Substance misuse in first-episode psychosis: 15-month prospective follow-up study*

D. WADE, S. HARRIGAN, J. EDWARDS, P. M. BURGESS, G. WHelan and P. D. McGorrry

Background Well-designed prospective studies of substance misuse in first-episode psychosis can improve our understanding of the risks associated with comorbid substance misuse and psychosis.

Aims To examine the potential effects of substance misuse on in-patient admission and remission and relapse of positive symptoms in first-episode psychosis.

Method The study was a prospective 15-month follow-up investigation of 103 patients with first-episode psychosis recruited from three mental health services.

Results Substance misuse was independently associated with increased risk of in-patient admission, relapse of positive symptoms and shorter time to relapse of positive symptoms after controlling for potential confounding factors. Substance misuse was not associated with remission or time to remission of positive symptoms. Heavy substance misuse was associated with increased risk of in-patient admission, relapse and shorter time to relapse.

Conclusions Substance misuse is an independent risk factor for a problematic recovery from first-episode psychosis.

Declaration of interest None.

Research on substance misuse in psychotic disorders has been hampered by methodological limitations including selection bias, lack of diagnostic rigour, failure to control for potential confounding variables and a lack of prospective follow-up studies (Blanchard et al., 2000; Murray et al., 2003). Prospective studies of first-episode psychosis can address these issues and improve our understanding of the risks associated with comorbid substance misuse and psychosis. A small number of prospective studies have reported that substance misuse is associated with a problematic recovery from recent-onset psychosis (Linszen et al., 1994; Strakowski et al., 1998; Sorbara et al., 2003). Consistent with the findings of these studies, our hypotheses for the current study were that substance misuse in first-episode psychosis would be associated with increased risk of in-patient admission, a longer time to remission of positive symptoms, and earlier and increased risk of relapse of positive symptoms.

METHOD

Participants Consecutive in-patient and out-patient admissions of individuals with first-episode psychosis were screened for the study between January and December 1997 at the Central East Area Mental Health Service (CEAMHS) and the Northern Area Mental Health Service (NAMHS), and between March and July 2001 at the Early Psychosis Prevention and Intervention Centre (EPPIC) in Melbourne, Australia. The CEAMHS and the NAMHS are generic mental health services for adults with serious mental illnesses and EPPIC is a specialist mental health service for youth with first-episode psychosis. The services provide comprehensive care within defined catchment areas and are funded by the state government. The inclusion criteria for the study were age 15–30 years, fluency in English, ability to give informed consent, and clear evidence of a functional psychotic disorder. The exclusion criteria were organic aetiology, learning disability, history of brain damage or epilepsy, and more than 6 months of prior treatment for a psychotic disorder. The aims of the study were fully explained to the participants, who provided written informed consent. The research and ethics committees of the North-Western Mental Health Program approved the study. In total 126 patients (EPPIC n = 71; CEAMHS n = 32; NAMHS n = 23) were recruited to the study. Twenty-three patients had missing data regarding the presence of any substance misuse during follow-up owing to their not being contactable at the 9-month or 15-month time point and were excluded from further analyses, leaving a sample of 103 patients (EPPIC n = 59, CEAMHS n = 25, NAMHS n = 19). For patients who were eligible for the study at EPPIC (n = 95), no significant difference was found between patients included (n = 59) and not included (n = 36) in the current analyses on demographic variables, psychotic disorder diagnosis or duration of untreated psychosis.

Measures and procedure A baseline assessment was completed at entry to treatment, and follow-up assessments were undertaken 3 months, 9 months and 15 months following the initial assessment. An updated version of the Royal Park Multidiagnostic Instrument for Psychoses (RPMIP; McGorry et al., 1990) was used to diagnose DSM–IV (American Psychiatric Association, 1994) psychotic disorders based on assessment at baseline and 3-month follow-up. Diagnoses were subsequently categorised as schizophrenia-spectrum psychosis (schizophrenia, schizophrheniform, schizoaffective or delusional) or other psychosis (bipolar, major depression, not otherwise specified, substance-induced or brief). The RPMIP was also used to estimate the duration of untreated psychosis in days, defined as the time from onset of psychotic symptoms to treatment entry. The Chemical Use, Abuse and Dependence Scale (CUAD; McGovern & Morrison, 1992) was used to diagnose DSM–III–R (American Psychiatric Association, 1987) substance misuse (criteria met for abuse or dependence) during the 15-month follow-up period. Substance misuse was assessed at the 9-month time point (for the interval between baseline and 9...
months) and the 15-month time point (for the interval between 9 months and 15 months). Substances assessed for included alcohol, amphetamine, benzodiazepine, cannabis, cocaine, hallucinogen, inhalant, opioid and phenycyclidine. Diagnoses of substance misuse were based on the 17 items rated ‘true’ or ‘false’ for each substance. Each item corresponds to a criterion of DSM–III–R substance abuse or dependence. Individual substance use severity scores are based on weighted scores from 1 to 4 for the 17 items rated ‘true’ for each substance. The sum of individual substance use severity scores provides a total substance use severity score. The higher total substance use severity score at the 9- or 15-month time point was used to calculate the total substance use severity score during the follow-up period. As in the study by Kavanagh et al. (2004), any misuse of substances other than alcohol or cannabis was defined as ‘other substance misuse’ and the presence of at least two of alcohol, cannabis or other substance misuse was defined as ‘poly-substance misuse’. Patients with substance misuse were grouped according to the pattern of substance misuse as follows: cannabis misuse; other but not cannabis misuse; or alcohol misuse only. Substance misuse was categorised as mild or heavy based on a median split of CUAD total substance use severity scores.

Remission and relapse of positive psychotic symptoms were the primary clinical outcomes. The Brief Psychiatric Rating Scale (BPRS; Lukoff et al., 1986) was used to rate remission and relapse of positive symptoms according to the following criteria: remission was defined as a score of 3 (mild) or less on all of the BPRS Psychotic sub-scale items (hallucinations, conceptual disorganisation, unusual thought content and suspiciousness) for at least 2 weeks; relapse was defined as a score of 4 (moderate) or more on any of the BPRS Psychotic sub-scale items. Participants diagnosed with other substance misuse were 45% (25 out of 54, missing data for 1 patient) met criteria for poly-substance misuse, 57% (31 out of 54, missing data for 1 patient) met criteria for cannabis misuse and 7% (4 out of 54, missing data for 1 patient) met criteria for hallucinogen misuse. The rate of poly-substance misuse was 30% (31 out of 102). Of the patients with a diagnosis of substance misuse met criteria for amphetamine and/or hallucinogen misuse. The rate of poly-substance misuse was 30% (31 out of 102). Of the patients with a diagnosis of substance misuse, 57% (31 out of 54, missing data for 1 patient) met criteria for poly-substance misuse. The varying denominator for these analyses is owing to missing data on misuse of some individual substances.

**RESULTS**

**Participants**

The mean age of the 103 patients was 21.6 (s.d. = 3.5) years. The patients were predominantly male (71%) and single (90%), and approximately a third (34%) of the patients had completed secondary school. The majority of patients were diagnosed with schizophrenia-spectrum psychosis (75%) and hospitalised during the first 3 months of treatment (76%). No significant difference was found between these 103 patients and the 23 patients excluded from analyses because of missing substance misuse data on demographic or clinical variables, including the rates of any or individual lifetime substance misuse at baseline.

**Rates of substance misuse**

Overall, 53% of patients (55 out of 103) were given a diagnosis of substance misuse during follow-up; these included cannabis 42% (43 out of 103), alcohol 30% (30 out of 100) and other substance misuse 17% (17 out of 98). Thirteen of the 17 patients diagnosed with other substance misuse met criteria for amphetamine and/or hallucinogen misuse. The rate of poly-substance misuse was 30% (31 out of 102). Of the patients with a diagnosis of substance misuse, 57% (31 out of 54, missing data for 1 patient) met criteria for poly-substance misuse. The varying denominator for these analyses is owing to missing data on misuse of some individual substances.

**In-patient admission**

The rates of in-patient admission for patients with and without a diagnosis of substance misuse were 45% (25 out of 55) and 15% (7 out of 48) respectively (Table 1). Logistic regression analyses showed that substance misuse was significantly associated with in-patient admission during follow-up and remained so after controlling for the effects of gender, age,
Psychotic disorder diagnosis, duration of untreated psychosis and medication compliance. The mean number of in-patient admission days was 12.0 (s.d. = 19.9, median = 0) for patients diagnosed with substance misuse compared with 1.4 (s.d. = 4.2, median = 0) for patients without substance misuse (Mann–Whitney U-test, Z = −3.6, P < 0.001). When patients were grouped according to the pattern of substance misuse, 21 out of 43 patients with cannabis misuse, 3 out of 5 patients with other substance misuse but not cannabis misuse, and 1 out of 7 patients with alcohol misuse only were hospitalised following the first 3 months of treatment.

**Remission of psychotic symptoms**

For patients who achieved remission of positive symptoms during follow-up (98 out of 103), the mean duration of remission (that is, the period from the time that remission criteria were first met to psychotic relapse or the end of follow-up) was 343.7 days (s.d. = 133.6, median = 386.0). The rates of remission during follow-up for patients with and without a diagnosis of substance misuse were 93% (51 out of 55) and 98% (47 out of 48) respectively. Univariate logistic regression analyses showed that the association between substance misuse and remission was not statistically significant (Table 1). Multivariate analyses were not undertaken owing to the small number of patients who did not achieve remission (n = 5).

**Time to remission of psychotic symptoms**

A Kaplan–Meier survival analysis showed no significant difference between patients with substance misuse (n = 55, 4 censored cases; median time to remission 39 days, 95% CI 22–56) and patients without substance misuse (n = 48, 1 censored case; median time to remission 41 days, 95% CI 31–51) on days to remission (log-rank test, χ² = 1.1, d.f. = 1, P = 0.300). A Cox regression analysis showed that substance misuse was not significantly associated with time to remission (hazard ratio 0.8, 95% CI 0.5–1.2, P = 0.277) after controlling for the effects of gender, age, psychotic disorder diagnosis, duration of untreated psychosis and medication compliance.

**Relapse of psychotic symptoms**

For patients who achieved remission during follow-up (n = 98), the rates of relapse of positive symptoms during follow-up for patients with and without substance misuse diagnosis were 51% (26 out of 51) and 17% (8 out of 47) respectively. Logistic regression analyses showed that substance misuse was significantly associated with relapse and remained so after controlling for the effects of gender, age, psychotic disorder diagnosis, duration of untreated psychosis and medication compliance (Table 1). When patients were grouped according to the pattern of substance misuse, 23 out of 40 patients with cannabis misuse and 2 out of 4 patients with other substance misuse but not cannabis misuse relapsed, compared with 1 out of 7 patients with alcohol misuse only.

**Time to relapse of psychotic symptoms**

For patients who achieved remission (n = 98), a Kaplan–Meier survival analysis showed that substance misuse was a significant risk factor for time to relapse (Fig. 1). Patients with a diagnosis of substance misuse (n = 51, 25 censored cases) had a significantly shorter time to relapse of psychotic symptoms compared with patients without substance misuse (n = 47, 39 censored cases; log-rank test, χ² = 12.7 d.f. = 1, P < 0.001). The median time to relapse for patients with substance misuse was 378 days (95% CI 271–485, mean = 359). The median time to relapse for patients without substance misuse could not be calculated because fewer than half the patients relapsed (mean 477 days). A Cox regression analysis showed that substance misuse remained significantly associated with shorter time to relapse after controlling for the effects of gender, age, psychotic disorder diagnosis, duration of untreated psychosis and medication compliance (hazard ratio 2.8, 95% CI 1.2–6.7, P = 0.021).

**Relationship between severity of substance misuse and in-patient admission, relapse and time to relapse**

Patients whose substance misuse was categorised as heavy (n = 27), mild (n = 28) and none (n = 48) were compared on rates of in-patient admission and relapse and time to relapse. The heavy substance misuse group had a higher rate of in-patient admission (52%; 14 out of 27) than the mild substance misuse group (39%; 11 out of 28), who in turn had a higher rate of in-patient admission than the no substance misuse group (15%, 7 out of 48). A univariate logistic regression analysis showed that patients with heavy or mild substance misuse were significantly more likely to be admitted for in-patient care than patients who did not misuse substances (Table 2). Heavy but not mild substance misuse...
remained significantly associated with in-patient admission after controlling for the effects of gender, age, psychotic disorder diagnosis, duration of untreated psychosis, and medication compliance (Table 2). Among patients whose disorder was in remission (n=98), those with heavy substance misuse had a higher rate of relapse (64%; 16 out of 25) than those with mild substance misuse (38%; 10 out of 26), who in turn had a higher rate of relapse than patients with no substance misuse (17%; 8 out of 47). A univariate logistic regression analysis showed that patients with heavy or mild substance misuse were significantly more likely to experience relapse compared with patients with no substance misuse. After adjusting for the effects of the covariates, heavy but not mild substance misuse was significantly associated with relapse (Table 2).

A Kaplan–Meier survival analysis showed that substance use severity was a significant risk factor for time to relapse (Fig. 2). Patients with heavy substance misuse (n=25, 9 censored cases; median 327 days, 95% CI 238–416) had a shorter time to relapse of psychotic symptoms than patients with mild substance misuse, a difference that just failed to reach statistical significance (n=26, 16 censored cases; log-rank test, \( \chi^2=3.8, \) d.f.=1, \( P=0.052 \)), and a significantly shorter time to relapse than patients with no substance misuse (n=47, 39 censored cases; log-rank test, \( \chi^2=19.2, \) d.f.=1, \( P<0.001 \)). Patients with mild substance misuse had a significantly shorter time to relapse than patients with no substance misuse (log-rank test, \( \chi^2=4.3, \) d.f.=1, \( P=0.038 \)). The median time to relapse for patients with mild and no substance misuse could not be calculated because fewer than half of these patients relapsed. A multivariate analysis showed that heavy substance misuse (hazard ratio 4.6, 95% CI 1.7–12.5, \( P=0.003 \)) but not mild substance misuse (hazard ratio 2.0, 95% CI 0.8–5.4, \( P=0.160 \)) was significantly associated with a shorter time to relapse compared with no substance misuse.

**DISCUSSION**

The aim of the study was to examine the potential impact of substance misuse on clinical outcome in individuals treated for first-episode psychosis. The findings supported the hypotheses that substance misuse is associated with increased risk of in-patient admission and earlier and increased risk of psychotic relapse. The hypothesis that substance misuse is associated with longer time to remission of positive symptoms was not supported.

**Strengths and limitations**

A range of methodological problems have affected the study of comorbid substance misuse and psychosis. Briefly, these problems include use of criteria to diagnose

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**Table 2** Associations between substance use severity and in-patient admission following the initial 3-month treatment period and relapse during the 15-month follow-up

<table>
<thead>
<tr>
<th>Substance use severity</th>
<th>In-patient admission (n=103)</th>
<th>Clinical outcome</th>
<th>Relapse (n=98)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>% (n)</td>
<td>OR (95% CI)</td>
<td>Adjusted OR (95% CI)</td>
</tr>
<tr>
<td>Heavy misuse</td>
<td>52 (14)</td>
<td>6.3 (2.1–19.0)**</td>
<td>5.7 (1.5–21.9)**</td>
</tr>
<tr>
<td>Mild misuse</td>
<td>39 (11)</td>
<td>3.8 (1.3–11.4)*</td>
<td>2.8 (0.8–9.8)</td>
</tr>
<tr>
<td>No misuse</td>
<td>15 (7)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

OR, odds ratio.
1. Adjusted for gender, age, psychotic disorder diagnosis (schizophrenia-spectrum or other psychosis), duration of untreated psychosis (log-transformed) and medication non-compliance.
2. Missing compliance data for 1 patient; n=102.
3. Missing compliance data for 1 patient; n=97.
4. Reference category.

*P < 0.05, **P < 0.01, ***P < 0.001.
reports from research interviewers that most patients were willing to discuss substance-related problems. Further, urine drug screens can only detect substance use within a limited period and cannot provide information about the functional impact of substance misuse necessary to make a diagnosis. Third, analysis of the independent effects of different types of substance misuse on outcome was not possible, given that more than half of the patients with a diagnosis of substance misuse met criteria for poly-substance misuse. The finding that 87% of patients with substance misuse met criteria for cannabis and/or other substance misuse tends to implicate these substances in the observed adverse effects of substance misuse. Descriptive analyses suggested that patients with alcohol misuse only were less likely to experience in-patient admission or relapse compared with patients reporting cannabis or other substance misuse. These findings are consistent with evidence for a stronger association between psychotic exacerbations and cannabis or stimulant misuse compared with alcohol misuse (Dixon, 1999). Fourth, the operational definition of remission (minimal positive symptoms for at least 2 weeks) may be criticised for the relatively low threshold for remission criteria to be met (Andreasen et al., 2005). However, 94 of the 98 patients in remission maintained their initial remission of positive symptoms for at least 8 consecutive weeks, which is similar to criteria used in other studies of first-episode psychosis (e.g. Lieberman et al., 1993; Amminger et al., 1997).

Comparison of current findings with other research

Several (but not all) prospective studies have reported associations between substance misuse and worse outcome in first-episode or recent-onset psychosis. Sorbara et al. (2003) found that drug misuse but not alcohol misuse in first-episode psychosis was associated with an increased risk of in-patient admission. It is worth noting, however, that 3 of the 13 alcohol misusers were also diagnosed with drug misuse in this study. Linszen et al. (1994) found that cannabis misuse in recent-onset psychosis was associated with earlier relapse and an increased risk of relapse of positive symptoms. Despite differences in the definitions of substance misuse and relapse, our study and Linszen et al. (1994) found similar rates of relapse in misusing (51% and 42% respectively) and non-misusing patients (17% in both studies). Sevy et al. (2001) did not find a link between substance misuse in first-episode psychosis and earlier relapse or an increased risk of relapse; this negative finding might have been owing to the analysis of effects of substance misuse diagnosed at initial presentation rather than during the follow-up treatment period.

In contrast to our findings, Strakowski et al. (1998) reported that substance misuse in first-episode affective psychosis was associated with a longer time to symptomatic remission. Differences in sample characteristics and methodology between the two studies may help to explain the discrepant findings. For example, Strakowski et al. (1998) recruited patients with bipolar or major depressive disorder with psychosis rather than a broad range of psychotic disorders, and operationally defined remission in terms of positive, negative and affective symptoms rather than positive symptoms alone. It is also feasible that the lack of association between substance misuse and remission of positive symptoms in our study might have resulted from variation in the severity of substance use following entry to treatment. That is, patients might have reduced or stopped their substance use immediately following entry to treatment in response to the onset of acute psychosis and/or subsequent treatment.
including in-patient admission. If this is correct, a short-term reduction in severity of substance use might have enabled substance misusers to achieve rapid remission while still meeting criteria for substance misuse during the follow-up period. Unfortunately, the collection of interval-based substance use data did not enable us to test this proposition.

The association between more severe substance use and in-patient admission as well as relapse of positive symptoms is consistent with a previous report of a dose–response relationship between frequency of cannabis misuse and relapse (Linszen et al., 1994). A dose–response relationship is consistent with a causal link between substance misuse and worse clinical outcome. The high rate of relapse following initiation of treatment for first-episode psychosis (Robinson et al., 1999), and the increased risk of chronicity (Wiersma et al., 1998) and higher costs (Almond et al., 1994) associated with relapse, suggest that relapse prevention should be a high priority in the treatment of early psychosis. A key challenge for effective relapse prevention programmes will be to develop and implement proven interventions for comorbid psychosis and severe substance misuse (Ley & Jeffery, 2003).

REFERENCES


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References

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