

Beneficial effects of agomelatine in experimental model of sepsis-related acute kidney injury

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ABSTRACT

BACKGROUND: Sepsis-related acute kidney injury (AKI) is a serious complication of sepsis. Problems persist regarding early diagnosis and treatment of AKI. The aim of the present study was to evaluate the efficacy of agomelatine, which is primarily known for its positive effects on depressive and anxiety disorders in sepsis-related AKI.

METHODS: Sepsis model was created with cecal ligation puncture (CLP). Rats were separated into 4 groups of 8 each: the control group, the sham-operated group, the CLP+saline group, and the CLP+agomelatine group. Agomelatine was administered intraperitoneally in doses of 20 mg/kg.

RESULTS: In the agomelatine group, reductions were observed in levels of tumor necrosis factor α (TNF- α), malondialdehyde (MDA), blood urea nitrogen (BUN), and creatinine, as well as in histological kidney scores, compared to the non-treated group. In addition, it was demonstrated that agomelatine treatment had positive effect on sepsis-induced morphological damage to renal and tubular tissues.

CONCLUSION: Agomelatine showed strong efficacy in sepsis-related AKI, demonstrated with histological and biochemical results in an experimental model. It is believed that antioxidant and pro-inflammatory effects of agomelatine are responsible for the improvement in kidneys.

Keywords: Acute kidney injury; agomelatine; cecal ligation puncture; sepsis.

INTRODUCTION

Sepsis is a complex clinical syndrome with high rates of morbidity and mortality.^[1] According to criteria of the American College of Chest Physicians (ACCP) and the Society of Critical Care Medicine (SCCM), the elementary definition of sepsis is “a complex immune reaction against a microorganism.”^[2] Sepsis is associated with multiple organ dysfunctions.^[3] Renal functions are particularly affected in the early stages of sepsis.^[4] Acute kidney injury (AKI) is a common and serious

complication that increases mortality and is characterized by rapid kidney failure due to micro- and macro-hemodynamic impairment and immune toxicity in kidney tissue cells.^[5,6] Pathophysiology of AKI is not clear, though multifactorial mechanisms containing ischemia–reperfusion injury, direct inflammatory injury, coagulation, endothelial cell dysfunction, and apoptosis are generally considered to be causes.^[7]

Agomelatine, N-[2-(7-Methoxy-1-naphthyl)ethyl]acetamide, is specified for the treatment of major depression, with dual effects on sleep problems and depressive disorders.^[8] It is a synthetic drug, an agonist of melatonin receptors (MT1 and MT2) and an antagonist of serotonin receptor (5-HT2C).^[9–11] Oxidative stress, which plays a key role in pathogenesis of sepsis, is a leading cause of organ dysfunction in particular.^[12] Melatonin at high doses is known for its strong antioxidant and anti-inflammatory activity.^[13] In several experimental studies, beneficial effects of melatonin have been demonstrated with sepsis models.^[13–16] Although the exact mechanism is not clear, it is thought that the antioxidant, anti-apoptotic, anti-inflammatory, and immunomodulating effects of melato-

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nin are responsible for good effects on sepsis.^[14] In addition, reports have indicated protective effects of melatonin on renal functions.^[17-19] However, exact pathophysiology of sepsis-related AKI remains unclear and, in spite of new treatment options, AKI continues to occur with high mortality.^[20]

The aim of the present study was to research effects of agomelatine, a melatonin analogue, on sepsis-induced AKI. Cecal ligation puncture (CLP) was performed on rats to create an animal model of sepsis, and agomelatine effects on renal functions were evaluated with biochemical and histopathological testing.

MATERIALS AND METHODS

Animals

Used in the present study were 42 male Sprague Dawley mature albino rats, each weighing 200–220 g. They were fed ad libitum and housed in pairs in steel cages in a temperature-controlled environment ($22\pm 2^\circ\text{C}$) with light/dark cycle of 12 hours each. Protocol was approved by the committee for animal research and the study strictly conformed to the animal experiment guidelines of the Committee for Human Care.

Drugs

All drugs were freshly prepared. Agomelatine (Valdoxan®, Servier Laboratories Ltd., Slough, UK) was dissolved in saline. Saline (0.9% NaCl) was used as control solution. All solutions were administered intraperitoneally (IP) in a volume of 1 mL/kg body weight.

Experimental Design

Rats were separated at random into 2 initial groups, and CLP was performed on 26 rats to induce a sepsis model. Ten rats (7 rats in the CLP+saline group and 3 in the CLP+agomelatine group) died in the first 24 hours after surgical procedure and were excluded from the study. No mortality occurred in the sham-operated group. Study groups were designed as follows:

- Group 1: Normal (non-operated and orally fed control, n=8)
- Group 2: Sham-operated (n=8)
- Group 3: CLP and 1 ml/kg 0.9 NaCl (saline) IP (n=8)
- Group 4: CLP and 20 mg/kg agomelatine IP (n=8)

Rats were anesthetized IP with injection of combination 80 mg/kg ketamine hydrochloride (Alfamine®; Alfasan International BV., Woerden, Holland) and 7 mg/kg xylazine hydrochloride (Alfazyne®; Alfasan International BV, Woerden, Holland).

Under aseptic conditions, 3-cm midline laparotomy was performed to expose the cecum with the adjoining intestine. The cecum was ligated tightly with a 3.0 silk suture at its base under the ileocecal valve and punctured once with a 22-gauge needle. The cecum was then gently squeezed to extrude a small amount of feces from the perforation site. The cecum

was returned to the peritoneal cavity, and the laparotomy incision was closed with 4-0 polyglactin 910 sutures. Following surgery, the animals were permitted a period to recover before being placed in their cages. In the sham group, under aseptic conditions, only laparotomy was performed; the cecum was neither ligated nor punctured. Rats were considered septic 5 hours after CLP.^[21] Treatments were performed within the first hour of surgical procedure. The study was concluded after 24 hours. The rats were euthanized with an overdose of pentobarbital sodium, and blood samples were collected by cardiac puncture for biochemical analysis.

Determination of BUN and Creatinine Levels

Blood urea nitrogen (BUN) and creatinine concentrations were determined spectrophotometrically, using an automated system of analysis. BUN and creatinine concentrations were expressed as mg/dL.

Determination of Plasma TNF- α Levels

Plasma TNF- α levels were measured using commercially available ELISA enzyme-linked immunosorbent assay kit (Quansys Biosciences, Logan, UT, USA). Plasma samples were diluted 1:2, and TNF- α was determined in duplicate, according to manufacturer's guide. Detection limit for TNF- α assay was <2 pg/mL.

Determination of Lipid Peroxidation

Lipid peroxidation was determined in plasma samples by measuring malondialdehyde (MDA) levels as thiobarbituric acid-reactive substances (TBARS).^[22] Briefly, trichloroacetic acid and TBARS reagent were added to the plasma samples, then mixed and incubated at 100°C for 60 minutes. After cooling on ice, the samples were centrifuged at 3000 rpm for 20 minutes, and the absorbance of the supernatant was read at 535 nM. MDA levels were expressed as nM, and tetraethoxypropane was used for calibration.

Histopathological Studies of Kidney

For histological and immunohistochemical studies, all animals were IP anesthetized with 40 mg/kg ketamine and 4 mg/kg xylazine, and were perfused with 200 mL of 4% formaldehyde in 0.1 M phosphate-buffered saline. Formalin-fixed kidney sections (4 μm) were stained with hematoxylin and eosin. All sections were photographed with Olympus C-5050 digital camera mounted on Olympus BX51 microscope.

Morphological evaluation was performed by computerized image analysis system (Image-Pro Express 1.4.5; Media Cybernetics, Inc., Rockville, MD, USA) on 10 microscopic fields per section, examined with $\times 20$ magnification by an observer blinded to the study group.

Kidney sections from all groups were evaluated semi-quantitatively to determine extent of tubular epithelial necrosis,

Table 1. Malondialdehyde, TNF- α , BUN and creatinine levels

	Normal group	Sham-operated group	CLP + saline group	CLP + 20 mg/kg agomelatine group
Malondialdehyde (nM)	65.3 \pm 3.2	71.2 \pm 3.5	218.9 \pm 7.5**	212.5 \pm 6.5##
TNF- α (pg/mL)	21.07 \pm 1.9	22.8 \pm 2.14	274.5 \pm 7.9**	119.9 \pm 5.8##
Plasma BUN content (mg/dL)	24.7 \pm 1.2	21.6 \pm 1.5	55.01 \pm 3.2**	37.8 \pm 3.6#
Plasma creatinine content (mg/dL)	0.35 \pm 0.02	0.37 \pm 0.03	0.79 \pm 0.03*	0.56 \pm 0.05#

Results are presented as mean \pm SEM. * p <0.01, ** p <0.000 (different from normal and sham-operated groups), ## p <0.000, # p <0.05 (different from CLP + saline Group). TNF- α : Tumor necrosis factor alpha; BUN: Blood urea nitrogen; CLP: Cecal ligation and puncture.

luminal necrotic debris, tubular dilatation, hemorrhage, and interstitial inflammation, rated as follows: 0-5% = score 0; 6-20% = score 1; 21-40% = score 2; 41-60% = score 3; 61-80% = score 4; and 81-100% = score 5.^[23,24]

Statistical Analysis

Data are presented as mean \pm standard error of the mean (SEM). Data analyses were performed using SPSS software for Windows (version 15.0; SPSS Inc., Chicago, IL, USA). Data were analyzed with non-parametric Mann-Whitney U test, and p values of 0.05 or less were considered statistically significant.

RESULTS

Serum TNF- α , MDA, BUN, and creatinine levels of all groups

are shown in Table 1. MDA is a predictor of lipid peroxidation, and in cases of sepsis, high levels indicate oxidative stress. Plasma MDA levels were markedly elevated in the CLP+saline group, compared to the normal and sham-operated groups (p <0.000). A significant decrease was observed in the CLP+agomelatine group, compared to the CLP+saline group (p <0.000). No differences were observed between the normal and sham-operated groups.

TNF- α , which causes harmful effects of inflammation, is a pro-inflammatory cytokine that can be used to determine severity of sepsis as organ dysfunction. In the present study, plasma TNF- α was also found to be significantly higher in the CLP+saline group, compared to the normal and sham-operated groups (p <0.000). A significant decrease in TNF- α was

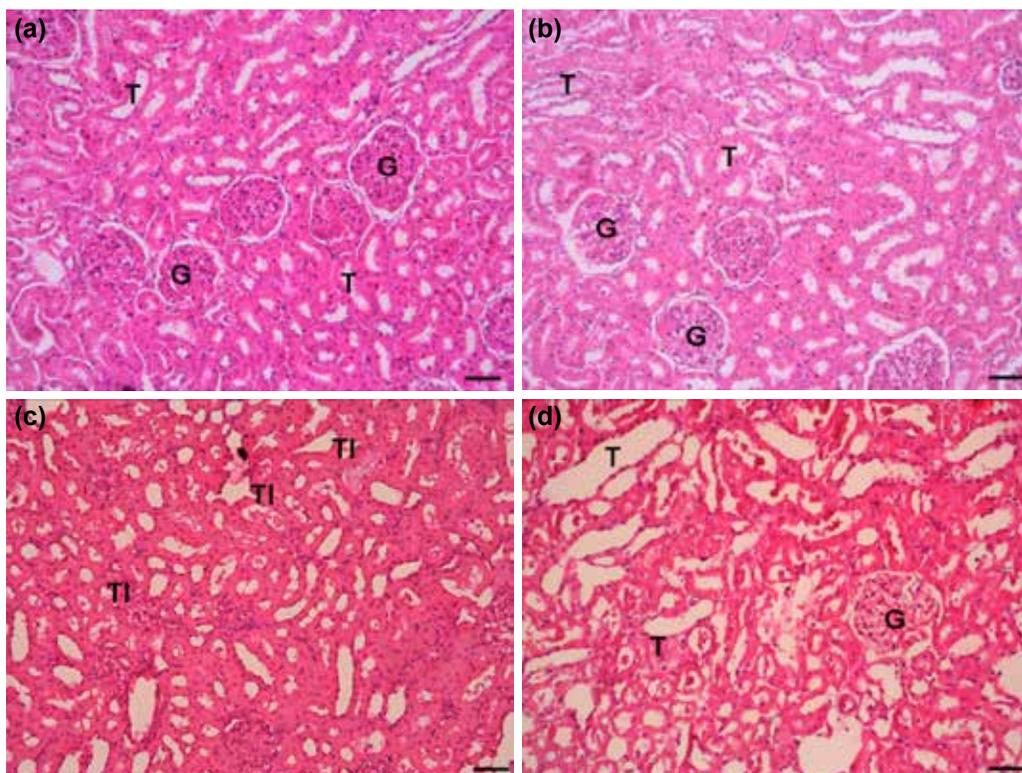


Figure 1. Kidney histopathology. (a) Kidney from normal group, H&E x10 magnification, renal tubules (T), (b) sham group showed minimal histopathological alteration, (c) CLP+saline group showed severe histopathological alteration related to tubular injury (TI). (d) CLP+agomelatine group showed decrease in tubular injury.

Table 2. Changes in histopathological kidney injury scores

	Normal group	Sham-operated group	CLP + saline group	CLP + 20 mg/kg agomelatine group
Tubularepithelial necrosis	0	0.5±0.18 [†]	3.3±0.18 ^{**}	2.1±0.22 ^{##}
Luminal necrotic debris	0	0.62±0.18 [†]	2.5±0.32 ^{**}	1.37±0.32 [#]
Tubular dilatation	0	0.37±0.19 [†]	3.1±0.22 ^{**}	1.8±0.35 [#]
Hemorrhage	0	0.5±0.18 [†]	3.0±0.3 ^{**}	1.9±0.22 ^{##}
Interstitial inflammation	0	0.6±0.26 [†]	3.25±0.5 ^{**}	1.75±0.36 [#]

Results are presented as mean ± SEM. [†]p<0.05 (different from normal group), ^{**}p<0.000 (different from normal and sham-operated groups), [#]p<0.05, ^{##}p<0.01 (different from CLP + saline group). CLP: Cecal ligation and puncture.

also observed in the CLP+agomelatine group (p<0.000). No differences were observed between the normal and sham-operated groups.

AKI was assessed by measurement of BUN and creatinine levels. BUN levels were compared among the 4 groups, with the highest in the CLP+saline group. BUN was significantly lower in the CLP+agomelatine group, compared to the CLP+saline group (p<0.05). Creatinine levels were markedly increased in the CLP+saline group, compared to the normal and sham groups (p<0.01). In the CLP+agomelatine group, a significant decrease was observed in creatinine levels, compared to the CLP+saline group (p<0.05). No difference in BUN or creatinine levels was observed in the normal or sham groups.

Histopathology of kidney tissue in all groups is shown in Fig. 1. Alteration of kidney tissue in sham group was minimal. Histopathological indicators of AKI (tubular epithelial necrosis, luminal necrotic debris, tubular dilatation, hemorrhage, and interstitial inflammation) were observed in the CLP+saline group. Tubular injury was observed in the CLP+agomelatine group, though to a lesser extent than in the CLP+saline group. Mean±SD of kidney tissue tubular epithelial necrosis scores were (0.5±0.18), (3.3±0.18), and (2.1±0.22); scores for luminal necrotic debris were (0.62±0.18), (2.5±0.32), and (1.37±0.32); scores for tubular dilatation were (0.37±0.19), (3.1±0.22), and (1.8±0.35); scores for hemorrhage were (0.5±0.18), (3.0±0.3), and (1.9±0.22), and scores for interstitial inflammation were (0.6±0.26), (3.25±0.5), and (1.75±0.36) in the sham, CLP+saline, and CLP+agomelatine groups, respectively (Table 2). A marked increase in tubular epithelial necrosis, luminal necrotic debris, tubular dilatation, hemorrhage, and interstitial inflammation was observed in the sham group, compared to the normal group (p<0.05). Scores of all histological parameters in the CLP+saline group were higher than those in the sham and normal groups (p<0.000). There was a significant decrease in scores of all histological parameters in the CLP+agomelatine group, compared to the CLP+saline group (p<0.05 for luminal necrotic debris scores, tubular dilatation scores, and interstitial inflammation scores; p<0.01 for tubular epithelial necrosis scores and hemorrhage scores).

DISCUSSION

The curative effect of agomelatine on the kidney was demonstrated in the present study with biochemical and histological parameters in an experimental model of sepsis-induced AKI. AKI occurs in nearly half of all septic patients and is associated with increased mortality.^[25] It has been suggested that AKI in sepsis is correlated with important destructive effects. While detection of acute lung injury is not easy in early stages, the authors suggested that it can indicate severity of sepsis.^[26]

Increased levels of BUN and creatinine were used to define AKI. Craciun et al. reported that creatinine is a predictor of glomerular filtration rate, which, though influenced by factors such as gender and age, can be used to detect AKI. It was also reported that BUN can indicate kidney function, but is less specific than creatinine. The authors suggested that high levels of BUN can reflect mortality of sepsis.^[27] In the present study, in addition to BUN and creatinine levels, histopathological changes in renal and tubular tissues were used to determine renal injury. Olguner et al. reported that an association between high kidney injury scores and sepsis has been histologically demonstrated.^[28] For example, the histological kidney score was found to be higher in the sepsis group than in the sham group of a study by Koca et al. These parameters were also used to evaluate an agent in AKI.^[29] In the present study, histological findings were similarly used in the diagnosis stage, as well as to evaluate effects of agomelatine on AKI.

Certain mechanisms are believed to contribute to the pathogenesis of AKI, which has yet to be clearly understood. In the present study, TNF-α levels were higher in the sepsis group than in the normal and sham groups. Luo et al. demonstrated that pro-inflammatory cytokines such as TNF-α increase with sepsis-related AKI. The authors suggested that the inflammation largely contributed to the pathogenesis of AKI.^[30] Similarly, Chancharoentana and et al. demonstrated that hypercytokinemia plays a more distinct role in sepsis-related AKI than in non-sepsis related AKI.^[31]

It has been determined that oxidative stress plays an important role in sepsis-related AKI.^[32] The results of the present

study support this conclusion. MDA, the end-product of lipid peroxidation, was higher in the sepsis group than in the normal and sham groups.

TNF- α , MDA, BUN, and creatinine levels were used to evaluate the effect of agomelatine on sepsis-induced AKI. Histological investigations and scores to determine kidney injury were used for further assessment. Levels of TNF- α , MDA, BUN, and creatinine were reduced, as were histological kidney injury scores, in the AKI group treated with agomelatine, compared to the AKI group that was not. Put simply, decrease in BUN and creatinine can indicate the positive effect of agomelatine on renal function. This suggestion is supported with histopathological kidney investigations.

Agomelatine is a melatonin analogue, known to be more potent than melatonin in antidepressant models.^[33] The effect of agomelatine on inflammation was studied by Molteni et al.^[34] Pro-inflammatory cytokines were found to be reduced in rats treated with agomelatine, a finding similar to that of the present study. The authors reported that agomelatine modified the expression of enzymes involved in the kynurenine pathway. It has been suggested that the kynurenine pathway is aggravated in cases of septic shock.^[35] Agomelatine may have effects beneficial to this pathway, in addition to its antioxidant and pro-inflammatory effects on sepsis-related AKI.

The efficacy of melatonin in sepsis models has been documented.^[13,14,16] Shang et al. reported that use of melatonin as a therapeutic agent in endotoxemic rats decreased incidence of acute lung injury by reducing lipid peroxidation, neutrophil infiltration, TNF- α release, and IL-10 production. It has been demonstrated that melatonin affects sepsis by reducing inflammation and inhibiting nuclear factor κ B (NF- κ B) activation.^[16] Srinivasan noted that Gitto et al. demonstrated positive effects of melatonin on oxidative stress in newborns with sepsis, utilizing MDA, 4-HDA concentration, and nitrate level. Melatonin was recommended because of its immunomodulatory, antioxidant, and anti-apoptotic effects on sepsis caused by multiple organ failure.^[14] Similarly, in a study by Lowes et al., melatonin was administered to rats with sepsis induced by lipopolysaccharide/peptidoglycan G (LPS/Pep G) in an acute model, and it was reported that melatonin protected mitochondria from oxidative stress and inflammation.^[13] In addition to those using sepsis models, other studies have demonstrated beneficial effects of melatonin on renal damage as ischemia-reperfusion injury or acute renal failure.^[11,18,36]

Agomelatine has generally been studied in relation to depressive and anxiety disorders, and, to the best of our knowledge, the present study is the first to research its effects on sepsis-related AKI. Agomelatine has effects similar to those of melatonin, due to its melatonergic receptors. Hence, the mechanism responsible for these beneficial effects is generally attributed to the antioxidant and anti-inflammatory effects of melatonin. Though it is outside the scope of the present

study, the authors suspect that agomelatine is more effective than melatonin, due to its impact on the kynurenine pathway in cases of sepsis-related AKI.

Conclusion

Results of the present study indicate that pro-inflammatory and antioxidant mechanisms play important roles in the pathogenesis of sepsis-induced AKI. While the mechanism is not entirely clear, the present data suggest that agomelatine may have beneficial effects on kidneys. Further studies are needed before agomelatine can be suggested as treatment of sepsis-related AKI.

Conflict of interest: None declared.

REFERENCES

1. Wang HE, Shapiro NI, Safford MM, Griffin R, Judd S, Rodgers JB, et al. High-sensitivity C-reactive protein and risk of sepsis. *PLoS One* 2013;8:e69232. [CrossRef](#)
2. Hecker A, Uhle F, Schwandner T, Padberg W, Weigand MA. Diagnostics, therapy and outcome prediction in abdominal sepsis: current standards and future perspectives. *Langenbecks Arch Surg* 2014;399:11–22.
3. Bozza FA, D'Avila JC, Ritter C, Sonnevile R, Sharshar T, Dal-Pizzol F. Bioenergetics, mitochondrial dysfunction, and oxidative stress in the pathophysiology of septic encephalopathy. *Shock* 2013;39 Suppl 1:10–6.
4. Seija M, Baccino C, Nin N, Sánchez-Rodríguez C, Granados R, Ferruelo A, et al. Role of peroxynitrite in sepsis-induced acute kidney injury in an experimental model of sepsis in rats. *Shock* 2012;38:403–10. [CrossRef](#)
5. Mayeux PR, MacMillan-Crow LA. Pharmacological targets in the renal peritubular microenvironment: implications for therapy for sepsis-induced acute kidney injury. *Pharmacol Ther* 2012;134:139–55. [CrossRef](#)
6. Legrand M, Dupuis C, Simon C, Gayat E, Mateo J, Lukaszewicz AC, et al. Association between systemic hemodynamics and septic acute kidney injury in critically ill patients: a retrospective observational study. *Crit Care* 2013;17:R278. [CrossRef](#)
7. Ademri C, Selmi V, Vitali L, Nosi D, Tani A, Thyron GD, et al. Expression and characterization of anionic components in the tubulointerstitial compartment of rat kidney during polymicrobial sepsis. *Acta Histochem* 2014;116:94–105. [CrossRef](#)
8. Jakovljević M. Agomelatine as chronopsychopharmaceutics restoring circadian rhythms and enhancing resilience to stress: a wishful thinking or an innovative strategy for superior management of depression? *Psychiatr Danub* 2011;23:2–9.
9. Boyce P, Barriball E. Circadian rhythms and depression. *Aust Fam Physician* 2010;39:307–10.
10. Fornaro M, Prestia D, Colicchio S, Perugi G. A systematic, updated review on the antidepressant agomelatine focusing on its melatonergic modulation. *Curr Neuropharmacol* 2010;8:287–304. [CrossRef](#)
11. Eser D, Baghai TC, Möller HJ. Agomelatine: The evidence for its place in the treatment of depression. *Core Evid* 2010;4:171–9.
12. Wheeler DS. Oxidative Stress in Critically Ill Children with Sepsis. *Open Inflamm J* 2011;4:74–81. [CrossRef](#)
13. Lowes DA, Webster NR, Murphy MP, Galley HF. Antioxidants that protect mitochondria reduce interleukin-6 and oxidative stress, improve mitochondrial function, and reduce biochemical markers of organ dysfunction in a rat model of acute sepsis. *Br J Anaesth* 2013;110:472–80.
14. Srinivasan V, Pandi-Perumal SR, Spence DW, Kato H, Cardinali DP. Melatonin in septic shock: some recent concepts. *J Crit Care* 2010;25:656.e1–6. [CrossRef](#)

15. Srinivasan V, Mohamed M, Kato H. Melatonin in bacterial and viral infections with focus on sepsis: a review. *Recent Pat Endocr Metab Immune Drug Discov* 2012;6:30–9. [CrossRef](#)
16. Shang Y, Xu SP, Wu Y, Jiang YX, Wu ZY, Yuan SY, et al. Melatonin reduces acute lung injury in endotoxemic rats. *Chin Med J (Engl)* 2009;122:1388–93.
17. Fariás JG, Zepeda AB, Calaf GM. Melatonin protects the heart, lungs and kidneys from oxidative stress under intermittent hypobaric hypoxia in rats. *Biol Res* 2012;45:81–5. [CrossRef](#)
18. Nava M, Romero F, Quiroz Y, Parra G, Bonet L, Rodríguez-Iturbe B. Melatonin attenuates acute renal failure and oxidative stress induced by mercuric chloride in rats. *Am J Physiol Renal Physiol* 2000;279:F910–8.
19. Aslaner A, Gunal O, Turgut HT, Celik E, Yildirim U, Demirci RK, et al. Effect of melatonin on kidney cold ischemic preservation injury. *Int J Clin Exp Med* 2013;6:794–8.
20. Liu B, Chen YX, Yin Q, Zhao YZ, Li CS. Diagnostic value and prognostic evaluation of Presepsin for sepsis in an emergency department. *Crit Care* 2013;17:R244. [CrossRef](#)
21. İşeri SO, Sener G, Saglam B, Gedik N, Ercan F, Yegen BC. Oxytocin protects against sepsis-induced multiple organ damage: role of neutrophils. *J Surg Res* 2005;126:73–81. [CrossRef](#)
22. Demougeot C, Marie C, Beley A. Importance of iron location in iron-induced hydroxyl radical production by brain slices. *Life Sci* 2000;67:399–410. [CrossRef](#)
23. Ashrafi F, Nematbakhsh M, Safari T, Talebi A, Nasri H, Khazaei M, et al. A combination of vitamin C and losartan for cisplatin-induced nephrotoxicity in rats. *Iran J Kidney Dis* 2012;6:361–5.
24. Eren Z, Coban J, Ekinci ID, Kaspar C, Kantarci G. Evaluation of the effects of a high dose of erythropoietin-beta on early endotoxemia using a rat model. *Adv Clin Exp Med* 2012;21:321–9.
25. Holthoff JH, Wang Z, Patil NK, Gokden N, Mayeux PR. Rolipram improves renal perfusion and function during sepsis in the mouse. *J Pharmacol Exp Ther* 2013;347:357–64. [CrossRef](#)
26. Bhargava R, Altmann CJ, Andres-Hernando A, Webb RG, Okamura K, Yang Y, et al. Acute lung injury and acute kidney injury are established by four hours in experimental sepsis and are improved with pre, but not post, sepsis administration of TNF- α antibodies. *PLoS One* 2013;8:e79037.
27. Craciun FL, Iskander KN, Chiswick EL, Stepien DM, Henderson JM, Remick DG. Early murine polymicrobial sepsis predominantly causes renal injury. *Shock* 2014;41:97–103. [CrossRef](#)
28. Olguner CG, Koca U, Altekin E, Ergür BU, Duru S, Girgin P, et al. Ischemic preconditioning attenuates lipid peroxidation and apoptosis in the cecal ligation and puncture model of sepsis. *Exp Ther Med* 2013;5:1581–8.
29. Koca U, Olguner ÇG, Ergür BU, Altekin E, Taşdoğan A, Duru S, et al. The effects of dexmedetomidine on secondary acute lung and kidney injuries in the rat model of intra-abdominal sepsis. *ScientificWorldJournal* 2013;2013:292687. [CrossRef](#)
30. Luo CJ, Zhang FJ, Zhang L, Geng YQ, Li QG, Hong Q, et al. Mesenchymal stem cells ameliorate sepsis-associated acute kidney injury in mice. *Shock* 2014;41:123–9. [CrossRef](#)
31. Chanchaoenthana W, Tiranathanagul K, Srisawat N, Susantitaphong P, Leelahavanichkul A, Praditpornsilpa K, et al. Enhanced vascular endothelial growth factor and inflammatory cytokine removal with online hemodiafiltration over high-flux hemodialysis in sepsis-related acute kidney injury patients. *Ther Apher Dial* 2013;17:557–63. [CrossRef](#)
32. Wang Z, Holthoff JH, Seely KA, Pathak E, Spencer HJ 3rd, Gokden N, et al. Development of oxidative stress in the peritubular capillary micro-environment mediates sepsis-induced renal microcirculatory failure and acute kidney injury. *Am J Pathol* 2012;180:505–16. [CrossRef](#)
33. Srinivasan V, Zakaria R, Othman Z, Lauterbach EC, Acuña-Castroviejo D. Agomelatine in depressive disorders: its novel mechanisms of action. *J Neuropsychiatry Clin Neurosci* 2012;24:290–308. [CrossRef](#)
34. Molteni R, Macchi F, Zecchillo C, Dell'aghi M, Colombo E, Calabrese E, et al. Modulation of the inflammatory response in rats chronically treated with the antidepressant agomelatine. *Eur Neuropsychopharmacol* 2013;23:1645–55. [CrossRef](#)
35. Zeden JP, Fusch G, Holtfreter B, Schefold JC, Reinke P, Domanska G, et al. Excessive tryptophan catabolism along the kynurenine pathway precedes ongoing sepsis in critically ill patients. *Anaesth Intensive Care* 2010;38:307–16.
36. Shih YC, Lee PY, Cheng H, Tsai CH, Ma H, Tarng DC. Adipose-derived stem cells exhibit antioxidative and antiapoptotic properties to rescue ischemic acute kidney injury in rats. *Plast Reconstr Surg* 2013;132:940e–51e. [CrossRef](#)

DENEYSEL ÇALIŞMA - ÖZET

Sepsis nedenli akut böbrek hasarında agomelatinin etkilerinin değerlendirilmesi

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AMAÇ: Sepsis nedenli akut böbrek hasarı sepsisin önemli komplikasyonlarından biridir. Sepsisin erken dönemlerinde gelişir ve mortaliteyi artırır. Hem erken tanı sürecinde hem de tedavi aşamasında hala problemler vardır. Bu çalışmanın amacı, genellikle anksiyete ve depresyonda kullanılan agomelatinin sepsis nedenli akut böbrek hasarındaki olası etkilerinin değerlendirilmesidir.

GEREÇ VE YÖNTEM: Sepsis modeli çekal ligasyon ve delme (CLP) tekniği ile oluşturuldu. Siçanlar her biri sekizerli dört gruba ayrıldı. İlk grup normal, ikinci grup sham grubu olarak belirlendi. Üçüncü grup sepsis modeli oluşturulup sadece salin verilen grup, dördüncü grup ise sepsis modelini takiben agomelatine uygulanan grup olarak belirlendi. Yirmi dört saatin sonunda böbrek doku örnekleri ve kanlar alınarak histopatolojik ve biyokimyasal yöntemlerle analizleri yapıldı.

BULGULAR: Çalışmada agomelatine uygulanan grupta sadece salin uygulanan gruba göre TNF- α , MDA, BUN, kreatin ve histolojik böbrek skorlaması daha düşük olarak bulundu. Aynı zamanda, histopatolojik değerlendirmede agomelatine tedavisinin böbrek dokularında sepsisin oluşturduğu hasarlanmayı düzelttiği görüldü.

TARTIŞMA: Bu çalışmada, agomelatine sepsis nedenli akut böbrek hasarında faydalı olduğu savunulmaktadır. Bu hem biyokimyasal hem de histolojik değerlendirmeler ile gösterilmiştir. Agomelatine anti-oksidan ve proenflamatuar etkilerinin böbreklerdeki bu düzelmeden sorumlu olduğu düşünülmektedir.

Anahtar sözcükler: Agomelatine; akut böbrek hasarı; CLP; sepsis.

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