

Assessment of the Serotonin and Norepinephrine Reuptake Blocking Properties of Duloxetine in Healthy Subjects

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Duloxetine is a dual inhibitor of norepinephrine (NE) and serotonin (5-HT) uptake. Initial trials conducted in depressed patients using regimens of 20 mg/day or less did not convincingly demonstrate its efficacy as an antidepressant. The aim of this study was to assess the effects of duloxetine on the 5-HT and NE reuptake processes in healthy human volunteers. Twenty-seven healthy young males without a history of psychiatric disorder were randomly assigned to four groups, each group receiving one of the following daily drug regimens: placebo, clomipramine (a potent 5-HT/NE reuptake blocker) 100 mg/day, duloxetine 20 mg/day, or duloxetine 60 mg/day. In order to assess the NE reuptake process, the pressor response to intravenous tyramine (4 and 6 mg) was measured. Determination of the whole blood 5-HT content was used to evaluate the 5-HT reuptake blockade. These measurements were performed at baseline and repeated after 7 and 14 days

of drug intake. Both duloxetine, at doses of 20 to 60 mg/day, and clomipramine significantly interfered with the 5-HT reuptake process, as demonstrated by marked decreases in blood 5-HT concentrations. However, the same doses of duloxetine, unlike clomipramine, failed to impede the usual increase in blood pressure that follows a tyramine intravenous infusion, indicating that clomipramine but not duloxetine blocked NE reuptake. At doses tested in a population of healthy volunteers, duloxetine acted as a selective 5-HT reuptake inhibitor, having no clear effect on the NE reuptake process. Nevertheless, given that the highest dose of duloxetine increased supine systolic blood pressure, it is possible that it represents the threshold regimen for NE reuptake inhibition.

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Most antidepressants used today increase the efficacy of the serotonergic (5-HT) and/or noradrenergic (NE) neurotransmission. Thus, modulation of the synaptic availability of these two neurotransmitters is thought to

be involved in the therapeutic effects of these drugs (Blier and de Montigny 1999; Delgado et al. 1999). Blockade of the reuptake of 5-HT and/or NE by tricyclic antidepressants (TCAs), or of 5-HT only by selective 5-HT reuptake inhibitors (SSRIs) leads to a marked improvement of depressive symptoms in about 70% of patients. About 30% of them remain unsatisfactorily improved despite adequate dosing, however. Treatments combining NE and 5-HT reuptake blockade have been reported to produce more robust antidepressant effects in selected groups of depressed patients (Nelson et al. 1991; Seth et al. 1992; Nelson 1998; Poirier and Boyer 1999). Development of molecules acting as dual inhibitors of 5-HT and NE reuptake might thus lead to

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faster-acting and more efficacious antidepressant treatments.

Duloxetine [LY248686; (+)-N-methyl-3-(1-naphthalenyloxy)-2 thiophenepropanamine], a compound now being investigated as an antidepressant, is endowed with the property of blocking both the 5-HT and NE reuptake processes (Wong et al. 1993; Engleman et al. 1995; Kasamo et al. 1996; Fuller et al. 1994; Kihara and Ikeda 1995; Wong 1998), while being devoid of significant affinities for other neuronal receptors (Wong et al. 1993). Animal studies have shown that, *in vitro* and *ex vivo*, duloxetine inhibits [^3H]5-HT and [^3H]NE in hypothalamic and cortical synaptosomes (Wong et al. 1993; Engleman et al. 1995), as well as in hippocampus slices (Kasamo et al. 1996). Radioligand-binding studies support this finding in that duloxetine antagonizes the binding of [^3H]paroxetine and [^3H]tomoxetine to the 5-HT and NE carriers, respectively (Wong et al. 1993; Engleman et al. 1995). Duloxetine prevents the p-chloroamphetamine-induced depletion of 5-HT in the rat brain (Wong et al. 1993; Engleman et al. 1995; Kasamo et al. 1996; Fuller et al. 1994), and also antagonizes the depletion of NE induced by 6-hydroxydopamine in mice hearts and in the hypothalamus of rats (Fuller et al. 1994). Microdialysis studies indicate that acute systemic duloxetine administration induces a large increase in extracellular levels of 5-HT and NE in the hypothalamus and cortex of rats (Engleman et al. 1995; Kihara and Ikeda 1995). Finally, using an electrophysiological approach, Kasamo et al. (1996) found that, when given acutely, duloxetine suppresses the spontaneous firing of both the dorsal raphe 5-HT and the locus coeruleus NE neurons, indicating an increased availability of these neurotransmitters in the immediate vicinity of their somatodendritic receptors regulating their firing activity.

In spite of the pharmacology of duloxetine compared to that of currently marketed antidepressants and of promising results in depressed subjects (Berk et al. 1997), controlled trials conducted in depressed patients have yet to convincingly demonstrate its efficacy as an antidepressant. In the largest controlled study reported to date for patients with major depression, duloxetine was administered at daily doses of 5, 10, and 20 mg, and was compared to placebo and to 150 mg/day of clomipramine (Joubert et al. 1997). A preliminary indication in that study is that duloxetine was not as effective as clomipramine. In contrast, venlafaxine (Effexor®) has been shown to block reuptake of both 5-HT and NE and to be an effective antidepressant (Yardley et al. 1990; Derivan et al. 1995; Nierenberg et al. 1994). It is possible that, in doses tested thus far, duloxetine has not adequately blocked both 5-HT and NE reuptake processes and as a result has not been found as efficacious as expected.

The main goal of this study was to examine the ef-

fects of duloxetine on the 5-HT and NE reuptake processes in healthy volunteers. The secondary objective was to compare the potency of different doses of duloxetine in blocking NE and 5-HT reuptake processes to that of placebo and the reference drug, clomipramine.

SUBJECTS AND METHODS

Study Subjects

Volunteers were recruited via a newspaper advertisement. Prior to their inclusion in the study, all volunteers underwent a structured clinical interview (SCID for non-patients), a physical examination, and laboratory tests consisting of a complete blood count, electrolytes, liver enzymes, urea and creatinine, urine drug screening, and an electrocardiogram. Participants met the following inclusion criteria: 1) male, between the ages of 18 and 40 years old; 2) normal body weight (body mass index not in the obese range); 3) no significant medical illness, including migraines, high blood pressure (BP > 140/90), hepatitis B or HIV; 4) not taking medication on a regular basis; and 5) no evidence suggestive of a past or present history of mental disorders.

Subjects were excluded from the study for any of the following reasons: 1) use of an investigational drug within 30 days of study entry; 2) orthostatic hypotension (fall in systolic BP > 10 mm Hg); 3) use of any psychotropic medication or illicit drugs; 4) regular use of any medication other than acetaminophen or vitamins and mineral supplements; 5) smoking more than 10 cigarettes per day; and 6) history of difficult phlebotomy (e.g., vasovagal reaction). Any significant abnormality upon physical examination or laboratory tests was also a reason for exclusion.

Randomization

Volunteers were randomly assigned to one of four treatment groups: placebo, clomipramine 100 mg/day, duloxetine 20 mg/day (low-dose), or duloxetine 60 mg/day (high-dose). Thirty subjects qualified for the study, but three dropped out because of adverse events (one in each of the following groups: placebo, clomipramine and duloxetine 60 mg/day). A total of twenty-seven subjects completed the protocol, and only these subjects were considered in the analysis.

Drug Regimens and Timing of Assessments

Each subject was randomly assigned to one of the four treatment arms. If assigned to either placebo or a duloxetine arm, neither the subject nor the investigator could tell whether the subject was given duloxetine or placebo, as the study drug capsules were identical in appearance. If assigned to the clomipramine arm, the sub-

ject was given clomipramine tablets which were identifiable as such. In all arms, subjects took study drug in divided doses (twice daily) for a total of 14 days. Subjects in the low-dose duloxetine arm started on the full dose and received the same dose for the entire study period. The doses were gradually escalated in the high-dose duloxetine and clomipramine arms in order to minimize unwanted side effects. Subjects in the high-dose duloxetine arm received 20 mg/day for three days, 40 mg/day for four days, and then 60 mg/day for the next seven days. Dose-escalation for the clomipramine group was as follows: 25 mg/day for three days, 50 mg/day for four days, and then 100 mg/day for the last seven days. Plasma levels of the drugs were determined by standard HPLC methods approved by Eli Lilly & Co (Sharma et al. 2000).

The assessment of NE and 5-HT reuptake processes was performed at baseline (day 0), on day 7 and on day 14. As both duloxetine and clomipramine have half-lives of less than 24 hours, it was considered that four days at the same dose would be sufficient to achieve near-steady-state plasma levels. To insure maximal biological response during the reuptake assessments, participants were instructed to take their last dose of medication 2 to 3 hours prior to the beginning of testing. The peak plasma level of duloxetine is reached after 2 to 5 hours (Sharma et al. 2000), while that of clomipramine is typically reached after 2 hours.

Assessment of 5-HT Reuptake

Platelet 5-HT content has become a widely used tool for studying the presynaptic serotonergic neurons. Blood platelets can take up, store and release 5-HT in a fashion very similar to that of 5-HT neurons, but lack the enzymatic system necessary to synthesize 5-HT and most of the 5-HT receptors subtypes (Pletscher 1968). Thus, any 5-HT present in platelets has been taken up by the transport system, and platelet 5-HT content is considered to provide an indirect indication of 5-HT reuptake activity by 5-HT neurons. Once 5-HT uptake has been effectively blocked and as it remains blocked, the platelet 5-HT content gradually decreases as older platelets which have previously taken up 5-HT are removed from the circulation and are replaced by new ones incapable of taking up 5-HT. Moreover, as a very large proportion (90–95%) of blood 5-HT is sequestered in the platelets, whole blood 5-HT content largely reflects platelet 5-HT content (Demet et al. 1978; Artigas et al. 1985; Zucker 1965).

In this study, whole blood 5-HT was measured by high performance liquid chromatography (HPLC) with electrochemical detection, basically as described by Artigas et al. (1985). Two indices were used to evaluate 5-HT reuptake: 1) depletion of blood 5-HT content, measured on day 7 and day 14, compared to the baseline level; and

2) that same 5-HT content of whole blood, this time divided by the platelet count, as measured at baseline and on day 14. Adjusting the whole blood 5-HT content with the platelet count in this fashion is expected to help reduce inter-individual and intra-individual variance. Flament et al. (1987) found that the mean level of 5-HT changed from 49 to 45 ng/ 10^8 platelets after five weeks of treatment with placebo, whereas it dropped to 2 ng/ 10^8 platelets with clomipramine treatment.

Assessment of NE Reuptake

Tyramine acts indirectly as a sympathomimetic agent. It is taken up in NE neurons by the NE transporter and then releases NE from its intraneuronal stores (Ishigooka et al. 1997). In turn, NE acts on postsynaptic adrenoceptors to cause an elevation of systolic BP. Pretreatment with desipramine, which blocks the high-affinity NE reuptake site, prevents the increase in BP that usually follows tyramine administration (Hoffman and Lefkowitz 1990) by preventing tyramine from entering the NE neurons. The tyramine test consists of measuring the increase in systolic BP of patients following a fixed dose of tyramine, or the dose of tyramine necessary to cause a predetermined increase in systolic BP, such as 30 mm Hg (Ghose et al. 1976). A decrease in the pressor response to the same doses of tyramine before a given treatment, or an increase in the dose of tyramine necessary to cause a given increase in systolic BP, may be considered an index of NE reuptake blockade.

Blockade by a drug of vascular postsynaptic α -adrenoceptors might also be expected to decrease tyramine pressor response (Ghose 1984). Some commercially available antihypertensive agents, such as prazosin, are known to act by blocking the α_1 -adrenoceptors and thereby producing vasodilation. Recently, however, Schäfers et al. (1997) reported that the α_1 -adrenoceptor antagonist doxazosin did not decrease tyramine pressor response, although the β_1 -adrenoceptor antagonist bisoprolol did. They proposed that the rise in systolic BP obtained with intravenous tyramine infused over 15 minutes reflects a β_1 -adrenoceptor-mediated positive inotropic effect rather than a vascular pressor response. Regardless of whether the pressor response to tyramine is mediated through α -adrenoceptors or β -adrenoceptors, if tyramine is prevented from entering NE nerve terminals, it should not be able to cause NE release and thus should not cause any change in blood pressure. In the present study, the pressor response to tyramine was considered to provide specifically an index of NE reuptake blockade by duloxetine, since duloxetine has no affinity for either α -adrenoceptors or β -adrenoceptors. The possible contributions of α - and β -adrenoceptor activity to the effects of clomipramine and other agents are addressed further in the discussion later in this paper.

The effects of two fixed doses of intravenous tyramine (4 and 6 mg) on systolic BP were assessed. The choice to measure blood pressure response to a fixed dose of tyramine (rather than to estimate the dose of tyramine necessary to produce a given change in blood pressure) was made with the expectation that duloxetine- and/or clomipramine-treated subjects would require large doses of tyramine to produce increases of ≥ 30 mm Hg in systolic blood pressure, such that medical complications might result. Blockade of NE reuptake was expected to manifest itself by a decrease in the elevation of BP that was shown prior to drug administration (effect of tyramine measured on day 0).

In response to the physiological challenge of standing up, NE plasma levels usually increase (Ghose and Turner 1975). Following a 5-week treatment with clomipramine, Flament et al. (1987) found an increased elevation of plasma NE levels on standing up, presumably due to an increased synaptic availability of NE secondary to NE reuptake blockade. In our study, to evaluate NE reuptake inhibition further, supine and standing NE plasma levels were measured by HPLC.

The following procedure was followed at each visit: the subject was connected to a Criticare 408 apparatus, which automatically registers the heart rate (HR) and blood pressure, and a catheter installed in one arm. Following a resting period of 30 minutes in supine position, during which HR and BP were recorded every 10 minutes, blood was drawn for 5-HT, NE, and study drug level measurements. The participant then stood up for 5 minutes, at which point BP and HR were recorded, and blood was once again drawn for NE level measurement.

After another 30-minute period in supine position, a test dose of 0.5 mg of tyramine was administered intravenously in order to rule out the presence of a pheochromocytoma which would not have been detected by the physical examination and laboratory tests. After 5 minutes, and if there was no significant elevation in blood pressure after the test dose, a dose of 4 mg of tyramine was administered. The BP and HR were then measured every 2 minutes for 10 minutes. This time period allowed us to detect the peak increase in BP, which usually occurs 2 minutes after the injection of tyramine, and provided sufficient time to let BP return to its baseline level. If the increase in BP did not reach 30 mm of Hg after the 4 mg dose of tyramine, an additional dose of 6 mg was given, and the BP and HR were measured as just described.

Evaluation and Statistical Analysis

The pressor response to tyramine was considered to be the difference between the systolic blood pressure measured 2 minutes after the tyramine infusion was complete, and that just prior to it. Serotonin reuptake data is

expressed in concentrations in 5-HT in pmol/L and in 5-HT content of 10^9 platelets compared to the baseline levels. Analysis of variance (ANOVA) for repeated measures (treatment \times time) was performed on the data to assess the effects of the different drug regimens on the following parameters: 1) concentrations of 5-HT; 2) ratio of 5-HT per 10^9 platelets at baseline and day 14; 3) pressor response to a load of 4 mg of tyramine; 4) pressor response to a load of 6 mg of tyramine; 5) NE concentrations; 6) supine and standing pulse; 7) supine and standing systolic BP; and 8) supine and standing diastolic BP. Post-hoc analyses were conducted where appropriate with Tukey's test. Linear regression analyses were conducted between the drug plasma levels and the 5-HT concentrations and the pressor response to tyramine. All values are expressed as means \pm SEM unless indicated otherwise. The level of significance was $p < .05$.

RESULTS

Subjects

A total of 27 healthy male volunteers completed the study. The average age of the subjects was 24 ± 1 year (mean \pm SEM). They weighted on average 72.7 ± 2 kg, and their average height was 174.2 ± 1.2 cm. Mean body mass index, calculated by dividing the weight (in kg) by the square of the height (in meters), was 24. There was no significant difference between subject groups for age, height, weight, or body mass index.

Effects on 5-HT Reuptake

The first 11 samples were lost during preparation. A total of 16 samples were available for analysis of 5-HT concentrations (Figure 1). The data for 5-HT collected for one of these subjects was withdrawn from all further analysis because the measured baseline 5-HT level was 195 pmol/ml of whole blood, a result which was clearly outside the range of two standard deviations (S.D. = 851 pmol/ml of blood) of the mean for the rest of the subjects at that point: 2086 pmol/ml of whole blood. An undetected technical error might be responsible for this aberrant result.

There was a significant interaction between treatment group and time of testing [$F_{6,22} = 6.82$; $p < .01$]. Simple main effect tests revealed that, while placebo had no significant effect on whole blood 5-HT concentrations [$F_{2,22} = 1$; $p > .05$], clomipramine treatment, as well as both duloxetine regimens, significantly decreased 5-HT levels [$F_{2,22} = 34.22, 30.67$, and 23.97 , respectively; $p < .0001$]. Indeed, 5-HT levels had significantly decreased from baseline as early as on day 7 for subjects receiving clomipramine and duloxetine 20 mg/day, and those levels did not decrease any further the

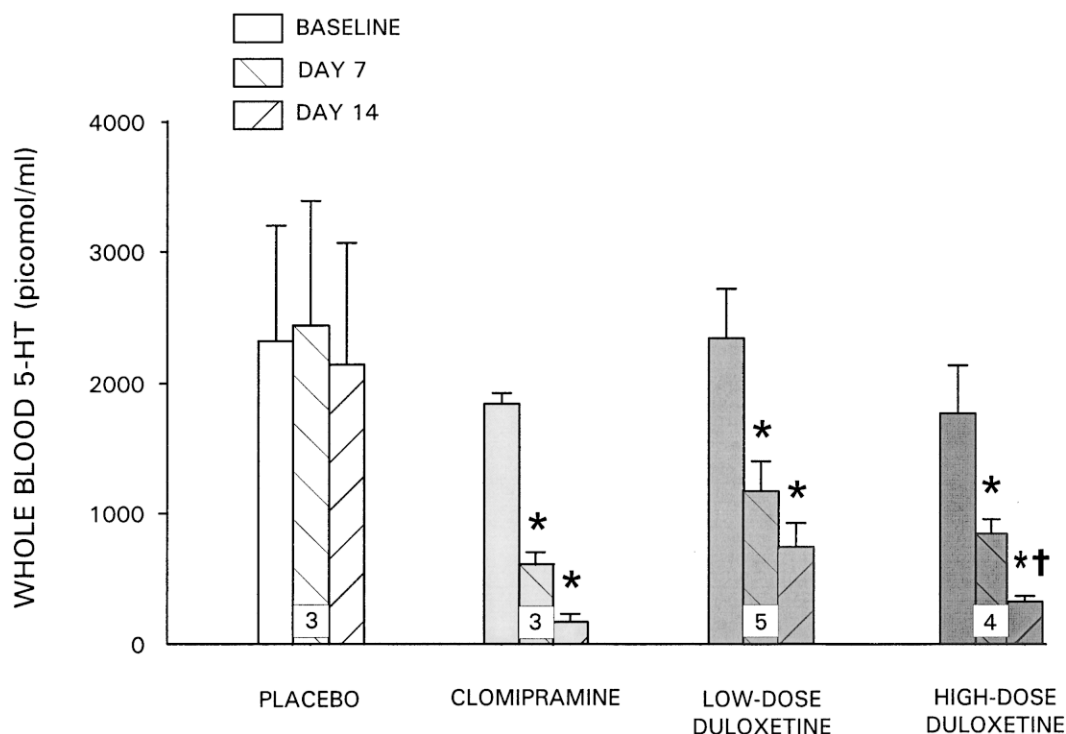


Figure 1. Mean (\pm SEM) whole blood 5-HT concentration (pmol/ml) per treatment group. Measures were obtained at baseline, after 7 days and after 14 days of drug administration. (*) indicates a significant change from baseline and (+) a significant change from day 7.

following week. Subjects on the duloxetine dose-escalation regimen (40 mg/day on day 7 and 60 mg/day on day 14) also had a significant decrease in whole blood 5-HT concentrations on day 7, but with this dose there was a significant further decrease in 5-HT levels during the following week of treatment, suggesting a dose-dependent response.

Similar analyses were conducted on the 5-HT content of 10^9 platelets at baseline compared to that on day 14. Results show that the 5-HT content of 10^9 platelets was significantly decreased from baseline in subjects taking clomipramine, duloxetine 20 mg/day, and duloxetine 60 mg/day [$F_{1,10} = 23.88, 19.61$, and 24.76 , respectively; $p < .01$], whereas placebo had no effect on that parameter [$F_{1,10} = 0.01$; $p > .05$].

Effects on NE Reuptake

NE Plasma Levels. There was no significant difference between groups in supine NE plasma levels [$F_{3,23} = 0.49$, $p = .69$] at any time of testing, nor was there a significant interaction between treatment group and time of testing [$F_{6,46} = 1.66$, $p = .15$]. Similarly, there was no difference in standing NE levels [$F_{3,23} = 1.59$, $p = .21$], and there was no interaction between treatment groups and time of testing [$F_{6,46} = 0.72$, $p = .63$]. Since these results did not reveal significant differences, the raw data are not presented.

Vital Signs. There was no significant difference between treatment groups with regard to standing systolic BP [$F_{3,23} = 0.30$, $p = .83$] and standing diastolic BP [$F_{3,23} = 1.13$, $p = .36$], nor was there an interaction between treatment groups and time of testing for those parameters [$F_{6,46} = 1.80$, $p = .12$ and $F_{6,44} = 0.99$, $p = .44$, respectively]. The standing diastolic BP value was unavailable for one subject at baseline, therefore the data for that parameter were removed from the analysis for standing diastolic BP.

There was, however, a significant interaction between time of testing and treatment group for the following parameters: supine pulse [$F_{6,46} = 4.36$, $p = .002$], standing pulse [$F_{6,46} = 2.89$, $p = .02$], supine systolic BP [$F_{6,46} = 2.22$, $p = .057$], and supine diastolic BP [$F_{6,46} = 2.90$, $p = .02$]. Simple main effect tests revealed that clomipramine significantly increased both supine and standing pulse, as well as the supine diastolic BP, on day 7 and day 14, when compared to baseline. However, there were no significant changes in those parameters from day 7 to day 14 in the subjects taking clomipramine. Clomipramine did not, at any time, significantly change the supine systolic BP, whereas volunteers receiving the high-dose duloxetine (60 mg/day) had a significant increase in supine systolic BP on day 14, when compared to baseline.

There was no significant difference in supine systolic BP in the high-dose duloxetine group between baseline

and day 7, when the dose was 40 mg/day, nor was there a difference on that parameter between day 7 and 14. High-dose duloxetine was not associated with significant changes in measurements which clomipramine had effects on, namely the supine pulse, standing pulse, and supine diastolic BP. Placebo and the lower dose of duloxetine had no significant effects on cardiovascular parameters. Cardiovascular measures are summarized in Table 1.

Tyramine 4 mg. The ANOVA for repeated measures showed that there was an interaction between the day of testing and the increase in systolic BP caused by the intravenous administration of 4 mg of tyramine [$F_{6,46} = 2.44$; $p < .05$] (Figure 2). A simple main effect test indicated that there was a difference in the increase in systolic BP over time in subjects taking clomipramine [$F_{2,46} = 5.06$; $p < .05$], but not in subjects receiving placebo and both duloxetine regimens [$F_{2,46} = 0.22, 0.14$, and 3.03 , respectively; $p > .05$]. Post-hoc analysis of the results indicated that the mean increase in blood pressure in subjects taking clomipramine was significantly reduced on days 7 and 14, when compared to baseline. However, there was no significant difference in the pressor response presented on day 7 and day 14 for these same subjects.

Tyramine 6 mg. Two subjects were removed from the statistical analysis for this parameter because they had shown an increase in systolic BP near or superior to 30 mm Hg after the administration of 4 mg of tyramine at

one point or another in the study, so they did not receive the 6 mg dose of tyramine on that visit. Thus, since some data were missing for them, it was deemed appropriate to remove them altogether from the statistical analysis. Similar analyses were performed on the measured pressor response to 6 mg of intravenous tyramine (Figure 3). As expected, results show that there was an interaction between the day of testing and the increase in systolic BP caused by the intravenous administration of tyramine [$F_{6,46} = 3.08$; $p < .05$]. Once again, a simple main-effect test indicated that there was a significant difference in systolic BP over time only in subjects taking clomipramine [$F_{2,46} = 8.85$; $p < .001$], but not in subjects receiving placebo, duloxetine 20 mg/day or duloxetine 60 mg/day [$F_{2,46} = 0.34, 0.89$, and 0.64 , respectively; $p > .05$].

Post-hoc analysis of the results indicated that the mean increase in blood pressure in subjects taking clomipramine was significantly lower on days 7 and 14 than at baseline. However, there was no significant difference between their pressor response on day 7 and that on day 14 ($p > .05$).

Drug Plasma Levels

The average clomipramine level was 34.9 ± 5.5 ng/ml of plasma ($n = 6$) on day 7 and 93.1 ± 24.2 ng/ml on day 14, whereas those of desmethylclomipramine were 21.0 ± 3.4 and 73.2 ± 11.6 ng/ml, respectively. Linear regression analysis showed that there was no signifi-

Table 1. Effects of the Different Drug Regimens on The Cardiovascular Parameters

Parameter	Group		Baseline	Day 7	Day 14
Systolic blood pressure (mm Hg)	Placebo	Supine	113 ± 4	116 ± 2	113 ± 5
		Standing	123 ± 5	118 ± 3	122 ± 6
	Clomipramine	Supine	115 ± 4	122 ± 5	121 ± 5
		Standing	122 ± 4	119 ± 2	127 ± 5
	Low dose	Supine	115 ± 5	115 ± 3	112 ± 3
		Standing	122 ± 5	122 ± 4	114 ± 3
	High dose	Supine	114 ± 2	115 ± 1	$121 \pm 2^*$
		Standing	120 ± 3	120 ± 4	127 ± 3
Diastolic blood pressure (mm Hg)	Placebo	Supine	59 ± 3	63 ± 4	58 ± 4
		Standing	66 ± 3	64 ± 3	67 ± 3
	Clomipramine	Supine	61 ± 3	$67 \pm 3^*$	$68 \pm 2^*$
		Standing	72 ± 3	72 ± 3	72 ± 2
	Low dose	Supine	62 ± 3	60 ± 2	57 ± 2
		Standing	72 ± 5	72 ± 3	62 ± 4
	High dose	Supine	61 ± 2	61 ± 2	62 ± 2
		Standing	66 ± 7	71 ± 3	71 ± 2
Pulse (beats/min)	Placebo	Supine	66 ± 2	71 ± 2	71 ± 4
		Standing	86 ± 4	88 ± 7	85 ± 4
	Clomipramine	Supine	66 ± 6	$78 \pm 7^*$	$81 \pm 7^*$
		Standing	90 ± 7	$110 \pm 7^*$	$108 \pm 8^*$
	Low dose	Supine	68 ± 5	60 ± 3	62 ± 3
		Standing	83 ± 6	75 ± 3	79 ± 3
	High dose	Supine	62 ± 3	62 ± 2	60 ± 2
		Standing	78 ± 4	82 ± 3	82 ± 5

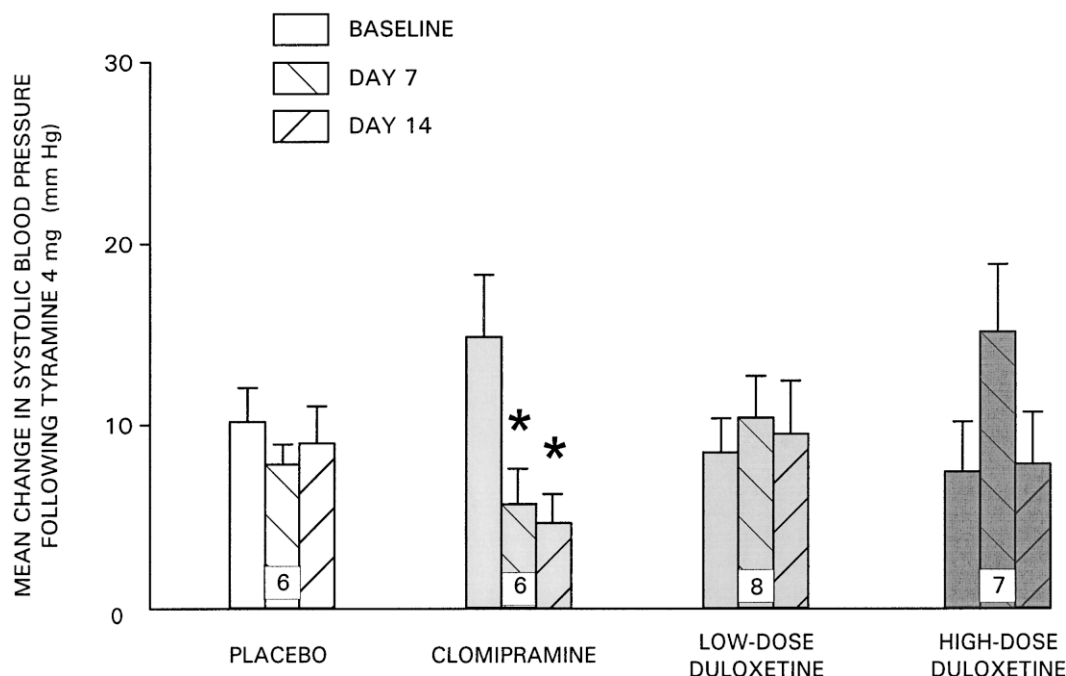


Figure 2. Mean (\pm SEM) increase in systolic BP following the intravenous administration of 4 mg of tyramine (in mm Hg), per treatment group. Systolic BP measured at baseline, after seven days, and after 14 days of drug administration. (*) indicates a significant change from baseline.

cant linear relationship between either clomipramine or desmethylclomipramine plasma levels and the pressor response to tyramine at any time during the study. Furthermore, there was no significant linear relationship between clomipramine levels and 5-HT concentrations.

In the subjects taking 20 mg of duloxetine per day, the average plasma level of the medication was 6.0 ± 2.1 ng/ml on day 7 and 8.6 ± 2.3 ng/ml on day 14. There was no significant difference in duloxetine plasma levels between day 7 and day 14 [$W(8) = 28$; $p > .05$] in that group. In subjects taking the duloxetine dose-escalation regimen, the drug concentration on day 7, when they reached the 40 mg/day dose, was 15.1 ± 1.9 ng/ml and that on day 14 was 27.9 ± 6.6 ng/ml. The difference in plasma concentration between day 7 and day 14 was not quite significant [$t(6) = -2.30$; $p = .06$] for that group of subjects either. There was no correlation between the duloxetine plasma level and the pressor response to tyramine, nor was there a linear relationship between the plasma level of the drug and the whole blood 5-HT level, at any of the different times of testing.

DISCUSSION

The aim of this study was to assess the effects of duloxetine on the 5-HT and the NE reuptake processes in healthy male subjects. Duloxetine, at doses of 20, 40, and 60 mg/day, significantly interfered with the 5-HT

reuptake processes, as demonstrated by the decrease in blood 5-HT concentrations. However, the same doses of duloxetine failed to impede the usual increase in blood pressure that follows a tyramine intravenous infusion, indicating that this drug did not alter the NE reuptake process. The tricyclic antidepressant clomipramine, known to potently inhibit both the NE and the 5-HT reuptake process, caused a significant decrease in whole blood 5-HT concentration and also markedly reduced the pressor response to tyramine. The effectiveness of clomipramine to decrease both the 5-HT concentration and the pressor response to tyramine supports the validity of the assays used in this study and the conclusion that, at doses tested on our population of healthy volunteers, duloxetine acted as a selective 5-HT reuptake inhibitor, while having no effect on the NE reuptake process.

There was a high between-subject variability in duloxetine plasma levels. This was not unexpected given that other commercially-available SSRIs show similar between-subject variabilities in plasma levels, with, for example, a five-fold difference in the plasma levels of the SSRI paroxetine at the standard effective dose of 20 mg/day.

The finding of a significant 5-HT reuptake blockade by duloxetine is in keeping with results from animal studies (Wong et al. 1993; Engleman et al. 1995; Kasamo et al. 1996; Fuller et al. 1994; Kihara and Ikeda 1995) and previous studies performed on platelets of human volunteers (Wong et al. 1993; Ishigooka et al. 1997). The

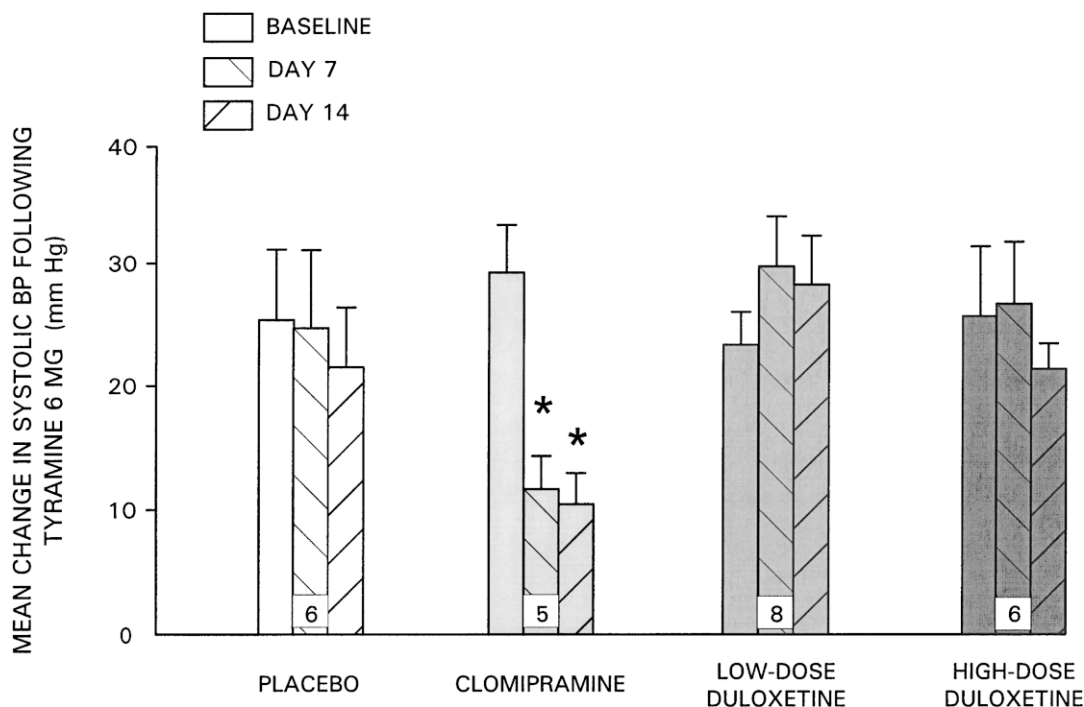


Figure 3. Mean (\pm SEM) increase in systolic BP following the intravenous administration of 6 mg of tyramine (in mm Hg) per treatment group. Systolic BP measured at baseline, after seven days, and after 14 days of drug administration. (*) indicates a significant change from baseline.

fact that there was no correlation between duloxetine or clomipramine plasma levels and 5-HT concentrations can be explained by the marked potency of these drugs to block 5-HT reuptake even at the lowest dose used. For example, in a previous study conducted in our laboratory using the same experimental paradigm, paroxetine, a SSRI with a high affinity for the 5-HT carrier, but not the much weaker SSRI nefazodone, blocked the 5-HT reuptake process (Debonnel et al. 1998). The lack of effect of nefazodone on the 5-HT reuptake process in this case, in addition to its weaker potency to block 5-HT reuptake, could be explained by its short half-life. Indeed, the activity of the 5-HT carrier probably returned to normal between the two daily administrations of nefazodone, thus precluding a depletion of the platelet stores of 5-HT.

As expected, clomipramine significantly decreased the pressor response to intravenous administration of tyramine. This effect, however, was less pronounced than that obtained with desipramine, which practically abolished the pressor response to tyramine in the abovementioned study (Debonnel et al. 1998). The *in vitro* inhibition of NE uptake caused by desipramine is stronger than that caused by clomipramine, but not that of its metabolite, desmethylclomipramine. The half-maximally effective drug concentration (the IC_{50} value) of desipramine is 1.2 nM on [3H]NE uptake in rat brain preparations, whereas that of desmethylclomipramine

is 0.5 nM and that of clomipramine is 60 nM (Baldessarini 1985).

Given that the mean plasma concentration of desmethylclomipramine was 73.2 ± 11.6 ng/ml (on day 14) and that of desipramine was lower in the previous study, it appears at first glance difficult to account for the lesser effect of the clomipramine treatment when compared to the abovementioned desipramine regimen (100 mg /day). Possibly, in comparison to desipramine, clomipramine's greater affinity for α_1 -adrenoceptors or its greater antagonism of muscarinic receptors, an action which Schäfers et al. (1997) suggest may enhance the pressor response to tyramine, may explain why clomipramine produced a lesser attenuation of the tyramine pressor response. Further studies in animals and humans using tricyclic and non-tricyclic NE reuptake inhibitors with different affinities for adrenergic and muscarinic receptors should help resolve this apparent discrepancy. These experiments are presently ongoing in our laboratory.

The lack of effect of duloxetine on the tyramine pressor response was an unexpected finding since animal studies have repeatedly shown that it blocks NE reuptake *in vivo* (Wong et al. 1993; Engleman et al. 1995; Kasamo et al. 1996; Fuller et al. 1994; Kihara and Ikeda 1995). This cannot be accounted for by an effect of duloxetine at α_1 - or β_1 -adrenoceptors or at muscarinic receptors since it has no affinity for these receptors

(Wong et al. 1993). Furthermore, a possible desensitization of peripheral β -adrenoceptors (which might be expected to attenuate pressor response to tyramine) by duloxetine and the tricyclic antidepressants is unlikely to have contributed a significant role in the present study because these drugs produce this adaptive change in the brain only after long-term administration (see Wong 1998). Therefore, there is the possibility that insufficient doses of duloxetine were used to block NE reuptake. Animal studies have demonstrated duloxetine to be only about twice as potent in blocking the reuptake of 5-HT as it is in blocking that of NE (Wong et al. 1993; Kasamo et al. 1996). Since the reuptake of 5-HT was inhibited in our human volunteers at the lowest dosage (20 mg/day), it seems unlikely that the 60 mg/day regimen was insufficient to block the NE carrier. Nonetheless, differences between animals and humans in relative potencies for blocking NE and 5-HT reuptake or in the absorption, distribution, and metabolism of duloxetine may be such that the 60 mg/day regimen was less than necessary to achieve a detectable amount of NE reuptake blockade.

Another possibility to explain the lack of efficacy of duloxetine to suppress the pressor response to tyramine comes from the study of venlafaxine, another putative dual uptake inhibitor of 5-HT and NE. In a previous study (Debonnel et al. 1998), venlafaxine in doses up to 300 mg/day had no effect on the pressor response to tyramine in healthy volunteers. One tentative explanation is that venlafaxine and duloxetine might interact with a different NE reuptake site, yet to be

identified, than tricyclic antidepressants, thereby producing differential effects on the indirectly-acting sympathomimetic tyramine. Indeed, in the case of venlafaxine, it has been shown that there is a marked discrepancy between its affinity with the NE carrier, using [3 H]-nisoxetine as a ligand, which is extremely low, and its great potency to block NE reuptake *in vivo*. This finding would support the idea of two distinct uptake sites for NE. This is, however, not the case with duloxetine, as duloxetine is endowed with both a high affinity for the NE carrier, using [3 H]-nisoxetine as a ligand, and a high potency to block NE reuptake *in vivo* (Béique et al. 1998, 1999).

An additional intriguing result comes from the study of depressed patients who underwent the same procedure as in the present study before and during a venlafaxine treatment (Debonnel et al. 1998). In the latter study, the pressor response to tyramine of patients receiving 225 mg/day of venlafaxine was significantly decreased after a 7-day titration, which stands in sharp contrast to the lack of effect of venlafaxine in our healthy volunteers, even if the dose used in the volunteers was higher (300 mg/day). Thus, it is possible that the NE reuptake process is differentially modulated in healthy volunteers and depressed patients. This hypothesis is supported by results from other investigators showing an increased sensitivity to tyramine in depressed patients, compared to control subjects (Coppen and Ghose 1978; Friedman 1978; Klimek et al. 1997). The biological basis for this difference remains to be elucidated, but a few possibilities can be envisaged. De-

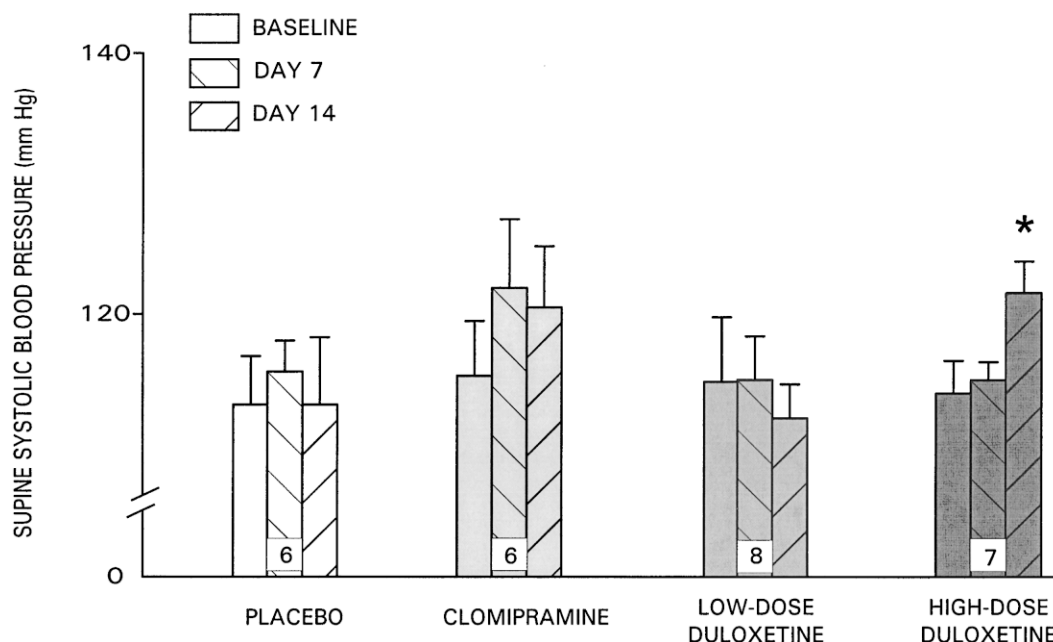


Figure 4. Systolic BP in mm Hg (mean \pm SEM) in supine position (prior to injection of tyramine) per treatment group. (*) indicates a significant change from baseline.

pressed patients might have fewer NE carrier sites; the physiological reuptake process of NE would then be easier to block. Indeed, a study comparing [^3H]-nisoxetine binding to NE carrier sites in brain tissue from depressed subjects to those of age-matched controls found that there was a decreased binding of the ligand in the midcaudal portion of the locus coeruleus in the depressed patients, indicating a decreased density of NE carriers (Klimek et al. 1997). Alternatively, the postsynaptic adrenoceptors mediating the effect of the released NE by tyramine may be more sensitive in depressed patients than in healthy volunteers. Finally, another explanation for the putative increased sensitivity of depressed patients to tyramine test is that the intraneuronal stores of NE may be greater in depressed patients than in volunteers.

Another interesting result is that there was no change in plasma NE levels, either supine or standing, in the volunteers. Those results stand in contrast to those of Flament et al. (1987) who found a significant increase in standing plasma NE levels in children with obsessive-compulsive disorders, following a five-week treatment with clomipramine. One possible explanation for this difference between their subjects and those of our present study is that the plasma levels of desmethyl-clomipramine, the metabolite of clomipramine with a high affinity for the NE reuptake site, attained in our study (73 ± 12 ng/ml) were lower than those achieved in the study by Flament et al. (208 ± 86 ng/ml). Nevertheless, some of our findings are in agreement with certain results reported by Flament et al. (1987): the increase in supine and standing pulse, as well as the increase in diastolic BP, in patients receiving clomipramine. The changes in these cardiovascular parameters may well represent a physiologic reaction to an increased availability of NE. None of these parameters were affected by the lower duloxetine regimen, which is in agreement with the lack of effect of this drug regimen on the pressor response to tyramine, suggesting no NE reuptake blockade. The higher dose of duloxetine (60 mg/day) did not alter any of the cardiovascular parameters except one: a significant increase in supine systolic BP on day 14 (Figure 4). Importantly, this increase was not observed on day 7, when the same subjects were receiving a lower dosage of duloxetine was used (40 mg/day). The possibility therefore remains that a dose of duloxetine of 60 mg/day may represent the threshold for NE reuptake blockade, since doses of 20 mg/day and 40 mg/day did not increase the supine systolic BP, while a regimen of 60 mg/day of duloxetine did. The fact that this parameter was not affected by the clomipramine regimen does not invalidate this interpretation. Indeed, the differential effects of duloxetine and clomipramine on supine systolic BP might be due to the blockade of α -adrenoceptors by the latter drug only (Baldessarini 1985).

In conclusion, the present study has demonstrated that duloxetine acts as a selective 5-HT reuptake inhibitor in non-depressed human male volunteers, and that, at doses up to 60 mg/day, it failed to inhibit the tyramine pressor response, as would be otherwise expected of a NE reuptake blocker. It would be interesting to determine whether daily dosages higher than 60 mg/day could suppress the tyramine pressor response in healthy volunteers, and whether there would be any difference observed in this pressor response in depressed subjects versus healthy subjects.

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