

Escitalopram versus paroxetine controlled release in major depressive disorder: a randomized trial

Taro Kishi¹
Yuki Matsuda¹
Shinji Matsunaga¹
Masatsugu Moriwaki^{1,2}
Yoichiro Otake²
Kaku Akamatsu³
Tomo Okochi⁴
Shigeki Hirano⁴
Toshihiko Funahashi⁵
Momoko Okuda⁶
Hideaki Tabuse⁶
Kiyoshi Fujita²
Nakao Iwata¹

¹Department of Psychiatry, Fujita Health University School of Medicine, ²Department of Psychiatry, Okehazama Hospital, Toyoake, ³Department of Psychiatry, Jindai Clinic, Nagoya, ⁴Department of Psychiatry, Toyota Memorial Hospital, ⁵Department of Psychiatry, Jindai Hospital, Toyota, Aichi, ⁶Department of Psychiatry, Holy Cross Hospital, Toki, Gifu, Japan

Correspondence: Taro Kishi
Department of Psychiatry, Fujita Health University School of Medicine, Toyoake, Aichi 470-1192, Japan
Tel +81 562 93 9250
Fax +81 562 93 1831
Email tarok@fujita-hu.ac.jp

Objective: There are no direct comparisons between escitalopram and paroxetine controlled release in patients with major depressive disorder (MDD).

Methods: We conducted a 24-week, rater-masked, randomized trial of escitalopram (5–20 mg/day) versus paroxetine controlled release (12.5–50 mg/day) in patients with MDD (UMIN000011191). Patients with the diagnosis of moderate-to-severe MDD (a 17-item Hamilton Rating Scale for Depression [HAM-D-17], with total score at baseline being ≥ 20) were recruited to participate in a parallel, randomized, controlled trial. The primary outcome for efficacy was an improvement in the 21-item HAM-D (HAM-D-21) total score at 24 weeks. The secondary outcomes were the response, remission, and discontinuation rates and the incidence of individual adverse events.

Results: A total of 88 patients with MDD (males, 61.4%; mean age, 40.8 ± 13.4 years) were recruited. The discontinuation rate was 58.0% (escitalopram, 55.8%; paroxetine controlled release, 60.0%). Both escitalopram and paroxetine controlled-release treatment groups exhibited significant reduction in the HAM-D-21 total score at 2, 4, 8, 12, and 24 weeks from the baseline. However, there were no significant differences in the HAM-D-21 total score, response rate, remission rate, and discontinuation rate at any time point between the groups. In addition, there were no significant differences in the incidence of any individual adverse events (eg, nausea, vomiting, and somnolence) between the treatment groups.

Conclusion: Our results suggest that escitalopram and paroxetine controlled release had similar efficacy and safety profiles in patients with MDD. One of the primary limitations of this study is the small sample size.

Keywords: escitalopram, paroxetine controlled release, major depressive disorder, Hamilton Rating Scale for Depression, antidepressant

Introduction

In 2009, the Meta-Analysis of New Generation Antidepressants Study² reported that clinically important differences for both efficacy and acceptability exist among commonly prescribed antidepressants in favor of escitalopram and sertraline. In 2012, escitalopram was demonstrated to have the highest probability of remission and is the most effective and cost-effective pharmacological treatment in a primary care setting.³ Escitalopram appears to be the best first-line antidepressant for treating major depressive disorder (MDD). In contrast, in 2010, paroxetine immediate release was the best-selling antidepressant in Japan.⁴

There were three randomized trials of escitalopram versus paroxetine immediate release in patients with MDD. Boulenger et al's study and Kasper et al's study reported that escitalopram is more effective and safer than paroxetine immediate release in the long-term treatment of patients with MDD.^{5,6} Baldwin et al reported that significantly

($P < 0.01$) more paroxetine immediate release was associated with a higher discontinuation rate compared with escitalopram.⁷ As the National Institute for Health and Care Excellence guidelines indicated, a higher incidence of discontinuation symptoms is observed for paroxetine immediate release than for other selective serotonin reuptake inhibitors (SSRIs).⁸ In order to overcome these drawbacks of paroxetine immediate release, paroxetine controlled release was developed in Japan in 2012 to improve general tolerability, particularly, gastrointestinal tolerability. Although there are no published data which demonstrate that paroxetine controlled release has a lower risk for producing discontinuation effects than paroxetine immediate release, one randomized trial showed that paroxetine controlled release is associated with low rates of early-onset nausea and dropout rates due to adverse events, which were comparable to those of placebo.⁹ However, there are no direct comparisons between escitalopram and paroxetine controlled release in patients with MDD. Therefore, we conducted a 24-week, rater-masked, randomized trial of escitalopram versus paroxetine controlled release in Japanese patients with MDD.

Methods

Subjects

This study was conducted from July 2013 to December 2015 at the Fujita Health University Hospital, Jindai Clinic, Jindai Hospital, Toyota Memorial Hospital, Holy Cross Hospital, and Okehazama Hospital. The trial was registered at the University Hospital Medical Information Network (UMIN) Clinical Trials Registry (UMIN000011191). Patients were diagnosed according to the *Diagnostic and Statistical Manual of Mental Disorders*, Fourth Edition, Text Revision criteria by the consensus of at least two experienced psychiatrists on the basis of structured interviews conducted using the Mini-International Neuropsychiatric Interview and a review of all medical records. All subjects met the following inclusion criteria: 1) age 20–70 years; 2) a 17-item Hamilton Rating Scale for Depression (HAM-D-17),¹ with total score at baseline being ≥ 20 ; 3) no neurologic or systemic diseases, including disturbance of hematopoiesis; 4) no history of electroconvulsive therapy within 6 months before enrollment; 5) absence of pregnancy; and 6) no dependence on any addictive substances other than nicotine in the past 5 years before enrollment. All subjects underwent laboratory blood testing and electrocardiography at the time of enrollment, and no patients were excluded because of a medical condition.

The clinical trial was described in detail, and written informed consent was obtained from all participants and

their guardians. This study was approved by the six ethics committees of the Fujita Health University Hospital, Jindai Clinic, Jindai Hospital, Toyota Memorial Hospital, Holy Cross Hospital, and Okehazama Hospital.

Procedures

All patients were randomly assigned to one of the two treatment groups by the central registration office. When the patients satisfied the inclusion and exclusion criteria for this multicenter trial, the randomization procedure was conducted by the authors in contact with a central registration office by telephone. The authors generated an allocation schedule before the start of the study using a random number table. The initial starting dose of escitalopram was 5–10 mg/day, with dose increases when needed in increments of 5–10 mg/day to a maximum of 20 mg/day. The initial starting dose of the paroxetine controlled release was 12.5 mg/day, with a dose increase of 12.5–50 mg/day. The subsequent antidepressant dosage was increased or decreased according to the patient's tolerance and therapeutic response. Lorazepam was used as an anxiolytic, and the hypnotics brotizolam and eszopiclone were permitted during our clinical trial (by prescription and for valid clinical reasons).

Evaluation of psychopathology, tolerability, and safety

The 21-item HAMD (HAM-D-21)¹ was completed at baseline and at 2, 4, 8, 12, and 24 weeks after starting administration or at discontinuation. Treatment-emergent adverse events were recorded at each time point, including immediately before the study, upon spontaneous complaints and through clinical observation using the Udvalg for Kliniske Undersogelser Side Effect Rating Scale.¹⁰

The raters who administered the psychometric tests were masked to the treatment group allocation, but the patients and physicians were aware of the treatment that was received. Raters consisted of psychologists who were not involved in any treatment decisions or the evaluation of side effects. Therefore, they were not required to know the pharmacologic treatment. The following measures were taken to maintain the masking of the raters. Electronic data from the trial were password protected, all case report forms were securely stored, and discussions about patients among the research teams were restricted. Moreover, the participants were reminded to avoid open discussion of the treatment assignment with the raters.

The primary outcome measure was the improvement of the HAM-D-21 total scores at week 24. The secondary outcomes included: the HAM-D-21 scores at 2, 4, 8, and

12 weeks; the HAMD-17 scores at 2, 4, 8, 12, and 24 weeks; the response rate at 2, 4, 8, 12, and 24 weeks (clinical response defined as $\geq 50\%$ reduction in the HAMD-17 or -21 total score from baseline to endpoint); the remission rate at 2, 4, 8, 12, and 24 weeks (clinical remission defined as the HAMD-17 total score ≤ 6 or HAMD-21 total score ≤ 7);¹¹ the discontinuation rate; and individual adverse events.

Statistical analysis

Modified intent-to-treat analysis was performed using the last observation carried forward method. Baseline continuous and categorical variables were compared between the treatment groups by an independent *t*-test and a chi-square test, respectively. Paired Student's *t*-tests were used to assess the changes in the HAMD-21 and HAMD-17 total scores from the baseline to endpoint. The comparison between the escitalopram and paroxetine controlled-release treatment groups was made by determining the change in the HAMD-21 (or HAMD-17) total score at the endpoint (2, 4, 8, 12, and 24 weeks) from baseline, based on an analysis of covariance model using the baseline HAMD-21 (or HAMD-17) score as covariates. We also used multiple logistic regression analyses to examine which antidepressant was associated with a higher response rate or remission rate (dependent variable: response rate [HAMD-21 or HAMD-17] or remission rate [HAMD-21 or HAMD-17]; independent variable: treatment group; and covariates: baseline HAMD-21 or HAMD-17

score, respectively). A Kaplan–Meier analysis was used to estimate the time to discontinuation for the two treatment groups, and the results were compared using a log-rank test. We also compared the change in body weight from baseline to endpoint between the escitalopram and paroxetine controlled-release treatment groups based on an analysis of covariance model using the baseline value as a covariate. Incidences of individual adverse events during the study were compared between the treatment groups using a chi-square test. The final dose of antidepressants was evaluated based on the fluoxetine equivalent.¹² The statistical power regarding the primary outcome for efficacy was also calculated ($\alpha=0.5$, <http://www.biostat.ucsf.edu/samplesize.html>).

Statistical analyses were performed using Statistical Package for the Social Sciences Statistics for Windows (Version 22.0; Corporation, Armonk, NY, USA) and JMP (JMP 12.2.1J; SAS Japan Inc., Tokyo, Japan). A *P*-value of <0.05 was considered statistically significant for all tests.

Results

The demographics and other characteristics of the patients are presented in Table 1. A total of 88 patients were recruited, all of whom were diagnosed with MDD at enrollment. Of these, 77.3% were first-episode patients and 61.4% were male, and the mean age was 40.8 ± 13.4 years. There were no significant differences in most variables between the treatment groups (Table 1).

Table 1 Baseline characteristics of the patients

Variable	Escitalopram (n=43)	Paroxetine controlled release (n=45)	Group difference (χ^2 test or <i>t</i> -test)	<i>P</i> -value
Male (%)	25 (58.1)	29 (64.4)	$\chi^2=0.369$	0.544
Age, years (mean \pm SD)	38.9 ± 12.4	42.5 ± 14.2	$t=-1.28$	0.205
First episode (%)	35 (81.4)	33 (73.3)	$\chi^2=0.819$	0.366
Educational history, years (mean \pm SD)	14.0 ± 1.88	13.6 ± 2.22	$t=-0.752$	0.454
Hospitalization (%)	0 (0.00)	1 (2.22)	$\chi^2=1.35$	0.245
Study center (FHU hospital, %)	19 (44.2)	26 (57.8)	$\chi^2=1.63$	0.202
Married (%)	25 (58.1)	19 (42.2)	$\chi^2=2.24$	0.135
Unemployed (%)	8 (18.6)	12 (26.7)	$\chi^2=0.819$	0.366
Current smoker (%)	9 (20.9)	9 (20.0)	$\chi^2=0.012$	0.914
Hamilton Depression Rating Scale-17 total scores (mean \pm SD)	23.7 ± 3.58	23.2 ± 3.78	$t=-0.605$	0.547
Hamilton Depression Rating Scale-21 total scores (mean \pm SD)	25.8 ± 4.24	25.1 ± 4.71	$t=-0.710$	0.480
Comorbid psychiatric illness (%) [*]	6 (14.0)	6 (13.3)	$\chi^2=0.007$	0.933
Body weight, kg (mean \pm SD)	65.0 ± 13.9	65.7 ± 12.3	$t=0.240$	0.811
Use of antidepressant (%)	0 (0.00)	0 (0.00)	Not applicable	
Use of hypnotic (%)	6 (14.0)	6 (13.3)	$\chi^2=0.007$	0.933
Use of anxiolytic (%)	0 (0.00)	1 (2.22)	$\chi^2=1.35$	0.245

Note: ^{*}Generalized anxiety disorder and/or panic disorder.

Abbreviations: FHU, Fujita Health University; SD, standard deviation.

Table 2 Discontinuation rate

Number of patients	Escitalopram	Paroxetine controlled release
Randomized	43	45
Completed the trial (%)	19 (44.2)	18 (40.0)
Discontinued the trial (%)	24 (55.8)	27 (60.0)
The reasons of discontinuation		
Inefficacy (%)	9 (20.9)	8 (17.8)
Adverse events (%)	8 (18.6)	11 (24.4)
Other reasons (%)	7 (16.2)	8 (17.8)

The discontinuation rate was 58.0% (escitalopram, 55.8% and paroxetine controlled release, 60.0%; Table 2). There was no significant difference in the mean times to discontinuation between the escitalopram and paroxetine controlled-release treatment groups (13.4 ± 10.1 vs 13.2 ± 9.92 weeks; $\chi^2_{(1)} = 0.0836$, $P = 0.773$). All 88 patients were included in the efficacy and safety analyses. The mean escitalopram and paroxetine controlled-release doses at endpoint were 13.6 ± 5.81 mg/day (fluoxetine equivalent, 30.2 ± 12.9 mg/day) and 24.8 ± 13.3 mg/day (fluoxetine equivalent, 29.2 ± 15.7 mg/day), respectively ($t_{(1)} = 0.0836$, $P = 0.773$).

Both escitalopram and paroxetine controlled-release treatments were associated with significant improvements in the HAMD-21 and HAMD-17 total scores at 2, 4, 8, 12, and 24 weeks (Table S1). However, there were no significant differences in the magnitude of the HAMD-21, and HAMD-17 total score decreases at any time between the escitalopram and paroxetine controlled-release treatment groups (Table 3). The statistical power with respect to the primary outcome was 43%. Also, there were no significant differences in the response rate at any time point between the treatment groups (Figures 1 and 2).

No patients had serious adverse events such as death, suicide attempt, and serotonin syndrome, and there were no significant differences in the incidence of individual adverse events between the groups (Table 4). Moreover, there were no significant differences in body weight at the endpoint between the treatment groups (Table 4).

Discussion

This is the first randomized trial involving a comparison between escitalopram and paroxetine controlled release in Japanese patients with MDD. There were no significant differences in the efficacy and safety outcomes between the escitalopram and paroxetine controlled-release treatment groups in this study. However, our study found that discontinuation rates due to adverse events in both treatment

Table 3 Group difference in Hamilton Depression Rating Scale scores at 2, 4, 8, 12, and 24 weeks

	Baseline score	Change in score at 2 weeks	F_{df}	P-value	Change in score at 4 weeks	F_{df}	P-value	Change in score at 8 weeks	F_{df}	P-value	Change in score at 12 weeks	F_{df}	P-value	Change in score at 24 weeks	F_{df}	P-value
Hamilton Depression Rating Scale-17 total scores																
Escitalopram	23.7 \pm 3.58	-9.40 \pm 6.57	$F_{1,86} = 2.00$, 0.161		-12.0 \pm 7.01	$F_{1,86} = 1.92$, 0.169		-12.8 \pm 7.36	$F_{1,86} = 1.03$, 0.314		-13.4 \pm 7.56	$F_{1,86} = 2.58$, 0.112		-14.5 \pm 7.73	$F_{1,86} = 3.10$, 0.082	
Paroxetine controlled release	23.2 \pm 3.78	-7.11 \pm 7.26			-9.58 \pm 7.76			-10.9 \pm 7.62			-10.6 \pm 7.86			-11.4 \pm 7.84		
Hamilton Depression Rating Scale-21 total scores																
Escitalopram	25.8 \pm 4.24	-10.0 \pm 7.18	$F_{1,86} = 1.62$, 0.206		-12.6 \pm 8.01	$F_{1,86} = 1.25$, 0.267		-13.8 \pm 8.01	$F_{1,86} = 0.775$, 0.381		-15.3 \pm 7.75	$F_{1,86} = 3.07$, 0.084		-15.6 \pm 8.16	$F_{1,86} = 1.02$, 0.315	
Paroxetine controlled release	25.1 \pm 4.71	-7.69 \pm 7.93			-10.3 \pm 8.63			-12.0 \pm 8.42			-12.0 \pm 8.98			-13.4 \pm 9.38		

Notes: Covariate: Hamilton Depression Rating Scale score at baseline. Data presented as mean \pm standard deviation.

Abbreviation: df; degrees of freedom.

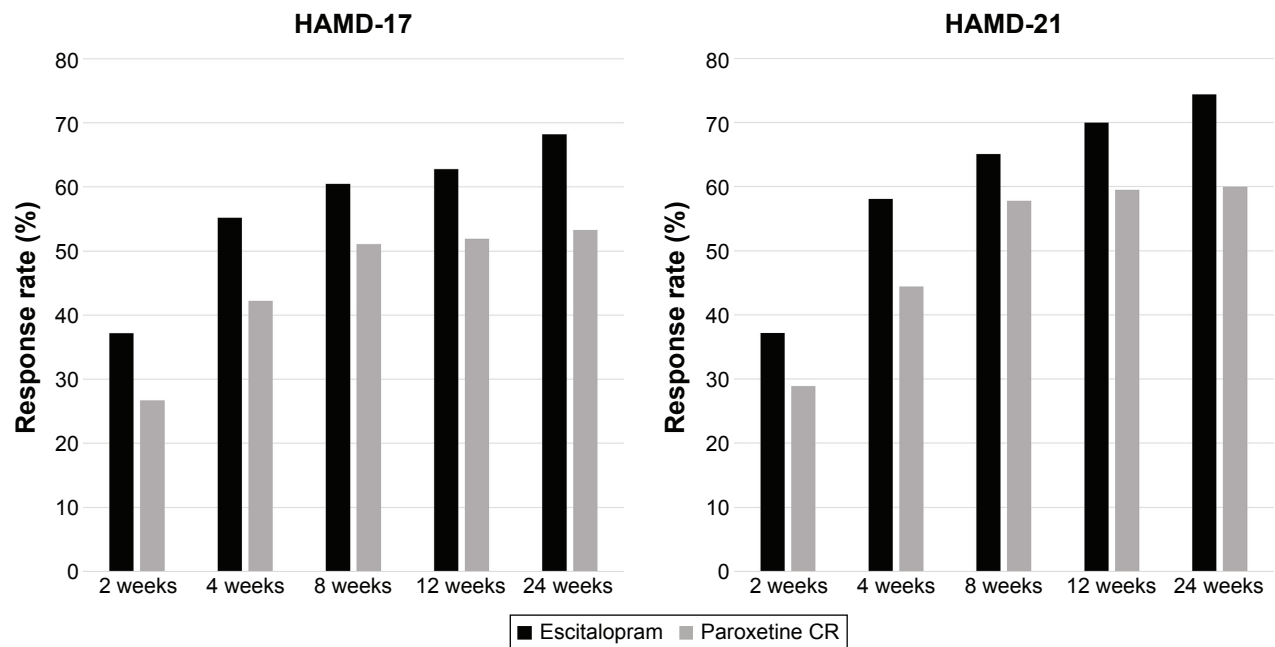


Figure 1 Response rate.

Abbreviations: CR, controlled release; HAMD, Hamilton Depression Rating Scale.

groups were relatively high compared with other studies on SSRIs.¹³ In a recent network meta-analysis, escitalopram emerged as the most efficacious agent among the SSRIs and was the best tolerated of the new-generation antidepressants (eg, agomelatine, duloxetine, escitalopram, fluvoxamine, fluoxetine, mirtazapine, paroxetine, sertraline, trazodone, and

venlafaxine) that were analyzed.¹³ We did not find significant differences in any efficacy outcome between the escitalopram and paroxetine controlled-release treatment groups in this study. The statistical power with respect to the primary outcome was 43%. Therefore, our results might be a statistical error due to an insufficient sample size.

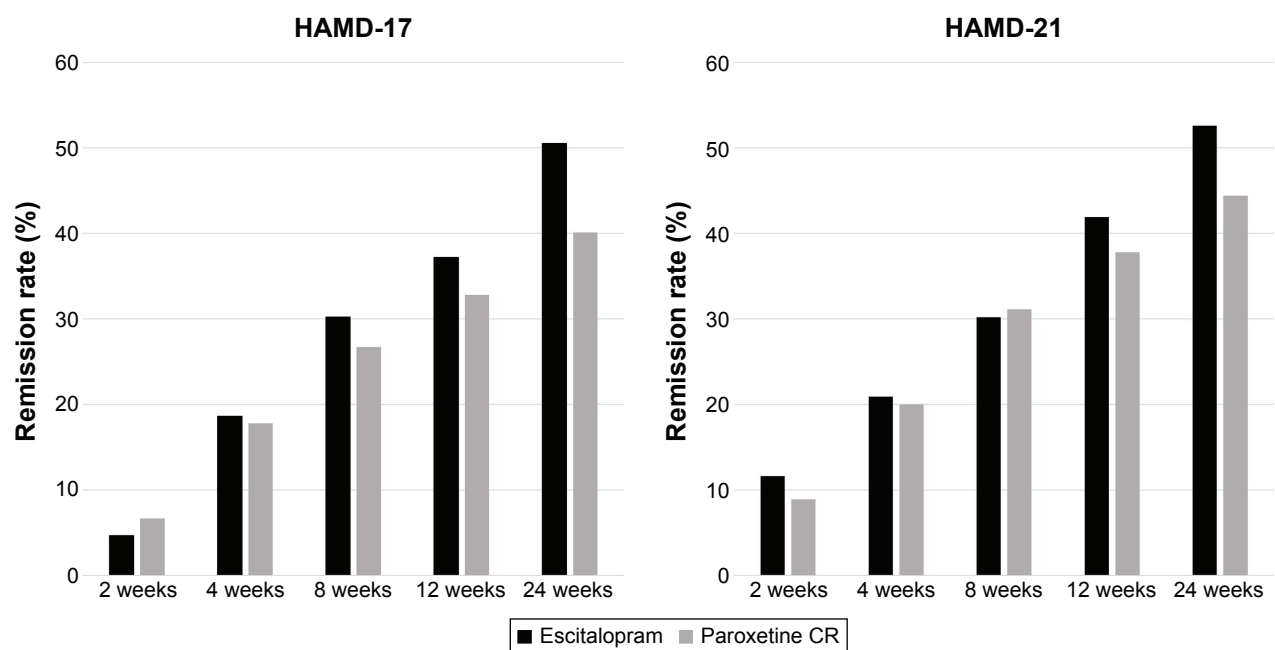


Figure 2 Remission rate.

Abbreviations: CR, controlled release; HAMD, Hamilton Depression Rating Scale.

Table 4 Adverse events observed in the study (incidence $\geq 5\%$)

Individual adverse events	Escitalopram (n=43)	Paroxetine controlled release (n=45)	Group difference (χ^2 test or analysis of covariance and P-value)
At least one adverse event (%)	26 (60.5)	29 (64.4)	$\chi^2=0.149$ and 0.700
Headache (%)	5 (11.6)	2 (4.44)	$\chi^2=1.59$ and 0.207
Somnolence (%)	12 (27.9)	8 (17.8)	$\chi^2=1.29$ and 0.256
Dizziness (%)	6 (14.0)	4 (8.89)	$\chi^2=0.562$ and 0.453
Anxiety (%)	9 (20.9)	4 (8.89)	$\chi^2=2.58$ and 0.108
Insomnia (%)	6 (14.0)	5 (11.1)	$\chi^2=0.162$ and 0.687
Decreased salivation (%)	3 (6.98)	5 (11.1)	$\chi^2=0.460$ and 0.498
Nausea/vomiting (%)	11 (25.6)	12 (26.7)	$\chi^2=0.013$ and 0.908
Diarrhea (%)	9 (20.9)	6 (13.3)	$\chi^2=0.901$ and 0.342
Constipation (%)	6 (14.0)	3 (6.67)	$\chi^2=1.29$ and 0.256
Loss of sexual desire (%)	2 (4.65)	3 (6.67)	$\chi^2=0.168$ and 0.682
Use of hypnotic (%)	6 (14.0)	6 (13.3)	$\chi^2=0.007$ and 0.933
Use of anxiolytic (%)	0 (0.00)	1 (2.22)	$\chi^2=1.35$ and 0.245
Weight gain (%)	1 (2.33)	3 (6.67)	$\chi^2=1.00$ and 0.317
Mean weight change (kg)*	0.853 \pm 3.05	0.502 \pm 2.11	$F_{1,86}=0.471$ and 0.494

Notes: *Covariate: body weight at baseline. Data presented as mean \pm standard deviation.

Limitations

One of the primary limitations of this study is that this study was not double blinded. The physicians responsible for assessing the adverse events were aware of which treatment was received by which patient. Although the raters were masked to the nature of the antidepressant treatment and were not involved in any side effect ratings or management decisions, it is possible that they were inadvertently informed of the group allocation by the patients. The second limitation is that this study did not include a placebo arm. The third limitation is the small sample size, particularly in the subgroup analysis (the statistical power with respect to the primary outcome was 43%). Therefore, we did not perform an HAMD item analysis. The fourth limitation is that we did not correct for multiple comparisons (eg, using Bonferroni's correction), because the application of a more stringent alpha level for secondary outcomes would have been too conservative in this small sample.^{14,15} The fifth one is that we did not count the number of patients who were screened for entry in the study. Finally, the intent-to-treat analysis using the last observation carried forward method may have influenced the results. The discontinuation rate was high, and thus, this method may yield a biased estimate of the treatment effect and underestimate the variability of the results.¹⁶ It is not clearly justified why this analysis, in particular, is appropriate and valid. However, if we had selected a complete analysis, our sample size would have been too small.

Conclusion

In conclusion, our findings suggest that although escitalopram and paroxetine controlled release are effective in patients with

MDD, discontinuation rates due to adverse events in both treatment groups were relatively high. However, because our study might be underpowered to detect significant differences in efficacy and safety outcomes between the escitalopram and paroxetine controlled-release treatment groups, further study using a larger sample size is required.

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Disclosure

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

Table S1 Endpoint change in Hamilton Depression Rating Scale scores at 2, 4, 8, 12, and 24 weeks from the baseline

Mean score at baseline	Mean score at 2 weeks	t_{df} , P-value	Mean score at 4 weeks	t_{df} , P-value	Mean score at 8 weeks	t_{df} , P-value	Mean score at 12 weeks	t_{df} , P-value	Mean score at 24 weeks	t_{df} , P-value
Hamilton Depression Rating Scale-17 total score: escitalopram										
23.7±3.58	14.3±5.97	$t_{42}=-9.38, <0.0001$	11.7±6.37	$t_{42}=-11.2, <0.0001$	10.9±6.71	$t_{42}=-11.4, <0.0001$	10.3±7.00	$t_{42}=-11.6, <0.0001$	9.23±7.24	$t_{42}=-12.3, <0.0001$
Hamilton Depression Rating Scale-17 total scores: paroxetine controlled release										
23.2±3.78	16.1±7.09	$t_{44}=-6.57, <0.0001$	13.6±7.63	$t_{44}=-8.28, <0.0001$	12.3±7.64	$t_{44}=-9.61, <0.0001$	12.6±7.67	$t_{44}=-9.03, <0.0001$	11.4±7.86	$t_{44}=-9.72, <0.0001$
Hamilton Depression Rating Scale-21 total scores: escitalopram										
25.8±4.24	15.7±6.53	$t_{42}=-9.15, <0.0001$	13.2±7.41	$t_{42}=-10.3, <0.0001$	12.0±7.58	$t_{42}=-11.3, <0.0001$	10.4±7.76	$t_{42}=-13.0, <0.0001$	10.1±7.97	$t_{42}=-12.6, <0.0001$
Hamilton Depression Rating Scale-21 total scores: paroxetine controlled release										
25.1±4.71	17.4±8.25	$t_{44}=-6.50, <0.0001$	14.8±8.82	$t_{44}=-8.00, <0.0001$	13.1±8.79	$t_{44}=-9.55, <0.0001$	13.1±9.27	$t_{44}=-8.95, <0.0001$	11.6±9.62	$t_{44}=-9.61, <0.0001$

Note: Data presented as mean ± standard deviation.

Abbreviation: df, degrees of freedom.

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