Neuroprotective Effects of Agmatine in Experimental Peripheral Nerve Injury in Rats: A Prospective Randomized and Placebo-Controlled Trial

Sıçanlarda Deneysel Periferik Sinir Yaralanmasında Agmatinin Nöroprotektif Etkileri: Prospektif, Randomize, Plasebo Kontrollü Çalışma

Aykut SEZER1, Bulent GUCLU1, Burak KAZANCI2, Murteza CAKIR3, Mustafa Kemal COBAN4

1Ministry of Health, Sevket Yılmaz Research and Training Hospital, Department of Neurosurgery, Bursa, Turkey
2Ufuk University, Faculty of Medicine, Department of Neurosurgery, Ankara, Turkey
3Ataturk University Hospital, Department of Neurosurgery, Erzurum, Turkey
4Ministry of Health, Erzurum Research and Training Hospital, Department of Neurosurgery, Erzurum, Turkey

Corresponding Author: Bulent GUCLU / E-mail: guclubulent@hotmail.com
INTRODUCTION
Peripheral nerve injury is a common health problem that often results in both social and economic losses for the victim (2). Peripheral nerve injury may occur as a result of trauma (blunt or penetrating) or acute compression and result in demyelination or axonal degeneration. Clinically, both demyelination and axonal degeneration result in disruption of the sensory and/or motor function of the injured nerve. Recovery of function occurs with the remyelination, axonal regeneration and reinnervation of the sensory receptors, muscle end plates, or both. No definitive treatment for peripheral nerve injuries exists. Functional recovery after peripheral nerve injury is often poor; however, unlike in the central nervous system, regeneration in the peripheral nervous system is possible (4). New therapies that protect injured peripheral neurons and enhance regeneration are needed. The purpose of this study was to demonstrate the activity of agmatine, an inducible nitric oxide synthase (iNOS) inhibitor and selective N-methyl-D-aspartate receptor (NMDAR) antagonist, on reducing tissue damage in distal part of trauma in an experimental rat peripheral nerve injury model.

MATERIAL and METHODS
I- Laboratory Study
This experimental study was conducted in the Animal Laboratory of the Pharmacology Department, Histology Department, Pathology Department, and Neurosurgery Department of Ataturk University / Erzurum / Turkey. The approval of the Ethics committee approval was obtained for the study from the Local Research Ethics Committee.

This study involved thirty male Sprague Dawley rats. Rats were divided into 5 groups (with 6 rats in each group). The rats were numbered with ear tags. The rats in all of the groups were sedated with intraperitoneal thiopental (Pental Sodyum, Ulagay, Istanbul, Turkey), and approximately 1.5-2 cm of sciatic nerves were removed for microscopic analysis and comparison. The sciatic nerves were then fixed in 4% formaldehyde for 5 days.

Sciatic nerve samples were taken proximal to the lesion, the lesion site, and distal to the lesion for all of the groups, and these parts were embedded in paraffin blocks. Three sections were taken from every block in order to make a better evaluation, and the sections were stained with Hematoxylin and Eosin. Injury scores were obtained by evaluating the sections with a light microscope. The sciatic nerve sections in the axonotmesis + placebo group (group 2) showed mild injury in the proximal site, and severe injury in the lesion and distal sites. The sciatic nerve sections in axonotmesis + 50 mg/kg agmatine group (group 3) showed mild injury in the proximal site and moderate injury in the lesion and distal sites. The sciatic nerve sections in the neurotmesis + placebo group (group 4) showed mild-moderate injury in the proximal and lesion sites, and severe injury in the distal site. The sciatic nerve sections in the neurotmesis +50 mg/kg agmatine group (group 5) showed mild injury in the proximal site and moderate injury in the lesion and distal sites. Damage was more prominent in the distal part of the nerves, and only the distal parts of the nerves were evaluated and compared in this study.

II- Statistical Examination
Axonolysis, axon degeneration, edema, hemorrhage, and inflammation were evaluated in the histopathologic examination of the distal sites of the sciatic nerves in groups 2, 3, 4, and 5 and were compared with the control group. Axonolysis, axon degeneration, edema, hemorrhage, and inflammation were graded as 0, 1, 2, or 3 according to a modified grading system described by Wahl et al. (22) (Table 1). A light microscope with both 20x and 40x magnification was used in evaluating the histopathologic specimens, and final grades of axonolysis, axon degeneration, edema, hemorrhage, and inflammation were obtained for all of the nerves. Statistical comparisons were conducted using the Independent Samples t Test, and the results were evaluated using the Mann-Whitney U test; p <0.05 was accepted as statistically significant.
RESULTS

I- Light Microscopic Evaluation

The sections taken from the control group (group 1) were evaluated under a light microscope and normal axon, vascular, and connective tissue structures were seen (Figure 1). Nerve sections of the distal parts of trauma in the axonotmesis + placebo group (group 2) revealed severe axonolysis, degenerated ghost axons, migrated inflammatory cells, and severe edema in the connective tissue when evaluated under a light microscope (Figure 2). Nerve sections of the distal parts of trauma in the axonotmesis + agmatine treatment group (group 3) displayed some intact axons, a moderate degree of axonolysis, degenerated ghost axons, and moderate edema when evaluated under a light microscope (Figure 3). Nerve sections of the distal parts of trauma in the neurotmesis + placebo group (group 4) showed severe axonolysis, degenerated ghost axons, degenerated myelin sheaths, and intense edema when evaluated under the light microscope (Figure 4). Nerve sections of the distal parts of trauma in the neurotmesis + agmatine group (group

### Table 1: Table Showing Criteria of Histopathologic Scoring Depending on Axonolysis, Axon Degeneration, Edema, Hemorrhage, and Inflammation

<table>
<thead>
<tr>
<th>Grade of damage</th>
<th>Percentage of damage</th>
<th>Score</th>
</tr>
</thead>
<tbody>
<tr>
<td>No damage</td>
<td>0%</td>
<td>0</td>
</tr>
<tr>
<td>Minor damage</td>
<td>25% &gt;</td>
<td>1</td>
</tr>
<tr>
<td>Moderate damage</td>
<td>25%-75%</td>
<td>2</td>
</tr>
<tr>
<td>Severe damage</td>
<td>75% &lt;</td>
<td>3</td>
</tr>
</tbody>
</table>

**Figure 1:** Light microscopic examination (HE staining) of sciatic nerve of control group showing normal axonal structure (A), normal peripheral nerve vascular structure (V), and normal structure of connective tissue (x40).

**Figure 2:** Light microscopic examination (HE staining) of the distal part of the sciatic nerve of the axonotmesis + placebo group showing axonolysis (Aks), a large number of ghost axons that are degenerated, inflammatory cells migrating (I) out of vessels, and edema (E) (x40).

**Figure 3:** Light microscopic examination (HE staining) of the distal part of the sciatic nerve of the axonotmesis + 50 mg /kg agmatine treatment group showing less axonolysis (Aks) and degenerated axons and less edema (E). Some normal axonal structure (A) is also seen (x20).

**Figure 4:** Light microscopic examination (HE staining) of the distal part of the sciatic nerve of the neurotmesis + placebo group showing intense edema (E), axonolysis (Aks), and large number of ghost axons that are degenerated (x40).
Peripheral nerve injury may result in demyelination or axonal degeneration, the self-destructive set of cellular and molecular processes by which degenerating axons and myelin are cleared after injury, is initiated by macrophages and Schwann cells. Molecular inflammatory mediators such as cytokines (IL-1, IL-6, IL-10, and TNF-alpha, among others), transcription factors (NF-kappaB, c-Jun), the complement system and arachidonic acid metabolites have been shown in various studies to modulate these processes (1). Furthermore, the alteration of spinal arachidonic acid turnover after peripheral nerve injury regulates regional glutamate uptake activity and glutamate homeostasis (20).

Treatment of peripheral nerve injuries is considered to be a challenging procedure. Treatment of these injuries varies depending on the type of the injury, place of the injury, condition of the surrounding structures, and associated injuries. Generally, the surrounding structures are minimally injured. Surgical and medical methods are used to treat peripheral nerve injury. The literature presents numerous experimental peripheral nerve injury models, and different materials were used to treat the injuries. We selected our model as to avoid injuring surrounding structures. In the treatment of peripheral nerve injuries, suturing with vinyl and silk was done first; however in light of the number of sutures and the negative effects of the sutures on recovery, fibrin glue and wrapping the injured nerve were also tried. Electrical stimulation, therapeutic ultrasound, low-dose radiation, and low-intensity laser were also tested in experimental models to produce nerve regeneration and functional recovery (8, 13, 15, 16, 17).

After peripheral nerve injury, some surgeons advise immediate repair, while others advise recommend waiting some weeks (Seddon). In the lesion site of a peripheral nerve injury, the decrease in oxygenation and changes in morphology increase arachidonic acid metabolism. Unstable endoperoxides collect in intracellular and extracellular areas. These substances are superoxidized and are converted to superoxide anion radicals. To begin recovery, these free oxygen radicals should be removed and should no longer be produced. The processes that occur in peripheral regeneration can be divided into the following major events: Wallerian degeneration, axon regeneration/growth, and nerve reinnervation. Wallerian degeneration, the self-destructive process by which degenerating axons and myelin are cleared after injury, is initiated by macrophages and Schwann cells. Molecular inflammatory mediators such as cytokines (IL-1, IL-6, IL-10, and TNF-alpha, among others), transcription factors (NF-kappaB, c-Jun), the complement system and arachidonic acid metabolites have been shown in various studies to modulate these processes (1). Furthermore, the alteration of spinal arachidonic acid turnover after peripheral nerve injury regulates regional glutamate uptake activity and glutamate homeostasis (20).
Agmatine [4-(aminobutyl)-guanidine-NH2-CH2-CH2-CH2-CH2-NH-(NH2)2] is a guanidinium compound formed by the decarboxylation of L-arginine by arginine decarboxylase, and found in abundance in bacteria, and plants (21) and in trace amounts in mammalians (9). Agmatine is a neurotransmitter-neuromodulator with both N-methyl-D-aspartate receptor (NMDAR)–antagonizing and nitric oxide synthase (NOS)–inhibiting activities. The compound is present in the brain in very low amount; however, its synthesis is greatly increased during brain development and after brain ischemia (6). Agmatine plays a role in the modulation of neurotransmission functions and interacts with various neurotransmitter receptors (7). Agmatine can also interfere with second messenger pathways by acting as an adenosine diphosphate (ADP)-ribose acceptor and inhibits ADP-ribosylation of proteins (14). As a result of these characteristics, agmatine has certain neuroprotective effects after neurotrauma. Treatment with exogenous agmatine has been found to be non-toxic and to exert significant neuroprotective effects in models of neurotoxic and ischemic brain injuries (5, 11, 12). Treatment with agmatine has also been found to be neuroprotective following excitotoxic spinal cord injury (3, 10, 23).

In summary, the present findings demonstrate that agmatine treatment can exert significant neuroprotective effects in the experimental model of peripheral nerve injury and suggests that this naturally occurring, non-toxic compound should be tried for therapeutic use after peripheral nerve injury. Limitations of this study are the small number of rats, lack of knowledge on the minimum effective dose, minimum duration of usage, and the best way of administration of agmatine. This is the first study demonstrating the activity of agmatine, an inducible NOS (iNOS) inhibitor and selective NMDAR antagonist, on reducing tissue damage in distal part of trauma in an experimental peripheral nerve injury in rats. However, further studies should be conducted prior to the use of agmatine as a drug.

The purpose of this study was to demonstrate the activity of agmatine, an inducible NOS (iNOS) inhibitor and selective NMDAR antagonist, on reducing tissue damage in distal part of trauma in an experimental peripheral nerve injury in rats. In the literature, agmatine was used early in the process of the nerve injury in all studies about the effects of agmatine in neural tissue. Therefore in our study we also used agmatine early in the process of the nerve injury. The results of our study indicate that intraperitoneally administrated agmatine (50 mg/kg) had antioxidant and antineurotoxic effects and positive effects on recovery in experimental peripheral nerve injury in rats. Agmatine reduced axonalysis, axon degeneration, and edema. This effect is thought to be caused by NMDA blockage of agmatine, blockage of the production of NO, which, in turn, blocks peroxidation and decreases apoptosis, enabling cell function to continue. Nevertheless, additional studies examining the effective dose, side effects, and duration of usage of agmatine are recommended.

REFERENCES


