The effect of extracorporeal shock wave lithotripsy on distribution of interstitial cells of Cajal in rabbit renal pelvis and proximal ureter

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1. Introduction
Urolithiasis in the pediatric population is increasing in frequency (1,2). Management in children is very similar to that in adults in recent years. Among several management options, extracorporeal shock wave lithotripsy (ESWL) has become established for the management of stones located in the calyces or the renal pelvis of up to 2 cm in diameter. Although the efficacy of ESWL in children is higher than in adults due to easy passage of stone fragments through shorter and more elastic ureters, it still has some adverse effects (3). Besides renal and extrarenal tissue damages, incomplete fragmentation, residual stone fragments, and obstruction of urinary flow may also be seen after ESWL treatment (4,5). It is reasonable to assume that clearance of stone fragments after ESWL depends on contractility of the ureter, but to the best of our knowledge, the effect of ESWL on the contractility of the ureter has not been demonstrated before.

2. Materials and methods
Six New Zealand rabbits weighing 2500–3000 g were included in the study. After being fasted overnight,
rabbits were anesthetized with intramuscular ketamine hydrochloride (50 mg/kg Ketalar, Eczacıbaşı, Turkey). Right kidneys were exposed to 3 sessions of ESWL with 1000 shock waves at 14 kV (total of 3000 shock waves) by using an electrohydraulic-type third-generation Stonelith V5 ESWL device (PCK, Turkey). Each session of ESWL was performed for 20 min (rate: 50 shock waves/min) and 48 h after the previous treatment. The ESWL was performed by the same investigator each time. Ultrasonographic probing was performed to localize the right kidney. Since the focus distance of the ESWL device is 130 mm, an elevating shelter was put under the subjects. During the treatment period the rabbits were kept warm and in the same conditions with standard feeding.

Right side ureters and renal pelvises were allocated as the ESWL group (EG, n = 6), whereas left sides were allocated as the control group (CG, n = 6). Both pelvies and proximal ureters were harvested on the seventh day after ESWL. Tissues were examined histopathologically for presence of edema, inflammation, congestion, hemorrhage, fibrosis, and vascularization. Mast cell tryptase and CD117 (c-kit) staining was performed for evaluation of ICC distribution in samples.

The data obtained from the experiments were analyzed with the Mann–Whitney U test and median test since the variables were nonparametric (not normally distributed) as confirmed by ANOVA test (SPSS 15.0). P-values of lower than 0.05 were considered significant. The experiments were performed in adherence to the Declaration of Helsinki and by approval of the Local Ethics Committee of Kırıkkale University (2011/11). This study was supported by the Kırıkkale University Scientific Research Council (2011/59).

2.1. Histopathological evaluations

The preserved tissues were fixed with 10% formalin. All segments were embedded in paraffin blocks after tissue processing. Tissues were sectioned in pieces of 4–5 µm and stained with routine hematoxylin and eosin staining. The specimens were examined under a light microscope (Olympus CX31; Mason Technology, Ireland) by the same pathologist blinded to the study.

For both the EG and the CG, presences of edema, inflammation, congestion, hemorrhage, fibrosis, and vascularization were evaluated. Histopathologic findings were graded semiquantitatively for each parameter separately as follows: grade 0, normal; grade 1; mild, grade 2; moderate, and grade 3; severe.

2.2. Immunohistochemistry

Sections of 5 µm were prepared from each paraffin block. Each section was deparaffinized in xylene and rehydrated in graded serial ethanol baths, and was then placed in 3% H₂O₂ for 5 min to block endogenous peroxidase activity. Antigen retrieval was performed using a citrate buffer solution (pH6) for CD117 and mast cell tryptase antibodies. After 3 rinses of 5 min each in phosphate-buffered saline, sections were incubated for 1 h with primary antibodies in an automatized immunohistochemical staining machine (Leica Bond Max, Leica Biosystems, Newcastle Ltd., UK). Anti-CD117 (CD117-R-7-CE, Novastra, Leica Biosystems) and antimat cell tryptase (PA0019, Novastra, Leica Biosystems) were used as primary antibodies. A Novastra Bond Polymer Refine Detection system including peroxide block, postprimary, polymer reagent, DAB chromogen, and hematoxylin counterstain was used to detect positive staining. Phosphate-buffered saline was used as the negative control.

2.3. Determination of ICC distribution

Cytoplasmic immunostaining for CD117 and mast cell tryptase were considered as positive immunoreactivity. Since double-staining could not be performed, immunohistochemically stained sections were evaluated separately. First, positive staining of mast cells with mast cell tryptase was evaluated. Cells with polygonal-shaped cytoplasm having rough granules that were stained positive with CD117 at the same location were considered as mast cells in the immunohistochemical examination. The other cells that were negative for mast cell tryptase, positive for CD117, and had spindle-shaped cytoplasm were estimated as ICCs. The ICCs were counted manually in renal pelvis and proximal ureter sections under light microscope.

3. Results

Histopathologically, tissue edema was increased in renal pelvises and inflammation was increased in ureters in the EG compared to the CG. However, no statistical difference was detected between exposed and unexposed samples (P > 0.05) (Table). There was no statistical difference between groups regarding other parameters including congestion, hemorrhage, fibrosis, and vascularization (P > 0.05).

CD117 staining was detected in the renal pelvis (n = 2) and ureter (n = 2) of the CG. These cells were considered as Cajal cells since they were able to be distinguished from mast cells with mast cell tryptase staining (Figures 1 and 2). However, none of the samples obtained from the EG showed CD117 staining. There was no statistical difference between groups regarding CD117-positive cells (P > 0.05) (Figure 3).

4. Discussion

The treatment of choice for UUT stones has been ESWL since the 1980s, when it was first introduced (1,7). The first report regarding ESWL in children revealed that the complications and safety were similar to those in adults (7). Further studies were performed demonstrating the effectiveness of ESWL in children, and then it became a treatment of choice for the stones located in calyces or renal pelvises with a diameter of up to 2 cm (1,2).
The effectiveness of ESWL is measured by the stone-free rate, which has been reported from 50% to 95% in children (1–3,8). Some cases require more than 1 session of ESWL (up to 3 sessions) for a better stone-free rate (1,2,8,9). Each session consists of 1800 to 2000 shock waves (and up to 4000) between 14 and 21 kV (2,3). The usage of ultrasound for localization will provide less radiation exposure. In the present study, we performed ESWL on rabbits with a similar dosage in 3 sessions, just as reported in the literature (8–10).

Although ESWL is an efficient treatment modality in the management of urolithiasis in children, it has some adverse effects (3–5). Some of these are renal and extrarenal tissue damages, skin bruising, hematuria, transient tubular function deterioration, incomplete fragmentation, residual stone fragments, and obstruction of urinary flow (2–5). Besides the several clinical reports in humans about complications of ESWL, some studies in animals have also reported on the acute and chronic effects of ESWL (4,5,11). Nevertheless, contractility, which has an effect on stone clearance, has not been demonstrated before. There is one study revealing that ureteric contraction frequency was increased by in vitro prostaglandin stimulation as detected in ESWL-treated patients (12). Some of the adverse effects of ESWL on urinary flow are thought to be transient and would resolve several days after ESWL (2–5). We detected alteration in the distribution of ICCs in the EG in the present study. Although there was no statistical difference between groups, this decrease in ICC distribution in the EG might be an indicator of altered ureteric peristalsis, which could be transient or permanent.

Figure 1. Mast cell tryptase expression on samples (DAB, 20×).

Figure 2. CD117-positive, spindle-shaped cells that are considered Cajal-like cells within the muscular layer of the renal pelvis and proximal ureter in the CG (DAB, 40×).

Figure 3. The comparison of ICC distribution in the pelvis and proximal ureter of the groups.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Edema</th>
<th>Inflammation</th>
<th>Congestion</th>
<th>Hemorrhage</th>
<th>Fibrosis</th>
<th>Vascularization</th>
</tr>
</thead>
<tbody>
<tr>
<td>CG-Renal pelvis</td>
<td>0 * (0–1)</td>
<td>1 (0.75–1)</td>
<td>1 (1–1)</td>
<td>0 (0–0.25)</td>
<td>0 (0–0)</td>
<td>0 (0–0)</td>
</tr>
<tr>
<td>EG-Renal pelvis</td>
<td>1 * (0–1.5)</td>
<td>1 (0.75–3)</td>
<td>1 (1–1)</td>
<td>0 (0–0.25)</td>
<td>0 (0–0.25)</td>
<td>0 (0–0)</td>
</tr>
<tr>
<td>CG-Ureter</td>
<td>0 (0–0)</td>
<td>0 β (0–0.25)</td>
<td>1 (1–1)</td>
<td>0 (0–0.25)</td>
<td>0 (0–0)</td>
<td>0 (0–0)</td>
</tr>
<tr>
<td>EG-Ureter</td>
<td>0 (0–0.25)</td>
<td>1 β (0–1)</td>
<td>1 (0.75–1)</td>
<td>0 (0–0)</td>
<td>0 (0–0)</td>
<td>0 (0–0)</td>
</tr>
</tbody>
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*, β: P > 0.05.
this propelling movement and the factors deteriorating it are not clear (6,13,14). This peristalsis occurs even after surgical denervation of ureter or isolation of the pyeloureteric system, indicating autonomous pacemaker activity (6,13,14). Some pacemaker potentials were able to be shown by electrophysiological studies in the renal pelvis in animal studies (6,14). After recognition of ICCs in the gut (15,16), several studies were performed to investigate the presence and distribution of pacemaker cells in the UUT. Some spindle-shaped c-kit-positive cells were detected in the UUT. They were differentiated from other c-kit-positive cells, such as mast cells, by additional staining with mast cell markers, and were called Cajal-like cells (6,13,14). These cells are usually found within smooth muscle layers of the UUT. The number of them is highest in the proximal UUT and decreases gradually in distal parts of ureter. However, the distribution of these cells differs between species. They are found less frequently in pig, for example. On the other hand, differentiation of them requires species-specific antibodies (6,13,14). In the present study, we could detect only a few Cajal-like cells. Difficulty in double immunohistochemical staining of ICCs in rabbit tissue was also experienced in previous studies (6). We could detect ICCs in only 2 pelvis and 2 ureter samples in the CG. Although there was no statistical difference between groups, we could not detect any ICCs in the EG. This result may be obtained not only due to the deleterious effect of shock waves to the ICCs and ureteric peristalsis, but also due to the difficulty in staining with rabbit-specific antibodies against ICCs. Therefore, future studies are needed to prove the adverse effect of ESWL on ICC distribution in the UUT with different animal models.

In conclusion, rabbit does not appear a good model for investigating ICCs. ESWL may cause histopathological alterations in the renal pelvis and ureter. Since it has not been statistically proven, the reduced contractility of the ureter after ESWL may not be attributed to altered distribution of ICCs in renal pelvis and ureter.

Acknowledgments
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References