Endogenous erythropoietin level and effects of exogenous erythropoietin in a rat model of blunt chest trauma-induced pulmonary contusion

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ABSTRACT

BACKGROUND: The present objective was to investigate endogenous erythropoietin (EPO) level and relationship to oxidative stress within the first 24 hours of blunt chest trauma-induced pulmonary contusion (PCn) in a rat model.

METHODS: Thirty-five rats were divided into 3 groups. In the baseline control group (BC, n=7), rats were uninjured and untreated. In the positive control group (PC, n=21) rats were injured but untreated. In the EPO-24 group (n=7), rats were injured and a single dose of intra-peritoneal EPO (5000 IU/kg) was administered immediately after lung injury. The PC group was divided into 3 subgroups: PC-6 (n=7), PC-12 (n=7), and PC-24 (n=7). The BC group was subjected to thoracotomy, and the right lung was harvested. The PC subgroups were euthanized at 6, 12, and 24 hours after injury, respectively. The EPO-24 group was euthanized at the 24th hour after injury. Lung samples were obtained, levels of malondialdehyde (MDA) and EPO were analyzed, and activities of superoxide dismutase (SOD) and catalase (CAT) were then measured in homogenized lung tissue samples. Histologic damage to lung tissue in the BC group, the EPO-24 group, and PC subgroup euthanized at the 24th hour after injury were scored by a single pathologist blinded to group assignment.

RESULTS: Mean MDA levels, as well as SOD and CAT activities, of the BC and EPO-24 groups were significantly lower than those of the PC group (p<0.005). Mean EPO concentration of the PC group was significantly higher than that of the BC group (p<0.005). Lung tissue damage scores measured at 24 hours after injury were significantly lower in the EPO-24 group than in the PC group (p<0.005).

CONCLUSION: In the present PCn rat model, EPO concentrations, as well as SOD and CAT levels, were high in lung tissue, when measured at 24 hours after PCn. When administered early after chest trauma, EPO significantly attenuated oxidative damage and tissue damage in the early phase, as assessed by biochemical markers and histologic scoring.

Keywords: Contusion; erythropoietin; lung injury; oxidative stress.
antioxidants have been widely tested as treat-ment for blunt chest trauma-related PCn, no standard pharmacological ther-
avy yet exists.[3–4]

Erythropoietin (EPO) is a hypoxia-inducible hematopoietic growth factor with antiapoptotic, antioxidant, anti-inflam-
atory, angiogenic, and neuroprotective effects. Hypoxia is the main stimulus for EPO production; in fact, EPO synthesis may increase 50–100-fold in severely hypoxic conditions.[5,4] EPO is induced via hypoxia-inducible factors such as HIF-1. Hypox-
ic preconditioning and non-hypoxic inflammatory mediators, including ROS, have been found to increase levels of HIF-1.[7,4] EPO has also been found to attenuate ischemia/reperfusion (I/R)-induced lung injury[9] as well as tracheobronchial and pulmonary type II epithelial in-jury following traumatic brain injury.[3]

In light of the results mentioned above, the present aim was to determine effects of EPO on lung tissue following blunt chest injury-induced PCn in a rat model.

**MATERIALS AND METHODS**

**Animals**

Following approval from the Ethics Committee on Animal Re-
search of the present medical faculty, 35 male Sprague-Daw-
ley rats, weighing 300–330 g each were selected and acclima-
tized for 10 days in the animal laboratory of the research center, receiving a standard diet and water ad libitum.

**Blunt Chest Trauma**

PCn was induced by method of Raghavendran et al.[10] A hol-
low, 300-g cylindrical weight encased in a vertical stainless steel tube was dropped from 40 cm, positioned on a Lexon plat-form resting on the rat's chest. The platform was sus-
pended on Teflon guides, in order to mini-mize friction and facilitate energy transfer to the animal. The platform was at-
tached to a plast-ic protective shield in direct contact with the lateral aspect of the rat. This precordial shield was de-
gined to protect the heart from contusion, directing impact energy to the lateral sec-tions of the chest. Impact energy, E (in joules) was calculated using the equation E=mgh, in which m signi-
ifies mass of the cylindrical weight (0.3 kg), g signifies gravitational accelera-tion (9.8 m/s^2), and h signifies height of the weight above the Lexon platform (0.4 m). Thus, total en-
ergy transferred to the chest wall of the rat was 1.17 J.

**Experimental Design**

Rats were divided into 3 groups. The baseline control group (BC, n=7) was uninjured and un-treated. The positive control group (PC, n=21) was injured but untreated, and the EPO-24 group (n=7) was injured and administered EPO. The PC group was divided into 3 subgroups: PC-6 (n=7), PC-12 (n=7), and PC-24 (n=7). Rats were anesthetized with 60 mg/kg of intra-
peritoneal ketamine hydrochloride (Ketalar; Eczacıbaşı AŞ, Istanbul, Turkey). Blunt chest trauma was administered. Im-
mediately after the trauma, EPO-24 rats were given a single dose of Eprex intraperitoneal EPO (5000 U/kg; Janssen-Cilag AG, Sihlbruggstrasse, Switzerland). Analgesia was provided by morphine sulphate (0.05 mg/kg) intraperitoneally admin-
istered. Following the procedure, all rats were transferred to their cages. The PC-6, PC-12, or PC-24 groups were eutha-
nized by decapitation 6, 12, or 24 hours after chest trauma, respectively. The BC and EPO-24 groups were euthanized by decapitation 24 hours after chest trauma. The right lung was harvested from rats in the BC, PC, and EPO-24 groups. The upper lobes of rats in the BC and EPO-24 groups, as well as those in the PC-24 subgroup were fixed in 10% formaldehyde for histopathological examination. The middle and lower lobes were stored at -8°C until biochemical and EPO assays were performed.

**Biochemical Analysis**

Levels of malondialdehyde (MDA), and activities of superox-
ide dismutase (SOD) and cata-

cate (CAT) were measured in the following fashion. Tissue samples were homogenized with 3 volumes of ice-cold 1.15% potassium chloride. Activities of antioxidant enzymes (CAT, SOD), and MDA levels were measured in the supernatant obtained after centrifugation at 14000 rpm. SOD activity was measured in the tissue samples according to the method de-scribed by Fridovich.[11] This method employs xanthine and xanthine oxidase to generate superoxide radicals, which react with p-iodonitrotetrazolium violet (INT) to form a red form-assan dye measured at 505 nm. The assay medium consisted of 0.01 M phosphate buf-
fer, CAPS (3-cyclohexylamino-1-propane sulfonic acid) buffer solution (50 mM CAPS, 0.94 mM EDTA, saturated NaOH) at a pH of 10.2, solution of substrate (0.05 mM xanthine, 0.025 mM INT), and 80 UL xanthine oxidase. SOD activity was ex-
pressed as U/mg protein.

CAT activity was determined by measuring the decrease in hydrogen peroxide concentration at 230 nm by the method of Beutler.[12] The assay medium consisted of 1-M Tris hydro-
chlo-ride, 5-mM buffer solution (pH 8.0), 10 mM H_2O_2, and the tissue sample, to make a final vol-ume of 1.0 mL. CAT ac-
activity was expressed as U/mg protein. Tissue sample protein concen-tration was measured with a spectrophotometer by the method of Lowry.[13] MDA levels in tissue samples were measured using the 2-thiobarbituric acid (TBA) test.[14] The reaction mixture contained a 0.1-mL sample, 0.2 mL of 8.1% sodium dodecyl sulphate, 1.5 mL of 20% acetic acid, and 1.5 mL of an 0.8% aqueous solution of TBA. The pH of the mix-
ture was adjusted to 3.5, and the volume was increased to 4.0 mL, by means of distilled water, and 5.0 mL of n-butanol and pyridine mixture (15:1; v/v) was then added. The mixture was shaken vigorously. After centrifugation at 4000 rpm for 10 min, absorbance of the organic layer was 532 nm.

**Measurement of Tissue EPO Levels**

Tissue EPO concentrations were measured with a commer-
cial kit using the ELISA method (Rat EPO ELISA kit, Cusabio Biotech Co. Ltd., Wuhan Hubei, China). Tissue homogenates were prepared as follows: 100 mg of tissue was rinsed with 1x phosphate buffered solution (PBS), and was homogenized in 1 mL of 1xPBS, then stored overnight at –20°C. After 2 freeze-thaw cycles were performed to break cell membranes, homogenates were centrifuged for 5 minutes at 5000 rpm. The supernate was immediately removed then tested.

Histopathological Evaluation

Tissue samples were fixed in 10% neutral buffered formalin solution and embedded in paraffin. At least 8 tissue sections of 5-μm thickness were obtained, then stained with hematoxylin–eosin and scored by a single pathologist blinded to group distribution. Each specimen was scored for congestion (vascular dilation), hemorrhage, interstitial edema, and alveolar col-lapse on a scale from 0–3, in which 0 signified absence of pathology (<5% of maximum pathology), 1 signified mild (5-10% of maximum pathology), 2 signified moderate (11–20% of maximum pathology), and 3 signified severe pathology (>21% of maximum pathology). The specimen was scored for inflammation as follows: 0 signified no extravascular leukocytes, 1 signified <10 leukocytes, 2 signified 10–45 leukocytes, and 3 signified >45 leukocytes. Total tissue damage was calculated using the scoring system described above.[15]

Statistical Analysis

Individual group biochemical parameters were checked for normalcy of distribution prior to statistical analysis using the Shapiro-Wilk test. The Kruskal-Wallis test was used for non-normally distributed variables, and Mann–Whitney U test was performed on biochemical data to examine differences among groups. Tissue damage scores were compared using the Mann–Whitney U test. Data were expressed as mean±SD. Results with p<0.005 were considered statistically significant.

RESULTS

Biochemical Findings

Test protocol was successfully implemented in all animals. Lung tissue MDA levels, SOD and CAT activities, EPO concentrations, and total tissue damage scores are listed in Table 1. All biochemical parameters were significantly different between the PC and EPO groups (p<0.005). MDA levels, and CAT and SOD activities were significantly lower in the BC and EPO groups, compared to the PC group (p<0.005). MDA levels and EPO concentrations were significantly higher in the PC-6, PC-12, and PC-24 subgroups, compared to the BC group (p<0.005; Fig. 1). MDA levels in the EPO group were similar to those of the BC group, suggesting that MDA levels returned to normal range (Table 1, Fig. 1).

Histopathological Findings

Congestion, edema, and alveolar collapse were more prominent in the PC-24 subgroup specimens, compared to the BC and EPO-24 group specimens. Total tissue damage scores correlated with MDA levels. Total lung tissue damage score of the EPO-24 group was lower than that of the PC-24 group (p<0.005; Fig. 1). MDA levels in the EPO group were similar to those of the BC group, suggesting that MDA levels returned to normal range (Table 1, Fig. 1).

DISCUSSION

Results of present study demonstrate that blunt chest trauma caused oxidative stress evidenced by biochemical changes. MDA levels were approximately 8 times higher in untreated traumatized rats, compared to non-traumatized and EPO-treated rats. EPO administration also resulted in lower levels of MDA, a marker of hypoxic tissue injury, compared to traumatized rats not administered EPO. Similar to the present results, Türüt et al.[15] and Basaran et al.[16] found significantly

Table 1. Biochemical parameters and tissue damage scores in groups (mean±SD)

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Groups</th>
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<tbody>
<tr>
<td></td>
<td>BC (baseline control, n=7)</td>
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<tr>
<td></td>
<td>PC (positive control, n=21)</td>
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<td></td>
<td>EPO-24 (erythropoietin treated, n=7)</td>
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<td></td>
<td>PC-6 (n=7)</td>
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<td></td>
<td>PC-12 (n=7)</td>
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<td></td>
<td>PC-24 (n=7)</td>
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<tr>
<td>MDA (nmol/mg protein)</td>
<td>4.5±1.8</td>
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<tr>
<td></td>
<td>32.3±5.8</td>
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<td></td>
<td>33.4±5.3</td>
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<td>38.4±13.4</td>
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<td></td>
<td>5.8±1.6</td>
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<td>CAT (U/mg protein)</td>
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<td></td>
<td>75.4±22.7</td>
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<tr>
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<td>71.5±18.3</td>
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<td></td>
<td>62.5±19.5</td>
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<td></td>
<td>35.2±15.1</td>
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<td>SOD (U/mg protein)</td>
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<td></td>
<td>13.0±1.5</td>
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<td>18.4±1.6</td>
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<td>21.5±5.6</td>
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<tr>
<td></td>
<td>12.4±1.6</td>
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<tr>
<td>Tissue erythropoietin (ng/mL)</td>
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<td>241.2±46.1</td>
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<td>239.1±58.2</td>
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<td>302.0±51.5</td>
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<tr>
<td>Tissue damage score</td>
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<td></td>
<td>10.5±2.1</td>
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<td>5.1±1.3</td>
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</table>

MDA: Malondialdehyde; CAT: Catalase; SOD: Superoxide dismutase.
Groups were baseline control = uninjured and untreated, positive control = injured but untreated, erythropoietin-treated = injured and given a single dose of IP erythropoietin just after lung injury. All rats of EPO-24 group were killed at hour 24 after chest trauma. Rats of PC-6, PC-12, and PC-24 groups were killed at hour 6, 12, and 24 after chest trauma, respectively.

a: p<0.005 vs baseline control and erythropoietin treated groups. b: p<0.005 vs positive control and erythropoietin treated groups. c: p<0.005 vs erythropoietin treated group.
higher MDA levels in traumatic lung contusion samples, as well as in lung samples following acute lung injury induced by sodium taurodeoxycholate (causing acute necrotizing pancreatitis). Increased EPO concentration in lung tissues was presently observed in the first 24 hours after PCn. Because EPO has anti-inflammatory and antioxidant effects,[17–20] levels were analyzed during the early phase of blunt chest trauma. It was determined that EPO administration attenuated oxidative stress and inflammation, as evidenced by biochemical and histological parameters. Presently administered was epoetin alfa, a 165 amino acid glycoprotein manufactured by recombinant DNA technology, which has the same biological effect as endogenous human EPO. It has a molecular weight of 30400 Da and is produced by mammalian cells into which the human EPO gene has been introduced.[5,6]

The primary present aim was to determine the EPO level in lung tissue and its relationship with oxidative stress in the first 24 hours of blunt chest trauma— the most critical period, in terms of morbidity and mortality. For these reasons, EPO concentrations were measured at 6, 12, and 24 hours of blunt chest trauma. To our knowledge, EPO levels have never before been reported in cases of blunt chest trauma, either in animal or human experimental models. It was presently determined that EPO concentrations were elevated in lung tissue of rats subjected to blunt chest trauma-induced PCn. Furthermore, it was found that EPO concentrations, as well as antioxidant enzyme (including SOD and CAT) levels, were increased, compared to controls.

EPO exerts its antioxidative effects either directly as a potent free-radical scavenger by the scavenging actions of sugar moieties, or indirectly by increasing the activities of antioxidant enzymes such as SOD, glutathione peroxidase, and CAT.[9,21] Reduced arterial oxygen associated with anemia or hypoxia is the predominant stimulus for EPO production, primarily in the kidneys. On the other hand, EPO is also synthesized in organs such as the liver, spleen, lung, bone marrow, and brain.[5,6] In the present preliminary study, only EPO concentrations in lung tissue were measured, as serum levels could have been affected by other sources of EPO. The present primary limitations were lack of vital signs and oxygenation status measurements. The present study was preliminary, including an evaluation of EPO levels in damaged lung tissue, as well as the effect of endogenous EPO in the first 24 hours of lung injury; the present results comprise only preliminary research. Further studies, with similar blunt chest trauma models, are being planned, and with similar parameters, including dosage and timing of administration. It is hoped that all limitations will be addressed in future studies.

Increase in antioxidant enzyme activities, such as SOD and CAT, may indicate failure to compensate for oxidative stress.[18,22] In addition, ROS contributes to intensified synthesis of antioxidant enzymes in tissues, the elevated activity of which may be an adaptive response to oxidative stress.[23]
Expression of CAT has been reported in alveolar type II cells and macrophages. Increase in CAT activity during hypoxia indicates presence of oxidative stress, and may be an adaptive response aimed to protect mitochondria from elevated levels of $H_2O_2$. CAT is an important intracellular antioxidant enzyme, detoxifying $H_2O_2$ to oxygen and water. A significant increase in CAT production in the PC group was presumably the result of $H_2O_2$ generation in response to hypoxia. SOD plays an important role in catalyzing the conversion of superoxide to $H_2O_2$ and $O_2$, which are then metabolized to water by CAT or other peroxidases. This enzyme has 3 isoforms, and all SODs are highly expressed in the lung tissue, vessels, and airways. However, extracellular SOD activity is reportedly much higher in the lung. Organisms can defend themselves against oxidative stress by increasing SOD activity as a protective mechanism. In the present study, CAT and SOD activity levels in EPO-treated animals were close to those of the BC group, which underwent no trauma.

In the present EPO-treated group, the restoration of antioxidant enzyme activity confirms the critical role of EPO in regulating oxidative stress due to blunt chest trauma, and points to its property as a direct, potent free-radical scavenger. Production and secretion of EPO are regulated according to the supply of oxygen to the tissues. Hypoxia is the main stimulus for EPO production; in fact, EPO synthesis may increase 50-fold in response to 100 to 70 ToR induces a transitory release of EPO. EPO most likely acts through a variety of pathways to protect lung tissue against damage from inflammation.

REFERENCES

5. Jelkman W. Erythropoietin after a century of research: younger than

In conclusion, endogenous lung parenchyma levels of EPO are elevated following PCn. Exogenous EPO administered after PCn reduced oxidative damage, evidenced histopathologically and biochemically. Antioxidant and cytoprotective treatment with compounds such as EPO may contribute to the health of lung tissues following PCn. Further studies of dosage and timing should be performed to clarify underlying mechanisms and maximize the benefit of the protective effect of EPO on lung parenchyma following traumatic injury. Conflict of interest: None declared.

PCn is associated with leukocyte-mediated secondary inflammatory response leading to capillary leak, alveolar edema, and protein extravasation. In addition, histopathological changes are related to severity of the injury. Leukocyte infiltration in the alveolar space and atelectasis have been observed at 24 hours post-contusion. In the present study, congestion, hemorrhage, inflammation, interstitial edema, and alveolar collapse were scored. Total tissue damage score demonstrated significant decrease with the use of EPO. This and other results emphasize the protective effect of EPO on the lungs by reducing inflammation, likely related to anti-inflammatory potential.

In the present study, CAT and SOD activity levels in EPO-treated animals were close to those of the BC group, which underwent no trauma.

Lung tissue is a major site of ROS production. Cells involved also include alveolar macro-phages, neutrophils, mast cells, type II pneumocytes, endothelial cells, smooth muscle cells, and lung fibroblasts. In inflammatory lung conditions, $H_2O_2$ is primarily produced in alveolar macrophages and neutrophils. Alveolar macrophages are directly acti-vated by low oxygen levels. For example, lowering partial pressure of oxygen from 100 to 70 ToR induces a translatory release of $H_2O_2$ into the supernatant of alveolar macrophage cul-tures. Protective properties of EPO have been demonstrated in models of ischemic and inflammatory injury in neuronal, vascular, cardiac, and intestinal tissues. However, little is known regarding expression of EPO in pulmonary tissues, or the potential role of EPO in pathological processes of the lung. Recently, EPO and its receptor were found to be expressed in respiratory epithelium. EPO was also found to protect epithelial cells from neutrophil-mediated apoptosis agents. Thus, EPO is thought to have anti-apoptotic and cytoprotec-tive properties in the respiratory epithelium.

Pretreatment with EPO appears to attenuate I/R-induced lung injury in rats. This ability is at least partially due to EPO's inhibition of the accumulation of polymorphonuclear neutrophils in lung tissue. In the present study, CAT activities in the EPO-treated group were similar to those of the BC group, suggesting a lack of polymorphonuclear neutrophil accumulation and/or lack of excessive intracellular $H_2O_2$ accumulation. At 24 hours after contusion in the present study, alveolar edema/congestion and leukocyte infiltration in the lung tissue significantly increased, but subsided following administration of EPO. EPO most likely acts through a variety of pathways to protect lung tissue against damage from inflammation.

Conflict of interest: None declared.

REFERENCES

5. Jelkman W. Erythropoietin after a century of research: younger than
Künt göğüs travmasıyla oluşturulan pulmoner kontüzyon şoc modelinde eritropoietin düzeyleri ve eksojen eritropoietinin etkileri

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AMAC: Şcanlarda oluşturulan künt göğüs travmasına bağlı akışçıklar kontüzyon modelinde, ilk 24 saat içinde endojen eritropoietin düzeylerinin oksidatif stresli ilkeyi ve eksojen eritropoietinin etkilerini araştırmaya amaçlamaktayız.

GEREÇ VE YöNTEM: Otuş başı sıcak üç gruba ayrıldı: Grup BC (bazal kontrolü, n=7), herhangi bir işlem yapılmadı; Grup PC (pozitif kontrol, n=21), şcanlarda kontüzyon oluşturulan fakat tedavi verilmedi; Grup EPO-24 (eritropoietin tedavi, n=7), şcanlarda kontüzyon oluşturuldu ve künt göğüs travmasından hemen sonra intraperitoneal (5000 UI/kg) tek doz bir doz eritropoietin verildi. Pozitif kontrol grubu her grupta yedisiş çıkarlan olacak şekilde üç alt gruba ayrıldı (PC-6, PC-12, PC-24). PC-6, PC-12, PC-24 grubundaki şcanlar, künt travmadan 6, 12 ve 24 saat sonra, EPO-24 ve BC grubundaki şcanlar ise travmadan 24 saat sonra olduruldu. Akışçık dokusunda malondialdehit (MDA) ve EPO düzeyleri, süperoksit dismutaz (SOD) ve katalaz (CAT) aktiviteleri ölçüldü. BC, EPO-24 ve EPO-24 grubundalarında histopatolojik incelemeyle total doku hasanın tespit edildi.

BULGULAR: Ortalama MDA düzeyleri, SOD ve CAT aktiviteleri BC ve EPO-24 grubu gruplarda PC grubuna göre düşüktü (p<0.005). PC-6, PC-12 ve PC-24 grubunun ortalaması EPO kontrası say安全管理 BC grubuna göre daha yüksek bulundu (p<0.005). Akışçık dokusunda skarnf EPO-24 grubundan PC-24 grubuna göre daha düşüktü (p<0.005).

TARTIŞMA: Eritropoietin konsantrasyonları, SOD ve CAT düzeyleri pulmoner kontüzyon sonrası ilk 24 saatte akışçık dokusuna yüksek olduğu saptandı. Göğüs travması sonrası akut dönemde uygulanan tek doz intraperitoneal eritropoietin (5000 UI/kg), travmanın erken döneminde oksidatif hasan ve doku zedelemesini azaltmaktaydı.

Anahtar sözcükler: Akışçık hasan; eritropoietin; kontüzyon; oksidatif stres.