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The evaluation of arterial blood pressure in anesthetized dogs with xylazine and ketamine

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ABSTRACT

Objective: It was aimed to investigate the effects of ketamine combination administered with xylazine used for general anesthesia in dogs on arterial blood pressure, heart rate, and body temperature.

Materials and Methods: In the study, a total of 20 dogs, 14 females and 6 males, from various breeds and ages 1 to 5, which undergone elective ovariohysterectomy or castration according to body weights. 1 mg/kg xylazine (xylazine hydrochloride, 23.3mg/ml, Xylazinbio 2% Bioveta®, Czechia) and 10 mg/kg ketamine (ketamine hydrochloride, 100 mg/ml, Ketazol 10% Richter® Pharma Ag, Austria) combination within a single injection with 21G needle were applied intramuscularly. The food and water access were ceased 12 hours before drug administration. Systolic and diastolic blood pressures, pulse measurements, and body temperatures were measured 3 times before and 5 times during anesthesia (at the 15th, 30th, 45th, 60th and 120th minutes of anesthesia) by using the AM6100 veterinary bedside monitor. Muff was placed to cover 1/3 of the proximal leg and for the artery to be recognizable by the microprocessor. Pulse rates were measured with electrodes connected to the device. Body temperature was measured by a rectal thermometer which was a part of the device. Measurements were taken before anesthesia was considered as control measurements.

Results: In systolic blood pressure, the recordings at 0, 15th, 30th mins have been found as statistically significant in relation to 45th, 60th, and 120th ($p<0.05$). In diastolic blood pressure, there were no significant differences recorded. The change between the preintervention and post-application has been found significant ($p<0.001$). Body temperature has shown a meaningful change in comparison to the starting point after the readings ($p<0.001$).

Conclusion: Eventually, decreases in blood pressure, heartbeat, and body temperature were observed for the dogs that have been anesthetized with the combination of xylazine-ketamine.

Keywords: Blood Pressure, Body Temperature, Diastolic, Dog, Heart Rate, Systolic

INTRODUCTION

Nowadays, anesthetics are used in immobilization of wild animals, surgical interventions, and during operations for pain relief, as well as during labor, diagnosis, and treatments of diseases with pain (Atasoy and Karadeniz, 2003). After the administration, differences are seen in body temperature, rate of respiration, heart rate, arterial

blood pressure, arterial pH, blood gasses, and hematologic values within the organism. (Allen et al., 1986; Hall et al., 2001; Koç and Sarıtaş, 2004).

Xylazine is the first α_2 agonist used in veterinary medicine for premedication (Greene and Thurman, 1988; Bilgili and Doğan, 1991). After parental application, while decrease in respiratory rate, bradycardia, hypothermia form is observed, long

term hypotension forming after intravenous application. (Samy and Othman, 1985; Börkü et al., 2005; Murrel and Hellebrekers, 2005; Lemke, 2007; Cardoso et al., 2014; Kellihan et al., 2015). In some animals, it causes a decrease in intraocular pressure by reducing aqueous humor production (Sinclair, 2003).

Ketamine is a dissociative anesthetic which is a phencyclidine derivative that constitutes dissociation from the surrounding environment like catalepsy (Booth, 1982; Allen et al., 1986; Hellyer, 1996). Depending on the dosage, analgesic, sedative, and anaesthetic effects can be observed (Aantaa and Scheinin, 1993). In recent years, antidepressant-like features were also discovered (Mihara et al., 2012; Franceschelli et al., 2015). When used alongside anaesthetics with depressive effects on the nervous system, hypoxia, hypercarbia and changes in body temperature can occur (Short et al., 1993). It causes increase in heart rate and blood pressure. Temporary respiration depression forms following anaesthesia, outflow hallucinations may take place. therefore, it is not suggested that it is applied on its own (Colby and Sanford, 1981; Booth, 1982; Haskins et al., 1985; Izci et al., 1993; Gülanber et al., 2001; Topal, 2005; Flecknel, 2016).

The side effects caused by the combination of xylazine-ketamine, which is often used in small animal medicine, are being underestimated. Complications are occurring especially upon usage in animals with an unhealthy cardiovascular system and at absence of essential precautions. In this study, the effect of xylazine-ketamine combination on blood pressure was investigated using the oscillometric technique.

MATERIALS and METHODS

A variety of dog breeds, 14 females and 6 males, aged between 1-5, weighed 21-45 kg, brought to the clinic for spaying and the method ovariohysterectomy, a castrating method, were used as study subjects. 1 mg/kg xylazine (xylazine hydrochloride, 23.3mg/ml, Xylazinbio 2% Bioveta®, Czechia) and 10 mg/kg ketamine (ketamine hydrochloride, 100 mg/ml, Ketamol 10% Richter®, Pharma Ag, Austria) combination within a single injection with 21G needle was applied intramuscularly. The food and water access were ceased for 12 hours prior to the practice. Before practice and during the 15th, 30th, 45th, 60th and 120th minutes of the practice their systolic and diastolic blood pressures, pulse rates and body

temperatures were recorded, to produce three repeat measures in total. Data recorded before practice was used as control measurements. Measurements were taken by a special veterinary use bed-side monitor (AM6100 Veterinary Monitor, Shanghai-China). Blood pressures were measured automatically by an oscillometer. According to the literature, muff was placed to cover 1/3 of the proximal leg and for the artery to be recognizable by the microprocessor. Pulse rates were measured with electrodes connected to the device. Body temperature was measured by a rectal thermometer which was a part of the device.

Statistical analysis

Before significance tests, Numeric data obtained was first evaluated by Shapiro-Wilk test to assess the normality of parametric test assumptions, and Levene test to assess homogeneity of the variances. Variance analysis was used for Dependent quantitative data which were repeated more than two times to test variations. In the case of determined differences, Post-hoc Analysis with Bonferroni correction was used to specify which repetition the differences resulted from. All statistical calculations were analyzed with a minimum of 5% error margin. Level of significance was determined as $p < 0.05$.

RESULTS

Changes in systolic and diastolic blood pressure, heart rate per minute and body temperature caused by xylazine-ketamine combination before and after application presented in Table 1.

Systolic blood pressure was significant at 0, 15 and 30 minutes after application in comparison with the measurements at 45, 60 and 120 minutes (Figure 1).

No significant change in diastolic blood pressure (Figure 2). Changes in heart rate were statistically significant ($p < 0.001$). Heart rate measurements showed a significant decrease after the application (at 45, 60 and 120 minutes) in comparison to before application ($p < 0.05$) (Figure 3).

Body temperature showed a significant fall in comparison to values before application ($p < 0.001$). Value obtained before application and at 30th minute are statistical meaningful in comparison to values obtained at 45th, 60th and 120th minutes ($p < 0.05$) (Figure 4).

Table 1. Data before and after the xylazine-ketamine combination application.

Parameter	Measurements (minutes)							P
	0	15	30	45	60	120		
Systolic Blood Pressure (mmHg)	167.417 ± 4.875 ^a	169.117 ± 3.857 ^a	176.233 ± 6.225 ^a	158.567 ± 4.523 ^b	160.167 ± 3.474 ^b	157.517 ± 3.711 ^b	0.010	
Diastolic Blood Pressure (mmHg)	106.283 ± 6.413	118.017 ± 5.189	126.033 ± 7.839	108.667 ± 5.835	112.267 ± 3.971	105.800 ± 5.995	0.051	
Heart rate (bpm)	90.950 ± 5.246 ^a	79.783 ± 5.346 ^{ab}	90.633 ± 4.557 ^a	73.433 ± 4.199 ^b	69.100 ± 4.641 ^b	72.767 ± 3.776 ^b	<0.001	
Body temp. (°C)	38.317 ± 0.121 ^a	37.710 ± 0.230 ^{ab}	37.925 ± 0.142 ^a	37.072 ± 0.198 ^{bc}	37.007 ± 0.203 ^{bc}	36.717 ± 0.187 ^c	<0.001	

^{a, b, c}: The difference between the groups is statistically significant shown with different letters in the same row (p<0.05).

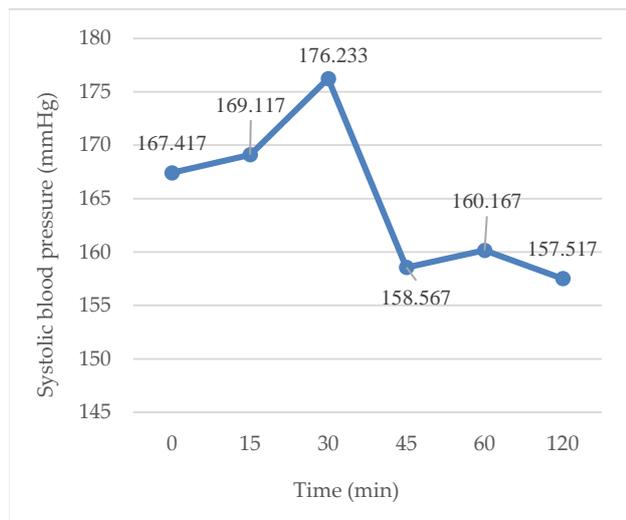


Figure 1. Changes in systolic blood pressure

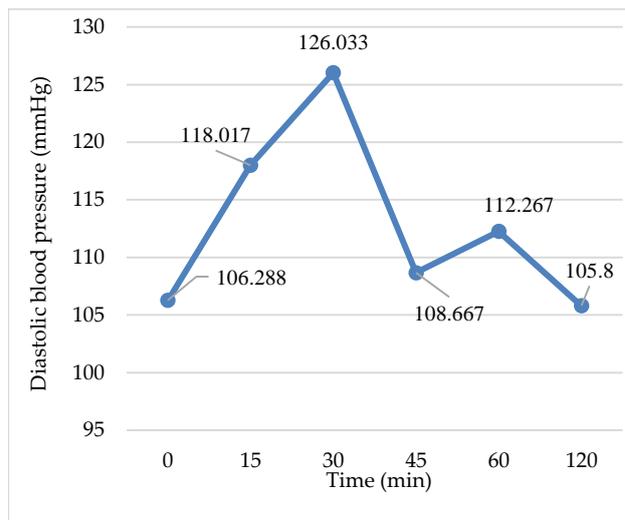


Figure 2. Changes in diastolic blood pressure

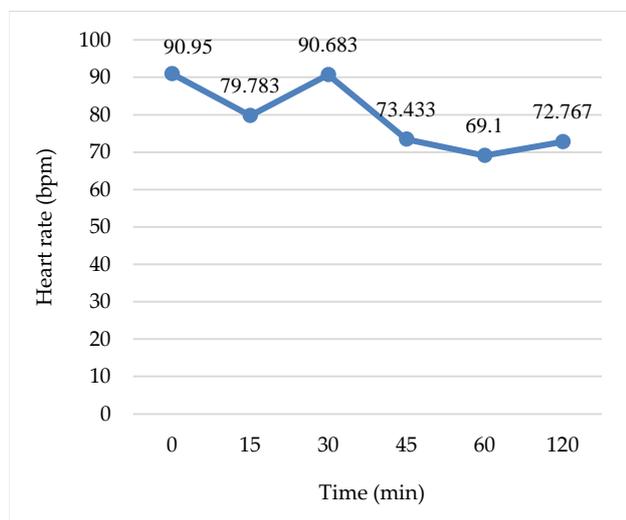


Figure 3. Changes in heart rate

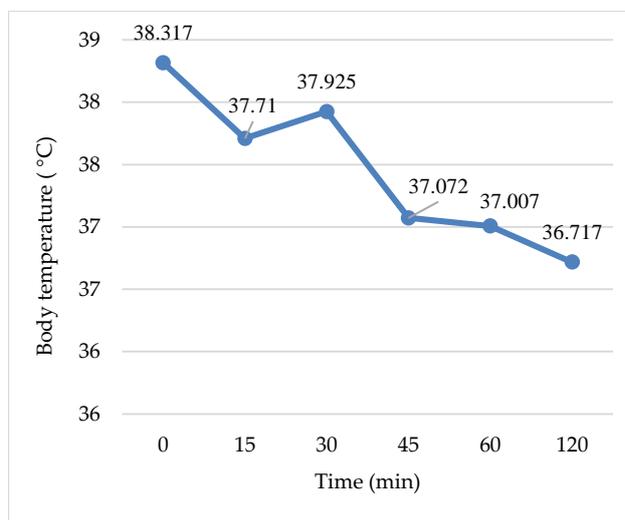


Figure 4. Changes in body temperature due to time

DISCUSSION

Physiological changes occur due to the effects of anesthetic medications in animals' system. In this study aimed at observing the effects of the

combination of xylazine- ketamine and determining any potential results of blood pressure on dogs.

Xylazine, causes temporary hypertension in arterial blood pressure (Samy and Othman, 1985; Borku et al., 2005). Ketamine causes rises in blood pressure and heart rate because of stimulating

cardiovascular system (Colby and Stanford, 1981; Haskins et al., 1985; Izci et al., 1993). Therefore, in our study it is thought that rise in the blood pressure during 15th and 30th minutes cause xylazine to cause temporary hypertension and ketamine stimulating cardiovascular system. In a study conducted by Koç et al., (2002), blood pressure was found to decrease time-dependently, and this decrease indicated that the depressive effect of xylazine in the cardiovascular system was greater than the stimulatory effect of ketamine. Data acquired from our study, drop in arterial blood pressure after 30th minute and reaching out to the minimum value at 120th minute, showed us that our study is compatible with Koç et al., (2002). During our study, significant decrease in heart rate at 60th minute supported literature information about xylazine's depressive effect on cardiovascular system causes a decrease in heart rate (Haskins et al., 1985; Samy and Othman, 1985; Allen et al., 1986; Izci et al., 1993; Koç et al., 2002). Between 15- and 30-minutes rise in heart rate was recorded which showed that the idea of ketamine, rising heart rate temporarily is compatible with literature which specifies the same thing (Colby and Stanford, 1981; Haskins et al., 1985; Samy and Othman 1985; Izci et al., 1993).

As a result of vasodilatation that is caused by xylazine's depressive effect on peripheral sympathetic system, fall in body temperature is indicated (Samy and Othman, 1985; Koç et al., 2002). It is also indicated that, drops in body temperature is because of thermoregulation center is being depressed because of ketamine being used with depressive effective anesthetics (Short et al., 1993). In this case, body temperature reaching its minimum value at 120th minute and showing a significant fall when compared to measurements taken between 0 and 30 minutes are all arise from xylazine depressing peripheral sympathetic system and ketamine depressing thermoregulation center. Facts that obtained from this study are compatible with literature information's (Samy and Othman, 1985; Short et al., 1993; Koç et al., 2002).

CONCLUSION

Consequently, dogs which were put under anesthesia with the combination of xylazine-ketamine have shown decreases in arterial blood pressure, heart rate and body temperature.

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Author's Contributions: The study was designed by BO and SG. Measurements were recorded by BO. BO and SG participated in the interpretation of the results. The draft and revision of the manuscript were done by BO and SG.

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