EXPERIMENTAL STUDY

The effects of epidural bupivacaine on ischemia/reperfusion-induced liver injury

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ABSTRACT

BACKGROUND: Several animal studies showed beneficial effects of thoracic epidural anesthesia (TEA) in hippocampal, mesenteric and myocardial IR injury (2–4). In this study, we investigated the effects of epidural bupivacaine on hepatic ischemia reperfusion injury in a rat model.

MATERIAL AND METHODS: Eighteen rats were randomly divided into three groups each containing 6 animals. The rats in Group C had sham laparotomy. The rats in the Group S were subjected to liver IR through laparotomy and 20 mcg/kg/h 0.9 % NaCl was administered to these rats via an epidural catheter. The rats in the Group B were subjected to liver IR and were given 20 mcg/kg/h bupivacaine via an epidural catheter. Liver tissue was harvested for MDA analysis, apoptosis and histopathological examination after 60 minutes of ischemia followed by 360 minutes of reperfusion. Blood samples were also collected for TNF-α, IL-1β, AST and ALT analysis.

RESULTS: The AST and ALT levels were higher in ischemia and reperfusion group, which received only normal saline via the thoracic epidural catheter, compared to the sham group. In the ischemia reperfusion group, which received bupivacaine via the epidural catheter, IL-1 levels were significantly higher than in the other groups. TNF-α levels were higher in the Groups S and B compared to the sham group. Bupivacaine administration induced apoptosis in all animals.

CONCLUSION: These results showed that thoracic epidural bupivacaine was not a suitable agent for preventing inflammatory response and lipid peroxidation in experimental hepatic IR injury in rats. Moreover, epidural bupivacaine triggered apoptosis in hepatocytes. Further research is needed as there are no studies in literature investigate the effects of epidural bupivacaine on hepatic ischemia reperfusion injury (Tab. 3, Fig. 3, Ref. 34). Text in PDF www.elis.sk.

KEY WORDS: thoracic epidural anesthesia, bupivacaine, ischemia reperfusion injury, liver.
Experimental groups

Rats were randomly divided to three groups: Group S (n = 6) was the sham group. Group IR (n = 6) was the control group. After the induction of general anesthesia, an epidural catheter was placed and 20mcg/kg/h of 0.9% NaCl was administered via this catheter. After this procedure, laparotomy was performed and IR was induced by the procedure explained below. Group IR-B (n = 6) was the Bupivacaine group. After the induction of general anesthesia and the administration of 20mcg/kg/h Bupivacaine (Marcain® 1 ml = 5mg Astra Zeneca, Istanbul, Türkiye) via an epidural catheter, IR was induced. General anesthesia was induced by 50 mg/kg Ketamin (Ketalar® 1 mL = 50 mg, Pfizer, Istanbul, Türkiye) and 2.5 mg/kg Xyslazin (1 mL = 20mg, Egevet, Izmir, Türkiye) The rats were hydrated via a 24 Gauge catheter placed into the tail vein.

Epidural catheterization

Epidural catheterization was performed using a microsurgical technique. Briefly, each rat was placed in the prone position. The lumbar vertebral column was flexed by placing a cylinder transversely under the lower abdomen. The fourth lumbar spinal process was exposed and cut. A small hole was drilled through the cranial subdural position of the catheter. Correct positioning of the epidural catheter was confirmed after the completion of the experiment by autopsy. All catheters were sutured to the fascia, tunneled and advanced to T6. A repeated negative aspiration test excluded the subdural position of the catheter. Correct positioning of the epidural catheter was confirmed after the completion of the experiment by autopsy. All catheters were sutured to the fascia, tunneled under the skin, exteriorized at the neck, and protected by a swivel device. The epidural catheter dead-space (28 L) was filled with bupivacaine 0.5% or saline 0.9% according to group assignment.

Induction of ischemia and reperfusion

A midline laparotomy was performed; hepatic artery in the left portal triad was identified and clamped with a microvessel clip (Harvard Apparatus, Inc., Holliston, MA). Epidural infusions started as continuous infusion (20 mcg/kg/h of saline 0.9% and 20 mcg/kg/h of bupivacaine 0.5%) and continued until the end of the reperfusion period. During the period of ischemia, the abdominal wall incision was kept closed to prevent heat and fluid loss. The swivel device allowed continuation of the infusion during the reperfusion period and unrestricted movement of animals in the cage. After 1 hour of ischemia, the vascular clip was removed from the hepatic artery in the left portal triad to permit reperfusion, return of pulse, and the reestablishment of pink color. During the reperfusion phase, the intestines were returned to the abdominal cavity, and the laparotomy incision was closed in layers. A reperfusion period was maintained for 360 min. After this period, 4–5 ml blood was collected via intracardiac puncture, then all rats were sacrificed and liver tissues were quickly removed.

Liver function tests

Serum levels of AST, ALT were measured with standard methods using commercial kits in Beckmann Olympus AU2700 auto-analyser.

Cytokine measurement

The blood samples were obtained and placed in microcentrifuge tubes, and the plasma was separated by centrifugation, immediately frozen, and stored at 80°C until the time of assay. Plasma TNF-α, and IL-1β levels were detected in a 96-well microtiter plate with a commercial enzyme-linked immunosorbent assay (ELISA) kit (eBioscience®) according to the manufacturer’s guidelines. All samples were tested in duplicate. The plate was read on ELX800 automated microplate readers at 450 nm. The concentrations of TNF-α, and IL-1β were calculated from a standard curve and expressed in picogram per milliliter (pg/mL). The lower limit of detection for ELISA was 8 to 16 pg/mL.

Processing and preparation of tissue

The tissue specimens were rapidly excised, washed in ice-cold normal saline, blotted, frozen in liquid nitrogen, and stored at −80°C until use. 10% (w/v) homogenation of liver tissues were made in Tris-HCl (0.1 M, pH 7.4) using an ice-chilled glass homogenizing vessel in a homogenizer fitted at 15 000 rpm. The suspended mixture was centrifuged at 1000 g for 10 min at 4°C in a refrigerated centrifuge.

Liver MDA activity assay

The extent of lipid peroxidation was determined as the concentration of thiobarbituric acid reactive substances (TBARS), according to the method of Okhawa et al (15). Briefly, 100 μL of liver homogenates or MDA standards were pipetted into test tubes containing 1.5 mL of 20% (w/v) glacial acetic acid (pH 3.5), 200 μL of 8.1% (w/v) sodium dodecyl sulphate (SDS), 1.5 mL of 0.8% (w/v) thiobarbituric acid (TBA) and 250 μL of distilled water. The test tubes were incubated at 95°C for 60 minutes with a marble on top of each test tube. After incubation, the test tubes were cooled and then centrifuged at 4000 × g for 10 minutes. The amount of malondialdehyde (MDA) formed was measured spectrophotometrically at 532 nm. 1,1,3,3-Tetraethoxypropane (TEP), a form of MDA, was used as standard in this assay. TBARS concentration was expressed as nmol of MDA per mg protein.

Histological determinations

All of the specimens were fixed in 10% buffered neutral formalin and embedded in paraffin. To visualize myocardial lesions at different levels, the entire heart was cut into four segments from apex to bottom. The segments were embedded in paraffin and 4-μm thickness cross-sections were cut from each segment.

The slides were stained with Hematoxylin-Eosin (Bio-optica, Milano, Italy) for the evaluation of the tissues' histological features. The slides were evaluated under a light microscope for apoptosis, inflammatory activity and IR injury.

Statistical analysis

Statistical analysis was performed by using the SPSS 17.0 packet program. Continuous variables were expressed as the mean ± SD. Values of p < 0.05 were accepted as significant. In order to identify normal or abnormal distributions, Shapiro-Wilk test was used. One way ANOVA was used in normally distributed indepen-
Results

Biochemical findings

We found significant differences between groups in terms of AST enzyme levels (p = 0.009). AST levels were significantly higher in the Group S than Group C (p = 0.001) (Tab. 1). Also, significant differences were found between groups in terms of ALT enzyme levels (p = 0.018). ALT levels were higher in the Group S than the Group C (p = 0.022) (Tab. 1). There were significant differences between the groups in terms of IL-1β levels (p < 0.0001). IL-1β levels in the Group S and B were significantly higher than in the Group C (p = 0.045, p < 0.0001, respectively) (Tab. 1).

We found significant inter-group differences in TNF-α levels (p = 0.001). TNF-α levels were significantly higher in the Group B than in the Group C and S (p = 0.001, p < 0.0001, respectively) (Tab. 1).

We couldn’t find any significant differences between study groups in terms of tissue MDA levels (p = 0.216) (Tab. 2).

Histopathological findings

We found normal hepatic tissue microanatomical characteristics in histological evaluation of specimens belonged to all study animals. In the group S, only in a single animal we found cellu-
lar damage findings including a congestion around portal fields, nuclear loss of hepatocytes, appearance of collapsed and irregular hepatocyte lines. In the group B, cellular damage findings were found in 3 animals. Additionally, in this group we found apoptotic objects in forms of cell groups or necrotic single cells. Irregular sinusoidal dilatations were partly identified around the central veins. Also mitotic figures – refer to regenerative hepatocyte activity - were identified. Histopathological findings are represented in Table 3. Also, microscopical appearances of specimens are showed in Figures 1, 2 and 3.

Discussion

In this study, we hypothesized that epidurally infused bupivacaine (0.5 %) would protect hepatic tissue against ischemia reperfusion injury via increasing splanchnic and hepatic perfusion that was proven in previous experimental model (16). However, our findings have suggested that epidural bupivacaine -in contrast to our hypothesis- resulted in an increased systemic inflammatory response, lipid peroxidation in liver tissue and apoptosis of hepatocytes.

An increased sympathetic nerve activity is one of the leading factor in hepatic injury and immune response. Triggered intrahepatic inflammation and liver injury secondary to increased sympathetic system activation were shown in different experimental studies, while decreased sympathetic activation was defined as a preventing factor in hepatic injury (9–11).

Thoracic epidural anesthesia (TEA) blocks sympathetic tone, denerves afferent sensorial nerves and results in regional anesthesia and analgesia. There is an increasing evidence that shows beneficial effects of TEA in various experimental in vitro, in vivo studies conducted for clinical entities include sepsis, IR injury, cardiovascular mortality, necrotizing pancreatitis, inflammatory processes (12–14, 17). However, in contrast to supporting data there are several reports that showed no beneficial effects of TEA in experimental models of inflammation and IR (18–21).

Wei et al (18) designed an in vitro experiment with activated human umbilical vein endothelial cells (HUVEC) in order to investigate different doses of lidocaine effect on inflammation. The authors reported that high dose lidocaine (0.5 mg/mL) reduced the IL-1β, IL-6 and IL-8 concentrations while low dose (0.005 mg/ ml) lidocaine couldn’t make any effective decrement in plasma concentrations of ICAM-1, IL-1β, IL-6 and IL-8. Similarly, we found that epidural bupivacaine at clinical doses couldn’t exert any anti-inflammatory effect that presented with increased plasma IL-1β levels.

Huang et al (20) investigated anti-inflammatory effects of bupivacaine on murine macrophages that incubated with lipopolysaccharide (LPS) alone (100 ng/ml) or LPS plus bupivacaine (1, 10 or 100 mikroM). They reported that bupivacaine significantly inhibited the COX-2 transcription and PGE (2) production in a dose-dependent manner. Also, they showed that bupivacaine inhibited TNF-alpha, IL-1beta, and IL-6 production triggered by LPS in a dose dependent manner. Another interesting finding of this study was not achieved a significant bupivacaine effect on endotoxin induced IL-10 production. In contrast to these findings, we found that epidurally administered bupivacaine caused an increased TNF-α and IL-1β levels in hepatic IR rat model.

In another in vitro study Kiefer et al (21) designed the Staphylococcus aureus added inflammation model; and they showed a reduced reactive oxygen species generation, CD11b expression and decreased granulocyte phagocytosis activity with lidocaine and bupivacaine treatments only at the highest concentrations (1846 μM and 770 μM respectively). According to these results of different investigations, we can state that different local anesthetic agents exert different anti-inflammatory effects in a dose dependent manner and these anti-inflammatory effects may be achieved only with the highest in vitro doses of agents.

Lipid peroxidation secondary to reactive oxygen species is one of the major factors that leads IR injury. Malondialdehyde is an end-product in lipid peroxidation (22). In our study, we couldn’t show any differences between the study groups in terms of tissue MDA levels. In contrast to our findings, Bedirli et al (23) showed strong anti-inflammatory effects of 20 mcL/h epidurally administered bupivacaine at 0.5% concentration in a mesenteric IR rat model. They showed that bupivacaine reduced plasma TNF-α, IL-6, IL-1β and tissue MDA levels significantly. Also, inflammation related apoptosis was significantly decreased with bupivacaine treatment.

Several studies showed a strong correlation between bupivacaine and cellular apoptosis via inhibition of complex I in mitochondria, impairing oxidative phosphorylation which resulted in a reduced ATP production and disrupting mitochondrial membrane potential (24, 25). Although the exact mechanism is not known, it is thought that local anesthetic induced apoptosis was related to mitogen activated protein kinase pathway (26, 27). Boselli et al (28) and Castro et al (29) showed a significant relationship between local anesthetic related neurotoxicity and cellular apoptosis. Similarly, Park et al (30) reported that bupivacaine induced schwann cell apoptosis, while in another study, Lu et al (31) showed a correlation between bupivacaine treatment and mitochondrial apoptosis. In our study, we found a significant hepatocyte apoptosis in bupivacaine group correlated to previous studies conducted by Park et al and Lu et al (30, 31).

Lee et al (32) investigated the effects of various local anesthetics (lidocaine 5 %, bupivacaine 1 –0.5 % and tetracaine 2.5 %) on systemic inflammation and renal functions in a renal IR injury model. They concluded that continuous infusion of bupivacaine 1% resulted in an impaired renal functions at 24th and 48th hours of IR. Additionally, in all local anesthetic groups, the apoptosis rates (lidocaine, bupivacaine and tetracaine respectively), mRNA -coding ICAM-1 and TNF alpha- expression were significantly higher than in the control group. In the bupivacaine 0.5% group, renal functions were protected in contrast to other study groups. Another interesting finding of this study was an unchanged IL-1β expression rate in all study groups. In our study, we couldn’t find any differences for IL-1β levels between the bupivacaine and control groups. However, apoptosis rates were significantly higher in the bupivacaine 0.5 % group than rates noted in sham and control groups.
Freise et al (33) investigated the effects of bupivacaine 0.5 % (15 mcg/h continuous infusion via thoracic epidural route) on rat caecal ligation and perforation induced sepsis model. The results of this study indicated that TEA treatment with bupivacaine 0.5 % was ineffective in the restoring sepsis induced sinusoidal vasoconstriction, sinusoidal leukocytes adhesion, increased TNF-α levels and increased serum ALT, AST levels. Another study conducted by Freise et al (34) showed ameliorated apoptosis rates in experimentally induced pancreatitis treated with continuous infusion of thoracic epidural bupivacaine (0.5 % concentration, 15 mcg/h infusion rate) in rats. In addition, they showed an ameliorated sinusoidal vasoconstriction with bupivacaine treatment. In contrast to this study, we showed high apoptosis rates in all bupivacaine treated (0.5 % concentration, 20 mcg/kg/h infusion rate) rats.

In summary, there are conflicting results regarding the thoracic epidural bupivacaine treatment in experimental IR injury and sepsis models designed for various organ systems. According to these data, we can state that one explanation of these different results may be the different experimental designs – drug concentration, IR injury modal, time interval between IR and scarification etc- of investigations. Additionally we believe that the obtained results may vary depend on the liver is primarily or secondarily affected organ. Future studies need to be conducted in order to clearly determine thoracic epidural bupivacaine effects on hepatic IR injury.

References


Received April 14, 2015.
Accepted September 20, 2015.