

## Morphometric Analysis of Dose-dependent Effect of Progesterone on Experimental Vasospasm-induced Rat Femoral Arteries

### Abstract

**Objective:** Our aim of this study was to determine effective doses of progesterone which has a vasodilatory effect during the early stage of vasospasm. Cerebral vasospasm (CV) is a predominant cause of morbidity and mortality which develops following subarachnoid hemorrhage (SAH). Etiopathogenesis of CV is multifactorial. Despite many previously performed studies on this issue, the mechanism by which blood and blood products in the subarachnoid space induce CV has not been clarified yet. **Materials and Methods:** In our study, we used “Rat Femoral Artery Vasospasm Model” introduced by Okada *et al.* Thanks to easy procurement and maintenance of rats. Rats were divided into four groups as: Group 1 ( $n = 8$ ; control group), Group 2 ( $n = 8$ ; vasospasm group), Group 3 ( $n = 8$ ; vasospasm + 3 mg/kg progesterone group), and Group 4 ( $n = 8$ ; vasospasm +15 mg/kg progesterone group). Progesterone which is an endogenously synthesized natural steroid was preferred in our study. Progesterone increases the production of vasodilatory epoxyeicosatrienoic acid by acting on its binding sites termed as pregnane X receptor. It decreases the intracellular influx of  $Ca^{2+}$  by blocking the functioning of L-type channels in smooth muscle cells. It manifests another vasodilatory effect by decreasing expression of  $TxA_2$  receptor. In our study, at the end of the 7<sup>th</sup> day, where the most intense vasospasm is seen, 1 cm pieces were excised from the femoral arteries and histopathologically examined under light microscope. **Results:** Vascular walls of three vasospasm-induced groups were relatively thicker when compared with the control group. Drug-treated groups were not different from each other. Vascular walls of the groups treated with lower and higher doses of the drug were thinner when compared with the vasospasm group without any statistically significant difference between groups ( $P > 0.05$ ). Luminal cross-sectional areas of the drug-treated groups did not differ from each other. Mean luminal cross-sectional areas of the control and the drug-treated groups were larger than that of the vasospasm group without any statistically significant intergroup difference ( $P > 0.05$ ). **Conclusion:** Based on the results of our study, progesterone did not exert protective effects on vascular wall thickness, while histopathological examination of luminal cross-sectional areas revealed its vasodilatory effects without any statistically significant difference between groups. Starting from the study results obtained, we think that its potential use as a preventive agent against the development of post-SAH CV requires conduction of multicentered, placebo-controlled, randomized, and double-blind studies.

**Keywords:** Progesterone, subarachnoid hemorrhage, vasodilation, vasospasm

### Introduction

Subarachnoid hemorrhage (SAH) is the detection of blood between pia mater and arachnoid. The presence of SAH in patients exposed to head trauma increases mortality rate two-fold. When all these factors are taken into consideration, head trauma is the most frequently seen cause of SAH. The most frequent cause of spontaneously developed SAH is an aneurysmal rupture. Besides, many factors which develop without any known etiology, and causative factors related to infectious, and toxic agents, vascular pathologies, bleeding

disorders, and brain tumors have been described.<sup>[1]</sup>

Some patients die secondary to severe clinical manifestations before the establishment of diagnosis, and sometimes misdiagnosis makes it impossible to estimate actual incidence of SAH. However, SAH is observed in nearly 10% of all cardiovascular diseases, and it is responsible approximately 25% of all deaths due to cerebrovascular mortality.<sup>[2]</sup>

Twelve percent of the patients who lost their lives after SAH because they did not receive any treatment. Twenty-five percent of the patients lost their lives within the

**Metin Kasap,  
Huseyin Canaz<sup>1</sup>,  
Gokhan Canaz<sup>2</sup>,  
Mehmet Tokmak<sup>3</sup>,  
Alper Bingul<sup>4</sup>,  
Ibrahim Alatas<sup>1</sup>**

*Department of Neurosurgery, Bakirkoy Dr. Sadi Konuk Training and Research Hospital, <sup>1</sup>Department of Neurosurgery, Florence Nightingale Hospital, Istanbul Bilim University, <sup>2</sup>Department of Neurosurgery, Haseki Training and Research Hospital, <sup>3</sup>Department of Neurosurgery, Faculty of Medicine, Medipol University, Istanbul, <sup>4</sup>Department of Neurosurgery, Kilis State Hospital, Kilis, Turkey*

**Address for correspondence:**  
Dr. Gokhan Canaz,  
Department of Neurosurgery,  
Haseki Research and Training  
Hospital, 34087 Fatih, Istanbul,  
Turkey.  
E-mail: gokhancanaz@gmail.  
com

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first 21 h, and another 40–60% of the cases existed within the first 30 days.<sup>[3]</sup>

Cerebrovascular spasm (cerebral vasospasm [CV]) is a narrowing of the large arteries of the basis cranii induced by accumulation of blood, blood degradation products, and other chemical substances in the cisterns leading to decreased perfusion in the distal segment of the artery.<sup>[4]</sup>

CV is the most important pathological condition which adversely affects mortality and morbidity.<sup>[2,5]</sup> Despite greater number of studies performed on this issue, its etiopathogenesis has not been revealed completely. Currently, medical treatments have not yielded satisfactory results. The main objective of the studies performed so far has been to reveal pathogenesis of SAH and develop effective treatment strategies which may limit the extent of vasospasm.

In this study, dose-dependent effects of an endogenous hormone (medroxyprogesterone acetate) without any serious side effects which conceivably might be an effective vasodilatory agent in the treatment of CV on rat femoral arteries were morphologically examined.

## Materials and Methods

In this study, surgical interventions were performed in Istanbul University Laboratory of Experimental Medicine Research Institute, Department of Experimental Animals Biology, and Biomedical Application Techniques. Light microscopic examinations were performed in Division of Pathology, Haseki Training and Research Hospital.

In this study, 32 Sprague-Dawley rats weighing 180–280 g were operated using microsurgical instruments and surgical microscope. As a vasospasm model “Rat Femoral Artery Vasospasm Model” suggested by Okada *et al.* was used.<sup>[6]</sup>

The study was approved by the faculty Animal Experiments Local Ethics Committee of Istanbul University Faculty of Medicine, Reference No. 2011/73. This research received no specific grant from any funding agency in the public, commercial, or not-for-profit sectors. We declare no conflict of interest in preparing this article.

Thirty-two rats were equally divided into four groups as follows:

- Group 1 ( $n = 8$  rats) control group; a silastic sheath was wrapped around femoral artery without application of vasospasm or drug therapy
- Group 2 ( $n = 8$  rats) vasospasm group; vasospasm of the femoral artery was performed without the application of progesterone
- Group 3 ( $n = 8$  rats) low-dose drug group; vasospasm of the femoral artery was induced, and progesterone at daily doses of 3 mg/kg was applied
- Group 4 ( $n = 8$  rats) high-dose drug group; vasospasm of the femoral artery was induced, and progesterone at daily doses of 15 mg/kg was applied.

For surgical intervention, the rats were successively anesthetized with percutaneous intraperitoneal 60 mg/kg ketamine HCl (Ketalar flacon 50 mg/ml, Eczacıbaşı, Turkey), and 10 mg/kg xylazine (Rompun 2% Bayer, Turkey), and laid in the supine position on the wooden blocks.

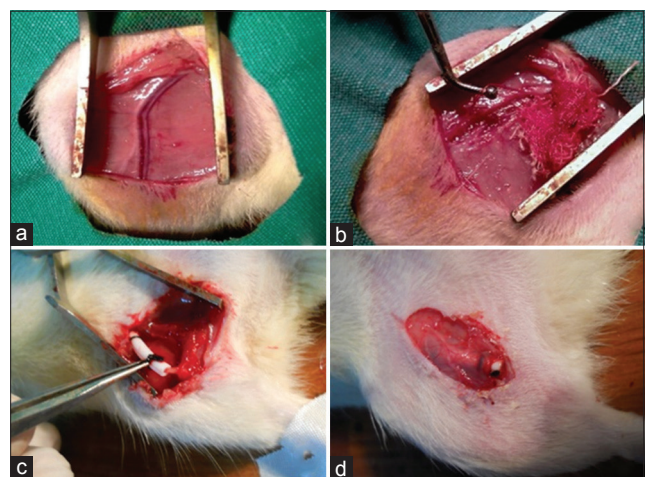
Inguinal regions of the rats were shaved and disinfected with povidone-iodine solution (Batticon solution 10%, Adeka, Turkey). A 3 cm long skin incision was made, and under microscopic examination, femoral neurovascular bundle was exposed [Figure 1a]. Avoiding traumatization of the femoral artery, femoral artery was dissected away from the femoral vein and nerve [Figure 1b]. A silastic sheath was wrapped around 1.5 cm segment of the femoral artery and sutured circumferentially [Figure 1c].

In Groups 2, 3, and 4, 1 cm proximal of the tip of the rat tails was excised and using insulin injector 0.1 cc blood sample was drawn and injected into silastic sheath wrapped around femoral artery to construct an SAH model. The tail was ligated proximal to the excised segment. Following hemostasis layers were closed in proper order.

Rats in all groups recovered normally from anesthesia under room temperature. The rats were fed with standard rat pellets in their cages heated to appropriate ambient temperature for 7 days. The day of surgery was considered as day 1, and once daily doses of 1 ml isotonic saline solution were administered through intramuscular route for 7 days.

The day of surgery was considered as day 1, and once daily doses of 1 ml isotonic saline solution containing 3 mg/kg and 15 mg/kg progesterone were administered through intramuscular route for 7 days in Groups 3 and 4, respectively.

At the end of the 7 days, the rats were reanesthetized with 60 mg/kg percutaneous intraperitoneal ketamine



**Figure 1:** (a) Neurovascular bundle. (b) Dissection and exposure of femoral artery. (c) Silastic sheath was wrapped around the artery so as to separate them from surrounding tissues. (d) Silastic sheath discerned under granulation tissue on postoperative 7 day

HCl (Ketalar flacon 50 mg/ml, Eczacıbaşı, Turkey), and 10 mg/kg xylazine (Rompun 2% Bayer, Turkey), and laid in the supine position on wooden blocks. The former incisions were opened, and the silastic sheath around femoral artery was approached [Figure 1d]. Silastic sheath was opened, and femoral arteries were exposed.

Meanwhile, the skin over the sternum was shaved, and following cleansing and disinfecting the surgical field with a povidone-iodine solution (Batticon solution 10%, Adeka, Turkey), sternum was dissected away from its attachments with ribs using dissection scissors, and thorax was explored [Figure 2]. The pericardium was opened, and the left ventricle was entered through a created puncture hole. Serum set was connected to the tip of the puncture catheter. With the aid of a large volume injector, 300 ml isotonic solution was delivered into the left ventricle under physiologic artery pressure. After circulation of the delivered solution within the vascular system, it was drained from the opened right atrium. Evacuation of all the delivered blood volume was ensured, and then perfusion procedure was initiated. For perfusion, a solution containing a mixture of 100 ml 0.003 M phosphate buffer (pH: 7.4) + 200 ml 4% formaldehyde and 1% gluteraldehyde was perfused as a bolus from a height of 1 m. Circulation of all delivered perfusion fluid through all vascular system and evacuation through excised right atrium were ensured.

At the end of this procedure, crystal clear perfusion fluid was drained wholly throughout the right ventricle. Nearly, 1.5 cm length femoral artery segment was excised, examined under a light microscope, and immersed in buffered 10% formaldehyde solution for analysis.

In the pathology laboratory from each vascular specimen, at least two 3 mm thick vascular segments were cut and immersed in buffered 10% formaldehyde solution. Then, cut sections were stored in cassettes. On the same day in the automated tissue processing device, the samples were subjected through processes of fixation with formaldehyde,



Figure 2: Exploration of thorax and heart

dehydration with grade alcohol, made transparent with xylene, and embedded in paraffin blocks. Tissue samples were left in paraffin for 24 h, and then taken out of tissue processing device, and embedded in paraffin blocks. Five-micron thick cut samples were placed on slides. Following deparaffinization procedure in the incubator at 60°C for 1 h, the slides were rinsed thrice with xylene solution. The specimens were treated with grade alcohol, rehydrated, and rinsed with water. The slides were stained with hematoxylin-eosin and examined under Nikon surgical microscope (Nikon Instruments Europe BV, Amsterdam / Netherlands) at  $\times 100$ ,  $\times 200$ , and  $\times 400$  magnification. Vascular wall thickness and luminal cross-sectional areas were measured using micrometers.

The light microscopic examination was performed in Division of Pathology, Haseki Training and Research Hospital. DP2-BSW program was used during measurements of vascular wall thickness and luminal cross-sectional areas.

## Results

### Histological changes

- Group 1: In the control group, light microscopic examination of the femoral artery revealed vessels with thin and smooth vascular endothelium, thin and uncurled internal elastic lamina, and concentrically arrayed smooth muscle cells [Figure 3a]
- Group 2: In the vasospasm group, light microscopic examination of the femoral artery disclosed markedly stenotic lumen and thickened vascular wall, disruption of the endothelial integrity, curling of internal elastic lamina, and vacuolization in the muscular layer [Figure 3b]
- Group 3: In the vasospasm plus low-dose progesterone group, in the light microscopic examination of the

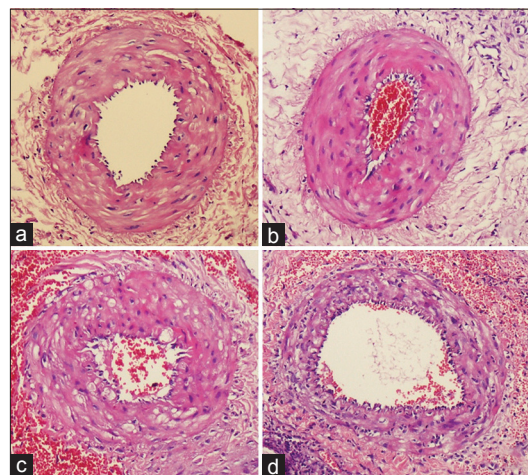


Figure 3: (a) Control group: Microscopic appearance of the vascular lumen under a light microscope ( $\times 200$ ). (b) Vasospasm group, light microscope. (c) Low-dose progesterone group under a light microscope. (d) High-dose progesterone group under light microscope

femoral artery, when compared with the vasospasm group, thinner vascular endothelium, patchy areas of curled internal elastic lamina, and concentrically arrayed smooth muscle cells were detected [Figure 3c]

- Group 4: In the vasospasm plus high-dose progesterone group, in the light microscopic examination of the femoral artery, similar to the vascular structure detected in the control group, thin and smooth endothelium, thin and occasionally curled internal elastic lamina, and concentrically arrayed smooth muscle cells were observed [Figure 3d].

### Morphometric analysis

Femoral arteries of four groups were harvested. Mean ( $\pm$ standard deviation [SD]) arterial wall thickness and luminal cross-sectional areas of femoral arteries were calculated. The values obtained were statistically analyzed.

When groups were compared as for mean arterial wall thickness, the smallest and the highest mean values were seen in the control and vasospasm groups, respectively. Mean values of low- and high-dose progesterone groups were closer to those of the vasospasm group.

When groups were compared as for mean values for luminal cross-sectional areas, the lowest and the highest mean values were detected in the vasospasm and control groups, respectively. Mean values of low- and high-dose progesterone groups were closer to those of the control group.

### Method of statistical analysis used

For the statistical evaluation of study data, Statistical Package for Social Sciences for Windows 18.0 (IBM, New York, USA) program was used.

In the evaluation of study data, in addition to descriptive statistical methods as means and SD, in the intergroup comparison of quantitative data among more than two groups, parameters with normal distribution were compared with “One-way ANOVA test” and in the detection of the group which caused intergroup difference “Tukey HSD test” was used.

In the comparison of quantitative data between two groups, parameters without normal distribution were compared using “Mann–Whitney U-test.” In the comparison of mean

values of more than two groups, “Kruskal–Wallis test” was used. The results were evaluated within 95% confidence interval and at a significance level of  $P < 0.05$ .

### Statistical values

In the study, four groups each consisting of eight cases were analyzed.

Maximum and minimum mean wall thicknesses were detected in the control and the vasospasm groups, respectively [Figure 4]. One-way variance analysis was used to determine if any difference between mean vascular wall thickness of the groups exists. Since one-way variance analysis could not yield homogenous results in assumptions as an alternative Kruskal–Wallis test was applied [Table 1]. As a result of statistical analysis, any statistically significantly difference between mean vascular wall thickness of the groups was not found ( $P = 0.948 > 0.05$ ).

Mean values for luminal cross-sectional areas of the groups were also analyzed [Table 2]. The highest and lowest mean values were noticed in the vasospasm and control groups, respectively [Figure 5]. Accordingly, the mean value for luminal cross-sectional area of the control group was observed higher than those of the other groups, while the relevant mean values in progesterone groups were nearly two-fold higher than those of the vasospasm group without any statistically significant intergroup difference because of increased SDs estimated in one-way variance tests ( $P = 0.618 > 0.05$ ).

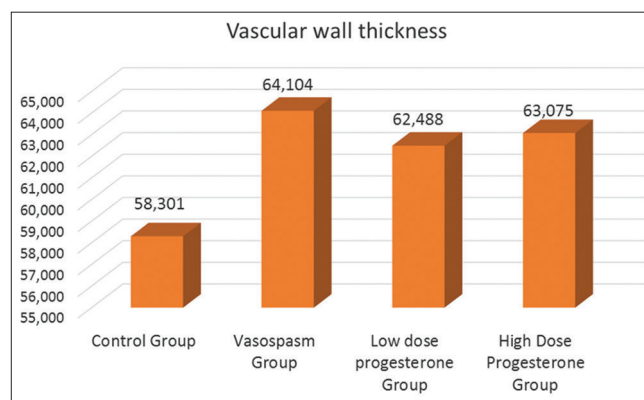


Figure 4: Median vascular wall thickness of the groups

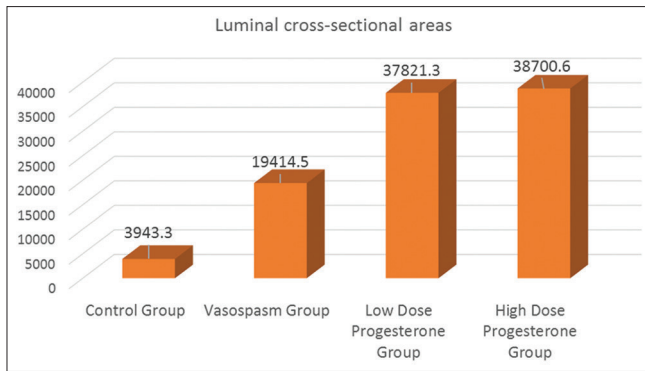
Table 1: Descriptive statistics related to distribution of vascular wall thickness among groups

	Mean $\pm$ SD			
	Control group	Vasospasm group	Low-dose progesterone group	High-dose progesterone group
Vascular wall thickness	58.301 $\pm$ 20.376	64.104 $\pm$ 10.640	62.488 $\pm$ 7.617	63.075 $\pm$ 10.527

SD – Standard deviation

Table 2: Distribution of luminal cross-sectional areas among groups

	Mean $\pm$ SD			
	Control group	Vasospasm group	Low-dose progesterone group	High-dose progesterone group
Luminal cross-sectional areas	39,436.30 $\pm$ 44,329.70	19,414.51 $\pm$ 22,655.36	37,821.30 $\pm$ 36,697.38	38,700.64 $\pm$ 33,128.05



**Figure 5: Mean luminal cross-sectional areas of the groups**

Excluding the control group, we concluded that mean values for luminal cross-sectional areas estimated in vasospasm and progesterone groups did not differ statistically significantly ( $P = 0.398 > 0.05$ ).

## Discussion

CV occurring following subarachnoid bleeding is one of the most important pathological events adversely affecting morbidity and mortality. CV can be defined as narrowing of cerebral vasculature caused by blood, metabolites, and chemical substances of blood which accumulate in the subarachnoid space.<sup>[7]</sup> Still, its pathophysiology has not been elucidated so far.

Its etiopathogenesis is multifactorial including main oxyhemoglobin and spasmogenic agents.

In the experimental studies, blood injected into subarachnoid space has induced CV. Still in clinical studies, computed tomography sections obtained immediately after aneurysmal bleeding revealed that amount of blood is the most important indicator of the development of late-term ischemic deficit.<sup>[8,9]</sup>

In a study performed by Okada *et al.*, the authors injected whole blood and washed erythrocytes and demonstrated that the most severe reaction had developed at the end of the 7<sup>th</sup> day. The outcomes seen in vasospasm and periadventitial blood were compared, and the consequences of femoral vasospasm observed in rats were similar to those found in their experiment as for the degree of severity, specificity, progression in time, and histological characteristics. Since silastic material applied around adventitia had no effect on vasospasm, lower cost, and easy maintenance of rats, “Rat Femoral Artery Vasospasm Model” used by Okada *et al.* was found to be applicable in our study.<sup>[6]</sup>

Cerebral arteries (CAs) were analyzed under light microscope immediately after SAH, and thickening of the vascular wall, internal elastic lamina, narrowing of the luminal diameter of CAs, disruption of endothelial cellular structure and integrity, vacuoli in smooth muscle, migration of proliferated myointimal cells into intima, and development of periadventitial inflammation associated with

perivascular axon loss were demonstrated.<sup>[10]</sup> Moreover, in our study, in compliance with literature findings, alterations in the vascular structure of CAs were observed.

Undoubtedly, mainly oxyhemoglobin and free oxygen radicals are responsible for the development of vasospasm occurring after SAH.<sup>[11-13]</sup> Oxidative stress stimulates hypertrophy and proliferation of smooth muscle cells, and accelerates apoptotic processes. These changes are associated with vascular contractile responses.<sup>[14]</sup> Correlations between the development of CV and increased levels of superoxide anions in the cerebrospinal fluid have been already reported. Oxyhemoglobin increases the amount of phospholipase C (PLC). PLC indirectly promotes the intracellular influx of  $Ca^{2+}$  leading to vasoconstriction. PLC also increases the levels of diacylglycerol resulting in increased amounts of protein kinase C. However, PKC decreases the production of NO and consequently prevents vasodilation.

Still, the most potent vasoconstrictor endothelin-1 activates RhoA/Rho kinase and protein kinase-C which reinforces contraction of cerebrovascular smooth muscles are under the influence of oxyhemoglobin.<sup>[15]</sup>

Previous studies demonstrated that progesterone decreases the expression of TxA2 receptor which is a potent vasoconstrictor, blocks L-type  $Ca^{2+}$  channels leading to decrease in intracellular  $Ca^{2+}$  content, and increases the production of vasodilatory agent epoxyeicosatrienoic acid. In our study, endogenous substance progesterone was used which was thought to have lesser side effects.

A histological and statistically significant difference was not detected between vasospasm and low-dose progesterone groups ( $P = 0.948 > 0.05$ ). Vascular walls of all these three groups were found to be thicker when compared with the control group.

Mean luminal cross-sectional areas of the vessels in the control and progesterone groups were found to be markedly higher relative to the vasospasm group. However, in statistical analysis because of higher SDs, the intergroup difference was deemed to be devoid of any significance ( $P = 0.618 > 0.05$ ).

These outcomes of our study, though statistically insignificant, have suggested a vasodilatory effect of progesterone on the cross-sectional area of the vascular lumen. Thanks to potent vasodilatory effects of progesterone, especially over  $Ca^{2+}$  metabolism, we think that its clinical use in the treatment of CV deserves merit, and consequently experimental and clinical studies should be maintained.

## Conclusion

Two different doses of progesterone which is an endogenous hormone with potent vasodilatory efficacy synthesized in the body without any known serious side

effect were analyzed in the experimental SAH model which was induced by injecting autologous blood on the periphery of rat femoral artery.

In our study, we used “Rat Femoral Artery Vasospasm Model” suggested by Okada *et al.* to obtain parameters of vascular wall thickness and cross-sectional areas of vascular lumens which were then subjected to statistical analysis.

Analysis of vascular wall thickness and luminal cross-sectional areas could not demonstrate any significant difference between Group 4 (15 mg/kg/day progesterone group) and Group 3 (3 mg/kg/day progesterone group).

Mean values for luminal cross-sectional areas of all three groups were higher than those of the vasospasm group without any statistically significant intergroup difference.

We think that progesterone's potential use as a preventive agent against the development of post-SAH CV requires conduction of multicentered, placebo-controlled, randomized, and double-blind studies.

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Nil.

#### Conflicts of interest

There are no conflicts of interest.

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