Group psychoeducation for stabilised bipolar disorders: 5-year outcome of a randomised clinical trial


Background
The long-term efficacy of psychological interventions for bipolar disorders has not been tested.

Aims
This study assessed the efficacy of group psychoeducation to prevent recurrences and to reduce time spent ill for people with bipolar disorders.

Method
A randomised controlled trial with masked outcome assessment comparing group psychoeducation and non-structured group intervention during 5-year follow-up. One hundred and twenty people with bipolar disorders were included in the study and 99 completed 5-year follow-up. Time to any recurrence, number of recurrences, total number of days spent ill, frequency and length of hospitalisations were the main outcome measures.

Results
At the 5-year follow-up, time to any recurrence was longer for the psychoeducation group (log rank=9.953, $P<0.002$). The psychoeducation group had fewer recurrences ($3.86 \text{ v. } 8.37, F=23.6, P<0.0001$) of any type and they spent less time acutely ill ($154 \text{ v. } 586 \text{ days}, F=31.66, P=0.0001$). The median number of days of hospitalisation per hospitalised participant was also lower for the psychoeducation group ($45 \text{ v. } 30, F=4.26, P=0.047$).

Conclusions
Six-month group psychoeducation has long-lasting prophylactic effects in individuals with bipolar disorders. Group psychoeducation is the first psychological intervention showing such a long-term maintained efficacy in people with bipolar disorders.

Declaration of interest
E.V. has served as consultant, advisor or speaker for the following companies: AstraZeneca, Bial, Bristol-Myers, Eli Lilly, Glaxo-Smith-Kline, Jannissen-Cilag, Lundbeck, Merck-Sharp and Dohme, Novartis, Organon, Otsuka, Pfizer Inc, Sanofi-Aventis, Servier and UCB Pharmaceuticals. F.C. has served as advisory or speaker for the following companies: Astra Zeneca, Eli-Lilly, Sanofi-Aventis, Tecnifar and Shire. Funding detailed in Acknowledgements.

The evidence for the use of psychoeducation and family-focused therapy as prophylactic adjuncts to medication in people with stabilised bipolar disorders is, at present, quite conclusive.\textsuperscript{1,2} Our group, in 2003, published a randomised controlled study of recurrence prevention using group psychoeducation, reporting efficacy of this intervention in preventing mania and hypomania, depression and mixed episodes, decreasing the rates of re-hospitalisation\textsuperscript{3} and enhancing adherence.\textsuperscript{4} Several psychosocial approaches have shown different degrees of efficacy in the short term.\textsuperscript{5–8} All the cited studies examined the efficacy of several psychological interventions at a maximum of 2-year follow-up. However, there are no data on the longer-term maintenance efficacy of such treatments.

The purpose of the present article is to report an additional 3 years of follow-up for our original trial, resulting in a total of 5-year follow-up. Moreover, we report data on time spent acutely ill, a very relevant issue regarding functioning and quality of life.\textsuperscript{9,10} To our knowledge, this is the first randomised masked clinical trial assessing the longer-term efficacy of a psychological intervention as a prophylactic adjunct to pharmacotherapy in bipolar disorders.

Method

Study design
This is a continuation study of a published randomised clinical trial.\textsuperscript{3} The first part of the study had a 2-year cut-off but in the present study we have extended the assessments up to 5 years. This was a parallel two group (experimental and control) randomised, single-masked trial with 20 weeks of treatment, carried out in the Bipolar Disorders Program of the Hospital Clinic of Barcelona (Barcelona, Spain), whose research and ethics committee approved the study. This programme merges clinical care, research and education, and focuses primarily on people with the most severe bipolar I and II disorders.

The study had two phases. The treatment phase consisted of 21 weeks of randomised treatment in which all participants received standard psychiatric care with standard pharmacological treatment. The experimental group ($n=60$) received additional group psychoeducation (21 sessions on a weekly basis) and the control group ($n=60$) met every week in groups of 8–12 without special instructions from the therapist (21 meetings on a weekly basis). This design was aimed to control the variability induced by the possible supportive effect of the group reunions themselves. To avoid interferences, participants agreed not to visit any psychologist outside the centre.

The follow-up phase comprised 5 years during which all participants continued receiving standard pharmacological treatment without psychological intervention in the study centre and were assessed monthly for several outcome measures. For statistical purposes we considered the whole time of the study after randomisation (5 years and 6 months) in our follow-up. After written informed consent was signed, participants were assessed at baseline and randomised, stratifying the groups by gender, age and number of previous episodes. Randomisation was carried out using a concealed block randomisation technique.
was made by a computerised random number generator, ensuring restricted randomisation allocation and was run by an independent statistician. All the statistical analysis was run considering a start the point where the intervention began. Significance was set at \( P=0.05 \).

**Treatment**

**Standard psychiatric care**

All participants were seen by two psychiatrists (E.V. or A.B.) every 4 weeks and were specifically told to go to the centre whenever they felt any change in their mood or any other problem such as insomnia. The psychiatrists had a minimum of 4 years of clinical and research experience in bipolar disorders.

Participants received pharmacological treatment following the treatment algorithms of the Barcelona Bipolar Disorders Program. Psychiatrists and assessors were masked to the nature of the treatment given to the participants, who were told not to tell their psychiatrists if they were receiving psychoeducative treatment. The psychiatrist was not allowed to provide formal psychotherapy or specific psychoeducation beyond good clinical practice.

**Participants**

One hundred and twenty people fulfilling DSM–IV criteria for bipolar disorder type I or II, aged 18–65 years, were recruited from participants with bipolar I and II disorders enrolled in the naturalistic prospective follow-up of the Bipolar Disorders Program of the Hospital Clinic at the University of Barcelona (Spain). Inclusion criteria were a lifetime diagnosis of bipolar disorder type I or II elicited by a trained psychiatrist (E.V. or A.B.); being euthymic (Young Mania Rating Scale (YMRS)) score above or equal to 20 on the YMRS and 12 on the HRSD–17 for manic recurrence; above or equal to 12 for hypomanic recurrence; above or equal to 6, Hamilton Rating Scale for Depression (HRSD–17) score above or equal to 8 for at least 6 months immediately prior to study entry; having sufficient data on the prior course of illness collected from a prospective follow-up of at least 24 months; and written consent to participate in the study. Exclusion criteria were current Axis II comorbidity (except mental retardation) was assessed by means of the Structured Clinical Interview for DSM–IV Axis I Disorders (SCID–I; ‘mental retardation’ (IQ < 70); organic brain damage; or deafness). Individuals currently receiving any kind of psychotherapy or enrolled in any pharmacological trial were also excluded. Axis II comorbidity (except mental retardation) was not considered as an exclusion criteria.

**Psychoeducation group**

Group psychoeducation was performed according to our manual. This programme consists on 21 sessions of 90 min, each aimed at improving four main issues: illness awareness, treatment adherence, early detection of prodromal symptoms and recurrences and lifestyle regularity. The programme was in groups of 8–12, conducted by two experienced psychologists (F.C. and A.M.-A.) who had previous experience with bipolar patients (3 years) and specific training on patients’ group management. The structure of each session consisted of a 30–40 min talk on the topic of the day, followed by an exercise related to the issue (e.g. drawing a life chart, writing a list of potential triggering factors) and a discussion. Participation was encouraged.

**Control group**

In addition to standard pharmacological treatment, participants assigned to the control group received an intervention consisting of 20 weekly group meetings of 8–12 participants with the same two psychologists (F.C. and A.M.-A.). The psychologists had instructions to avoid giving any psychoeducational feedback.

Examples of this non-psychoeducative attitude could be answers like ‘What do you think?’ or ‘What is the group opinion towards this comment?’ when a question was posed to the therapist.

**Assessments**

All participants were assessed monthly by the study psychiatrists and every 2 weeks by a research assistant who had instructions to contact the psychiatrist if a recurrence was suspected. The psychiatrist and research assistant were masked to treatment and participants were requested not to reveal significant details.

All participants in this study had been enrolled in the naturalistic prospective follow-up of the Bipolar Disorders Program of Barcelona for at least 2 years. This follow-up includes assessment of recurrences, symptom checking and treatment registration, and is performed every 2 months. Baseline assessment includes the administration of the SCID–I, SCID–II, YMRS, HRSD–17, and the Holmes & Rahe inventory for stressful life-events, which are also repeated every 2 months or whenever a new episode was suspected by the psychiatrist in charge of the participant.

**Psychoeducational medication and reasons for changes were recorded. The number of hospitalisations, reasons for admission and the total number of days that the participant remained hospitalised were also recorded. Adherence was assessed by the combination of an adherence-focused interview with the individual, an adherence-focused interview with significant first-degree relatives or a partner and by analyses of plasma concentrations of mood stabilisers. This method has been extensively explained elsewhere.**

**Main outcome measures**

**Time to recurrence**

The primary outcome measure was time to recurrence. This was defined as the time (days) elapsed between baseline and the emergence of a new acute episode according to DSM–IV criteria and scores above or equal to 20 on the YMRS for manic recurrence; above or equal to 12 for hypomanic recurrence; above or equal to 17 on the HRSD–17 for depressive recurrence; and above or equal to 20 on the YMRS and 12 on the HRSD–17 for mixed recurrence.

**Number of recurrences**

The number of recurrences, separating for type of episode (manic, hypomanic, mixed or depressive) was also recorded as a secondary outcome measure.

**Time spent ill**

Another secondary outcome measure was time spent ill. We obtained this data by prospectively registering the number of days that a participant fulfilled criteria for a specific episode.

**Participant flow and drop out**

Overall, 120 euthymic out-patients were recruited at the Bipolar Disorders Program of Barcelona; 60 were randomly assigned to psychoeducation and 60 to the control group. Group rules, which were provided equally to both experimental and control participants, included the possibility of group exclusion for those individuals missing five sessions. Within the experimental group, 44 individuals out of 60 (73.3%) fully adhered to the psychoeducation programme. Sixteen people (26.6%) withdrew from the group for a number of reasons, the most common being manic recurrence (eight participants) followed by depressive recurrence (four participants) emerging during group intervention. Full participant flow is shown in Fig. 1.

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Out of 60 people in the control group, 7 (11.6%) withdrew from the weekly meetings but continued with the psychiatrist visits, the pharmacological treatment and the follow-up phase procedures. The total number of participants who withdrew was higher among the experimental group \((P < 0.05)\). There were no significant differences in the number of sessions not attended: people in the control group did not attend a mean of 3.75 (s.d.=0.87) sessions compared with 3.38 (s.d.=1.16) sessions in the treatment group.

Twenty-one participants out of 120 (25.2%) dropped out from the 5-year follow-up and had to be considered as lost to follow-up. Of these, 11 belonged to the control group and 10 to the psychoeducation group. Hence, the number of non-completers per group was almost equal.

**Baseline characteristics of treatment groups**

Both groups were comparable at baseline regarding clinical (Table 1) and therapeutic variables. Each group consisted of 38 women and 22 men with a similar number of previous episodes.

### Table 1 Comparison of clinical characteristics of participants at study entry

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Control group ((n=60))</th>
<th>Psychoeducation group ((n=60))</th>
<th>Statistical test(^a) (d.f.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male gender, (n) (%)</td>
<td>22 (36.7)</td>
<td>22 (36.7)</td>
<td>(z=0) (1)</td>
</tr>
<tr>
<td>Diagnostic subtype, bipolar I: (n) (%)</td>
<td>48 (80)</td>
<td>52 (86.7)</td>
<td>(z=0.96) (1)</td>
</tr>
<tr>
<td>Psychotic features, yes:(^b) (n) (%)</td>
<td>42 (70)</td>
<td>47 (78.3)</td>
<td>(z=1.08) (1)</td>
</tr>
<tr>
<td>Attempted suicide, yes:(^b) (n) (%)</td>
<td>19 (31.7)</td>
<td>21 (35)</td>
<td>(z=0.15) (1)</td>
</tr>
<tr>
<td>Axis II comorbidity, yes: (n) (%)</td>
<td>22 (36.7)</td>
<td>15 (25)</td>
<td>(z=1.91) (1)</td>
</tr>
<tr>
<td>Age, years: mean (s.d.)</td>
<td>34.26 (7.80)</td>
<td>34.03 (9.32)</td>
<td>(t=0.15)</td>
</tr>
<tr>
<td>Age at onset, years: mean (s.d.)</td>
<td>23.25 (7.55)</td>
<td>22.26 (6.69)</td>
<td>(t=0.75)</td>
</tr>
<tr>
<td>Total number of episodes, mean (s.d.)</td>
<td>8.81 (6.60)</td>
<td>10.31 (10.55)</td>
<td>(t=-0.93)</td>
</tr>
<tr>
<td>Number of hospitalisations, mean (s.d.)</td>
<td>2.01 (2.12)</td>
<td>1.81 (1.78)</td>
<td>(t=0.56)</td>
</tr>
</tbody>
</table>

\(^a\) \(P\) not significant in all instances.

\(^b\) Lifetime history.

**Statistical analysis**

Recurrence free curve analysis was performed using Kaplan–Meyer’s survival analysis. Cox proportional hazards regression analysis was used to assess the association between number of previous episodes and time to recurrence, independent of other predictors. The following covariates were included in every analysis: age at onset, number of previous episodes and number of previous hospitalisations. Comparison of baseline characteristics of the sample were made by a \(\chi^2\)-test for categorical variables such as gender, diagnostic subtype, polarity of first episode, history of rapid cycling, seasonal pattern, psychotic features, history of suicide attempts, Axis II comorbidity and type of treatment received using the Fisher \(z\)-test when needed and the \(t\)-test for quantitative variables. The separate analysis at follow-up of the number of participants who relapsed for each condition was by a \(\chi^2\)-test. The comparison of the mean number of recurrences during the treatment and the follow-up phase was made using the ANCOVA model including age at onset, number of previous episodes and number of previous hospitalisations as covariates. Statistical significance was set at \(P < 0.05\).
Results

Time to recurrence

Survival analysis of participants remaining in remission is shown in Fig. 2. The event curves for the control and treatment groups were significantly different for time to any recurrence (log rank=9.953, P<0.002). Number of previous episodes was not associated with time to recurrence (95% CI 0.013–0.461, P=0.49 by Cox model).

Number of recurrences

Over the 5-year follow-up period, people in the psychoeducation group had less recurrences than those in the non-psychoeducation group (3.86 v. 8.37, F=23.6, P<0.0001). When analysing each type of episode (mania, hypomania, mixed or depression), individuals in the psychoeducation group had a lower number of recurrences. When comparing effect sizes (Cohen's d) between the 2-year and the 5-year assessments, we could see how effect sizes did not decrease with the passing of time. This was true for any episode (0.79 v. 0.87), mania (0.40 v. 0.57), hypomania (0.27 v. 0.42) and mixed episodes (0.34 v. 0.61). As for depression, the effect size at 5 years was slightly inferior to the 2-year effect size but still in the range of large effect (0.91 v. 0.80).

Time spent ill

People in the psychoeducation group spent much less time acutely ill than those in the non-psychoeducation group. This is mainly owing to the dramatic differences accounting for time spent with depression, but it is also true for every type of episode (Table 2).

Hospitalisations

During the treatment phase, 8 individuals out of 60 (13.3%) in the control group were hospitalised owing to recurrence compared with 9 out of 60 (16.1%) in the treatment group (non-significant difference). At the end of the 5-year follow-up period, 24 individuals out of 60 (40%) in the control group had been hospitalised v. 17 out of 56 (30.4%) in the treatment group (P=0.278).

The number of hospitalisations at the 5-year follow up for the psychoeducation group was 0.24 (s.d.=0.52) admissions v. 0.59 (s.d.=0.96) in the control group (F=5.34, P=0.023). The median number of days of hospitalisation per hospitalised participant was also lower for people in the psychoeducation group (45 v. 30) (F=4.26, P=0.047).

Treatment adherence

As for treatment adherence, both groups were comparable at entry, with rates of poor adherence reaching 25.9% (15 individuals out of 58) in the control group and 30.5% (18 individuals out of 59) in the psychoeducation group at baseline (χ²=0.312, P=0.577). At 5-year follow-up there were no differences regarding adherence, with rates of poor adherence of 14.3% and 14% respectively (χ²=0.002, P=0.967).

Discussion

Although several psychological interventions tested on people with bipolar disorders report short-term prophylactic efficacy (up to 1 year), the use of 6-month group psychoeducation is the first one to show efficacy in the long term. In fact, other psychological therapies may lose effect in the middle term. This may be the case for cognitive–behavioural relapse prevention therapy as this intervention showed prophylactic efficacy in the short term, 1 year, but had no significant effect in relapse reduction after 2 years other than the number of days being ill. Thus, the present study might change the way in which psychological interventions for bipolar disorders are considered, as one of the traditional criticisms that psychotherapy for severe mental disorders has received in the past is that its effect may progressively disappear once the intervention has finished. In contrast, our study shows how a time-limited, single intervention, without the help of any booster session, is able to bring about a major improvement in the outcome of bipolar disorders in the long term (5 years). Thus, group psychoeducation may enhance behavioural and attitudinal changes that seem to be maintained over the years.

The study has several limitations. Generalisability is limited by the exclusion of individuals with severe comorbidities and substance misuse, and by the fact that this was a single-site trial. The outcome was measured as the presence of full episodes and subthreshold presentations were not included in the definition. Masking may also be an issue in this sort of study, as some participants might have made comments to the psychiatrists that would indirectly indicate their treatment allocation.

All our 5-year follow-up data confirm the 2-year follow-up findings: the number of bipolar episodes is much lower for the psychoeducation group over time. In the long term, group psychoeducation may prevent all sorts of episodes including mania. Interestingly enough, when comparing 2-year and 5-year follow-up, prevention of mania seems to be more powerful in the longer term (Cohen's d effect size=0.4 v. 0.57), probably as a result of the fact that changes concerning regularity of habits and, especially, early detection, may be more noticeable in the longer term. Psychoeducation goes far beyond a mere transmission of information and should be considered as disease-management training more than anything else. According to this view, it is completely logical that its effects are more noticeable with the passing of time (Cohen's d effect size for prevention of all sorts of episodes=0.79 v. 0.87).
On the other hand, it should be emphasised that the psychoeducation programme and the subsequent follow-up was highly accepted by our participants, as most of them did not withdraw and engaged in the follow-up visits, as demonstrated by the fact that just 20% of the participants had to be considered as lost to follow-up at the 5-year end-point.

It is worth mentioning the effect of psychoeducation on hospitalisations. We could not find any difference regarding the number of participants who required hospitalisation, either at 2- or at 5-year follow-up. However, a significant difference was found regarding the number of hospitalisations per individual, lower for people in the psychoeducation group, which may mean that through psychoeducation we could be avoiding the ‘revolving door’ phenomena (the frequent rehospitalisation of a subgroup of people). This effect has not been described by any other psychological intervention in bipolar disorders. Simon et al8 reported how their ‘systematic care program’ for bipolar disorders had a significant effect on reducing the frequency and severity of mania at a worthy cost but could not find any efficacy on hospitalisations.

We have observed a significant effect of group psychoeducation on time spent ill, whereas individuals in the non-psychoeducation group spent up to 30% of their time acutely ill; for people in the psychoeducation group, this was approximately just 8%. Results for the control group were slightly different from those reported by Judd et al17,18 as people in that study spent between 47% and 54% of time being ill; depending on the bipolar subtype, this difference was somewhat predictable if we keep in mind that we were using quite narrower recurrence criteria. But what is more striking is the benefit regarding time to keep in mind that we were using quite narrower recurrence criteria. Furthermore, in a secondary analysis of 2-year data, people with comorbid Axis II disorders significantly improved with psychoeducation.24 In a recent trial, cognitive–behavioural therapy was not effective in preventing further episodes in people with bipolar disorders in general, and particularly in those with comorbidities.25 The improved understanding of the disease and increase of illness awareness may be another factor underlying the positive effects of psychoeducation, especially regarding empowerment for decision-taking, which may include dose increase of certain medications as soon as some warning sign is detected, if the psychiatrist is not available at that point.13 The better the knowledge about the disease is, the more likely are the patients to seek appropriate help. Simon et al8 reported that ‘short-term (up to 5 years) in bipolar disorders, significantly reducing mood and reported mood changes day in day out during a 3-week period after the session devoted to mood checking. Most of these changes were transient, but they deserve careful monitoring; too often, psychotherapies are seen as free of adverse events and this might be another factor underlying the positive effects of psychoeducation.

<table>
<thead>
<tr>
<th></th>
<th>Mean (s.d.)</th>
<th>Psychoeducation group (n=50)</th>
<th>Mean (s.d.)</th>
<th>Fisher’s F</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>398.55 (364.17)</td>
<td>364.17 (165.47)</td>
<td>35.46</td>
<td>&lt;0.001</td>
<td></td>
</tr>
<tr>
<td>Mania</td>
<td>62.17 (114.1)</td>
<td>26.1 (42.31)</td>
<td>4.39</td>
<td>0.035</td>
<td></td>
</tr>
<tr>
<td>Hypomania</td>
<td>60.35 (106.41)</td>
<td>13.8 (24.92)</td>
<td>8.84</td>
<td>0.004</td>
<td></td>
</tr>
<tr>
<td>Mixed episode</td>
<td>66.29 (100.39)</td>
<td>20.54 (41.1)</td>
<td>9.12</td>
<td>0.003</td>
<td></td>
</tr>
<tr>
<td>Depression</td>
<td>586.45 (510.91)</td>
<td>153.72 (184.38)</td>
<td>36.49</td>
<td>0.000</td>
<td></td>
</tr>
</tbody>
</table>
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