Paliperidone ER in the Treatment of Borderline Personality Disorder: A Pilot Study of Efficacy and Tolerability

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Antipsychotics are recommended for the treatment of impulsive dyscontrol and cognitive perceptual symptoms of borderline personality disorder (BPD). Three reports supported the efficacy of oral risperidone on BPD psychopathology. Paliperidone ER is the metabolite of risperidone with a similar mechanism of action, and its osmotic release reduces plasmatic fluctuations and antidopaminergic effects. The aim of this study is to evaluate efficacy and safety of paliperidone ER in BPD patients. 18 outpatients with a DSM-IV-TR diagnosis of BPD were treated for 12 weeks with paliperidone ER (3–6 mg/day). They were assessed at baseline, week 4, and week 12, using the CGI-Severity item, the BPRS, the HDRS, the HARS, the SOFAS, the BPD Severity Index (BPDSI), and the Barratt Impulsiveness Scale (BIS-11). Adverse events were evaluated with the DOTES. Paliperidone ER was shown to be effective and well tolerated in reducing severity of global symptomatology and specific BPD symptoms, such as impulsive dyscontrol, anger, and cognitive-perceptual disturbances. Results need to be replicated in controlled trials.

1. Introduction

Borderline personality disorder (BPD) is characterized by a pervasive pattern of instability in interpersonal relationships, self-image, and affects as well as impulsive dyscontrol that begins by early adulthood and appears in a variety of contexts [1].

Although psychotherapy plays a significant role in the treatment of BPD, focusing on maladaptive personality traits and interpersonal relationship patterns, pharmacotherapy is indicated by the treatment guidelines of the American Psychiatric Association [2, 3] to manage vulnerability traits, state symptoms, and acute relapses.

The availability of second-generation antipsychotics with a favourable tolerability profile has offered recent treatment options in the management of BPD patients. In particular, atypical antipsychotics are associated with fewer extrapyramidal adverse effects, a lower risk of tardive dyskinesia, and an improvement in cognitive functions [4–6].

Newer antipsychotics promise to be also more efficacious than traditional neuroleptics, probably due to their dual mechanism of action, targeting dysfunctions of both serotonergic and dopaminergic systems. These drugs produce their effects primarily on cognitive perceptual disturbances, such as transient paranoid or dissociative symptoms but also on symptoms of depression, anxiety, impulsivity, and aggressiveness.

Three reports supported the efficacy of oral risperidone on a large spectrum of BPD psychopathology, including affective instability, disturbed relationships, impulsivity, aggressiveness, hostility, and cognitive-perceptual impairment [7–9]. Paliperidone ER (9-hydroxyrisperidone) is a benzisoxazole derivative and the main active metabolite of risperidone. The efficacy and tolerability profile of this molecule is modulated both by drug-membrane interaction and drug-receptor action [10]. Paliperidone is an antagonist at dopamine receptor type 2 and serotonin receptor subtype 2A, with a greater affinity for 5HT2A receptor blockade relative to D2 receptor blockade. This drug has also some affinity for H1-histaminergic receptors and for α1- and α2-adrenergic receptors, whereas it lacks significant interactions with muscarinic receptors and with β1- and β2-adrenergic receptors [11–13]. Paliperidone does not undergo significant hepatic metabolism and is unlikely to induce drug-drug interactions.
Paliperidone extended release (ER) is compounded using osmotic-controlled dose-release system, a technology that reduces plasma level fluctuations of the immediate-release formulation [24–26]. This mechanism is expected to provide several advantages: a stable plasmatic level allows to avoid initial dosage adjustments and lowers the risk of antiparkinsonian adverse effects. Another benefit is represented by once-daily administration, that simplifies treatment regimen and may improve patients’ adherence.

Several clinical trials evaluated effectiveness of paliperidone ER in treating schizophrenia [15, 22, 27, 28]. In particular, recent studies ascertained significant improvements in positive symptoms, negative symptoms, depression and anxiety, uncontrolled hostility/excitement, and disorganized thoughts [27, 29]. At the moment, no data regarding the use of paliperidone ER in the treatment of personality disorders have been published.

The present paper is a pilot study with the aim of testing the efficacy and tolerability of paliperidone ER in patients with borderline personality disorder.

2. Materials and Methods

Consecutive outpatients aged between 18 and 60 years, who received a diagnosis of borderline personality disorder, were included in this study. Patients attended the Service for Personality Disorders, Unit of Psychiatry 1, Department of Neurosciences, University of Turin, Italy. Diagnosis of borderline personality disorder was made by an experienced clinician (S.B.) using DSM-IV-TR criteria [1] and performing the Structured Clinical Interview for DSM-IV Axis II Disorders [30].

Patients were required to fulfill exclusion criteria: a lifetime diagnosis of dementia, delirium, or other cognitive disorders; schizophrenia or other psychotic disorders; bipolar disorders; a co occurring major depressive episode; substance/alcohol abuse in the last six months; hyperprolactinemia at baseline.

Administration of psychotropic medications and/or psychotherapy in the three months before recruitment was excluded too.

Female patients in childbearing age were excluded if they were not using adequate birth control methods (according to the judgment of the clinician).

Each patient participated voluntarily in the study after providing a written informed consent. Declaration of Helsinki guidelines were followed, and approval by local Ethics Committee was obtained.

Patients included in the study were treated for 12 weeks with open-label paliperidone ER 3–6 mg/day. No other psychotropic drug or psychological intervention was allowed during the trial.

Participants in the study were repeatedly assessed (at baseline, week 4, and week 12) using the following assessment instruments:

1. the Clinical Global Impression Scale-Severity item (CGI-S) [31];
2. the Brief Psychiatric Rating Scale (BPRS) [32];
3. the Hamilton Depression Rating Scale (HAM-D) [33];
4. the Hamilton Anxiety Rating Scale (HAM-A) [34];
5. the Social Occupational Functioning Assessment Scale (SOFAS) [35];
6. the Barratt Impulsiveness Scale-version 11 (BIS-11) [36];
7. the Borderline Personality Disorder Severity Index (BPDSI) [37].

The BPDSI is a semistructured clinical interview assessing frequency and severity of BPD-related symptoms. The interview consists of eight items scored on 10-point frequency scales (0 = never; 10 = daily), including “abandonment”, “interpersonal relationships”, “impulsivity”, “parasuicidal behavior”, “affective instability”, “emptiness”, “outbursts of anger”, “dissociation and paranoid ideation”, and one item scored on a 4-point severity scale, concerning “identity”. The BPDSI showed excellent reliability coefficients and good validity indices according to Arntz et al. [37].

The BIS-11 is a 30-item self-report questionnaire assessing the personality trait of impulsivity on a 4-point Likert Scale [38]. Higher scores for each item indicate higher levels of impulsivity. Twelve items are reverse-scored, in order to avoid response sets. The BIS-11 showed adequate reliability and construct validity in both US [39] and Italian [40] samples.

Assessment was performed by an investigator (P.B.) who was unaware of the dosing strategy. Prior to this study, the interviewer received training sessions on the BPDSI.

Adverse effects were assessed using the Dosage Record and Treatment Emergent Symptoms Scale (DOTES) [41].

Serum prolactin level was measured at baseline, after four and twelve weeks of treatment. Blood samples were collected in fasting patients two hours after they woke up. Hyperprolactinemia was defined as a level of serum prolactin ≥ 20 ng/mL in males and ≥ 25 ng/mL in females [42, 43]. Body weight was measured at baseline and endpoint. Weight gain at least 7% of baseline was considered significant [44].

Statistics were performed on the scores of each rating scale with the analysis of variance (ANOVA) for repeated measures with Bonferroni correction for multiple comparisons (software system SPSS, version 17.0, SPSS Inc., Chicago, Ill., 2008). P values were considered significant when ≤ 0.05.
3. Results

Initial sample was made of eighteen patients. They were five males and thirteen females, with a mean age ± SD = 24.3 ± 5.4. Fourteen patients (77.8%) completed the trial period. Four patients (1 male and 3 females, 22.2%) dropped out in the first four weeks of treatment: three patients due to noncompliance, one female patient for hyperprolactinemia. The fourteen patients who completed the trial had a mean age of 25.2 ± 4.9 years; they were four males and ten females. The mean ± SD daily dose of paliperidone ER was 4.8 ± 1.5 mg/day.

Results of ANOVA with Bonferroni correction applied to rating scales scores are reported in Tables 1, 2, and 3.

A statistically significant improvement was found for CGI severity item (P = 0.001), BPRS mean score (P = 0.001), SOFAS mean score (P = 0.001), BIS-11 mean score (P = 0.005), BPDSI total score (P = 0.001), and items “impulsivity” (P = 0.001), “outbursts of anger” (P = 0.01), and “dissociative symptoms/paranoid ideation” (P = 0.002). On the contrary, there were no significant changes of HAM-D and HAM-A mean score and BPDSI items “abandonment,” “interpersonal relationships,” “identity,” “parasuicidal behaviour,” “affective instability,” and “emptiness.”

As far as paliperidone tolerability is concerned, only one case of severe side effect (hyperprolactinemia) inducing treatment discontinuation was observed in the initial group of eighteen subjects (5.6%). Adverse effects recorded in the final sample of fourteen patients were mild to moderate and included insomnia (n = 3, 21.4%), gastrointestinal symptoms (n = 2, 14.3%), agitation (n = 1, 7.1%), and extrapyramidal symptoms (tremor) (n = 1, 7.1%). Six patients (42.9%) had
at least one adverse effect during treatment period. No cases of significant weight gain (≥7% of baseline) were recorded. Mean weight gain ± SD was 0.7 ± 0.8 kg.

4. Discussion

Findings of this pilot study of paliperidone ER in patients with borderline personality disorder showed significant clinical and functional improvements, as measured with CGI severity item, BPRS, SOFAS, Barratt Impulsiveness Scale, BPDSI total score, and items “impulsivity,” “outbursts of anger,” and “dissociative symptoms/paranoid ideation.”

Because of the lack of trials of paliperidone in the treatment of BPD, these results can only be compared with data concerning the effects of other second-generation antipsychotics in this personality disorder. A particular interest is raised by clinical trials of risperidone, a drug with a chemical structure very similar to paliperidone (actually paliperidone is the active metabolite of risperidone).

The significant changes of the rating scale scores after treatment, in particular the reduction of the CGI-S, BPRS, and total BPDSI scores and the increase of the SOFAS score, indicated that paliperidone ER was efficacious on a large spectrum of BPD psychopathology. The improvement of global symptomatology and social functioning was reported by previous open and controlled studies concerning BPD treatment with atypical antipsychotics, such as risperidone [8, 45], clozapine [46, 47], olanzapine [48, 49], quetiapine [50–56], and aripiprazole [57–60].

In addition to the improvement of global psychopathology, paliperidone ER produced in our patients significant effects on specific symptom dimensions, such as dyscontrol of impulsivity and cognitive perceptual distortions. Measures with the clinician-rated BPDSI and the self-report BIS-11 were concordant and indicated that paliperidone was effective on impulsive-behavioral dyscontrol symptoms and outbursts of anger. Efficacy of second generation antipsychotics on this cluster of symptoms was already reported by several trials of olanzapine [48, 61, 62] and quetiapine [50–53, 55, 56]. Data concerning the effect on impulsivity of other antipsychotics are less consistent. This effect was never reported in trials of clozapine, and it was found in two pilot studies of risperidone [9, 45] and in one pilot study of aripiprazole in treatment refractory BPD patients [60]. Controlled trials of these two drugs did not list improvement of impulsive behaviors among their findings [7, 58]. Discordant data on this issue, particularly concerning risperidone, suggest the need to replicate our initial data on paliperidone, before drawing any conclusion for clinical practice.

Measures with the BPDSI item “dissociation and paranoid ideation” in our sample provided initial evidence that paliperidone ER was effective in treating cognitive and perceptual disturbances, such as ideas of reference, illusions, and dissociative symptoms. The efficacy of second-generation antipsychotics on cognitive and perceptual symptoms was considered in rather few studies of BPD patients. It was reported in several trials with clozapine [46, 47, 63], in two studies with risperidone [7, 9], one with olanzapine [64], one with ziprasidone [65], and one with aripiprazole [60]. Further investigations are needed to confirm this antipsychotic effect in BPD.

Concerning tolerability, adverse effects in the final sample of BPD patients were insomnia, gastrointestinal disturbances, agitation, and extrapyramidal symptoms. Side effects were mild to moderate in severity. Only one case

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Table 3: Results of analysis of variance (ANOVA) for repeated measures with Bonferroni correction for multiple comparisons, performed with BPDSI items “parasuicidal behaviour,” “affective instability,” “emptiness,” “outbursts of anger,” and “dissociation and paranoid ideation” and with BIS-11 score.

| Variable                      | T0   | SD  | SE  | P    |
|-------------------------------|------|-----|-----|------|
| Parasuicidal behaviour        | T0   | 3.46| 2.05| 0.48 | 0.101|
|                               | T1   | 2.78| 2.03| 0.48 | 0.101|
|                               | T2   | 2.08| 1.53| 0.36 | 0.083|
| Affective instability         | T0   | 5.19| 1.26| 0.30 | 1.000|
|                               | T1   | 4.54| 0.87| 0.20 | 0.101|
|                               | T2   | 4.42| 0.86| 0.20 | 0.101|
| Emptiness                     | T0   | 6.41| 1.73| 0.41 | 0.101|
|                               | T1   | 6.22| 1.62| 0.38 | 0.101|
|                               | T2   | 6.45| 1.43| 0.34 | 0.101|
| Outbursts of anger            | T0   | 7.38| 1.05| 0.25 | 0.101|
|                               | T1   | 6.77| 0.72| 0.17 | 0.101|
|                               | T2   | 6.38| 0.72| 0.20 | 0.101|
| Dissociation and paranoid ideation | T0   | 5.98| 1.04| 0.25 | 0.101|
|                               | T1   | 5.85| 1.04| 0.25 | 0.101|
|                               | T2   | 4.84| 1.18| 0.28 | 0.101|
|                               | T0   | 71.17| 7.79| 1.83 | 0.101|
|                               | T1   | 65.17| 6.31| 1.49 | 0.101|
|                               | T2   | 62.28| 6.43| 1.52 | 0.101|
of hyperprolactinemia induced treatment discontinuation. It is noticeable that we found a low incidence of hyperprolactinemia, compared with previous reports in patients with schizophrenia [20–22, 66]. No cases of significant weight gain were recorded, in contrast with a recent estimate of 8% in patients with schizophrenia or bipolar disorder [44]. A possible explanation for these differences is that we used low to moderate doses of paliperidone (3–6 mg/day). Usually, doses of paliperidone in patients with schizophrenia were higher, and some authors retained that hyperprolactinemia induced by paliperidone was dose related [21, 67]. The pattern of side effects in our sample was generally concordant with investigations of paliperidone in the treatment of schizophrenia [15, 27, 29, 67–69].

5. Conclusions

Initial findings were promising and indicated that low to moderate doses of the new antipsychotic paliperidone ER were efficacious in reducing severity of global symptomatology and social impairment and in treating two specific clusters of BPD symptoms, impulsive behavioral dyscontrol, and cognitive-perceptual disturbance. The level of tolerability recorded in our patients was favourable, with low incidence of hyperprolactinemia.

Limitations of the present study were the small sample size and the lack of a control group. However, this paper was aimed to provide initial data on the use of paliperidone ER in BPD treatment and promote larger trials with a double-blind controlled design.

Conflict of Interests

The authors have no relevant affiliations or financial involvement with any organization or entity with a financial interest in or financial conflict with the subject matter or materials discussed in the paper.

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