

Effect of milnacipran on suicidality in patients with mild to moderate depressive disorder

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Abstract: The presence of suicidal manifestations (thoughts and behavior) was studied in a cohort of 30 patients with mild to moderate depression during a 6-week treatment with the serotonin-norepinephrine reuptake inhibitor, milnacipran. At baseline mild suicidal thoughts were present in 46.7% of patients, the mean Hamilton Depression Rating Score (HDRS₁₇) was 23.9 ± 1.8 and the mean suicidality score on the Beck Scale for Suicidal Ideation (BSS) was 4.9 ± 4.9 . Suicidal thoughts decreased progressively throughout the study in parallel with other depressive symptoms. At no time during treatment was there any indication of an increased suicidal risk. Notably, the items retardation and psychic anxiety on the HDRS₁₇ decreased in parallel. This may possibly explain the lack of any “activation syndrome”, which is occasionally observed at the early stages of therapy with some antidepressants and may be linked to a temporary increase in suicidal ideation. To our knowledge this is the first detailed report of suicidality during treatment with milnacipran.

Keywords: suicidality, milnacipran, depression

Introduction

There is a clear association between suicide and depression. Mood disorders, principally major depressive disorder and bipolar disorder, are associated with about 60% of all suicides.¹⁻⁶ An often-quoted figure is that about 15% of patients with major depression will eventually die by suicide.^{7,8} This estimation is, however, derived largely from studies of severely depressed inpatients. More recently a more conservative lifetime suicide risk in the general population of depressed patients of about 6% has been calculated.⁹ This discrepancy has been erroneously interpreted by many primary care physicians and certain psychiatrists as evidence that overt suicidal behavior occurs only in severe depression. This notion can lead to inappropriate management of less severely depressed patients.

The more recent clinical literature clearly supports the idea that suicidality (thoughts and behavior) is associated with any type of depressive disorder and any degree of severity.^{10,11} Antidepressant therapy, which is effective in managing the symptoms of major depressive disorder, is expected to reduce or prevent suicidal behavior associated with depression.^{12,13} This hypothesis has been questioned, however, by large multi-study analyses that have suggested that, at least in certain patients groups, antidepressants may enhance or even induce suicidal ideation.^{14,15} In 2004 the US Food and Drug Administration warned of the possible enhancement of suicidal ideation in pediatric and adult patients receiving modern antidepressants especially at the beginning of therapy

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and at dose increase. This interpretation is controversial and the results of other meta-analyses^{16,17} have not supported the enhancement of suicidal ideation in adult patients with short-term antidepressant treatment compared to placebo. The latest analyses indicated that patients receiving SSRIs and tricyclic antidepressants (TCAs) do not show significantly more suicidality than patients receiving placebo.^{6,18} Indeed several studies have shown that suicide rates have decreased since the introduction of modern antidepressants supporting the hypothesis that the use of antidepressants may prevent suicide.^{19,20}

In the light of this controversy, the effect of an antidepressant, especially a recently introduced one, on suicidal ideation clearly remains an important question. Thus, in addition to assessing the general efficacy and tolerability of milnacipran (Ixel[®]), a serotonin and norepinephrine reuptake inhibitor (SNRI)²¹ recently introduced into Russia, the primary aim of the present investigation was to observe the effects of the drug on the occurrence and intensity of suicidal thoughts and behavior in patients with mild to moderate depressive disorders and to assess whether suicidality decreased in parallel with other depressive symptoms.

Methods

The investigation was a 6-week open label study in patients enrolled from an outpatient clinic (Psychiatric Hospital N12) in Moscow. Inclusion criteria were: age 16 to 65 years, a diagnosis of mild to moderate depression which could be either a single depressive episode, recurrent depressive disorder, bipolar disorder, dysthymia, cyclothymia, continuous depressive reaction within an adaptation disorder or other mood disorder. Exclusion criteria included severe depression with psychotic manifestations; schizophrenia and schizoaffective disorder; abuse of alcohol and other psychoactive agents; pronounced manifestations of psychorganic syndrome (or chronic brain disorder), including paroxysmal disorders; severe or decompensated somatic pathology requiring continuous polypharmacotherapy; pregnancy and lactation; participation in clinical trials within 30 days prior to start-up of the present study; and contradictions to milnacipran treatment. All diagnoses were based on ICD-10 criteria.

The study was approved by the local ethics committee and was carried out in accordance with the Declaration of Helsinki. All patients gave their written informed consent.

In addition to a standard psychiatric and general clinical examination (including urinalysis, hematology and

biochemistry, electrocardiogram), the 17-item Hamilton Depression Rating Scale (HDRS₁₇), Clinical Global Impression (CGI), Beck Depression Inventory (BDI), Beck Scale for Suicidal Ideation (BSS)²² and the Udvalg for Kliniske Undersøgelser Side Effect Rating scale (UKU-side-effects) were used.

Milnacipran was administered initially, for 2 to 3 days, at 25 to 50 mg/day depending on the severity of depressive symptoms. If the drug was well tolerated the dose was increased to 100 mg/day (50 mg twice daily).

The statistical significance of changes from baseline during the study were determined using paired Student's t-test and Wilcoxon nonparametric test using the statistical and analytical software package STATISTICA. The significance level was set as $P < 0.05$

Results

Thirty patients were enrolled in the study. Demographic characteristics of the patients are shown in Table 1. Women

Table 1 Patients' clinical demographics

		N	%
Female/Male		26/4	86.7/13.3
Mean age (yr)		41.4 ± 9.6	
Highest education level	Higher	18	60
	Secondary	10	33.3
	Primary	2	6.7
Professional activity	Employed	18	60
	Unemployed	12	40
Marital status	Married	12	40
	Unmarried	18	60
Diagnosis (ICD-10)	Single major depressive episode	18	60
	Recurrent major depressive disorder	12	40
Type of depression	Apathetic	10	33.3
	Anxious	6	20
	Asthenic	6	20
	Hysterical	6	20
Severity of current episode	Hypochondriac	2	6.7
Total	Mild	4	13.3
	Moderate	26	86.7
Single episode	Mild	2	
	Moderate	16	
Recurrent episode	Mild	2	
	Moderate	10	

comprised 86.6% of the cohort and the mean age of patients was 41.4 ± 9.5 (all values are given as mean \pm standard deviation). In general the educational level was high with 60% of patients having a higher (university) education and 33.3% a secondary education. Only 40% of patients were married.

The majority of patients (60%) had a diagnosis of a single depressive episode. Most patients (86.6%) at baseline were considered to be moderately depressed. The baseline HDRS₁₇ was 23.9 ± 1.8 and BDI was 37.7 ± 3.9 . The apathetic form of depression was the most frequent while anxious, asthenic and hysteric forms were also common. The hypochondriac form was only found in 2 patients.

Two patients withdrew from the study, one at the end of the first week of treatment because of adverse events (nausea) and the other at the end of the 4th week of treatment due to lack of efficacy of the treatment.

Antidepressant efficacy

The efficacy of the antidepressant treatment was demonstrated by the steady reduction of the mean HDRS₁₇ score

throughout the study (Figure 1). Statistically significant differences compared to baseline occurred from the first week of therapy. At endpoint (after 6 weeks treatment) mean HDRS₁₇ was 9.5 ± 4.4 and 60% of patients were treatment responders (reduction of baseline HDRS₁₇ of at least 50%). BDI scores showed a similar progressive reduction with a significant difference ($P < 0.05$) from baseline occurring from the third week of treatment. The mean BDI score at endpoint was 21.4 ± 6.1 (data not shown).

The mean baseline score on the CGI (Severity) scale was 3.9 ± 0.4 indicating a moderate severity. This value decreased progressively with a significant difference ($P < 0.05$) from baseline occurring after 3 weeks of treatment. At endpoint the mean CGI (Severity) score was 2.1 ± 0.3 , indicating borderline depression (data not shown).

Suicidality

At baseline mild suicidal ideation was found in 14 patients (46.7%). No current or previous suicidal attempts, planning or preparation were noted in any patient.

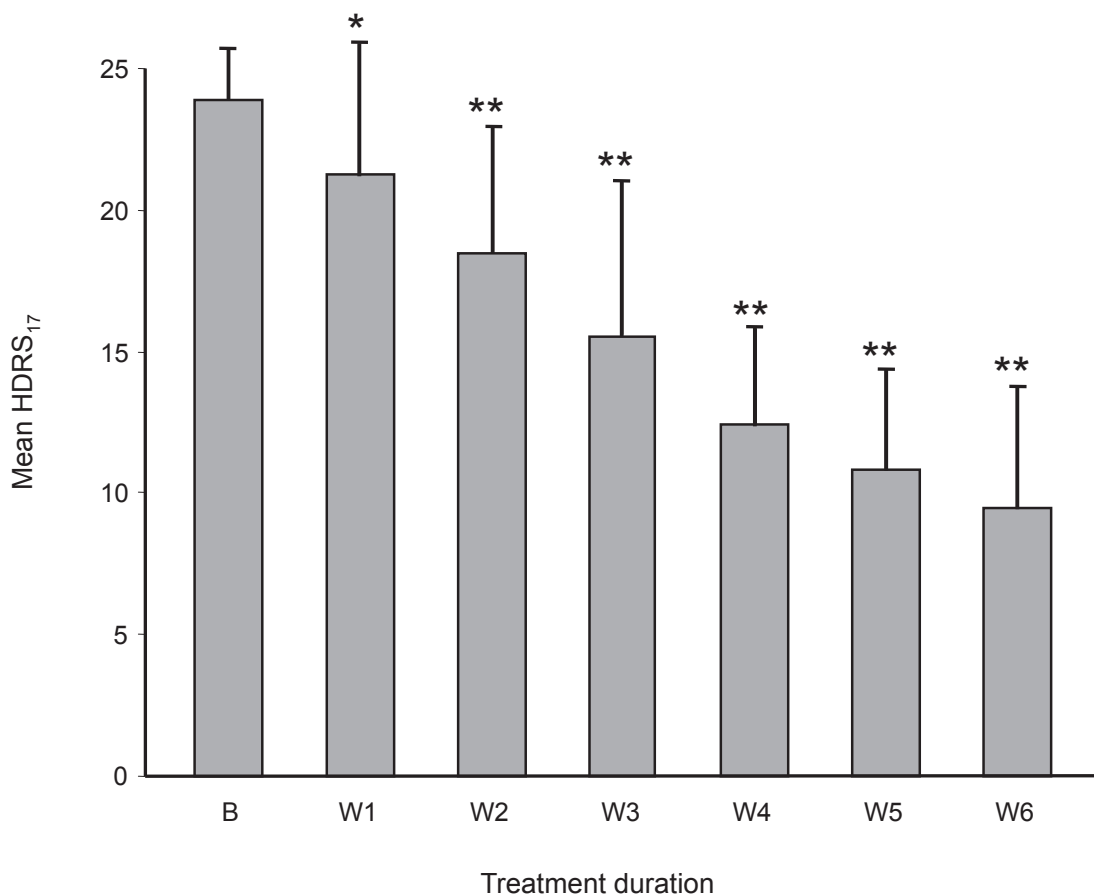


Figure 1 Reduction of mean global HDRS₁₇ scores during the study.

Notes: * $P < 0.05$; ** $P < 0.01$ compared to baseline values.

Abbreviations: B, baseline; W1 to W6, number of weeks of treatment.

A total of 57 suicidal manifestations were found during the study (Table 2). The most common suicidal manifestations were thoughts of “inanity of existence”, thoughts that death could resolve existing sufferings, suicidal thoughts and unwillingness to live.

More rarely, different signs of suicidality were reported: visualization of own suicide, death or funeral, as well as (supraliminal) death instinct. Only in a few patients did these suicidal ideations develop into either ideational compulsions (compulsive thoughts and concepts of death and funerals) or suicidal fantasies. Suicidal compulsive thoughts and concepts developed in patients with asthenic and hypochondriac depression, and suicidal fantasies in patients with hysteric depression. No other relationships, including those between suicidal phenomena and patients personality characteristics were detected.

The entry criteria of this study allowed the recruitment of patients with a single depressive episode, recurrent depressive disorder, chronic depression, bipolar disorder, dysthymia or cyclothymia. Unfortunately subanalysis of the different types of depression was not possible due to the small numbers of each subtype.

Mean baseline BSS score was 4.9 ± 4.9 , indicating a mild level of suicidality in this group of patients. Suicidality decreased rapidly with a significant reduction observed from the second week of treatment (Table 3). After 4 weeks of treatment, mean BSS scores were close to zero indicating an absence of any suicidal activity in treated patients.

Table 2 Suicidality (thoughts and behavior) reported during the study

	Number of patients reporting	% of all suicidal thoughts and behavior
Inanity of existence thoughts	11	19.4
Thoughts of death	9	15.8
Suicidal thoughts	7	12.3
Unwillingness to live	6	10.5
Visualizations of suicide	5	8.8
Visualizations of death	4	7
Visualizations of funerals	4	7
Death instinct	4	7
Suicidal compulsive thoughts	2	3.5
Funerals compulsive thoughts	2	3.5
Suicidal fantasies	2	3.5
Death compulsive thoughts	1	1.7
Total occurrences of suicidal thoughts or behavior ^a	57	100

^aSome patients reported more than one suicidal manifestation.

None of the patients showed any increase in BSS score during milnacipran treatment indicating an absence of any “activation syndrome”, sometimes observed at the early stages of therapy with certain antidepressants.¹⁰

Analysis of the “suicidal thoughts/attempts” item of the HDRS₁₇ at baseline showed a score of 1 in 10 patients (33.3%) and of 2 in one patient. The number of patients with scores 1 or 2 decreased progressively throughout the study. At endpoint only 3 patients had a score of 1, all others scoring zero.

The HDRS₁₇ items of “psychic anxiety” and “retardation” showed a similar progressive reduction throughout the study. The rate of decrease of the two items was approximately parallel (Figure 2). At the end of the study anxiety was reduced by 82% and retardation by 88%. The anxiety/retardation ratio remained essentially constant during milnacipran therapy confirming the absence of any “activation syndrome”.¹⁰

Adverse events

Adverse events (AEs) with milnacipran therapy were scored with the UKU Side Effect Rating in all patients. A total of 48 AEs were recorded during the study (Table 4). No AE was scored with a severity rating greater than 2, indicating that they did not significantly affect the patients’ daily life. The most frequent AEs, which occurred in more than 5 patients, were nausea, anxiety, tachycardia and dizziness. The majority of AEs occurred within the first week of treatment with milnacipran and tended to resolve spontaneously from the beginning of the second week. Seventy percent of all AEs had resolved by the third week of treatment.

Discussion

The cohort of the present study was a relatively classical mild to moderately depressed population although the proportion of women (86%) was somewhat higher

Table 3 Evolution of mean global Beck Scale for Suicidal Ideation scores during the study

	Mean (SD)	P ^a
Baseline	4.87 (4.91)	
Week 1	4.52 (4.38)	NS
Week 2	2.00 (2.83)	0.008
Week 3	1.10 (2.10)	0.0007
Week 4	0.20 (0.33)	0.0001
Week 5	0.10 (0.33)	>0.0001
Week 6	0.10 (0.33)	>0.0001

^aSignificance compared to baseline values.

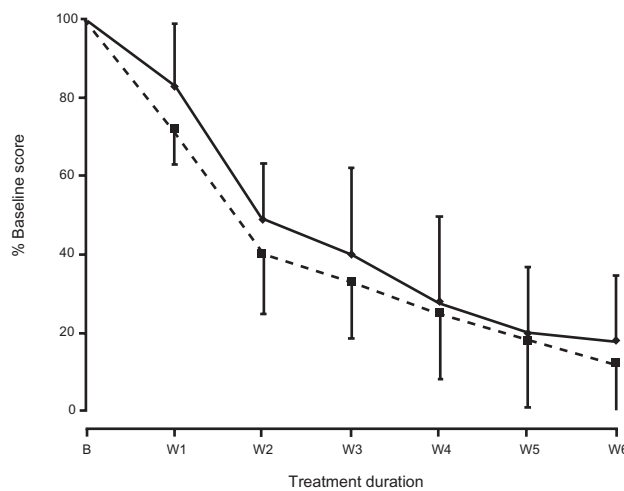


Figure 2 Reduction of mean "anxiety" and "retardation" scores during the study. **Notes:** "Anxiety" was the score of item 10 of the HDRS₁₇, "Retardation" was the score of item 8 of the HDRS₁₇. **Abbreviations:** B, baseline; W1 to W6, number of weeks of treatment; Solid line, "anxiety" item; broken line, "retardation" item.

than that commonly reported. At baseline nearly 50% of patients reported some form of suicidal ideation although none reported previous suicidal attempts, planning or preparation. Almost all of the suicidal manifestations observed during the study were limited to the cognitive sphere (thoughts, concepts). In addition, in the majority of cases, suicidal manifestations occurred only occasionally, and the suicidal fantasies, reported in only 2 patients, were transient (up to 1 to 2 hours per day over several days).

The wide prevalence of different suicidal manifestations in patients with mild to moderate depressive disorders shown here is important since, even in absence of a behavioral component they are considered to be a potent predictor of future suicidal behavior.^{2,13,23}

Regular patient assessment with HDRS₁₇, CGI and BDI showed a progressive regression of depressive symptoms throughout the study. There were few AEs, all of them mild as assessed by the UKU Side Effect Rating. Only one patient withdrew because of side effects.

The progressive regression of suicidal thoughts during milnacipran therapy observed at weekly testing showed no enhancement of suicidal risk at early stages of antidepressant pharmacotherapy. On the contrary, the administration of milnacipran led to a progressive and almost total reduction of all suicidal manifestations. The drug's effects on important aspects of the clinical profile of depression such as anxiety and retardation appeared to be well balanced. This may have contributed to the lack of "activation syndrome" which has been associated increased suicide risk.¹⁰ Even the occurrence of AEs such as anxiety during milnacipran therapy did not negatively affect the general suicidality of patients.

This study suffers from several weaknesses, namely the small cohort studied, its open nature and the relatively short observation period (6 weeks). To our knowledge this is the first study to have analyzed suicidality and depressive symptoms in parallel during the treatment with milnacipran. A meta-analysis of suicidal behaviors and ideation in clinical trials in depression with another SNRI, duloxetine, has also demonstrated the absence of increased risk of suicidal behaviors or ideation during treatment with the SNRI compared with placebo.²⁴

From this small open study there is a clear suggestion that mild suicidal thoughts can be relatively common even in mild to moderately depressed patients. During treatment with milnacipran suicidal thoughts regressed progressively in parallel with other depressive symptoms. At no time was there any noticeable increase in suicidal thoughts even in patients who reported anxiety as an

Table 4 Adverse events

	Number of adverse events recorded					
	Week 1	Week 2	Week 3	Week 4	Week 5	Week 6
Number of patients	29	29	29	28	28	28
Nausea	15	10	2	0	0	0
Anxiety	10	7	5	2	2	2
Tachycardia	8	5	1	0	0	0
Dizziness	7	3	0	0	0	0
Sweating	5	5	5	4	2	1
Somnolence	2	2	0	0	0	0
Abnormal ejaculation	1	1	1	1	0	0
Total number of adverse events	48	33	14	7	4	3

adverse event. This study clearly warrants replication in a larger cohort which should be followed for a longer period of time.

Acknowledgments and disclosures

We thank Dr Mike Briley for his help in correcting the manuscript. After completion of the study and its analysis, Pierre Fabre Médicament, manufacturers of milnacipran, offered to support the publication of the study with an unconditional educational grant. They played no role in the interpretation of the data or in writing the manuscript.

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