Effect of the duration of bladder overdistention on renal function and morphology in rats

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Received January 5, 2013; Accepted March 14, 2013

DOI: 10.3892/etm.2013.1028

Abstract. The aim of this study was to investigate the influence of the duration of bladder overdistention (DOBO) on kidney structure and function in rats. Bladder overdistention was induced in male Sprague-Dawley rats by an infusion of saline. Forty rats were divided into five groups: DOBO 1, 2, 4 or 8 h and the control. Renal function was evaluated using serum creatinine (Scr) and blood urea nitrogen (BUN). Apoptotic indices and morphologic changes of the kidney were detected by terminal deoxynucleotidyl transferase-mediated dUTP nick end labelling (TUNEL) staining and transmission electron microscopy (TEM). Compared with the control, rats undergoing 2, 4 or 8 h of overdistention showed significant, time-dependent increases in Scr (12.375 vs. 23.125, 34.375 and 51.500 µmol/l, respectively), BUN (6.980 vs. 18.689, 25.184 and 32.079 mmol/l, respectively), renal size (1.041 vs. 1.472, 1.484 and 1.634 cm², respectively) and renal pelvis separation (0.000 vs. 0.223, 0.320, 0.308 and 0.277 cm, respectively; P<0.01). In the rats, 2, 4 and 8 h of overdistension elicited time-dependent increases in the blood flow rate in the main renal artery (MRA; 44.827 vs. 49.082, 59.688 and 67.123 cm²/sec, control vs. DOBO 2, 4 and 8 h), the interlobar renal artery (IRA; 32.095 vs. 39.16 and 51.745 cm²/sec, control vs. DOBO 4 and 8 h) and the segmental renal artery (SRA; 21.171 vs. 24.355 and 25.358 cm²/sec, control vs. DOBO 4 and 8 h; P<0.01). TUNEL results showed that prolonged overdistention increased the apoptotic index of renal cells significantly (1.15, 1.77, 3.40, 5.34 and 13.91% for control and DOBO 1, 2, 4 and 8 h, respectively; P<0.01) and TEM indicated that prolonged overdistention resulted in ultrastructural injuries of increased severity. DOBO plays a significant role in the functional and structural impairment of the rat kidney. With increasing duration, the hemodynamic changes, cell apoptosis and ultrastructural injuries of the kidney are more evident, all of which may contribute to the increasingly serious impairment of renal function and morphology.

Introduction

Acute urinary retention (AUR) is a common complication in patients with benign prostatic hyperplasia (BPH) (1,2). Previous studies have shown that >10% of men >70 years old experience at least one episode of AUR over a 5-year period and this risk increases to one-third of men over a 10-year period (3).

Urinary retention induces an increase of intravesical pressure, which may affect renal function and morphology. Chronic urinary retention may impair renal function and lead to terminal kidney failure within a few years (4). Whether AUR results in an impairment of renal function and morphology requires investigation. A previous study observed that AUR affects glomerular and tubular renal function, which results in elevated urinary albumin excretion. Following AUR, glomerular permeability and tubular damage persists in the majority of patients (5). Data concerning renal function and morphology, however, remain scarce for AUR.

In clinical practice, the majority of patients with AUR are treated with catheterization (6). However, the durations of AUR prior to intervention differ markedly among patients due to differences in medical history and pain tolerance. Whether different AUR durations result in different impairments of renal function or morphology remains unclear; little has been published in the literature. In the current study, a rat model was used to investigate the effect of the duration of bladder overdistention (DOBO) on renal function and morphology, which may

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Key words: acute urinary retention, overdistention, rat, serum creatinine, terminal deoxynucleotidyl transferase-mediated dUTP nick end labelling, ultrastructure
Materials and methods

Animal model. Studies were performed on male Sprague-Dawley rats weighing 200-250 g (n=40). All rats received a standard diet, water ad libitum and were housed in a 12 h light/dark cycle. All animal care and experimental protocols were in accordance with the guidelines of Zhejiang University (Hangzhou, China). The 40 rats were allocated to five groups: the sham-operated control and DOBO 1, 2, 4 and 8 h groups (each n=8). Rats were anesthetized with urethane (1.0 g/kg i.p.) and anesthesia was maintained by supplementary injections of the same anesthetic. Once anesthetized, the rats were shaved for kidney Doppler ultrasound. The rat bladder was identified with a low midline abdominal incision. After emptying the bladder, the foreskin was ligated using 3-0 silk thread. A 24-G catheter was inserted into the apex of the bladder dome. The catheter was connected to an infusion pump, then 37°C 0.9% saline was infused (0.1 ml/min) until the total volume reached 1 ml. This was twice the mean bladder capacity of 0.5 ml established in preliminary experiments. The status of overdistention was maintained for 1, 2, 4 and 8 h, respectively. In the rats of the control group, the bladder was exposed and punctured but no saline was infused.

Doppler ultrasound detection. Kidney ultrasound was applied to all rats 0.5 h after the overdistention was relieved. The kidney length, width and cortex thickness were measured. Renal size was calculated using the following formula: Renal size (cm³) = renal width (cm) x cortex thickness (cm)/6. At the same time, the thicknesses of the cortex and hydrenephrosis levels were measured. Furthermore, Doppler ultrasound with a V4 MHz transducer was used to calculate the resistant index (RI) of the main renal artery (MRA), inter lobular artery (ILA) and segmental renal artery (SRA). The RI was calculated with the following formula: RI = (peak systolic shift - minimum diastolic shift)/peak systolic shift (4).

Renal function test. Blood samples were collected and stored at 4°C until examination. Serum creatinine (Scr) and blood urea nitrogen (BUN) were measured by the Department of Clinical Chemistry using an enzymatic method.

Transmission electron microscopy (TEM). To observe the ultra-structure of the bladder tissues, sections were stained with uranyl acetate and plumbic citrate and examined using TEM (TECNAI 10, Philips, Amsterdam, The Netherlands).

Statistical analysis. All results are expressed as the mean ± standard error of the mean (SEM). *P<0.05 vs. control group, †P<0.05 vs. 1 h DOBO group, ‡P<0.05 vs. 2 h DOBO group.

Table I. Effect of the duration of bladder overdistention (DOBO) on renal size, degree of separation of the renal pelvis and the thickness of the renal cortex.

| Group          | Renal size (cm³) | Separation of the pelvis (cm) | Thickness of the renal cortex (cm) |
|----------------|------------------|-------------------------------|-----------------------------------|
| Control (n=8)  | 1.041±0.159      | 0.000±0.000                   | 0.263±0.037                       |
| 1 h DOBO (n=8) | 1.146±0.162      | 0.223±0.059*                  | 0.259±0.058                       |
| 2 h DOBO (n=8) | 1.472±0.145a     | 0.320±0.029b                  | 0.276±0.032                       |
| 4 h DOBO (n=8) | 1.484±0.193c     | 0.308±0.044c                  | 0.251±0.052                       |
| 8 h DOBO (n=7) | 1.634±0.155c     | 0.277±0.028c                  | 0.256±0.035                       |

Data are reported as the mean ± standard error of the mean (SEM). *P<0.05 vs. control group, †P<0.05 vs. 1 h DOBO group, ‡P<0.05 vs. 2 h DOBO group.

Results

Survival. All rats survived and data were collected, with the exception of one rat in the 8 h overdistention group.

Doppler ultrasound detection. DOBO had significant effects on renal volume, the degree of separation of the pelvis and cortical thickness (Table I). Compared with the control, the renal size in the rats with 2, 4 or 8 h DOBO was significantly increased (P<0.05). Compared with the DOBO 1 h group, the DOBO 4 and 8 h groups demonstrated significant increases in renal size (P<0.05). A significant difference was also observed between the DOBO 2 h group and the DOBO 4 and 8 h groups (P<0.05). Furthermore, rats with a DOBO of 2, 4 or 8 h showed a significantly higher degree of separation of the pelvis than the controls (P<0.05); significant differences were also
observed between the DOBO 1 h group and the 2, 4 and 8 h groups (P<0.05). Cortical thicknesses showed no significant differences among the groups, however. Different overdistension durations had different effects on the blood flow of the renal arteries (Fig. 1). Blood flow in the MRA, IRA and SRA increased with distention time (P<0.01). The RIs of the MRA and SRA were significantly higher in the rats with 4 or 8 h DOBO than in the other groups (P<0.01, Table II).

Renal function test. Rats with 2, 4 or 8 h DOBO showed significant, time-dependent increases in SCr and BUN levels compared with sham-operated controls (P<0.01, Table II).

TUNEL assay. Increasing DOBO also promoted apoptosis within renal tissues. Compared with controls, the rats with 4 or 8 h DOBO showed significant increases in the apoptotic index (1.15 vs. 1.77, 3.4, 5.34 and 13.91%, respectively; P<0.01) and this effect was time-dependent. *P<0.01 vs. control group, †P<0.01 vs. 1 h DOBO group, ‡P<0.01 vs. 2 h DOBO group.

TEM. Electron microscopy revealed that increasing DOBO led to increasingly aggravated cellular and tissue injury. Evidence of injury included mitochondrial breakdown, tubular vacuolization and dilation, mesangial proliferation, necrosis,
Figure 5. Representative microscopic findings indicating the effect of the duration of bladder overdistention (DOBO) on renal ultrastructure. (A) Control group (magnification, x2,550). (B) 1 h DOBO group (magnification, x2,550). (C) 4 h DOBO group (magnification, x1,850). (D) 8 h DOBO group (magnification, x1,850). Electron microscopy revealed that an increase in the DOBO led to increasingly aggravated cellular and tissue injury. Evidence of injury included mitochondrial breakdown, tubular vacuolization and dilation, mesangial proliferation, necrosis, podocyte swelling and inosculcation and clearly visible slit pores (arrows).

Discussion

It is well known that BPH is a progressive disease, with risk of urinary retention and renal insufficiency (5). Renal impairment caused by BPH is usually a chronic disorder, which takes several years, even decades to develop (9). In the current study, it was identified that rats with bladder overdistention had significant, time-dependent reductions in SCr and BUN levels. Renal hemodynamic alterations and the consequences of progression to irreversible renal injury following unilateral ureteral obstruction (UUO) have been studied in detail (6). In animals with UUO, both renal blood flow (RBF) and glomerular filtration rate (GFR) decrease and remain depressed without intervention (7). The degree of damage to renal function depends on the duration of UUO and the species being examined (1,12,13). In the present study, the main ultrastructural changes were mitochondrial breakdown and disorganized podocytes; these and all other negative ultrastructural changes were further exacerbated by increasing durations of overdistention.

Based on these results and those reported previously, bladder overdistention affects not only the structure and function of the bladder but also of the kidney. Thus, more attention should be paid to the effects of AUR on the upper urinary tract. Clinical practice should be expanded to include monitoring of renal function in patients with AUR, especially in cases of prolonged AUR. These results indicate that the bladder should be decompressed as quickly as possible and the duration of AUR should be shortened to protect renal function in the treatment of AUR. Certainly, overdistention of the bladder is not the same as AUR and clinical circumstances are more complicated than animal experiments. Other factors such as comorbidity, age and physical conditions also contribute to the severity of impairment and the recovery of renal function following AUR.

In the present study, there was no recovery of renal function after the overdistention was relieved. How the duration of overdistention affects the recovery of renal insufficiency should be studied further.

DOBO plays an important role in functional and structural impairment of the kidney. Different overdistention durations lead to different severities of impairment of the rat kidney. With increasing duration, the hemodynamic changes, cell apoptosis and ultrastructural injuries of the kidney are more evident, all of which may contribute to more serious impairment of renal function and morphology.

Acknowledgements

The authors gratefully acknowledge Mrs. Li Wang from the Department of Electron Microscopy, Zhejiang University.
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