Efficacy of quetiapine in patients with bipolar I and II depression: a multicenter, prospective, open-label, observational study

Jong-Hyun Jeong¹
Won-Myong Bahk¹
Young Sup Woo¹
Ho-Jun Seo¹
Seung-Chul Hong¹
Duk-In Jon²
Kyung Joon Min³
Bo-Hyun Yoon⁴

¹Department of Psychiatry, College of Medicine, The Catholic University of Korea, Seoul, ²Department of Psychiatry, College of Medicine, Hallym University, Anyang, ³Department of Neuropsychiatry, College of Medicine, Chung-Ang University, Seoul, ⁴Naju National Hospital, Naju, Korea

Purpose: To evaluate and compare the therapeutic efficacy of quetiapine in bipolar I and II depression patients in the clinical setting.

Patients and methods: This was an 8-week, multicenter, open-label, observational study for bipolar depression. The dosage of quetiapine was flexible, and concomitant medications were permitted on clinician's judgments. A total of 1097 patients were enrolled, and 764 bipolar depression patients who exhibited good therapeutic compliance ($>75\%$ compliance rate) were analyzed.

Results: Clinical Global Impression – Bipolar scale and Montgomery–Asberg Depression Rating Scale scores were significantly improved at weeks four and eight compared with the baseline scores. At the end of the 8-week study, the response rate was 58.9%, and the remission rate was 42.1%. However, there were no significant differences in the response and remission rates between bipolar I and II disorder (BD-I and BD-II) patients (response rate 60.1% versus 56.3%; remission rate 44.5% versus 37.0%). Montgomery–Asberg Depression Rating Scale score at baseline ($\beta = 0.612$, $P < 0.001$), duration of current episode ($\beta = -0.152$, $P = 0.001$), and presence of remission on previous episode ($\beta = 0.111$, $P = 0.012$) were significantly associated with improvements in depressive symptoms. Fatigue (16.0%), somnolence (14.9%), and manic/hypomanic switching (0.6% at week four, 0.3% at week eight) were observed throughout the study period.

Conclusion: The results of this study suggest that quetiapine improves depressive symptoms in BD-I and BD-II patients with a minimal incidence of manic switching. The therapeutic efficacy of quetiapine increased with time. Quetiapine could be an effective and safe modality for the treatment of BD-I and BD-II.

Keywords: bipolar depression, quetiapine, therapeutic efficacy, observational study

Introduction

The lifetime prevalence of bipolar disorder (BD) is approximately 3%–5%, and 80%–90% of all BD patients experience depressive symptoms.^{1,2} In patients with BD, depressive periods tend to last longer than manic periods.³ In addition, a study that examined the symptoms experienced by BD patients over a 6-month period reported that approximately 60% of BD patients experienced depressive symptoms, which is two-fold higher than the incidence of manic symptoms.⁴ In other words, the period of depressive symptoms lasts longer, and the incidence of depressive symptoms is higher than that of manic symptoms. However, bipolar depression does not respond well to treatment; it frequently progresses to recurrent episodes of mood disorder, making it difficult to treat. The most widely used psychopharmacologic strategies for the treatment of bipolar depression include mood stabilizer monotherapy, a combination of mood stabilizers, and combination therapy with antidepressants. However, most recently,

Correspondence: Won-Myong Bahk
Department of Psychiatry, The Catholic University of Korea, 62, Yeoido-dong, Yeongdeungpo-gu, Seoul, Korea
Tel +82 2 3779 1051; +82 2 3779 1735
Fax +82 2 780 6577
Email wmbahk@catholic.ac.kr

monotherapy with atypical antipsychotics, combination therapy with antidepressants, and combination therapy with mood stabilizers have been newly recommended.^{5–10}

Atypical antipsychotics are approved for the treatment of schizophrenia and psychotic disorders. However, due to their mechanism of action, they have been widely used to improve depressive symptoms in patients with psychotic disorders and BD.⁷ In particular, quetiapine has been known to exert antidepressant effects in patients with psychotic disorders and depressive symptoms.¹¹ In two recent large-scale, double-blind, randomized controlled trials, quetiapine monotherapy was proven to have significantly higher antidepressant effects than placebo on bipolar I and II disorder (BD-I and BD-II), rapid cycling BD, and depression with anxiety symptoms. Moreover, the risk of inducing manic symptoms with quetiapine monotherapy was not higher than that with placebo. These results indicate that quetiapine could be used as a first-line drug for the treatment of bipolar depression.^{12,13} Quetiapine was the first antipsychotic to be approved as a monotherapeutic agent for bipolar depression, not only by the US Food and Drug Administration but also by the Korean Food and Drug Administration.

However, the previous data were obtained using highly selective patients with many prescription restrictions, which could be different from actual clinical situations. In addition, no published large-scale data have evaluated the efficacy of quetiapine in BD patients in clinical settings.

Therefore, this observational study was conducted to evaluate the therapeutic efficacy of quetiapine, which was administered for 8 weeks to patients with BD-I, most recent episode depressed, and BD-II, most recent episode depressed.

Material and methods

Subjects

This multicenter study was conducted at 98 sites including university hospitals and mental hospitals in Korea from June 1, 2007 to February 28, 2008. The study subjects were older than 18 years, and all subjects were diagnosed with BD-I, most recent episode depressed, or BD-II, most recent episode depressed, according to the diagnostic criteria of the Diagnostic and Statistical Manual of Mental Disorder, Fourth Edition, Text Revision (DSM-IV-TR).¹ The study procedure was explained to the patients who participated in this study, and only patients who submitted a written consent form were allowed to participate. Because this study was an observational study conducted in an actual clinical setting, special exclusion criteria were absent, excluding rapid

cycling, and even patients with chronic diseases were allowed to participate in the study, provided they were stable based on their past history and physical examination. However, patients with severe, unstable, or acute diseases as well as those with chronic physical illnesses were excluded from the study. The study protocol was approved by an institutional review board.

Medication administration

In this study, the dose of quetiapine was flexible depending on the clinical situation. In addition, the administration of concomitant medications was not restricted. Furthermore, even medications that had been continuously administered under stable conditions without a change in dose for the treatment of physical illnesses were allowed during the study period. To evaluate therapeutic compliance, the actual uptake of drugs was examined at weeks four and eight.

Evaluation of therapeutic efficacy and compliance

The Clinical Global Impression – Bipolar (CGI-BP) scale and Montgomery–Asberg Depression Rating Scale (MADRS) were used to evaluate therapeutic efficacy.^{14,15} Therapeutic efficacy was evaluated prior to the administration of quetiapine and after 4 and 8 weeks of administration.

As the main outcome measure of therapeutic efficacy, the CGI-BP and MADRS scores were compared at baseline and after 8 weeks of drug administration. In addition, the response rate ($\geq 50\%$ reduction in the total MADRS value) and remission rate (≤ 10 in the total MADRS value) were examined at the 4- and 8-week time points. In addition, differences in therapeutic efficacy between the BD-I and BD-II patients were examined. In addition, the numbers of patients in each CGI-BP subgroup at baseline and after 4 and 8 weeks were compared.

To evaluate therapeutic compliance, the subjective level of therapeutic compliance was examined after 4 and 8 weeks of drug administration. The patients were divided by compliance rate as follows: $>75\%$, $50\%–75\%$, $25\%–50\%$, and $<25\%$.

Data analysis

Of the patients who completed the baseline evaluation prior to drug administration, patients who took quetiapine more than once and patients who were evaluated at least once after drug administration were included in the intent-to-treat group. The data were analyzed by the last observation carried forward method. Regarding the statistical methods, repeated

measures of analysis of variance, Cochran–Mantel–Haenszel test, and the multiple linear regression test were applied. The independent *t*-test and Chi-squared test were used for comparisons of BD-I and BD-II patients, and all statistical analyses were performed with SPSS® 18.0 (IBM Corporation, Armonk, NY). The level of statistical significance was set at $P < 0.05$.

Results

Demographic data and clinical characteristics

A total of 1097 BD patients were recruited for participation in this study, and 1058 patients (96.4%) completed the entire study. To elucidate the true efficacy of quetiapine, patients who exhibited poor therapeutic compliance (<75% compliance rate) after 4 or 8 weeks were excluded ($n = 333$). Thus, the study finally included 764 BD patients (mean age: 42.7 ± 15.6 years) consisting of 526 patients with BD-I (mean age: 41.1 ± 14.7 years) and 238 patients with BD-II (mean age: 46.3 ± 16.9 years); the difference in age between the BD-I and BD-II groups was significant ($P < 0.001$).

The mean duration of current depressive episode was 6.3 ± 8.8 weeks and duration of current episode was 7.9 ± 12.1 weeks in BD-II patients, which was significantly longer than in BD-I patients (5.6 ± 6.7 weeks; $P = 0.006$).

Psychiatric comorbidity was detected in 3.8% of the subjects (BD-I 2.7% versus BD-II 6.3%; $P = 0.015$). Approximately 33% of BD patients ($n = 250$) were admitted (BD-I 35.7% versus BD-II 26.1%; $P = 0.008$), and mean number of admissions was 3.1 ± 1.9 (BD-I 3.1 ± 1.9 versus BD-II 3.2 ± 2.1 ; not significant) (Table 1).

Quetiapine dose and level of psychotropic concomitant medications

The mean dose of quetiapine was initially 197.5 ± 182.5 mg/day, and it was gradually increased to 349.0 ± 231.5 mg/day by week eight. During all observation periods, the dose of quetiapine was significantly higher in the BD-I patients than in BD-II patients ($P < 0.001$) (Table 2).

The combined administration of psychotropic agents was abundant in BD patients, and the difference in total concomitant medications administered between the BD-I and BD-II groups was not significant. Nonetheless, mood stabilizers (61.0% versus 35.7%; $P < 0.001$) were administered more frequently in BD-I patients, but anxiolytics (22.8% versus 31.5%; $P = 0.001$), antidepressants (12.4% versus 42.47%; $P < 0.001$), and hypnotics/sedatives (8.0% versus 13.4%;

Table 1 Demographic data and clinical characteristics of patients with bipolar I and II disorder

	Total (n = 764)	BD-I (n = 526)	BD-II (n = 238)	P-value
Sex				
Male	298 (39.0)	210 (39.9)	88 (37.0)	NS
Female	466 (61.0)	316 (60.1)	150 (63.0)	NS
Age (years)	42.7 ± 15.6	41.1 ± 14.7	46.3 ± 16.9	<0.001*
Psychiatric comorbidity	29 (3.8)	14 (2.7)	15 (6.3)	0.015**
Admission state	250 (32.7)	188 (35.7)	62 (26.1)	0.008**
Duration of current episode (weeks)	6.3 ± 8.8	5.6 ± 6.7	7.9 ± 12.1	0.006*
Previous history of mood episode	563 (73.7)	401 (76.2)	162 (68.1)	0.018**
Illness duration (years)	17.2 ± 17.1	17.7 ± 16.5	16.1 ± 18.5	NS
Previous history of admission	61.1	69.4	40.7	<0.001**
Number of admissions	3.1 ± 1.9	3.1 ± 1.9	3.2 ± 2.1	NS
Previous psychotic features	47.2	57.1	22.8	<0.001**

Notes: * $P < 0.05$ by independent *t*-test; ** $P < 0.05$ by Chi-squared test; data expressed as *n* (%) or mean $\pm$ standard deviation.

Abbreviations: BD-I, bipolar I disorder; BD-II, bipolar II disorder; NS, not significant.

$P = 0.018$) were more frequently coadministered to BD-II patients (Table 3).

Evaluation of therapeutic efficacy

The CGI-BP scores obtained at baseline, week four, and week eight were 4.3 ± 0.9 , 3.5 ± 0.9 , and 2.7 ± 0.9 , respectively. The values were significantly lower at week four and eight than at baseline ($P < 0.001$). However, the difference between BD-I and BD-II patients was not significant (Figure 1).

Moreover, when the entire patient group was divided according to disease severity and the subgroups were further divided according to CGI-BP score, the number of patients with a higher level of symptom severity was significantly

Table 2 Daily dosages of quetiapine in patients with bipolar I and II disorder

	Total (n = 764) (%)	BD-I (n = 526) (%)	BD-II (n = 238) (%)	P-value*
Day 1	197.5 ± 182.5	219.0 ± 189.8	149.6 ± 155.4	<0.001
Week 1	250.5 ± 208.9	280.7 ± 216.5	183.3 ± 173.5	<0.001
Week 2	301.2 ± 232.9	336.5 ± 241.2	222.4 ± 191.4	<0.001
Week 4	327.7 ± 228.0	359.8 ± 232.7	256.0 ± 199.7	<0.001
Week 8	349.0 ± 231.5	376.1 ± 239.4	289.1 ± 201.0	<0.001

Notes: * $P < 0.001$ by independent *t*-test; data expressed as mean $\pm$ standard deviation.

Abbreviations: BD-I, bipolar I disorder; BD-II, bipolar II disorder.

Table 3 Concomitant psychotropic medications in patients with bipolar I and II disorder

	Total (n = 764)	BD-I (n = 526)	BD-II (n = 238)	P-value
Total number of subjects	559 (73.2)	182 (76.5)	377 (71.7)	NS
Mood stabilizers	406 (53.1)	85 (35.7)	321 (61.0)	<0.001*
Anxiolytics	195 (25.5)	75 (31.5)	120 (22.8)	0.011*
Antidepressants	166 (21.7)	101 (42.4)	65 (12.4)	<0.001*
Antipsychotics	82 (10.7)	29 (12.2)	53 (10.1)	NS
Hypnotics and sedatives	74 (9.7)	32 (13.4)	42 (8.0)	0.018*
Number of concomitant medications				
One	288 (37.7)	207 (39.4)	81 (34.0)	NS
Two	189 (24.7)	118 (22.4)	71 (29.8)	0.028**
Three	72 (3.4)	50 (9.5)	22 (9.2)	NS
Four or more	10 (1.3)	2 (0.4)	8 (3.4)	0.001*

Notes: * $P < 0.001$ by Chi-squared test; ** $P < 0.05$ by Chi-squared test; data represents n (%).

Abbreviations: BD-I, bipolar I disorder; BD-II, bipolar II disorder; NS, not significant.

decreased after 4 and 8 weeks compared to that at baseline ($P < 0.001$). Such differences were also significant when the groups were compared after 4 and 8 weeks ($P < 0.001$; Table 4).

When comparing the severity of symptoms at baseline, week four, and week eight by MADRS, there was a statistically significant decrease in severity for both the total MADRS value and every subscale score ($P < 0.001$). In addition, the severity of symptoms was not significantly different between BD-I and BD-II patients at baseline or after 4 and 8 weeks (Table 5).

The response rates ($\geq 50\%$ reduction in the total MADRS value) at week four and eight were 25.5% and 58.9, respectively ($P < 0.001$), and the remission rates

Figure 1 Changes of clinical global impression - bipolar version scores in patients with bipolar depression. (n = 764).

Notes: † $P < 0.001$ by repeated measure analysis of variance in BD-I; * $P < 0.001$ by repeated measure analysis of variance in BD-II; there was no significance (independent t-test) between BD-I and BD-II.

Abbreviations: BD-I, bipolar I disorder; BD-II, bipolar II disorder; CGI-BP, clinical global impression-bipolar version.

Table 4 Changes of subgroup of subjects in Clinical Global Impression – Bipolar scale in patients with bipolar I and II disorder (n = 764)

CGI-BP subgroup	Baseline	4 weeks	8 weeks	P-value*
Normal, not ill	0 (0.0)	7 (0.9)	41 (5.4)	<0.001 ^{a-c}
Minimally ill	0 (0.0)	82 (10.8)	287 (37.6)	
Mildly ill	143 (18.7)	319 (42.1)	295 (38.6)	
Moderately ill	325 (42.5)	257 (33.9)	114 (14.9)	
Markedly ill	208 (27.2)	74 (9.8)	22 (2.9)	
Severely ill	70 (9.2)	17 (2.2)	4 (0.5)	
Very severely ill	18 (2.4)	1 (0.1)	1 (0.1)	

Notes: * $P < 0.001$ by Cochran–Mantel–Haenszel test; ^acomparison of baseline and 4 weeks; ^bcomparison of 4 weeks and 8 weeks; ^ccomparison of baseline and 8 weeks; data expressed as n (%).

Abbreviation: CGI-BP, Clinical Global Impression – Bipolar scale.

(≤ 10 in the total MADRS value) were 14.9% and 42.1%, respectively ($P < 0.001$). However, there was no difference in the response and remission rates between BD-I and BD-II patients (Figure 2).

Factors affecting the reduction of depressive symptom severity

To determine the factors independently associated with MADRS reductions at week eight, age, illness duration, number of hospitalization, MADRS score at baseline, number of concomitant medications, duration of current episode, presence of remission on previous episode, and use of antidepressants were added in the regression model. MADRS score at baseline ($\beta = 0.612$, $P < 0.001$), duration of current episode ($\beta = -0.152$, $P = 0.001$), and the presence of remission on previous episode ($\beta = 0.111$, $P = 0.012$) were significantly associated with improvements of depressive symptoms (Table 6).

Table 5 Changes of total Montgomery–Asberg Depression Rating Scale scores and subscores (n = 764)

	Baseline	4 weeks	8 weeks	P-value*
Total scores	30.1 ± 10.6	20.5 ± 9.9	14.0 ± 9.5	<0.001
Apparent sadness	3.1 ± 1.2	2.3 ± 1.1	1.5 ± 1.1	<0.001
Reported sadness	3.3 ± 1.2	2.3 ± 1.2	1.5 ± 1.1	<0.001
Inner tension	3.2 ± 1.3	2.3 ± 1.2	1.6 ± 1.1	<0.001
Reduced sleep	3.2 ± 1.4	1.8 ± 1.3	1.2 ± 1.1	<0.001
Reduced appetite	2.8 ± 1.4	1.8 ± 1.3	1.2 ± 1.2	<0.001
Concentration difficulties	3.1 ± 1.3	2.3 ± 1.2	1.7 ± 1.1	<0.001
Lassitude	3.1 ± 1.3	2.2 ± 1.2	1.6 ± 1.1	<0.001
Inability to feel	2.9 ± 1.3	2.1 ± 1.2	1.5 ± 1.2	<0.001
Pessimistic thoughts	3.1 ± 1.4	2.1 ± 1.3	1.5 ± 1.2	<0.001
Suicidal thought	2.3 ± 1.6	1.5 ± 1.3	0.9 ± 1.1	<0.001

Notes: * $P < 0.001$ by repeated measure analysis of variance; data represented as mean ± standard deviation.

Figure 2 Response and remission rates by Montgomery-Asberg Depression Rating Scale score in bipolar I and II disorder patients. (n = 764).

Notes: * $P < 0.001$ by Chi square test between week 4 and 8 in BD-I; † $P < 0.001$ by Chi square test between week four and eight in BD-II; ‡ $P < 0.001$ by Chi square test between week four and eight in BD-I and II; there was no significance (Chi square test) between BD-I and II.

Abbreviations: BD-I, bipolar I disorder; BD-II, bipolar II disorder; BD-I and II, bipolar I and II disorder.

Therapeutic compliance and the adverse events

To evaluate therapeutic compliance, the subjective level of therapeutic compliance was examined, and the results revealed compliance rates exceeding 75% after 4 weeks for 834 patients (76.0%). In addition, after 8 weeks, the compliance level exceeded 75% for 827 patients (75.4%). In excluded patients (n = 333), 30.6% of BD patients (n = 102) were admitted (BD-I 28.1% versus BD-II 35.8%; $P = 0.108$). In addition, there was no significant difference in compliance between the BD-I and BD-II groups after 4 and 8 weeks (Table 7).

In total, 176 of 1097 BD patients (16.0%) felt fatigued, and 165 patients (14.9%) reported somnolence during the study period. The mean weight change throughout the study was 1.4 ± 2.6 kg.

Manic/hypomanic episodes were observed in seven cases (three cases/four cases; 0.6%) and three cases (three cases/none; 0.3%) at week four and eight, respectively. Only four

Table 6 Factors associated with Montgomery-Asberg Depression Rating Scale score reductions at week eight (n = 764)

Variables	β	t	P-value
Age	0.076	1.766	0.078
Illness duration	0.007	0.162	0.871
Hospitalization number	0.005	0.116	0.907
MADRS score at baseline	0.612	14.266	<0.001*
Number of concomitant medication	0.060	1.392	0.165
Duration of current episode	-0.152	-3.472	0.001*
Presence of remission on previous episode	0.111	2.524	0.012**
Use of antidepressants	0.041	1.144	0.253

Notes: * $P < 0.05$ by multiple linear regression; ** $P < 0.05$ by dummy variable regression.

Abbreviation: MADRS, Montgomery-Asberg Depression Rating Scale.

of the patients who exhibited manic/hypomanic episodes had symptom severities exceeding moderately ill according to the CGI-BP score after 4 weeks, and only one patient had to terminate quetiapine therapy. In addition, the symptoms could only be controlled by adjusting the dose of quetiapine and concomitant medications in all of the remaining cases.

Discussion

Active treatment of depressive symptoms may be very important in patients with BD because many BD patients attempt to commit suicide during the depressive episode or mixed episode including depressive symptoms, and the risk of committing suicide is 15-fold greater in BD patients than in the general population.¹⁶ Generally, in patients with bipolar depression, the use of atypical antipsychotics was limited to patients exhibiting psychotic symptoms. However, the use of atypical antipsychotics has been accepted as a new treatment modality because recent studies have demonstrated the therapeutic efficacy of atypical antipsychotics in treating bipolar depression, regardless of the accompanying psychotic symptoms.^{12,13,17–20} In addition, the Korean Medication Algorithm Project for BD 2006 (KMAP-BP 2006) recommended the combined use of atypical antipsychotics and mood stabilizers in the treatment of BD, regardless of the presence or absence of psychotic symptoms.¹⁰

In the current study, treatment with the atypical antipsychotic quetiapine after 4 and 8 weeks resulted in noticeable improvements in depressive symptoms in both BD-I and BD-II patients compared to the baseline findings. In addition, when disease severity was classified according to the CGI-BP score, the number of patients with more severe symptoms was decreased after both 4 and 8 weeks of treatment. In particular, such a change was observed between 4 and 8 weeks of treatment, which suggests that the therapeutic effectiveness of quetiapine continuously increased for at least 8 weeks. The improvement of depressive symptoms was consistently observed on every subscale of MADRS, and this result is consistent with the results of previous double-blind, randomized controlled trials.^{12,13} In addition, the rates of response and remission continuously increased for 8 weeks, resulting in values of 58.9% and 42.1%, respectively. These results can be considered to support the results of previous studies and indicate that quetiapine may be effective in the treatment of bipolar depression.^{12,13,21}

BD-II patients were more chronically ill, they had a greater duration of illness and spent more time in partial remission and less time in full remission.²² In the current study, however, there were no differences in the response

Table 7 Compliance of quetiapine treatment in patients with bipolar I and II disorder

	Week 4			Week 8			P-value
	Total (n = 1097)	BD-I (n = 743)	BD-II (n = 354)	Total (n = 1097)	BD-I (n = 743)	BD-II (n = 354)	
<25%	15 (1.4)	11 (1.5)	4 (1.1)	29 (2.6)	16 (2.2)	13 (3.7)	NS
25%–50%	48 (4.4)	31 (4.2)	17 (4.8)	43 (3.9)	27 (3.6)	16 (4.5)	NS
50%–75%	200 (18.2)	141 (19.0)	59 (16.7)	198 (18.1)	132 (17.8)	66 (18.6)	NS
>75%	834 (76.0)	560 (75.4)	274 (77.4)	827 (75.4)	568 (76.5)	259 (73.2)	NS

Note: Statistical analysis was done by Cochran–Mantel–Haenszel test and Chi-squared test.

Abbreviations: BD-I, bipolar I disorder; BD-II, bipolar II disorder; NS, not significant.

and remission rates measured by MADRS between the BD-I and BD-II groups throughout the entire observation period. These results were concurrent with the results of a previous naturalistic study indicating that the treatment outcome of depressive symptoms was not different between the BD-I and BD-II patients.²³ This suggested that quetiapine could be used effectively in both BD-I and BD-II patients.

The mean initial daily dose of quetiapine was 197.5 mg, and this was higher than the 50 mg/day dose of quetiapine monotherapy administered in previous studies.^{12,13} In addition, the percentage of patients treated with a combination of psychotropic agents was 73.2%, which was relatively high. This discrepancy is thought to be due to the fact that previous studies were conducted in outpatients, whereas 32.7% of the subjects in the current study were inpatients. In addition, 81.3% of the patients had a level of disease severity greater than moderately ill, and thus patients with relatively severe symptoms were more abundant in this study than in previous studies. Furthermore, previous studies were conducted as randomized controlled trials, and the primary aim of the current study was to observe patients in an actual clinical setting. The mean daily dose of quetiapine was increased slowly from the beginning of administration to 327.7 mg and 349.0 mg after 4 and 8 weeks, respectively, which is in agreement with the results of previous studies demonstrating that therapeutic efficacy was significant with daily doses of 300 mg or 600 mg.^{12,13,21}

On the other hand, throughout the entire period of observation in the current study, the dose of quetiapine was lower in the BD-II patients than in the BD-I patients; however, there was no difference in treatment outcomes. One reason may be that the ratio of combination therapy was higher in BD-II patients than in BD-I patients, and this combination therapy might be countervailing the lower dose of quetiapine.

In fact, it was found that – with the exception of mood stabilizers – combination therapy with antidepressants, anxiolytics, and hypnotics was administered more frequently

to BD-II patients. This is thought to be due to the fact that the diagnosis of psychiatric comorbidity is more common in BD-II patients,²² and in the current study, psychiatric comorbidity was more prevalent in patients with BD-II. However, there was no difference in the number of patients administered combination therapy between the two disease groups (71.7% versus 76.5%; $P = 0.186$), and the number of concomitant medications ($\beta = 0.060$, $P = 0.165$) and the use of antidepressants ($\beta = -0.041$, $P = 0.253$) did not affect MADRS changes.

Evaluation of therapeutic compliance in BD patients has been considered difficult.²⁴ BD patients tend to evaluate their own therapeutic compliance as being high, and the pill count method or the measurement of drug concentration also has limitations in the accurate evaluation of therapeutic compliance.²⁵ In the current study, the subjective level of therapeutic compliance was evaluated after 4 and 8 weeks of treatment, and a compliance rate exceeding 75% was observed for approximately 70% of the study population; thus, the rate of therapeutic compliance was higher than in previous studies.^{26,27} This is thought to be attributable to several factors, including the fact that the current study was an observational study allowing for the dose of quetiapine to be adjusted freely, the lack of restriction on the use of concomitant medications, the large percentage of inpatients in the study population, and the relatively short period of observation.

It has been reported that in the treatment of BD, quetiapine tends to exert antidepressant effects with mild adverse events. In the current study, the level of adverse events was not evaluated systemically, but the rates of fatigue, somnolence, and dropout were 16.0%, 14.9%, and 3.6%, respectively. This result was comparable to those of previous studies.^{11,28,29} However, the detected dropout rate was extremely low compared to the rate of 16.0% observed in the monotherapy group administered a comparable daily dose (300 mg). This arose from the lack of strict restriction in the protocol.

Conversely, one of the difficulties encountered in the treatment of bipolar depression is manic/hypomanic switching. In the current study, the incidences of manic/hypomanic episodes after 4 and 8 weeks were 0.6% and 0.3%, respectively, which were lower than those reported previously.^{12,13,21} Although the current study was an open-label observational study, the research protocol was not strict, and the study was conducted with a relatively short period of observation; these results suggest that quetiapine could be used safely for the treatment of bipolar depression from the viewpoint of adverse events.

This was a large-scale study to examine the therapeutic efficacy of quetiapine in patients with BD-I and BD-II depression in an actual clinical setting. However, this study has several limitations. A structural interview was not performed. It was an open-label observational study permitting the flexible adjustment of the dose of quetiapine and the use of concomitant medications, and thus the control of variables contributing to the improvement of symptoms was difficult. Also, the investigation of adverse events was not systematic in this study. However, the use of a detailed checklist for adverse events was beyond the scope of this paper. Therefore, additional studies for the safety of quetiapine will be needed.

Conclusion

The results of this study demonstrated that the treatment of BD-I and BD-II depression with quetiapine for 8 weeks had significant antidepressant effects in an actual clinical setting, and the therapeutic efficacy improved with time, with minimal manic switching. In addition, the clinical improvement was similar in both BD-I and BD-II patients. However, the mean dosage of quetiapine was lower in BD-II patients throughout the observational periods, and the combination therapy with antidepressants, anxiolytics, and hypnotics was more frequently administered to BD-II patients.

Acknowledgment

This study was funded by AstraZeneca.

Disclosure

The authors report no conflicts of interest in this work.

References

- American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders, 4th Edition, Text Revision: DSM-IV-TR®*. Washington, DC: American Psychiatric Association; 2000.
- Shantry BS. Bipolar disorder: an update. *Neurochem Int*. 2005;46(4):273–279.
- Judd LL, Akiskal HS, Schettler PJ, et al. The long-term natural history of the weekly symptomatic status of bipolar I disorder. *Arch Gen Psychiatry*. 2002;59(6):530–537.
- Kupfer DJ, Frank E, Grochocinski VJ, Cluss PA, Houck PR, Stapf DA. Demographic and clinical characteristics of individuals in a bipolar disorder case registry. *J Clin Psychiatry*. 2002;63(2):120–125.
- Sachs GS, Printz DJ, Kahn DA, et al. The expert consensus guideline series: medication treatment of bipolar disorder 2000. *Postgrad Med*. 2000;Spec No:1–104.
- Min KJ, Shin YC, Jon DI, Yoon BH, Kwon JS, Bahk WM. Korean medication algorithm for bipolar disorder (III): depressive episode. *Korean J Psychopharmacol*. 2003;14(4):336–346.
- Post RM, Calabrese JR. Bipolar depression: the role of atypical antipsychotics. *Expert Rev Neurother*. 2004;4(6 Suppl 2):S27–S33.
- Yatham LN, Kennedy SH, O'Donovan C, et al. Canadian Network for Mood and Anxiety Treatments (CANMAT) guidelines for the management of patients with bipolar disorder: consensus and controversies. *Bipolar Disord*. 2005;7(Suppl 3):5–69.
- Yatham LN, Kennedy SH, O'Donovan C, et al. Canadian Network for Mood and Anxiety Treatments (CANMAT) guidelines for the management of patients with bipolar disorder: update 2007. *Bipolar Disord*. 2006;8(6):721–739.
- Min KJ, Bahk WM, Seo JS, et al. Korean medication algorithm for bipolar disorder 2006 (III): depressive episode. *Korean J Psychopharmacol*. 2006;17:436–448.
- Sajatovic M, Mullen JA, Sweitzer DE. Efficacy of quetiapine and risperidone against depressive symptoms in outpatients with psychosis. *J Clin Psychiatry*. 2002;63(12):1156–1163.
- Calabrese JR, Keck PE Jr, Macfadden W, et al. A randomized, double-blind, placebo-controlled trial of quetiapine in the treatment of bipolar I or II depression. *Am J Psychiatry*. 2005;162(7):1351–1360.
- Thase ME, Macfadden W, Weisler RH, et al. Efficacy of quetiapine monotherapy in bipolar I and II depression: a double-blind, placebo-controlled study (the BOLDER II study). *J Clin Psychopharmacol*. 2006;26(6):600–609.
- Spearing MK, Post RM, Leverich GS, Brandt D, Nolen W. Modification of the Clinical Global Impressions (CGI) Scale for use in bipolar illness (BP): the CGI-BP. *Psychiatry Res*. 1997;73(3):159–171.
- Montgomery SA, Asberg M. A new depression scale designed to be sensitive to change. *Br J Psychiatry*. 1979;134:382–389.
- Harris EC, Barraclough B. Suicide as an outcome for mental disorders. A meta-analysis. *Br J Psychiatry*. 1997;170:205–228.
- Vieta E, Gasto C, Colom F, et al. Role of risperidone in bipolar II: an open 6-month study. *J Affect Disord*. 2001;67(1–3):213–219.
- Montgomery SA. Dopaminergic deficit and the role of amisulpride in the treatment of mood disorders. *Int Clin Psychopharmacol*. 2002;17(Suppl 4):S9–S15.
- Tohen M, Vieta E, Calabrese J, et al. Efficacy of olanzapine and olanzapine-fluoxetine combination in the treatment of bipolar I depression. *Arch Gen Psychiatry*. 2003;60(11):1079–1088.
- Sachs G, Sanchez R, Marcus R, et al. Aripiprazole in the treatment of acute manic or mixed episodes in patients with bipolar I disorder: a 3-week placebo-controlled study. *J Psychopharmacol*. 2006;20(4):536–546.
- Young AH, McElroy SL, Bauer M, et al. A double-blind, placebo-controlled study of quetiapine and lithium monotherapy in adults in the acute phase of bipolar depression (EMBOLDEN I). *J Clin Psychiatry*. 2010;71(2):150–162.
- Judd LL, Akiskal HS, Schettler PJ, et al. The comparative clinical phenotype and long term longitudinal episode course of bipolar I and II: a clinical spectrum or distinct disorders? *J Affect Disord*. 2003;73(1–2):19–32.
- Mantere O, Suominen K, Valtonen HM, et al. Differences in outcome of DSM-IV bipolar I and II disorders. *Bipolar Disord*. 2008;10(3):413–425.
- Colom F, Vieta E. Non-adherence in psychiatric disorders: misbehaviour or clinical feature? *Acta Psychiatr Scand*. 2002;105(3):161–163.

25. Vieta E. Improving treatment adherence in bipolar disorder through psychoeducation. *J Clin Psychiatry*. 2005;66(Suppl 1):24–29.
26. Maarbjerg K, Aagaard J, Vestergaard P. Adherence to lithium prophylaxis: I. Clinical predictors and patient's reasons for nonadherence. *Pharmacopsychiatry*. 1988;21(3):121–125.
27. Colom F, Vieta E, Martinez-Aran A, Reinares M, Benabarre A, Gasto C. Clinical factors associated with treatment noncompliance in euthymic bipolar patients. *J Clin Psychiatry*. 2000;61(8):549–555.
28. Bowden CL, Grunze H, Mullen J, et al. A randomized, double-blind, placebo-controlled efficacy and safety study of quetiapine or lithium as monotherapy for mania in bipolar disorder. *J Clin Psychiatry*. 2005; 66(1):111–121.
29. McIntyre RS, Brecher M, Paulsson B, Huizar K, Mullen J. Quetiapine or haloperidol as monotherapy for bipolar mania – a 12-week, double-blind, randomised, parallel-group, placebo-controlled trial. *Eur Neuropsychopharmacol*. 2005;15(5):573–585.

Neuropsychiatric Disease and Treatment

Dovepress

Publish your work in this journal

Neuropsychiatric Disease and Treatment is an international, peer-reviewed journal of clinical therapeutics and pharmacology focusing on concise rapid reporting of clinical or pre-clinical studies on a range of neuropsychiatric and neurological disorders. This journal is indexed on PubMed Central, the 'PsycINFO' database and CAS.

Submit your manuscript here: <http://www.dovepress.com/neuropsychiatric-disease-and-treatment-journal>

The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit <http://www.dovepress.com/testimonials.php> to read real quotes from published authors.