

Additive effects of childhood abuse and cannabis abuse on clinical expressions of bipolar disorders

M. Aas^{1,2*†}, B. Etain^{3,4,5,6†}, F. Bellivier^{4,5,6,7,8}, C. Henry^{3,4,5,6,9}, T. Lagerberg², A. Ringen², I. Agartz^{1,10}, S. Gard^{5,11}, J.-P. Kahn^{5,12}, M. Leboyer^{3,4,5,6,9}, O. A. Andreassen^{1,2,6} and I. Melle^{1,2}

¹KG Jebsen Centre for Psychosis Research, Institute of Clinical Medicine, University of Oslo, Norway; ²Division of Mental Health and Addiction, Oslo University Hospital, Oslo, Norway; ³AP-HP, Hôpital H. Mondor – A. Chenevier, Pôle de Psychiatrie, Créteil, France; ⁴Inserm, U955, Créteil, France; ⁵Fondation Fondamental, Créteil, France; ⁶ENBREC, European Network of Bipolar Research Expert Centres (ENBREC), Paris, France; ⁷AP-HP, GH Saint-Louis – Lariboisière – Fernand Widal, Pôle Neurosciences, Paris, France; ⁸Université Paris 7, Denis Diderot, Paris, France; ⁹Université Paris Est, Faculté de Médecine, Créteil, France; ¹⁰Department of Psychiatric Research, Diakonhjemmet Hospital, Oslo, Norway; ¹¹Hôpital Charles Perrens, Centre Expert Trouble Bipolaire, Service de Psychiatrie Adulte, Pôle 3-4-7, Bordeaux, France; ¹²Service de Psychiatrie et Psychologie Clinique, Université de Lorraine et CHU de Nancy, Hôpitaux de Brabois, Vandoeuvre Les Nancy, France

Background. Previous studies of bipolar disorders indicate that childhood abuse and substance abuse are associated with the disorder. Whether both influence the clinical picture, or if one is mediating the association of the other, has not previously been investigated.

Method. A total of 587 patients with bipolar disorders were recruited from Norway and France. A history of childhood abuse was obtained using the Childhood Trauma Questionnaire. Diagnosis and clinical variables, including substance abuse, were based on structured clinical interviews (Structured Clinical Interview for DSM-IV Axis I disorders or French version of the Diagnostic Interview for Genetic Studies).

Results. Cannabis abuse was significantly associated with childhood abuse, specifically emotional and sexual abuse ($\chi^2=8.63$, $p=0.003$ and $\chi^2=7.55$, $p=0.006$, respectively). Cannabis abuse was significantly associated with earlier onset of the illness ($z=-4.17$, $p<0.001$), lifetime history of at least one suicide attempt ($\chi^2=11.16$, $p=0.001$) and a trend for rapid cycling ($\chi^2=3.45$, $p=0.06$). Alcohol dependence was associated with suicide attempt ($\chi^2=10.28$, $p=0.001$), but not with age at onset or rapid cycling. After correcting for possible confounders and multiple testing, a trend was observed for an interaction between cannabis abuse and childhood abuse and suicide attempt (logistic regression: $r^2=0.06$, $p=0.039$). Significant additive effects were also observed between cannabis abuse and childhood abuse on earlier age at onset ($p<0.001$), increased rapid cycling and suicide attempt (logistic regression: $r^2=0.03-0.04$, $p<0.001$). No mediation effects were observed; childhood abuse and cannabis abuse were independently associated with the disorder.

Conclusions. Our study is the first to demonstrate significant additive effects, but no mediation effects, between childhood abuse and cannabis abuse on increased clinical expressions of bipolar disorders.

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Key words: Bipolar disorder, cannabis abuse, childhood abuse, clinical characteristics.

Introduction

Bipolar disorders are highly co-morbid with other psychiatric disorders and medical conditions which are linked to both the development of, and the course of, the disease. This is particularly true for bipolar disorders and substance abuse, including alcohol and drug abuse, which have been found to be associated with earlier age at onset (Lin *et al.* 2006; Javard *et al.*

2011; Lagerberg *et al.* 2011), suicide attempts, mixed episodes, as well as greater disability (Agrawal *et al.* 2011). The study by Agrawal *et al.* (2011) showed that patients with a bipolar disorder are 6.8 times more likely to report a lifetime history of cannabis abuse than healthy controls. The findings above are supported by a study by Post (2010) indicating that drug abuse may increase sensitization and vulnerability to recurrent episodes, thus driving illness progression in bipolar disorder. Indeed, a recent study by Pettinati *et al.* (2013) showed that the lifetime prevalence rate of any bipolar disorder combined with any substance use disorder is 47.3%. A recent trial for co-morbid affective mood disorder and alcohol dependence again showed a strong link between these factors, demonstrating that combining treatment of both

* Address for correspondence: M. Aas, Ph.D., KG Jebsen Centre for Psychosis Research, Division of Mental Health and Addiction, Institute of Clinical Medicine, Bygg 49, Ullevål sykehus, PO Box 4956 Nydalen, 0424 Oslo, Norway.

(Email: monica.aas@medisin.uio.no)

† These authors contributed equally as joint first authors.

depression and alcohol dependence reduced depressive symptoms and excessive drinking simultaneously (Pettinati *et al.* 2010).

Individuals reporting substance abuse in adulthood also report high levels of childhood trauma experiences (Fergusson *et al.* 2008; Shin *et al.* 2009). More generally, childhood abuse has also been associated with severe mental disorders, such as schizophrenia and bipolar disorder, both with increased frequency (Read *et al.* 2005; Etain *et al.* 2010; Mondelli *et al.* 2010; Aas *et al.* 2011), and been linked to clinical characteristics of the disorder (Daruy-Filho *et al.* 2011). In bipolar disorders childhood abuse is associated with a more severe clinical expression shown by earlier onset of the illness (Garno *et al.* 2005), a rapid cycling course (Garno *et al.* 2005), more psychotic features (Bebbington *et al.* 2004; Janssen *et al.* 2004; Shevlin *et al.* 2007), a higher number of lifetime mood episodes (Nemeroff, 2004; Brown *et al.* 2005) as well as suicide ideation and attempts (Alvarez *et al.* 2011), though little is known about the mechanisms behind these associations.

As shown above, previous studies of bipolar disorders indicate that both childhood abuse and substance abuse are associated with the disorder, demonstrated by the high frequency of co-morbidity for both, as well as increasing clinical severity. The question is whether they both influence the clinical picture in bipolar disorders, or whether one is mediating the association of the other. That individuals reporting childhood abuse tend to show an increased risk of substance abuse in adulthood compared with individuals without childhood abuse (Fergusson *et al.* 2008; Shin *et al.* 2009) indicates possible interaction effects. It is also possible that associations between cannabis abuse and clinical characteristics observed in bipolar disorder, such as earlier age at onset, suicide attempts, and more severe illness (Agrawal *et al.* 2011; Lagerberg *et al.* 2011) are not directly linked to cannabis, but rather to the underlying effect of childhood abuse on clinical characteristics. Indeed, a study in patients with schizophrenia shows that the relationship between cannabis abuse and earlier age at onset disappeared when correcting for childhood trauma, indicating that childhood trauma, and not cannabis abuse, is driving this association (Houston *et al.* 2011).

To complicate this further, it is hypothesized that cannabis abuse elicits psychotic symptoms in people who otherwise would not develop a psychotic disorder. This group of people could be regarded as a selection of patients with good pre-morbid function (Ringen *et al.* 2013). This group would probably have a better prognosis than the group of patients with other causes of their symptoms, e.g. neurodevelopmental abnormalities (Ringen *et al.* 2008). Cannabis

abuse may, therefore, first reduce age at onset and worsen clinical characteristics in already vulnerable patients, and, second, increase the risk of developing the illness in patients with relatively good pre-morbid function and/or possible less genetic loading for the disorder.

The current study is, to our knowledge, the first to investigate the relationship between substance abuse (cannabis and alcohol), childhood abuse and clinical expressions in bipolar disorder. Our hypotheses are as follows: (1) childhood abuse will be related to increased substance abuse in bipolar disorder; (2) substance abuse will be associated with increased clinical expression, such as earlier age at onset, rapid cycling, lifetime history of at least one suicide attempt, and mood episodes; (3) interaction, or additive effects, will be observed between childhood abuse, substance abuse and clinical expressions, in the direction of earlier age at onset, history of suicide attempts, rapid cycling and more mood episodes in patients with both childhood abuse experiences and cannabis abuse; and (4) mediation effects will be observed between substance abuse, childhood trauma and clinical expression of bipolar disorders.

Method

Participants

A total of 418 patients from France were included at three French university affiliated departments of psychiatry (Paris/Créteil, Bordeaux and Nancy). Also, 169 patients from Norway were included as part of an ongoing study of severe psychiatric disorders (Thematically Organized Psychosis Study) and recruited from psychiatric in- and out-patient units at three major hospitals in Oslo. For inclusion in the study, all patients had to have a diagnosis of a bipolar disorder (type I, II or not otherwise specified) according to Diagnostic and Statistical Manual of Mental Disorders, fourth edition (DSM-IV) criteria (APA, 1994). The Norwegian project was approved by the Regional Committee for Medical Research Ethics and the Data Inspectorate. The local institutional review boards approved the French study. Written informed consent for participation in the study was obtained from all participants.

The total sample is presented in Table 1. Before merging the two samples for the analyses, comparison analyses were conducted between the two sites (see online Supplementary Table S1). A total of 13% reported cannabis abuse, and there were no differences between the two sites. Also, no differences between the groups were observed for cannabis dependence. The two samples were similar for Childhood Trauma

Table 1. Patients' demographic and clinical characteristics (n=587)

Characteristics	
Mean age, years (s.d.)	40.6 (13.6)
Gender, <i>n</i>	
Male	234
Female	353
Ethnicity, <i>n</i> (%)	
Caucasian	567 (96.6)
Other	20 (3.4)
Mean age at onset, years (s.d.)	24.2 (10.1)
Bipolar I, <i>n</i> (%)	425 (72.4)
Bipolar II, <i>n</i> (%)	126 (21.5)
Bipolar not otherwise specified, <i>n</i> (%)	36 (6.1)
Suicide attempt, <i>n</i> (%)	211 (36.4)
Rapid cycling, <i>n</i> (%)	114 (20.8)
Mean number of depressive episodes (s.d.)	4.8 (5.5)
Mean number of manic and hypomanic episodes (s.d.)	4.61 (6.0)
Psychotic episodes, <i>n</i> (%) ^a	247 (49.4)
Cannabis abuse, <i>n</i> (%)	75 (12.9)
Alcohol dependence, <i>n</i> (%)	53 (9.1)
Childhood trauma severity (moderate to severe intensity) ^b	
Physical abuse, yes (%)	55 (9.5)
Sexual abuse, yes (%)	104 (18.0)
Emotional abuse, yes (%)	156 (26.9)
Physical neglect, yes (%)	73 (12.5)
Emotional neglect, yes (%)	180 (31.3)

s.d., Standard deviation.

^a Having at least one psychotic episode during the course of the illness.^b 98.8% (*n*=580) of the patients completed the physical abuse subscale; 99.0% (*n*=581) completed the sexual abuse subscale; 98.8% (*n*=580) completed the emotional abuse subscale; 98.3% (*n*=577) completed the physical neglect subscale; 98.8% (*n*=580) completed the emotional neglect subscale.

Questionnaire (CTQ) total score (France: 41.74, s.d.=12.88 *versus* Norway 44.03, s.d.=16.43, *z*=0.88, *p*=0.37). Differences between the samples were found concerning alcohol dependence (Norway: 14% *versus* France: 7%). The French sample was also older and had a longer duration of illness than the Norwegian sample (see online Supplementary Table S1). The difference regarding lifetime alcohol dependence between the samples remained significant even when age at interview was used as a covariate. χ^2 Analysis showed no significant association between cannabis abuse (yes, no) and diagnosis (bipolar I, bipolar II and bipolar not otherwise specified) [$\chi^2=2.29$, degrees of freedom (df)=2, *p*=0.32]. Lastly, no significant difference was observed between abuse-dose score

(no trauma, one trauma, two traumas, three traumas) and subtypes of bipolar disorders ($\chi^2=7.38$, df=6, *p*=0.29).

Clinical assessment

In each country, clinical assessment was carried out by trained psychiatrists, medical doctors and clinical psychologists. The French patients were interviewed with the French version of the Diagnostic Interview for Genetic Studies (DIGS; Nurnberger *et al.* 1994), which provides lifetime DSM-IV Axis I diagnoses. French patients were euthymic at inclusion (i.e. having Mania Rating Scale and Montgomery–Asberg Depression Rating Scale scores of no more than 5). Several clinical variables were collected with the DIGS: lifetime substance abuse and dependence, history of rapid cycling (defined as in the DSM-IV criteria) and history of mixed episodes during the course of the disorder. 'Suicide attempt' was measured by the presence of at least one experience of lifetime suicide attempt.

A similar approach was taken in the Norwegian sample using the Structured Clinical Interview for DSM-IV Axis I disorders (SCID-I). It is an ongoing debate whether borderline personality disorder belongs to the bipolar spectrum (Coulston *et al.* 2012). We know from the literature that borderline patients, similar to bipolar patients, report high levels of childhood trauma (Coulston *et al.* 2012), as well as emotional dysregulation. In our study, the patients had to meet a DSM-IV SCID-I diagnosis of bipolar disorder to be included in the study, independent of possible personality disorders they may also have had.

For both samples, age at onset was defined as the age when the subject first met DSM-IV criteria for a major depressive, manic, hypomanic, or mixed episode. Duration of illness was calculated by current age at time of assessment minus age at onset.

Comparisons of DIGS and DSM criteria have shown excellent reliability scores. A two-phase test-retest (within-site, between-site) reliability study was carried out for DSM, third edition revised (DSM-III-R) criteria-based major depression, bipolar disorder, schizophrenia and schizo-affective disorder. Reliabilities using algorithms were excellent (0.73–0.95), except for schizo-affective disorder, for which disagreement on estimates of duration of mood syndromes relative to psychosis reduced reliability (Nurnberger *et al.* 1994). As we only investigated bipolar disorders this was not relevant for our study. Although the study by Nurnberger *et al.* (1994) compared DIGS with an earlier version of the DSM than the one we used in this study (DSM-III-R *versus* DSM-IV) there is no reason to think that the reliability score should be any different for DSM-IV used in our study.

CTQ

To measure childhood adverse events we used the CTQ, a retrospective questionnaire enquiring about traumatic experiences in childhood with answers ranging from 'never true', 'rarely true', 'sometimes true', 'often true' to 'very true', yielding scores for a total score as well as five subscores: physical, emotional and sexual abuse, and physical and emotional neglect (Bernstein *et al.* 1994). Reliability and validity of the CTQ have been demonstrated previously (Bernstein *et al.* 1994). In this study the short versions (28-item version) of the CTQ translated into Norwegian (Aas *et al.* 2012), or previously validated in French, were used (Paquette *et al.* 2004). We used the moderate-severe cut-off scores to dichotomize each subscore, to classify subjects as having/not having a history of childhood trauma in that category, as suggested by Bernstein & Fink (1998).

Statistical analyses

All statistical analyses were performed with the packages PASW Statistics 18 (release 18.0.1) and R (version 2.12.0). Continuous variables are presented as mean values and standard deviations. χ^2 Tests were used to investigate childhood trauma dichotomized using the moderate-severe cut-off scores, to classify subjects as having/not having a history of childhood trauma, as suggested by Bernstein & Fink (1998). For substance abuse, we particularly focused on the presence of lifetime DSM-IV cannabis abuse (yes/no), although separate analyses investigating lifetime alcohol DSM-IV dependence (yes/no) were also included. We chose here to focus on alcohol dependence to capture the most severe form of alcohol use. Alcohol dependence has also been more linked to psychopathology, compared with alcohol abuse. It is also less related to different social norms of different countries regarding drinking, and therefore more suitable to study across study sites (Rehm *et al.* 2005). As the clinical variables were not normally distributed, non-parametric tests, such as the Mann-Whitney and Kruskal-Wallis tests, were used to investigate relationships between substance abuse and clinical variables. For the interaction and additive analyses, binary logistic regression analyses, together with multiple linear regression analyses, were performed. As the age at onset variable was skewed, it was log transformed before being added into these analyses. *Post-hoc* analyses were conducted controlling for possible confounders such as gender, as well as duration of illness. Bipolar subdiagnoses and research sites (France/Norway) were also added into the models, but later taken out as they did not have any significant influence and did not improve the model. We used a three-step multiple regression

analysis to investigate if childhood trauma mediated the relationship between cannabis abuse and symptom severity measures (Baron & Kenny, 1986). In step 1, we investigated variables measuring clinical expression (age at onset, rapid cycling, mood episodes, suicide attempt and so on) one at a time, and childhood abuse as the independent variable. In step 2 we investigated variables measuring clinical expression as the dependent variable, again one at a time, and cannabis abuse (here as the mediator) as the independent variable. In step 3, we investigated variables measuring clinical expression, one at a time (as the dependent variable), with childhood abuse as an independent variable whilst controlling for the presence of cannabis abuse (i.e. both childhood abuse and cannabis abuse entered as independent variables at the same time). For the interaction analyses we used regression analyses with both childhood trauma and cannabis abuse as independent variables, together with the interaction term cannabis abuse $\times$ childhood trauma. For the additive effects between cannabis abuse and childhood abuse, the data were grouped into three groups accordingly: group one, 'no cannabis'; group two, 'cannabis abuse and no childhood trauma'; and group three, 'cannabis abuse and childhood trauma'.

To avoid type 1 error we corrected for the number of childhood trauma variables tested [physical abuse, sexual abuse, emotional abuse, and abuse dose (none to three types of abuse: physical, sexual or/and emotional)], and thus used a *p* value of 0.01 as a marker of statistical significance after multiple testing. As this was a hypothesis-driven study, with *a priori* ideas on directions, we decided this was sufficient enough without the potential of losing important data.

Results

Childhood trauma and substance abuse

Cannabis abuse was significantly associated with childhood abuse, specifically, emotional abuse ($p=0.003$), sexual abuse, ($p=0.006$), and childhood trauma abuse-dose effect ($p=0.002$) (see Table 2). A statistically significant correlation between cannabis abuse and alcohol dependence was also observed. This was statistically significant also after correcting for group site [binary logistic regression: Nagelkerke $r^2=0.03$, $p=0.003$, $\exp(B)=2.75$. No statistical significant associations were observed between alcohol dependence and childhood abuse.

Substance abuse and clinical characteristics of bipolar disorders

Cannabis abuse was significantly associated with earlier age at onset ($p<0.001$), history of lifetime suicide

Table 2. Childhood trauma and substance abuse in bipolar disorder patients^a

Variable	Emotional abuse			Physical abuse		Sexual abuse		Abuse-dose effect			
	Yes		No	Yes	No	Yes	No	None	One	Two	Three
Cannabis abuse, %	Yes	19.7	10.4	22.2	11.9	21.4	11.3	9.8	15.0	21.7	31.8
	No	80.3	89.6	77.8	88.1	78.6	88.7	90.2	85.0	78.3	68.2
Alcohol dependence, %	Yes	9.8	8.8	14.8	8.3	6.7	9.4	8.5	7.8	13.3	4.5
	No	90.2	91.2	85.2	91.7	93.3	90.6	91.5	92.2	86.7	95.5
		$\chi^2=8.63$, df=1, $p=0.003^*$		$\chi^2=4.62$, df=1, $p=0.03$		$\chi^2=7.55$, df=1, $p=0.006^*$		$\chi^2=14.52$, df=3, $p=0.002^*$			
		$\chi^2=0.14$, df=1, $p=0.71$		$\chi^2=2.59$, df=1, $p=0.11$		$\chi^2=0.74$, df=1, $p=0.39$		$\chi^2=2.22$, df=3, $p=0.53$			

df, Degrees of freedom.

^a Percentage of patients who both abused cannabis and scored positive on childhood trauma was based on moderate to severe cut-off scores from Bernstein & Fink (1998), and cannabis abuse only; the percentage of patients having both cannabis abuse and childhood trauma is greater compared with the group with cannabis abuse only. Similar procedures were performed for alcohol dependence.

* Significant after correcting for the number of trauma subtypes.

attempt (yes/no) ($p=0.001$), as well as a trend for rapid cycling ($p=0.06$; see Table 3 and Fig. 1). No significant associations were observed between cannabis abuse and number of depressive, (hypo)manic, or mixed episodes. Alcohol dependence was significantly associated with a history of lifetime suicide attempt ($p=0.001$). Alcohol dependence was also associated with increased number of depressive episodes ($p=0.03$); however this was no longer statistically significant after correcting for multiple testing. No significant associations were observed between alcohol dependence, (hypo)manic or mixed episodes, rapid cycling or age at onset.

Associations between childhood trauma, cannabis abuse and clinical characteristics

After correcting for multiple testing, a trend for interaction ($p=0.039$) was observed between sexual abuse, cannabis abuse, and a history of at least one suicide attempt (see Table 4 and Fig. 2). Although this was only at trend level after correcting for multiple testing, the Nagelkerke r^2 improved when the interaction term was added into the model, indicating that cannabis abuse and sexual abuse were not only independently associated with a history of at least one suicide attempt, but also that they interacted. Adding sites and diagnosis into the model did not change the results (data not shown here). No other significant interactions were observed between childhood trauma, cannabis abuse and clinical expression. Statistically significant additive effects were observed for cannabis abuse and childhood abuse on the severity of several clinical characteristics. For *age at onset*, a statistically significant additive effect for sexual abuse and cannabis abuse on earlier age at onset ($p<0.001$) was observed, with the earliest age at onset in the group reporting both cannabis abuse and childhood trauma [mean=18.6 (s.d.=7.3) years, compared with no cannabis or childhood abuse: mean 25.0 (s.d.=10.5) years, see Fig. 3]. Similar findings were observed for emotional abuse and physical abuse ($p<0.001$ and $p<0.001$, respectively). A significant additive effect of cannabis and childhood abuse on *rapid cycling* was also found. Binary logistic regression analysis indicated a significantly higher frequency of reports of rapid cycling in the group with both cannabis and childhood abuse compared with all other groups, specifically for sexual abuse [odds ratio (OR) 1.63, 95% confidence interval (CI) 1.11–2.38, $p=0.013$, $r^2=0.02$] and emotional abuse (OR 1.61, 95% CI 1.13–2.30, $p=0.003$, $r^2=0.02$). Binary regression analyses also showed statistically significant additive effects for cannabis abuse and childhood abuse and report of a history of at least one *suicide attempt*, specifically with regard to sexual abuse (OR 2.13, 95% CI

Table 3. Substance abuse and clinical characteristics of bipolar disorders

	Cannabis abuse		Alcohol dependence	
	Yes	No	Yes	No
Mean age at onset, years (s.d.) ^a	19.6 (5.8) $z = -4.17, p < 0.001^*$	25.0 (10.5)	23.3 (9.9) $z = -0.92, p = 0.36$	24.4 (10.0)
Number of depressive episodes (s.d.) ^a	4.4 (5.3) $z = -0.28, p = 0.78$	4.8 (5.5)	6.3 (6.2) $z = -2.13, p = 0.03$	4.7 (5.5)
Number of (hypo)manic episodes (s.d.) ^a	4.7 (5.9) $z = -1.13, p = 0.26$	4.5 (6.0)	5.7 (7.3) $z = -0.09, p = 0.93$	4.5 (5.9)
Suicide attempt, % ^b	53.3 $\chi^2 = 11.16, df = 1; p = 0.001^*$	33.5	56.6 $\chi^2 = 10.28, df = 1, p = 0.001^*$	33.9
Rapid cycling, % ^b	18.4 $\chi^2 = 3.45, df = 1, p = 0.06$	11.8	25.0 $\chi^2 = 0.63, df = 1, p = 0.43$	20.3
Mixed episodes, % ^b	12.0 $\chi^2 = 0.11, df = 1, p = 0.74$	13.1	26.0 $\chi^2 = 0.20, df = 1, p = 0.65$	23.2

s.d., Standard deviation; df, degrees of freedom.

^a Mann-Whitney test.

^b χ^2 test.

* Significant after correcting for number of trauma subtypes.

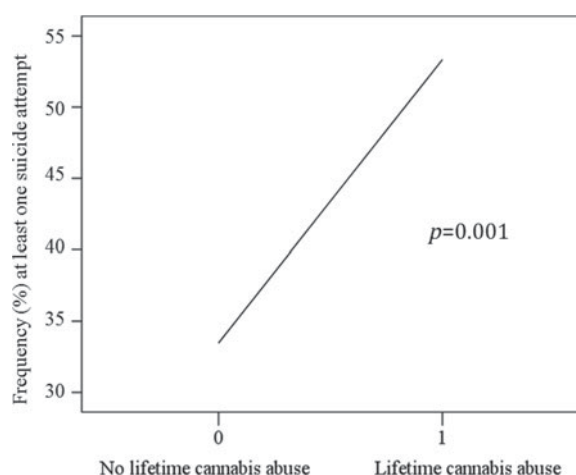


Fig. 1. Cannabis abuse and suicide attempt (wish to die). Analysis by χ^2 : 'no lifetime cannabis abuse', $n=368$; 'lifetime cannabis abuse', $n=211$.

1.49–3.14, $p < 0.001$, $r^2 = 0.04$) and emotional abuse (OR 1.88, 95% CI 1.34–2.63, $p < 0.001$, $r^2 = 0.03$), as well as a trend effect for physical abuse (OR 1.00, 95% CI 1.00–1.01, $p = 0.08$, $r^2 = 0.01$). Results also remained statistically significant ($p \leq 0.01$) after correction for the presence of alcohol dependence.

No additive effects were observed for cannabis abuse and childhood abuse, and increased number of mood episodes. *Post-hoc* regression analyses conducted for all additive analyses, correcting for gender and duration of illness, showed no changes in the results above (data not shown). No mediation effects were observed. Both childhood abuse and cannabis abuse were

independently significantly associated with the clinical expressions of bipolar disorders when entered into the model at the same time (for a history of at least one suicide attempt, see Table 4; for other clinical variables, data not shown).

Discussion

Our study is the first to demonstrate a trend for an interaction effect, as well as statistically significant additive effects, between childhood abuse and cannabis abuse on aspects of the clinical expression of bipolar disorders. Patients with both a lifetime history of cannabis abuse and childhood trauma had a higher frequency of reporting at least one lifetime suicide attempt, lower age at onset, and increased frequency of rapid cycling, suggesting an increased clinical severity of the bipolar disorder. Our study also demonstrates that while childhood sexual abuse and emotional abuse are significantly associated with cannabis use, they do not appear to be associated with the risk for alcohol dependence. When controlling for cannabis abuse, childhood abuse was still significantly associated with symptom load in bipolar disorder, indicating that cannabis abuse moderates, but does not mediate, the relationship between childhood abuse and increased severity of the clinical expression of bipolar disorders; similar findings were observed when childhood abuse was entered as the mediator. Based on the significant correlation between cannabis use and alcohol dependence, and the significant association between alcohol dependence and a history of

Table 4. Multivariate regression analyses, investigating childhood trauma, cannabis abuse and suicide attempt

Variable	Step 1		Step 2	
	OR (95% CI)	<i>p</i>	OR (95% CI)	<i>p</i>
Suicide attempt ^a				
Cannabis abuse	0.47 (0.29–0.78)	0.003	0.16 (0.05–0.53)	0.003
CTQ sexual abuse	0.50 (0.32–0.76)	0.002	0.16 (0.05–0.53)	0.002
Cannabis x sexual abuse	–		3.97 (1.07–14.69)	0.039
Nagelkerke (<i>r</i> ²)		<i>r</i> ² =0.048		<i>r</i> ² =0.059

OR, Odds ratio; CI, confidence interval; CTQ, Childhood Trauma Questionnaire.

^a Logistic regression; step 1 includes cannabis abuse and sexual abuse; step 2 includes cannabis abuse, sexual abuse, and the interaction term 'cannabis abuse×sexual abuse'.

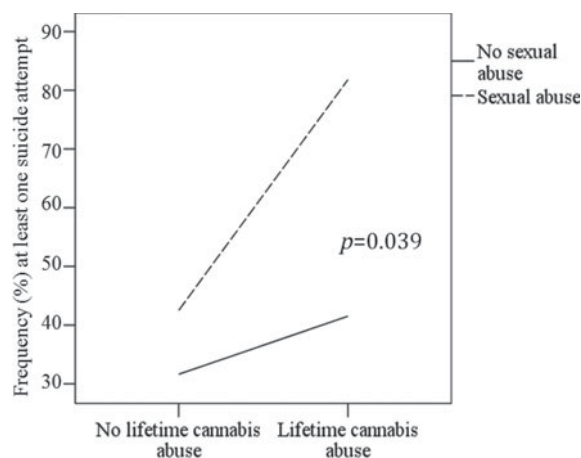


Fig. 2. Interaction between childhood sexual abuse, cannabis abuse and a history of at least one suicide attempt. Analysis by binary logistic regression: 'no lifetime cannabis abuse', *n*=368; 'lifetime cannabis abuse', *n*=211; 'no sexual abuse', *n*=475; 'sexual abuse', *n*=104. Sexual abuse was dichotomized into trauma *versus* no trauma based on moderate to severe cut-off scores of Bernstein *et al.* (1998).

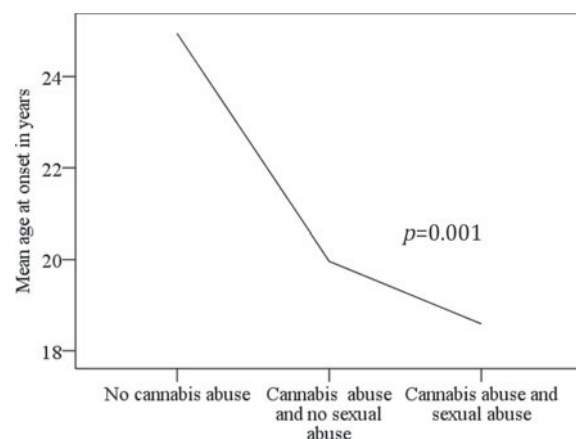


Fig. 3. Additive effect of cannabis abuse and childhood sexual abuse and lower age at onset. Analysis by the Kruskal–Wallis test: 'no lifetime cannabis abuse', mean 24.95 (S.E.M.=0.47) years, *n*=499; 'lifetime cannabis abuse and no sexual abuse', mean 19.96 (S.E.M.=0.70) years, *n*=53; 'lifetime cannabis abuse and sexual abuse', mean 18.59 (S.E.M.=1.57) years, *n*=22.

at least one suicide attempt, we performed additional analyses controlling for alcohol dependence. Our results suggest that cannabis abuse is associated with increased risk of at least one suicide attempt, independently of alcohol dependence. Our data further indicate that cannabis abuse has a stronger effect on symptom levels in bipolar disorder than alcohol dependence.

The results of this study also confirm previous findings of a relationship between childhood abuse and substance abuse (Fergusson *et al.* 2008; Shin *et al.* 2009), and between substance abuse and increased symptom severity in bipolar disorders (Lin *et al.* 2006; Javaid *et al.* 2011; Lagerberg *et al.* 2011). Our study also confirms findings reported by Fergusson *et al.* (2008) and Shin *et al.* (2009) that individuals

reporting a history of childhood abuse tend to show an increased risk of substance abuse in adulthood compared with individuals without childhood abuse. One of the main questions of the present study was whether childhood abuse and cannabis abuse independently influence the clinical picture, or if the effect disappeared when adding both factors at the same time as independent variables in a multivariate regression analysis. As both childhood abuse and cannabis abuse were independently significantly associated with the clinical expression of bipolar disorders when entered into the model at the same time, our data do not support mediation effects. However, our findings support a trend for interactions and statistically significant additive effects. The significant additive effects indicate that childhood abuse and cannabis abuse may have separate effects on bipolar disorders, and

the presence of both proportionally increases the risk for a more severe illness progression. Research in this area is still sparse, and future research is needed to further explore the mechanisms behind these associations. Our data support a role of childhood abuse, as well as cannabis abuse, in the development of clinical characteristics in bipolar disorder. It further highlights the importance of assessing both childhood abuse and cannabis abuse in bipolar disorder research. These should also be systematically included in the clinical assessment of the patients.

A possible mechanism behind the increased risk for a co-occurrence of childhood abuse and cannabis abuse may be linked to their suggested opposite effects on the hypothalamic–pituitary–adrenal (HPA) axis (Heim *et al.* 2000, 2008; van Leeuwen *et al.* 2011). Childhood trauma has been linked to long-term hyperactivation of HPA activation (Heim *et al.* 2000, 2008), while cannabis abuse has been linked to a reduction in HPA activity (van Leeuwen *et al.* 2011). Cannabis, or substance use, in individuals with childhood abuse could therefore be viewed as a form of self-medication aiming to ‘regulate’ the HPA axis.

Further studies investigating mechanisms behind the effects of substance abuse and childhood abuse on clinical expression in bipolar disorder are needed. A possible target for future research could be, for example, investigations of the effect of childhood abuse on impulsivity (Etain *et al.* 2013), as a possible link between trauma and substance misuse and other risk behaviours.

Limitations

As in most clinical studies of this phenomenon, data on childhood abuse were obtained retrospectively with the inherent weakness of retrospective reporting designs. However, the retrospective collection of childhood abuse data in patients with severe mental disorders has been found to be a valid and reliable source of obtaining information in previous studies (Fisher *et al.* 2011). Our study also fulfills the quality criteria described by Fisher & Hosang (2010), with the limitation that in the Norwegian sample no standardized assessment, but rather clinical judgment, was conducted to decide whether the patients were in a suitable mood to reliably fill in the CTQ form. All patients from the French sample were systematically assessed and defined as euthymic at inclusion. Another limitation should be acknowledged; it is supposed that cannabis abuse pre-existed a history of suicide attempts. However, this has not been rigorously verified in our sample. Also, although we assume that childhood abuse pre-existed cannabis abuse, we cannot exclude the possibility that some

individuals had a cannabis abuse history prior to childhood abuse experiences. Lastly, as already mentioned, we observed a higher rate of alcohol dependence in the Norwegian sample than in the French sample, as similarly observed in the overview article by Rehm *et al.* (2005). As opposed to this, a recent report by the World Health Organization showed higher alcohol consumption per adult in France than in Norway (WHO, 2011). In our sample different instruments were used to assess alcohol dependence (DIGS in the French sample and SCID-I in the Norwegian sample). It could be that the threshold to detect alcohol dependence is higher in the DIGS than the SCID-I, or it could be that alcohol dependence is linked not only to the amount of alcohol consumed, but also to drinking style, such as binge drinking, which is more frequent in Norway.

Conclusion

Our results show a trend for interaction, as well as statistically significant additive effects, between childhood abuse and cannabis abuse on the clinical expression of bipolar disorders. These results indicate that both childhood abuse and substance abuse (and in particular cannabis abuse) should be taken into account when investigating mechanisms behind the clinical expression and prevalence of bipolar disorders. If confirmed, these results have implications for our understanding of the pathophysiological mechanisms, and treatment implications, in bipolar disorders.

Supplementary material

For supplementary material accompanying this paper visit <http://dx.doi.org/10.1017/S0033291713002316>.

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Declaration of Interest

None.

References

- Aas M, Dazzan P, Fisher HL, Morgan C, Morgan K, Reichenberg A, Zanelli J, Fearon P, Jones PB, Murray RM, Pariante CM (2011). Childhood trauma and cognitive function in first-episode affective and non-affective psychosis. *Schizophrenia Research* **129**, 12–19.
- Aas M, Djurovic S, Athanasiu L, Steen NE, Agartz I, Lorentzen S, Sundet K, Andreassen OA, Melle I (2012). Serotonin transporter gene polymorphism, childhood trauma, and cognition in patients with psychotic disorders. *Schizophrenia Bulletin* **38**, 15–22.
- Agrawal A, Nurnberger JI Jr, Lynskey MT (2011). Cannabis involvement in individuals with bipolar disorder. *Psychiatry Research* **185**, 459–461.
- Alvarez MJ, Roura P, Osés A, Foguet Q, Sola J, Arrufat FX (2011). Prevalence and clinical impact of childhood trauma in patients with severe mental disorders. *Journal of Nervous and Mental Disease* **199**, 156–161.
- APA (1994). *Diagnostic and Statistical Manual of Mental Disorders*, 4th edn, revised. American Psychological Association: Washington, DC.
- Baron RM, Kenny DA (1986). The moderator–mediator variable distinction in social psychological research: conceptual, strategic, and statistical considerations. *Journal of Personality and Social Psychology* **51**, 1173–1182.
- Bebbington PE, Bhugra D, Brugha T, Singleton N, Farrell M, Jenkins R, Lewis G, Meltzer H (2004). Psychosis, victimisation and childhood disadvantage: evidence from the second British National Survey of Psychiatric Morbidity. *British Journal of Psychiatry* **185**, 220–226.
- Bernstein DP, Fink L (1998). *Childhood Trauma Questionnaire: A Retrospective Self-Report*. Harcourt Brace and Company: San Antonio.
- Bernstein DP, Fink L, Handelsman L, Foote J, Lovejoy M, Wenzel K, Sapareto E, Ruggiero J (1994). Initial reliability and validity of a new retrospective measure of child abuse and neglect. *American Journal of Psychiatry* **151**, 1132–1136.
- Brown GR, McBride L, Bauer MS, Williford WO (2005). Impact of childhood abuse on the course of bipolar disorder: a replication study in U.S. veterans. *Journal of Affective Disorders* **89**, 57–67.
- Coulston CM, Tanious M, Mulder RT, Porter RJ, Malhi GS (2012). Bordering on bipolar: the overlap between borderline personality and bipolarity. *Australian and New Zealand Journal of Psychiatry* **46**, 506–521.
- Daruy-Filho L, Brietzke E, Lafer B, Grassi-Oliveira R (2011). Childhood maltreatment and clinical outcomes of bipolar disorder. *Acta Psychiatrica Scandinavica* **124**, 427–434.
- Etain B, Mathieu F, Henry C, Raust A, Roy I, Germain A, Leboyer M, Bellivier F (2010). Preferential association between childhood emotional abuse and bipolar disorder. *Journal of Traumatic Stress* **23**, 376–383.
- Etain B, Mathieu F, Liqueur S, Raust A, Cochet B, Richard JR, Gard S, Zanouy L, Kahn JP, Cohen RF, Bougerol T, Henry C, Leboyer M, Bellivier F (2013). Clinical features associated with trait-impulsiveness in euthymic bipolar disorder patients. *Journal of Affective Disorders* **144**, 240–247.
- Fergusson DM, Boden JM, Horwood LJ (2008). The developmental antecedents of illicit drug use: evidence from a 25-year longitudinal study. *Drug and Alcohol Dependence* **96**, 165–177.
- Fisher HL, Craig TK, Fearon P, Morgan K, Dazzan P, Lappin J, Hutchinson G, Doody GA, Jones PB, McGuffin P, Murray RM, Leff J, Morgan C (2011). Reliability and comparability of psychosis patients' retrospective reports of childhood abuse. *Schizophrenia Bulletin* **37**, 546–553.
- Fisher HL, Hosang GM (2010). Childhood maltreatment and bipolar disorder: a critical review of the evidence. *Mind and Brain* **1**, 75–85.
- Garno JL, Goldberg JF, Ramirez PM, Ritzler BA (2005). Impact of childhood abuse on the clinical course of bipolar disorder. *British Journal of Psychiatry* **186**, 121–125.
- Heim C, Newport DJ, Heit S, Graham YP, Wilcox M, Bonsall R, Miller AH, Nemeroff CB (2000). Pituitary–adrenal and autonomic responses to stress in women after sexual and physical abuse in childhood. *Journal of the American Medical Association* **284**, 592–597.
- Heim C, Newport DJ, Mletzko T, Miller AH, Nemeroff CB (2008). The link between childhood trauma and depression: insights from HPA axis studies in humans. *Psychoneuroendocrinology* **33**, 693–710.
- Houston JE, Murphy J, Shevlin M, Adamson G (2011). Cannabis use and psychosis: re-visiting the role of childhood trauma. *Psychological Medicine* **41**, 2339–2348.
- Janssen I, Krabbendam L, Bak M, Hanssen M, Vollebergh W, de Graaf R, van Os J (2004). Childhood abuse as a risk factor for psychotic experiences. *Acta Psychiatrica Scandinavica* **109**, 38–45.
- Javadi N, Kennedy JL, De Luca V (2011). Ethnicity and age at onset in bipolar spectrum disorders. *CNS Spectrums* **16**, 127–134.
- Lagerberg TV, Sundet K, Aminoff SR, Berg AO, Ringen PA, Andreassen OA, Melle I (2011). Excessive cannabis use is associated with earlier age at onset in bipolar disorder.

- European Archives of Psychiatry and Clinical Neuroscience* **261**, 397–405.
- Lin PI, McInnis MG, Potash JB, Willour V, MacKinnon DF, DePaulo JR, Zandi PP** (2006). Clinical correlates and familial aggregation of age at onset in bipolar disorder. *American Journal of Psychiatry* **163**, 240–246.
- Mondelli V, Dazzan P, Hepgul N, Di FM, Aas M, D'Albenzio A, Di NM, Fisher H, Handley R, Marques TR, Morgan C, Navari S, Taylor H, Papadopoulos A, Aitchison KJ, Murray RM, Pariante CM** (2010). Abnormal cortisol levels during the day and cortisol awakening response in first-episode psychosis: the role of stress and of antipsychotic treatment. *Schizophrenia Research* **116**, 234–242.
- Nemeroff CB** (2004). Neurobiological consequences of childhood trauma. *Journal of Clinical Psychiatry* **65** (Suppl. 1), 18–28.
- Nurnberger JI Jr, Blehar MC, Kaufmann CA, York-Cooler C, Simpson SG, Harkavy-Friedman J, Severe JB, Malaspina D, Reich T** (1994). Diagnostic Interview for Genetic Studies. Rationale, unique features, and training. NIMH Genetics Initiative. *Archives of General Psychiatry* **51**, 849–859.
- Paquette D, Laporte L, Bigras M, Zoccolillo M** (2004). Validation of the French version of the CTQ and prevalence of the history of maltreatment [in French]. *Santé mentale au Québec* **29**, 201–220.
- Pettinati HM, O'Brien CP, Dundon WD** (2013). Current status of co-occurring mood and substance use disorders: a new therapeutic target. *American Journal of Psychiatry* **170**, 23–30.
- Pettinati HM, Oslin DW, Kampman KM, Dundon WD, Xie H, Gallis TL, Dackis CA, O'Brien CP** (2010). A double-blind, placebo-controlled trial combining sertraline and naltrexone for treating co-occurring depression and alcohol dependence. *American Journal of Psychiatry* **167**, 668–675.
- Post RM** (2010). Mechanisms of illness progression in the recurrent affective disorders. *Neurotoxicity Research* **18**, 256–271.
- Read J, van Os J, Morrison AP, Ross CA** (2005). Childhood trauma, psychosis and schizophrenia: a literature review with theoretical and clinical implications. *Acta Psychiatrica Scandinavica* **112**, 330–350.
- Rehm J, Room R, van den Brink W, Jacobi F** (2005). Alcohol use disorders in EU countries and Norway: an overview of the epidemiology. *European Neuropsychopharmacology* **15**, 377–388.
- Ringen PA, Lagerberg TV, Birkenaes AB, Engn J, Faerden A, Jonsdottir H, Nesvag R, Friis S, Opjordsmoen S, Larsen F, Melle I, Andreassen OA** (2008). Differences in prevalence and patterns of substance use in schizophrenia and bipolar disorder. *Psychological Medicine* **38**, 1241–1249.
- Ringen PA, Melle I, Berg AO, Agartz I, Spigset O, Simonsen C, Sundet K, Andreassen OA** (2013). Cannabis use and premorbid functioning as predictors of poorer neurocognition in schizophrenia spectrum disorder. *Schizophrenia Research* **143**, 84–89.
- Shevlin M, Dorahy MJ, Adamson G** (2007). Trauma and psychosis: an analysis of the National Comorbidity Survey. *American Journal of Psychiatry* **164**, 166–169.
- Shin SH, Edwards EM, Heeren T** (2009). Child abuse and neglect: relations to adolescent binge drinking in the national longitudinal study of Adolescent Health (AddHealth) Study. *Addictive Behaviors* **34**, 277–280.
- van Leeuwen AP, Creemers HE, Greaves-Lord K, Verhulst FC, Ormel J, Huizink AC** (2011). Hypothalamic–pituitary–adrenal axis reactivity to social stress and adolescent cannabis use: the TRAILS study. *Addiction* **106**, 1484–1492.
- WHO** (2011). *Global Status Report on Alcohol and Health*. World Health Organization: Geneva.