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Sensory signalling effects of tegaserod in patients with irritable bowel syndrome with constipation

Running title: Sensory Signalling Effects of Tegaserod in IBS-C

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Abstract

Tegaserod relieves overall and multiple individual constipation-predominant IBS (IBS-C) symptoms. However, mechanisms for the relief of abdominal pain/discomfort are not well understood. The effects of tegaserod on rectal sensitivity to distension were measured by the nociceptive flexion RIII reflex, as evidenced by spinal hyperexcitability (i.e. increase or facilitation of the RIII reflex), in IBS-C patients. A randomized, double-blind, placebo-controlled, parallel study was performed in 30 women with IBS-C. The effects of slow ramp rectal distension on the RIII reflex, recorded from the lower limb, were measured before (D1) and after 7 days (D8) of placebo ($n = 15$) or 6 mg tegaserod bid ($n = 15$). Pressure–volume and sensation–volume relationships were measured during distension, and patients reported their IBS symptoms daily. On D1, rectal distension facilitated the RIII reflex in both treatment groups. On D8 versus D1 these facilitatory effects were significantly lower ($P < 0.001$, ANOVA) after tegaserod (mean reduction: $-30.3 \pm 11.9\%$) than placebo (mean reduction: $-10.1 \pm 12.9\%$). No significant changes in the volume–sensation relationship or differences in compliance were observed with tegaserod or placebo. In conclusion, tegaserod reduces the facilitatory effects of rectal distension on the RIII reflex in women with IBS-C.

Keywords: abdominal pain, constipation-predominant irritable bowel syndrome, IBS-C, RIII reflex, tegaserod

Irritable bowel syndrome (IBS), one of the most common functional gastrointestinal disorders, is characterized by recurrent abdominal pain/discomfort and abnormal bowel function.¹ Pain and discomfort may be a consequence of hyperexcitability of nociceptive systems (i.e. central sensitization),² which could explain the hypersensitivity to rectal distension consistently reported in most IBS patients.³ This new pathophysiological concept highlights the need for standardized and reproducible methods to evaluate visceral sensitivity in humans. We therefore developed a reflexological technique based on the modulation of a somatic nociceptive cutaneomuscular flexion reflex (RIII reflex) during gut distension as a possible tool for objective assessment of visceral sensitivity in humans.⁴⁻⁹

The RIII reflex is a polysynaptic spinal reflex elicited by electrical stimulation of a cutaneous sensory nerve (usually the sural nerve at the ankle) that can be recorded from a flexor muscle on the ipsilateral limb (usually the biceps femoris).¹⁰ It is considered to be a reliable index of spinal transmission of nociceptive signals, and indeed, its threshold and amplitude are related closely to those of concomitant painful cutaneous sensations evoked by electrical stimulation.¹⁰ In healthy subjects, we demonstrated that gastric or rectal distensions induced a reduction of the RIII reflex response and that the change in the response was correlated with both the volume of the distension and the visceral sensation.^{4,6,7} Using this methodology we demonstrated that, in about two thirds of IBS patients, slow ramp rectal distension induced an increase (“ie facilitation”) of the RIII reflex response.⁹ These facilitatory effects, which probably reflect alterations of pain modulatory mechanisms, can therefore be used as an objective and quantifiable index of the hyperexcitability of spinal nociceptive processes in a subgroup of IBS patients. We also showed that this electrophysiological approach is of value for investigating the mechanisms of action of pharmacological agents.^{5,7,8}

The efficacy of tegaserod, a 5-HT₄ receptor agonist, in the treatment of IBS with constipation (IBS-C) in women has been confirmed in several large randomized controlled

trials.¹¹⁻¹⁶ In addition, we have previously analysed the effects of tegaserod on the modulation of the RIII response induced by rectal distension in healthy women, and showed that it may act directly on nociceptive processes.⁸

Here we report a double-blind, placebo-controlled study, evaluating the effects of tegaserod on modulation of the RIII reflex induced by rectal distension in women with IBS-C.

MATERIAL AND METHODS

Participants

Following approval from the local Ethics Committee, the study was conducted from 11 July 2003 to 29 October 2004 and included 32 women, aged 18 to 60 years, who provided written informed consent.

Patients were eligible if they met IBS-C (ROME II) criteria.¹ Specifically, they had to experience abdominal pain/discomfort with two of the following characteristics for at least 12 weeks (not necessarily consecutive) of the previous 12 months: 1) relief with defecation; 2) onset associated with a change in stool frequency; 3) onset associated with a change in stool form. Each patient underwent a complete clinical evaluation to exclude organic disease. Women were excluded if they had undergone abdominal surgery other than appendectomy, and only women with a normal total colonoscopy within the previous 5 years were included in the study.

Analgesics, antispasmodics, laxatives and antidiarrhoeal agents were stopped at least 7 days before the start of the experimental protocol. IBS-C criteria were confirmed during a 7-day baseline period, during which patients completed a daily questionnaire about their abdominal pain/discomfort (0 = none to 6 = very severe), the number of bowel movements, and stool consistency (1 = watery to 7 = very hard). At the end of the baseline period, the patients responded 'Yes' or 'No' to the question 'Did you have satisfactory relief of your overall symptoms during the last week?' Only patients who reported at least 4 days with abdominal

discomfort or pain score ≥ 3 (i.e. at least moderate pain), mean stool consistency ≥ 4 (i.e. neither loose nor hard) and answered 'No' to the question about the overall relief were included in the study.

Nociceptive flexion reflex (RIII reflex)

The RIII reflex was elicited from the lower limb and recorded using an entirely computerized system (PhysioLab System, Notocord, Igny, France) as previously described.^{4,6-9} Briefly, the subjects were placed in the lateral decubitus position. The sural nerve was electrically stimulated at a frequency of 0.17 Hz (10 stimulations/min) via a pair of surface electrodes placed 2 cm apart on degreased skin over the retromalleolar path of the nerve. Each electrical stimulation consisted of a train of five constant-current pulses of 1-ms duration. Electromyographic responses were recorded from the ipsilateral biceps femoris via a pair of surface electrodes placed 2 cm apart on degreased skin over the muscle. The RIII response was identified as a multiphasic signal appearing between 90 and 180 ms after each stimulation. After amplification, each reflex response was digitized, full-wave rectified, and integrated. The RIII response was thus scored as the integrated surface area of the signal. The RIII reflex threshold in each patient was defined as the average minimal current that elicited the reflex response. The intensity of electrical stimulation of the sural nerve was then adjusted to 20% above the threshold and kept constant during the pre-distension (control), distension and post-distension periods of each experimental sequence (see below).

Rectal distension

An oversized spherical polyvinyl bag (10 cm in diameter; infinite compliance until maximal volume of 600 mL) was mounted on the tip of a double-lumen polyvinyl tube (12 Fr), folded tightly, lubricated and inserted into the rectum. The distal attachment site was 4 cm from the anal

verge. The proximal opening of the tube was linked to an electronic barostat (INRA, Toulouse, France) that allowed controlled inflation and deflation of the balloon with air, and continuous monitoring and recording of the volume and pressure inside the balloon. When in place, the balloon was unfolded by slowly injecting air under controlled pressure (< 20 mmHg) and then completely deflated. After a 20-min rest period, the barostat was used to inflate the balloon at a constant rate of 40 mL/min (slow ramp distension).

Experimental design

Each patient participated in two experimental sessions, separated by 1 week. After verification of inclusion and exclusion criteria, control pre-treatment measurements were taken on the first experimental day (D1) after a 12-hour fast. The experimental session began with the determination of the RIII reflex threshold. Rectal distension was then performed up to the pain threshold (i.e., minimal distension eliciting a pain sensation). The RIII reflex responses were measured continuously for 3 minutes before the distension (i.e., pre-distension control period), during rectal distension, and for 3 minutes following the distension. The sensations elicited by rectal distension were scored as described below. At the end of the first experimental session, the patients were randomized to receive either 6 mg tegaserod twice daily (bid) or placebo for the following 7 days. During these 7 days they were asked to record their symptoms daily (abdominal pain and discomfort, number of bowel movements and stool consistency), in a diary similar to that used for the pre-inclusion period. On Day 8 (D8), at the end of the treatment, the last tablet of tegaserod or placebo was ingested 1 hour before the second experimental session, which was performed as described for D1. During this experimental session the investigators were blinded to the patients' D1 thresholds and reflex responses

Sensation elicited by rectal distension

Before the experiments, participants were informed of the visceral sensations they might experience. The sensation elicited by the rectal distension was graded from 0 to 6 using a validated verbal questionnaire: 0 = no perception; 1 = initial perception; 2 = sensation of gas; 3 = sensation of stool; 4 = urge to defecate or onset of discomfort; 5 = moderate pain; and 6 = intense or unbearable pain.^{6,7,9} Subjects reported their sensations at fixed intervals (every 50 mL of distension). Distensions were stopped at the pain threshold (i.e., score 5). If intense or unbearable pain (score 6) was experienced during any level of distension, the experiment was ended immediately.

Data analysis

The primary endpoint was the effect of treatment on the RIII reflex response during rectal distension. Secondary endpoints were the pressure–volume and sensation–volume relationships, and the mean symptom scores and stool consistency scores after treatment.

For each distension on D1 and D8, the RIII responses were averaged at 1-minute intervals in the pre- and post-distension periods and at every 50 mL during the distension. All the reflex responses are expressed as a percentage of the mean value for the control pre-distension period. If the pain threshold was reported before 600 mL, the RIII value recorded at the maximal tolerated volume was reported for subsequent volumes to standardize data (last observation carried forward). Pressure was recorded at fixed intervals (every 50 mL of distension) during slow ramp distension.

Intergroup changes from pre-treatment (D1) to post-treatment (D8) were tested by analysis of variance (ANOVA) or the Wilcoxon rank sum test. ANOVA included treatment and distension volumes as factors. A significance level of 0.05 (two-sided) was used in all cases. Wilcoxon rank sum and Fisher's exact tests were used to assess the homogeneity of the treatment

groups. Adverse events were summarized by body system and by individual adverse events. Severity and relation to study medication were noted. No interim analysis was performed.

RESULTS

Most of the IBS-C patients selected for the study were followed for IBS symptoms by one of the physicians of the gastroenterology department; the others were recruited after advertising in a local paper. The 32 IBS-C patients were screened on D1, of whom two were excluded because they could not tolerate the electrical stimulation. Thus, 30 women with IBS-C (mean age: 42.9 ± 9.5 years) were randomized to receive either placebo ($n = 15$) or tegaserod ($n = 15$) and completed the study. Demographic and clinical characteristics were similar in both groups (Table 1).

Effects of tegaserod and placebo on the modulation of RIII reflex by rectal distension

The RIII reflex threshold was not significantly different between the tegaserod (8.0 ± 0.4 mA) and placebo (7.5 ± 0.5 mA) groups on D1, and was not significantly altered by treatment on D8 (8.2 ± 0.6 vs 7.8 ± 0.5 mA, respectively). Sural nerve stimulation at 1.2 times threshold elicited a fairly stable RIII response and evoked a moderate sensation of pain of the pinprick type in every patient; this procedure was well tolerated throughout the experiments.

On D1, the RIII reflex was increased (i.e., facilitation) during rectal distension in both groups, but the magnitude of the facilitation was significantly higher ($P < 0.05$, ANOVA) in the tegaserod group ($134.3 \pm 43.9\%$) than in the placebo group ($112.8 \pm 50.9\%$). On D8, compared with D1, the facilitation of the RIII response was significantly more reduced ($P < 0.001$, ANOVA) by tegaserod treatment (mean reduction = $-30.3 \pm 11.9\%$) than by placebo (mean reduction = $-10.1 \pm 12.9\%$). The reduction of facilitation was consistent

throughout the distension period (Figure 1). An individual example of the effects of tegaserod on the facilitation of the RIII reflex induced by rectal distension is presented in Figure 2.

The difference between the two treatment groups in the mean facilitation effects on D1 was not due to a difference in the magnitude of facilitation itself, but to the unequal distributions of patients presenting facilitation or inhibitory effects. Rectal distension increased the RIII reflex in 20 patients (eight in the placebo group and 12 in the tegaserod group), and decreased the RIII reflex in 10 patients (seven in the placebo group and three in the tegaserod group). Consequently, the data were further analysed by comparing treatment within the subgroups of patients with facilitation and inhibition of the RIII reflex on D1 (Figures 3a and c). The magnitudes of facilitation and inhibition were similar before tegaserod and placebo. On D8 in the subgroup of patients with inhibition, the inhibitory effects were not significantly altered by tegaserod or placebo (Figure 3d). In contrast, in the subgroup of patients with facilitation, facilitation was significantly reduced ($P < 0.001$) by tegaserod compared with placebo (Figure 3b).

Pressure–volume and sensation–volume relationships

The pressure–volume relationship was similar in the two groups on D1 and was not significantly affected by the treatment (Figure 4).

The sensation–volume relationship on D1 was not statistically different in the two treatment groups; the maximal volume of distension tolerated was 363 ± 123 mL in the tegaserod group and 286 ± 131 mL in the placebo group. This was not significantly modified on D8 (change from Day 1 to Day 8 was -70.0 ± 82 mL and -73.3 ± 121 mL in the tegaserod and placebo groups, respectively).

IBS symptoms

Mean IBS symptom scores (abdominal pain/discomfort, frequency of bowel movements, and mean stool consistency) on D1 did not differ significantly between the placebo and tegaserod groups (Table 1). The mean abdominal pain score was significantly lower after treatment with tegaserod than placebo (2.45 ± 1.29 vs 3.06 ± 0.73 , $P < 0.05$). No significant change was observed in the frequency of bowel movements or mean stool consistency scores.

Safety

During the study, three of the 30 patients enrolled reported adverse events; two in the tegaserod group and one in the placebo group. In the tegaserod group, one patient experienced abnormal bowel sounds and another experienced nausea, headache and dry mouth. In the placebo group, one patient reported nausea. No serious adverse event was observed during the study.

DISCUSSION

This placebo-controlled study demonstrates that tegaserod reduces the pathological facilitation of the RIII flexion reflex induced by rectal distension in women with IBS-C. The threshold of the RIII reflex and the elastic properties of the rectum were not altered by tegaserod, suggesting possibly a selective interaction with spinal processing of sensory visceral information. These effects, probably the consequence of a reduction of spinal hyperexcitability, were associated with a significant decrease in abdominal pain/discomfort scores and, therefore, may contribute to the analgesic action of tegaserod.

Phase II and Phase III studies show that tegaserod effectively provides relief from overall and individual IBS symptoms, including a decreased abdominal pain, improved stool consistency, and increased bowel movement frequency with an acceleration of colonic transit

time for women with IBS-C.¹¹⁻¹⁶ However, the mechanisms of the analgesic action of tegaserod are not clearly established. In decerebrate cats, tegaserod dose-dependently inhibited the firing of rectal afferents during distension.¹⁷ A direct inhibition of mucosal mechanoreceptors may be a likely effect of tegaserod as it is rapidly absorbed and its actions mimic those of endogenous serotonin released from enterochromaffin cells, stimulating intrinsic sensory neurons in the intestinal mucosa.¹¹ The unchanged visceral pain threshold observed here is consistent with this hypothesis. Also, the unchanged RIII reflex threshold in this study suggests that tegaserod does not have general analgesic effects, but may act selectively on sensory visceral processes.

We found that slow rectal distension facilitated the RIII reflex in approximately two thirds of IBS patients. This proportion was similar to that observed in our previous study using the same methodology but with a different cohort of IBS patients.⁹ Thus, our findings are consistent with IBS being a heterogeneous clinical entity resulting from more than one pathophysiological mechanism.

The abnormal increase in the RIII reflex during rectal distension may reflect a true hyperexcitability of the spinal viscerosomatic nociceptive processes (i.e., central sensitization), which could contribute to the pathophysiology of chronic abdominal pain in a subgroup of IBS patients. The normal RIII reflex threshold in patients in this study suggests that any such spinal hyperexcitability was not secondary to a direct alteration of the properties of spinal nociceptive neurons, but is the consequence of a failure of segmental and/or heterosegmental (i.e., descending) systems that modulate the spinal transmission of nociceptive signals. Interestingly, studies using other approaches also concluded that IBS symptoms were the result of abnormalities in pain modulation.¹⁸ The lower facilitation of the RIII reflex induced by rectal distension after tegaserod treatment may result from action on pain modulatory systems. The role of serotonergic systems in the descending modulation of the spinal transmission of nociceptive signals has been established in animals, but is less well

described in humans.¹⁹ However, a specific pain modulatory system, Diffuse Noxious Inhibitory Controls (DNIC), initially described in the rat,²⁰ has also been demonstrated in humans and involves serotonergic systems.^{21–24}

An interesting finding with potential clinical significance in the present study is that tegaserod preferentially exhibits effects in patients presenting with a facilitation of the RIII reflex, but not in those presenting with a inhibition of the RIII reflex, during slow ramp rectal distension on D1 before the treatment period. These differential effects are again consistent with IBS being a heterogeneous condition, consisting of different subgroups of patients who respond differently to treatment. Identification of such subgroups may allow therapeutic strategies to be tailored to individual patients and thereby significantly improve treatment outcomes. Classification of patients according to a measurable neurophysiological index might be of particular value because the clinical criteria currently used, such as the ROME II criteria, do not discriminate between subgroups of patients according to pain mechanisms.²⁵ Most Phase II and III clinical studies are performed with subgroups of IBS patients defined by their altered bowel habits (i.e., diarrhoea or constipation). These criteria have no direct relationship with pain mechanisms, and this could explain the relatively small magnitude of the analgesic effects generally reported by Phase II and III trials of IBS treatments. Our findings suggest that analysis of the effects of rectal distension on the RIII reflex may enable identification of subgroups of patients with spinal hyperexcitability that may respond differently to treatment. This interesting hypothesis needs to be tested with other clinically active pharmacological agents.

In conclusion, this study showed that tegaserod had a true objective effect on visceral sensitivity by reducing the facilitatory effects of rectal distension on the RIII reflex in a large subgroup of IBS-C patients.

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Amy Wagner is an employee of, and owns stocks and shares in, Novartis Pharma AG, Switzerland. Yolande Loria is an employee of Novartis Pharma SA, France.

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Table 1 Baseline demographic and clinical characteristics of the patients

Variables	Tegaserod (n = 15)	Placebo (n = 15)
Age (yr)		
Mean (SD)	44.9 (7.4)	41.0 (11.1)
Range	29.0–56.0	22.0–54.0
Race (%)		
Caucasian	80.0	100.0
Black	13.3	0
Asian	6.7	0
Height (cm)		
Mean (SD)	161.7 (5.8)	162.9 (6.3)
Range	154.0–176.0	151.0–170.0
Weight (kg)		
Mean (SD)	60.4 (15.0)	62.7 (12.1)
Range	41.0–88.0	44.0–92.0
Body mass index (kg/m ²)		
Mean (SD)	23.0 (5.3)	23.6 (4.1)
Range	17.1–33.3	17.2–31.8
Abdominal pain (0 = none, 6 = very severe)		
Mean (SD)	3.06 ± 0.73	3.35 ± 0.81
Bowel movements		
Mean (SD)	0.73 ± 0.47	0.87 ± 0.58
Stool consistency (1 = watery, 7 = very hard)		
Mean (SD)	4.63 ± 1.22	5.06 ± 1.16

SD = standard deviation.

FIGURE LEGENDS

Figure 1 Overall effects of tegaserod (black circles) and placebo (white circles) on the facilitation of the RIII reflex during rectal distension.

Data (mean \pm standard deviation [SD]) are expressed as % change from baseline (D8 minus D1) during rectal distension (0–600 mL). Reduction in the facilitation of the RIII reflex was significantly greater with tegaserod than placebo ($P < 0.001$, analysis of variance [ANOVA]). Data recorded at each minute of the 3 minutes immediately following distension are also shown (P1–P3).

Figure 2 An example of RIII reflex recordings during slow ramp distension for an individual patient before and after 1 week of treatment with tegaserod.

Each bar represents a single reflex response, expressed as a percentage of the control value (i.e., mean value recorded during each minute of the 3-minute control period prior to distension [C1–C3]). The distension is indicated by arrows. Facilitation of the RIII reflex induced by rectal distension was lower on D8 than on D1. P1–P3 refers to each minute of the 3-minute period following distension.

Figure 3 Effects of tegaserod and placebo on the modulation of the RIII reflex during rectal distension according to RIII reflex profile on D1.

Data (mean \pm standard deviation [SD]) are shown for patients with facilitation (squares) before (a) and after treatment (b) for patients with inhibition (triangles) before (c) and after treatment (d). Tegaserod induced a significant ($P < 0.001$) reduction of the facilitatory effects (b). In contrast, the inhibitory effects (d) are not altered by tegaserod. No significant changes were observed after placebo.

Figure 4 Pressure–volume relationships before (a) and after (b) 1 week of treatment with placebo (white) or tegaserod (black).

The pressure–volume curves were not significantly altered by treatment.

LIST OF ABBREVIATIONS USED

ANOVA	analysis of variance
bid	twice daily
D1	Day 1 (first experimental day)
D8	Day 8
DNIC	Diffuse Noxious Inhibitory Controls
FDA	Food and Drugs Administration
IBS	irritable bowel syndrome
IBS-C	IBS with constipation
SD	standard deviation

Figure 1

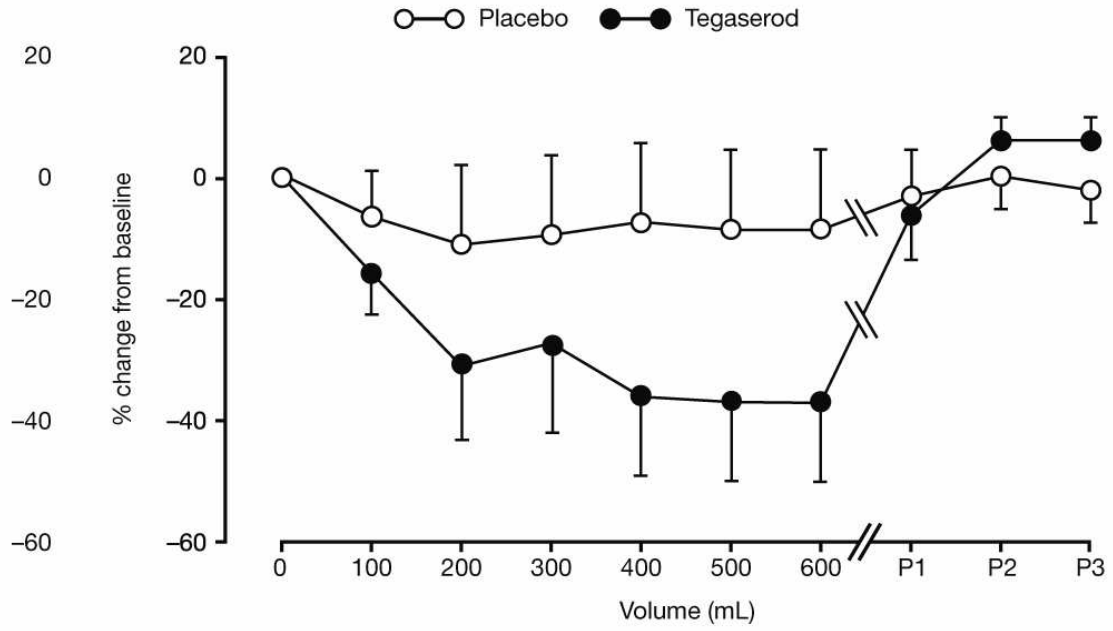


Figure 2

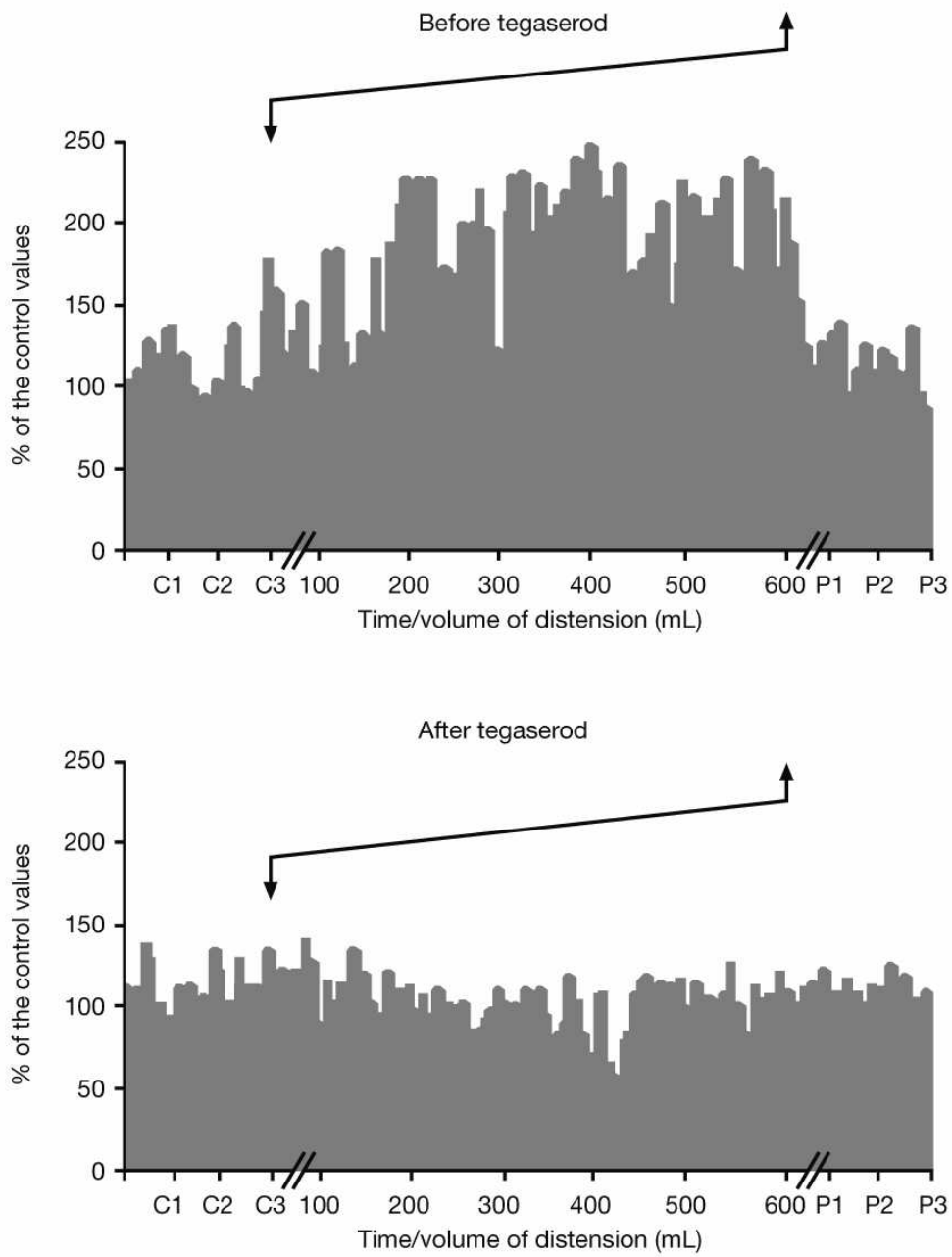


Figure 3

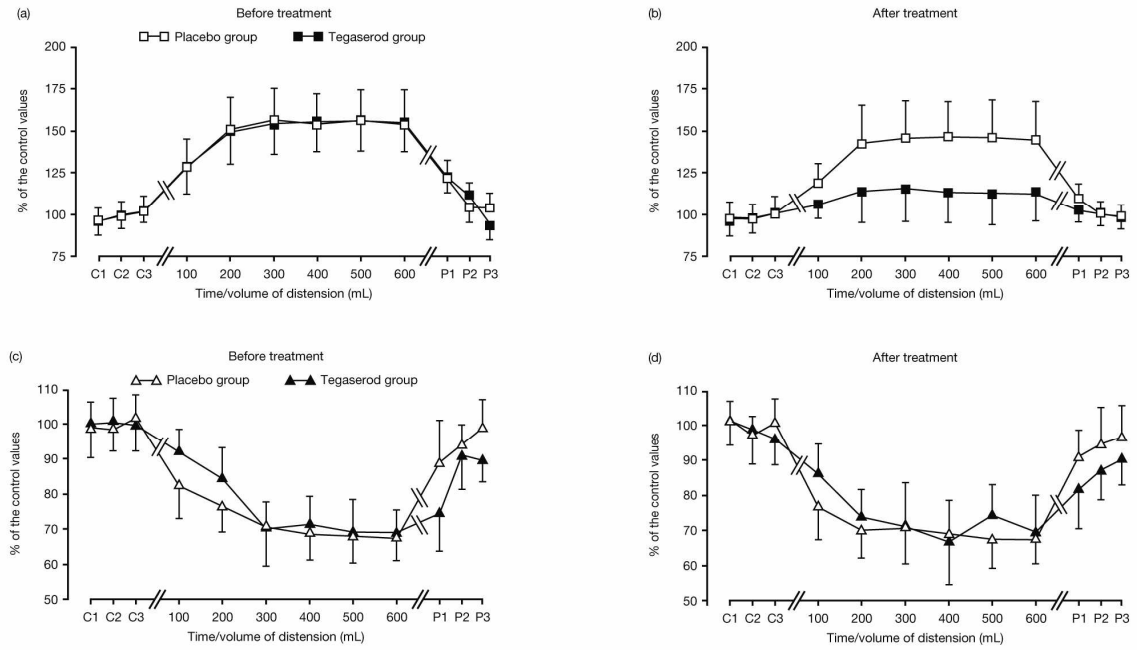


Figure 4

