(9):1119–27.
27. Darke S, Hall W, Wodak A, Heather N, Ward J. Development and validation of a multi-dimensional instrument for assessing outcome of treatment among opiate users: the Opiate Treatment Index. *Br J Addict.* 1992;87(5): 733–42.
28. Daigre C, Roncero C, Rodriguez-Cintas L, Ortega L, Lligona A, Fuentes S, et al. Adult ADHD screening in alcohol-dependent patients using the Wender-Utah Rating Scale and the adult ADHD Self-Report Scale. *J Atten Disord.* 2015;19(4):328–34. doi:10.1177/1087054714529819.
29. van de Glind G, van den Brink W, Koeter MW, Carpentier PJ, van Emmerik-van Oortmerssen K, Kaye S, et al. Validity of the Adult ADHD Self-Report Scale (ASRS) as a screener for adult ADHD in treatment seeking substance use disorder patients. *Drug Alcohol Depend.* 2013;132(3):587–96. doi:10.1016/j.drugalcdep.2013.04.010.
30. Perry AE, Olason DT. A new psychometric instrument assessing vulnerability to risk of suicide and self-harm behaviour in offenders: Suicide Concerns for Offenders in Prison Environment (SCOPE). *Int J Offender Ther Comp Criminol.* 2008;53(4):385–400. doi:10.1177/0306624X08319418.
31. Handelsman L, Cochrane KJ, Aronson MJ, Ness R, Rubinstein KJ, Kanof PD. Two new rating scales for opiate withdrawal. *Am J Drug Alcohol Abuse.* 1987;13(3):293–308.
32. Raffa JD, Grebely J, Tossonian H, Wong T, Viljoen M, Khara M, et al. The impact of ongoing illicit drug use on methadone adherence in illicit drug users receiving treatment for HIV in a directly observed therapy program. *Drug Alcohol Depend.* 2007;89(2-3):306–9.
33. Gryczynski J, Schwartz RP, Salkever DS, Mitchell SG, Jaffe JH. Patterns in admission delays to outpatient methadone treatment in the United States. *J Subst Abuse Treat.* 2011;41(4):431–9. doi:10.1016/j.jsat.2011.06.005.
34. Cone EJ. Oral fluid results compared to self reports of recent cocaine and heroin use by methadone maintenance patients. *Forensic Sci Int.* 2012; 215(1-3):88–91. doi:10.1016/j.forsciint.2011.01.046.
35. Peles E, Kreek MJ, Kellogg S, Adelson M. High methadone dose significantly reduces cocaine use in methadone maintenance treatment (MMT) patients. *J Addict Dis.* 2006;25(1):43–50. doi:10.1300/J069v25n01_07.
36. Dobler-Mikola A, Hattenschwiler J, Meili D, Beck T, Boni E, Modestin J. Patterns of heroin, cocaine, and alcohol abuse during long-term methadone maintenance treatment. *J Subst Abuse Treat.* 2005;29(4): 259–65. doi:10.1016/j.jsat.2005.08.002.
37. Daigre C, Roncero C, Grau-Lopez L, Martinez-Luna N, Prat G, Valero S, et al. Attention deficit hyperactivity disorder in cocaine-dependent adults: a psychiatric comorbidity analysis. *Am J Addict.* 2013;22(5):466–73. doi:10.1111/j.1521-0391.2013.12047.x.
38. Gudjonsson GH, Sigurdsson JF, Sigfusdottir ID, Young S. An epidemiological study of ADHD symptoms among young persons and the relationship with cigarette smoking, alcohol consumption and illicit drug use. *J Child Psychol Psychiatry.* 2012;53(3):304–12.
39. Wilson JJ, Levin FR. Attention deficit hyperactivity disorder (ADHD) and substance use disorders. *Curr Psychiatry Rep.* 2001;3(6):497–506.
40. Karila L, Petit A, Lowenstein W, Reynaud M. Diagnosis and consequences of cocaine addiction. *Curr Med Chem.* 2012;19(33):5612–8.
41. Kosten TR, Wu G, Huang W, Harding MJ, Hamon SC, Lappalainen J, et al. Pharmacogenetic randomized trial for cocaine abuse: disulfiram and dopamine beta-hydroxylase. *Biol Psychiatry.* 2013;73(3):219–24. doi:10.1016/j.biopsych.2012.07.011.
42. Carroll KM, Nich C, Shi JM, Eagan D, Ball SA. Efficacy of disulfiram and Twelve Step Facilitation in cocaine-dependent individuals maintained on methadone: a randomized placebo-controlled trial. *Drug Alcohol Depend.* 2012;126(1-2):224–31. doi:10.1016/j.drugalcdep.2012.05.019.

43. Oliveto A, Poling J, Mancino MJ, Feldman Z, Cubells JF, Pruzinsky R, et al. Randomized, double blind, placebo-controlled trial of disulfiram for the treatment of cocaine dependence in methadone-stabilized patients. *Drug Alcohol Depend.* 2011;113(2-3):184–91. doi:10.1016/j.drugalcdep.2010.07.022.
44. Rawson RA, Huber A, McCann M, Shoptaw S, Farabee D, Reiber C, et al. A comparison of contingency management and cognitive-behavioral approaches during methadone maintenance treatment for cocaine dependence. *Arch Gen Psychiatry.* 2002;59(9):817–24.
45. Epstein DH, Hawkins WE, Covi L, Umbricht A, Preston KL. Cognitive-behavioral therapy plus contingency management for cocaine use: findings during treatment and across 12-month follow-up. *Psychol Addict Behav.* 2003;17(1):73–82.
46. Darker C, Sweeney B, El Hassan H, Kelly ASOC, Smyth B, et al. Non-attendance at counselling therapy in cocaine-using methadone-maintained patients: lessons learnt from an abandoned randomised controlled trial. *Ir J Med Sci.* 2012;181(4):483–9. doi:10.1007/s11845-011-0803-4.
47. Penberthy JK, Ait-Daoud N, Vaughan M, Fanning T. Review of treatment for cocaine dependence. *Curr Drug Abuse Rev.* 2010;3(1):49–62.
48. Lopez A, Becona E. Depression and cocaine dependence. *Psychol Rep.* 2007;100(2):520–4.
49. Dakwar E, Nunes EV, Bisaga A, Carpenter KC, Mariani JP, Sullivan MA, et al. A comparison of independent depression and substance-induced depression in cannabis-, cocaine-, and opioid-dependent treatment seekers. *Am J Addict.* 2011;20(5):441–6. doi:10.1111/j.1521-0391.2011.00148.x.
50. Afshar M, Knapp CM, Sarid-Segal O, Devine E, Colaneri LS, Tozier L, et al. The efficacy of mirtazapine in the treatment of cocaine dependence with comorbid depression. *Am J Drug Alcohol Abuse.* 2012;38(2):181–6.
51. Winstanley EL, Bigelow GE, Silverman K, Johnson RE, Strain EC. A randomized controlled trial of fluoxetine in the treatment of cocaine dependence among methadone-maintained patients. *J Subst Abuse Treat.* 2011;40(3):255–64. doi:10.1016/j.jsat.2010.11.010.
52. Oliveto A, Poling J, Mancino MJ, Williams DK, Thostenson J, Pruzinsky R, et al. Sertraline delays relapse in recently abstinent cocaine-dependent patients with depressive symptoms. *Addiction.* 2012;107(1):131–41.
53. Moeller FG, Schmitz JM, Steinberg JL, Green CM, Reist C, Lai LY, et al. Citalopram combined with behavioral therapy reduces cocaine use: a double-blind, placebo-controlled trial. *Am J Drug Alcohol Abuse.* 2007;33(3):367–78.
54. Donny EC, Brasser SM, Bigelow GE, Stitzer ML, Walsh SL. Methadone doses of 100 mg or greater are more effective than lower doses at suppressing heroin self-administration in opioid-dependent volunteers. *Addiction.* 2005;100(10):1496–509.
55. Bond AJ, Reed KD, Beavan P, Strang J. After the randomised injectable opiate treatment trial: post-trial investigation of slow-release oral morphine as an alternative opiate maintenance medication. *Drug Alcohol Rev.* 2012;31(4):492–8.
56. Chiasson JP, Stavro K, Rizkallah E, Lapierre L, Dussault M, Legault L, et al. Questioning the specificity of ASRS-v1.1 to accurately detect ADHD in substance abusing populations. *J Atten Disord.* 2012;16(8):661–3.
57. Darke S. Self-report among injecting drug users: a review. *Drug Alcohol Depend.* 1998;51(3):253–63. discussion 67-8.

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