ABSTRACT

BACKGROUND AND OBJECTIVES: Tramadol blocks somatosensory potentials in vitro and may be associated to local anesthetics to improve analgesic quality. This study aimed at evaluating whether tramadol changes lidocaine motor block regression in two different concentrations.

METHOD: Male Wistar rats weighing 250 to 300 g were submitted to sciatic nerve block guided by percutaneous nerve stimulation. Animals were distributed in four groups (n = 5 per group): 2% lidocaine (GI), 0.5% lidocaine (GII), 2% lidocaine/1.25 tramadol (GIII), 0.5% lidocaine/1.25 tramadol (GIV). Partial and total motor block regression times were evaluated.

RESULTS: All animals had total motor block when awakening from anesthesia, which has totally regressed during the observation period. Total regression time of 2% lidocaine was 41 ± 1.71 minutes, 0.5% lidocaine was 25.26 ± 0.83 minutes, 2% lidocaine/tramadol was 46.06 ± 0.88 minutes and 0.5% lidocaine/tramadol was 36.15 ± 1.18 minutes. The association of 0.5% lidocaine and 1.25 mg tramadol was more effective as compared to 0.5% lidocaine alone. Data are presented in mean ± mean standard error (mse), considering significant p < 0.05 using ANOVA followed by Tukey test.

CONCLUSION: Tramadol has effects similar to local anesthetics and, when used as adjuvant of lidocaine, prolongs motor block duration in rats.

Keywords: Lidocaine, Nervous block, Tramadol.

INTRODUCTION

Tramadol (1-RS, 2RS)-2-[(dimethyl-amine)-methyl]-1-(3 methoxyphenyl)-cyclohexanol hydrochloride is a central action drug sold as a racemic mixture of two enantiomers [(+) and (-) tramadol]. The methyl group in the phenolic part of the molecule is responsible for the opioid agonist activity and its affinity for μ receptors is approximately 6 thousand times lower than morphine, 100 times lower than dextropropoxyphene and 10 times lower than codeine1. After systemic administration, tramadol is demethylated by the P450 cytochrome system in o-desmethyl-tramadol (M1), active metabolite with agonist activity in...
μ receptors 200 times higher than the original molecule. It has pharmacological effect similar to local anesthetics, blocking action potential conduction in isolated nerves. By spinal route, it suppresses spinal cord somatosensory potentials and after perineural injection it induces total motor block. A previous study has described different motor block intensities by perineural tramadol in sciatic nerve of rats, where motor block induced with 5 mg tramadol was similar to motor block induced with 2% lidocaine.

This study aimed at evaluating the possibility of tramadol potentiating duration and intensity of motor block induced by 0.5% and 2% perineural percutaneous lidocaine in the sciatic nerve of rats. Parameter was motor block regression time.

METHOD
Twenty male Wistar rats weighing 250 to 300 g were placed in pairs in cages with 12-hour light-dark cycles. Water and food were supplied ad libitum. Animals were supplied by the Central Vivarium of the School of Medicine, University of São Paulo (FMUSP) and experiments were carried out in FMUSP's LIM-08, after adaptation to the study environment for 30 minutes.

Anesthetic technique: animals were placed in a closed chamber where 4% isoflurane in oxygen was supplied by gauged vaporizer for anesthetic induction – anesthesia was maintained with 1% isoflurane via facial mask to allow sciatic nerve block.

Sciatic nerve block: after being anesthetized, right femur greater trochanter was located by palpation and a 2.5 cm needle without bevel (B Braun, Germany) connected to a Stimulation nerve stimulator (B Braun, Brazil) to locate the sciatic nerve. The needle was introduced at 1 mm from right femoral shaft, in the notch located between greater trochanter and ischial tuberosity and was then directed to the ischium. A ground electrode was fixed to the right ear and the needle was connected to an electric cable which supplied initial current with 0.6 mA intensity able to promote right thigh muscle contraction, which was progressively increased the greater the proximity to the sciatic nerve. Current was decreased to 0.2 mA observing the muscle contraction response pattern and, with the help of a Hamilton syringe, 50 μL lidocaine (Groups I and II) or tramadol/lidocaine (Groups III and IV) were administered by the previously filled needle lateral extension (15 μL). After drug administration the extension was washed with 15 μL saline.

Motor block evaluation: time for progressive and total motor block regression was recorded and was characterized by observing animals’ gait. Values from zero to 3 were attributed to the following criteria: 0 = total absence of motor block, unchanged gait; 1 = minimum motor block, normal gait however with paw inversion; 2 = moderate motor block, animals traction the paw using thigh muscles, but paw is flaccid and does not support the plantar aspect on surface; 3 = total motor block, totally flaccid paw.

Experimental design: animals were distributed in four different study groups. The first group (GI) was submitted to sciatic nerve block with 2% lidocaine. The second group (GII) received 0.5% lidocaine. The third group (GIII) received 2% lidocaine and 1.25 mg tramadol. The fourth group (GIV) received 0.5% lidocaine associated to 1.25 mg tramadol. Motor block duration and intensity were observed in the four groups. Tramadol concentration was based on a previous study, which has observed the presence of minimum motor block with 2.5% tramadol concentration without changing flinching ability.

Statistical analysis
Data are shown as mean ± MSE (mean standard error) of 5 animals from each group and were analyzed by ANOVA followed by Tukey test for multiple comparisons with the help of GraphPad Prism version 5.0 software, considering significant p < 0.05.

This study was approved by the Research Ethics Committee, School of Medicine, University of São Paulo (protocol 051/2002).

RESULTS
Sciatic nerve was blocked according to the technique described by Sousa et al., in compliance with IASP ethical recommendations for the study of conscious animals. Emergence time of rats after isoflurane withdrawal was similar for the four groups (2.13 ± 0.18, 2.33 ± 0.25, 2.58 ± 0.22, 2.47 ± 0.31 minutes for 2% lidocaine, 0.5% lidocaine, 2% lidocaine/tramadol and 0.5% lidocaine/tramadol) respectively GI, GII, GIII, GIV.

In all groups, animals had total motor block at emergence (degree 3).

Total motor block regression time of 2% lidocaine (41 ± 1.71 minutes) was significantly better than of 0.5% lidocaine (25.26 ± 0.83 minutes) (p < 0.05). Tramadol associated to lidocaine has prolonged total motor block regression time of 0.5% lidocaine (36.16 ± 1.19 versus 25.26 ± 0.83 minutes, of tramadol / 0.5% lidocaine and of 0.5% lidocaine, respectively) (p < 0.05), but did not significantly change 2% lidocaine total motor block regression time (46.06 ± 2.88 versus 41 ± 1.71 minutes, for tramadol / 2% lidocaine and 2% lidocaine, respectively) (p > 0.05) (Graph 1).

Graph 1 – Lidocaine-induced motor block regression time (in minutes) in sciatic nerve of rats.
The addition of 1.25 mg tramadol to 0.5% lidocaine solution has significantly increased the duration of the effect of 0.5% lidocaine alone (p < 0.05). ANOVA followed by Tukey test. Motor block regression time from degree 3 to degree 2, however, was longer in the group receiving 2% lidocaine associated to tramadol (6.85 ± 0.36 minutes) as compared to 2% lidocaine alone (4.73 ± 0.21 minutes) (p < 0.05) (Table 1).

## DISCUSSION

The administration of anesthetic solutions close to peripheral nerves or neural complexes induces longer anesthesia as compared to neuraxial blocks, depending on physical agent characteristics and the presence or not of vasoconstrictors. Combinations of local anesthetics and adjuvants, such as epinephrine, ketamine, neostigmine, clonidine and dexmedetomidine aim at prolonging analgesia time, allowing the use of local anesthetics in low concentrations and subsequent decrease of drug noxious effects.

Our study has evaluated duration and total regression of motor block induced by different lidocaine concentrations in mixed peripheral nerve, made up of sensory and motor fibers of different diameters with different sensitivities to local anesthetics. According to our results, it is clear that tramadol added to lidocaine has prolonged motor block duration. Previous studies have reported significant better quality of analgesia with tramadol associated to ropivacaine and intra-articular bupivacaine, as well as with lidocaine for brachial plexus block in orthopedic surgeries in humans; however, authors have not mentioned motor block duration. Our study has shown that motor block induced with 0.5% lidocaine, but not with 2% lidocaine, was prolonged by the addition of tramadol.

Such effect might have been caused by tramadol action on perineural adrenergic fibers, prolonging lidocaine action on sciatic nerve fibers. However, the most likely possibility for such effect is the action of tramadol on the kinetics of sodium channel, where it decreases neural excitability, the mechanism of which is possibly different from lidocaine and not totally clear. In addition, in vivo, tramadol blocks neural somatosensory potentials conduction and has local anesthetic effect as effective as 2% prilocaine. When directly applied on the sciatic nerve, it dose-dependently blocks neural conduction with lower potency as compared to lidocaine, being critical the proximity of neural sheaths for the synergistic effect.

We have not found significant increase in total duration of 2% lidocaine effect in this model. However, animals receiving tramadol associated to 2% lidocaine had deep neural block (degree 3) for a longer time as compared to those submitted to 2% lidocaine alone, which could be understood as indirect measurement of motor block intensity. One hypothesis for this phenomenon might have been the participation of tramadol in the early conduction blockade stage, where blockade duration is limited by drug potency.

Clinically, the association of tramadol to loco-regional anesthesia might decrease surgical stress and improve postoperative recovery in humans, due to the physiological advantages of such techniques and to the possibility of early hospital discharge. However, there have been limitations to the technique and the models of our study because it was impossible to measure blockade onset time.

## CONCLUSION

Current results allow concluding that tramadol may be used as adjuvant for lidocaine, prolonging motor block recovery time in rats, possibly by mechanisms similar to local anesthetics.

## REFERENCES

ERRATUM

Evaluation of the addition of tramadol on lidocaine-induced motor block regression time. Experimental study in rats

Avaliação da adição do tramadol sobre o tempo de regressão do bloqueio motor induzido pela lidocaína. Estudo experimental em ratos

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In the above mentioned article, graph 1 was published with error.

Graph 1 – Lidocaine-induced motor block regression time (in minutes) in sciatic nerve of rats.

Best regards,
Angela Maria Sousa