Comparison Of Silodosin (8mg) Versus Tamsulosin (0.4mg) In The Medical Expulsive Therapy

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Abstract - Background: Ureteral stones account for 22% of all urinary tract stones with 68% of them being located in the distal ureter. Conservative management strategies such as observation or medical expulsive therapy (MET) using pharmacological agents to facilitate spontaneous passage of ureteral stones have gained popularity in the management of ureteral stones during the recent years. Objectives: To compare the efficacy of silodosin (8 mg) versus tamsulosin (0.4mg), both in terms of the stone expulsion rate and the time to stone expulsion. Patients and Methods: A prospective and randomized controlled study was conducted in the department of Urology Ibn Sina Medical College, Dhaka, Bangladesh. Starting from October 2018, till September 2019; a total of 96 patients (M=56; F=40) who were between age group of 18-50 years, who had unilateral, non-impacted, uncomplicated middle or lower and loweror middle ureteral stones which were ≤ 1cm and ≤0.9cm were enrolled in a prospective study and they were randomized into two groups. Group 1 received tamsulosin (0.4mg), and group 2 received silodosin (8mg) for a maximum period of 4 weeks. The patients were followed up weekly or biweekly with imaging studies. Results: Four patients in Group A and six patients in Group B were lost to follow-up, with 86 patients remaining for per-protocol analyses. No significant differences were found between the groups with respect to age, stone size, or stone location. Spontaneous stone expulsion was observed in 26 of 44 patients (59%) in Group A and in 34 of 42 patients (80%) in Group B (P=0.027). The primary endpoint was the stone expulsion rate, and the secondary endpoint was the time to stone expulsion. Stone expulsion rate was observed in 59% of patients in group 1 and in 80% of patients in group 2, which was statistically significant. There was also significant difference between groups with regard to mean time to stone expulsion. Conclusion: Silodosin was more effective than tamsulosin with regard to

stone expulsion rate and with a less mean time to stone expulsion.

Keywords—Ureteric Stone, Silodosin, Tamsulosin, Medical Expulsive Therapy.

I Introduction

Ureteral stones account for 22 % of all urinary tract stones with 68 % of them being located in the distal ureter. Urinary stones have afflicted humankind since antiquity, with the earliest literary quotations to stone disease, describing symptoms and prescribing treatments to dissolve the stone, are observed within the medical texts of Asutu in Mesopotamia between 3200 and 1200 BC [1]. Conservative management strategies such as observation or medical expulsive therapy (MET) using pharmacological agents to facilitate spontaneous passage of ureteral stones have gained popularity in the management of ureteral stones during the recent years. Evidence on the association of stone size with spontaneous stone passage rates is scarce. Ureteral stones occupy an important place in daily urological practice, that causing an acute attack of ureteral colic by obstructing the ureter [2]. Of all urinary tract stones, ureteral stones are 20% and 70% of these stones are located in the distal portion of the ureter [3]. There has been a paradigm shift in the management of the ureteral calculi in the past decade, with the introduction of minimally invasive techniques and newer drugs (4). An excellent results with recent studies have reported with the medical expulsion therapy for the distal ureteral calculi, with alpha 1 blockers [5].

Treatment Methods Of Ureterolithiasis:

1. Observation (also termed "watchful waiting" and "expectant management").

- 2. Medical expulsive therapy (MET).
- 3. Shock wave lithotripsy (SWL).
- 4. Ureteroscopy (URS).

5. Percutaneous Antegradeureteroscopy (PAURS).

6. Laparoscopic surgery.

7. Open surgery [6, 7].

Indications For Active Removal Of Ureteral Stones:

1. Stones with low likelihood of spontaneous passage {for example: stones associated with ureteric stricture, stones <1 cm} [6].

2. Persistent pain despite adequate analgesic medication [7].

3. Persistent obstruction [7].

4. Renal insufficiency (bilateral obstruction, or single kidney, renal failure) [6].

5. The patient's employment (machinery, pilots) [4].

Medical Expulsive Therapy (MET):

Patients who have newly diagnosed ureteric stones who have no indication for active removal of ureteric stones (listed above) and of less than 10 mm in size, may be offered appropriate medical therapy to facilitate stone passage during observation (Medical Expulsive Therapy) i.e. {the administration of drugs to facilitate stone passage} [6]. There is growing evidence that (MET) can be efficacious [6]. And the use of (MET) has become an accepted practice [2]. Meta-analyses have shown that patients with ureteral stones mange with nifedipineor a-blockers are more likely to pass stones with less episodes of ureteiccolic than those not tacking such therapy [5, 8]. Tamsulosin, An adrenergic antagonist (α_{1A} , α_{1D}) is effective medical agents and the most popular one, which is used for the expulsive therapy (probably because of its lack of a need for dose titration upon initiation of treatment and the excellent tolerability) [6]. Silodosin. recently introduced selective а α1Aadrenoceptor antagonist, has shown promising results with fewer side effects and a better efficacy [9].

Factors Affecting Medical Expulsion Therapy:

1. Stone size: MET is less likely to increase the stone-free rate, due to the high likelihood of spontaneous passage of stones up to ~5 mm, [10-13].

2. Stone location: The vast majority of trials have investigated distal ureteral stones [4]. One randomized clinically controlled trial (RCT) has assessed the effect of tamsulosin on spontaneous passage of proximal ureteral calculi 5-10 mm [14]. The main effect was to encourage stone migration to a more distal part of the ureter 14].

3. Medical expulsive therapy after extracorporeal shock wave lithotripsy (SWL): Clinical studies and several meta-analyses have shown that MET after SWL for ureteral or renal stones can expedite expulsion and increase stone free rate and reduce analgesic requirements [15-17].

4. Medical expulsive therapy after ureteroscopy: Medical expulsive therapy following holmium: YAG laser lithotripsy increases stone free rates and reduce colic episodes [18]. 5. Duration of medical expulsive therapy treatment: Most studies have had duration of 30 daysor1 month [19].

Alpha 1 adrenoceptors (ARs) are a class of proteins belonging to the G protein-coupled receptor familv [20]. Molecular heterogeneity in aladrenoceptors has been widely documented by gene cloning technologies and three different subtypes have been cloned, according to the indications of the International Union of Pharmacology, pharmacologically characterized and named α_{1A} , α_{1B} , and α_{1D} [21]. The distribution of these α1 adrenoceptors in human ureter was studied using quantitative real-time PCR and $\alpha 1$ adrenoceptors was found that each ureteral region was endowed with mRNA encoding α_1 adrenoceptors subtypes, although with differences in terms of the amount expressed and receptor distribution [21]. The α_{1A} subtype accounted for about 38% of total adrenoceptors [22]. The α_{1D} subtype mRNA was highly expressed in each ureteral accounting for about 54% of total region. adrenoceptors mRNA [22]. The α_{1B} subtype accounted for about 8% [21]. In the proximal and middle ureter, the distribution of adrenoceptors was α 1D $\geq \alpha$ 1A> α 1B, like that of the total ureter [22]. The α_{1D} subtype expression was significantly higher than the α_{1A} subtype expression. In the distal ureter, the distribution of adrenoceptors was $\alpha 1D > \alpha_{1A} > \alpha_{1B}$ [22].The distal ureter expressed the highest amount of αı Dadrenoceptors subtype [22]. Alpha_{1A}adrenoceptors that is primarily located in the human prostate, bladder base, bladder neck, prostatic capsule and prostatic urethra. Silodosin is a highly selective for these receptors. Blockade of these alpha1A-adenoceptors causes smooth muscle in these tissues to relax [23]. Silodosin has been demonstrated in vitro that the alpha_{1A}: alpha_{1B} binding ratio of silodosin is (162:1) which is extremely high [24]. It has a substantially lower affinity for alpha1Badrenoceptors that are primarily located in cardiovascular system (23). Tamsulosin exhibits selectivity for both alpha_{1A} and alpha_{1D} receptors over the alpha1Badrenoceptor subtype [26]. These three AR subtypes have a distinct distribution pattern in human tissue (22). Tamsulosin hydrochloride is an alpha1adrenoceptor (AR) blocking agent used for the treatment of lower urinary tract symptoms (LUTS) associated with benign prostatic hyperplasia (BPH) [25]. Whereas approximately 70% of the alpha1receptors in human prostate are of the alpha1A subtype, the human bladder contains predominantly the alpha_{1D} subtype while blood vessels express predominantly alpha_{1B} subtype [27].

II Patients And Methods

A prospective and randomized controlled study was conducted in the department of Urology Ibn Sina Medical College, Dhaka, Bangladesh. Starting from October 2018, till September 2019; a total of 96 patients (F=56; M=40) who were in the age group (17–60) years, and hadnon–impacted, unilateral, uncomplicated loweror middle ureteral stones which were ≤0.9cm, were enrolled in a prospective study and they were randomized into two groups. Group 1 received tamsulosin (0.4mg), and group 2 received silodosin (8mg) for a maximum period of 4 weeks. The patients were followed up weekly or biweekly with imaging studies.

The Study Exclusion criteria:

- 1. Diabetes Mellitus.
- 2. Urinary tract infection.
- 3. Severe hydronephrosis.
- 4. Hypotension.
- 5. Ureteral strictures.
- Multiple stones. 6.
- Solitary kidney. 7.
- Current use of any type of alpha-blocker. 8.
- Asthma and gastrointestinal ulcers. 9

10. Stones larger than 10 mm in greatest dimension.

The sample size of the study was arbitrarily determined. The patients were diagnosed by unenhanced computed tomography (CT) scans and re-evaluated with ultrasonography, plain X-ray and unenhanced (CT) scans whenever they were necessary. The stone size was calculated on the CT scan by using a digital ruler and the greatest dimension of the stone was taken into consideration as the stone size.

All the patients provided informed written consents and they were properly informed about the study in which they would be enrolled. The patients were randomly allocated into two treatment groups of 48 patients each. The patient demographics in the two groups, in terms of the size of the stones in the two groups, their locations in terms of the laterality and their locations in the ureter. Group A received tamsulosin (0.4 mg) daily, whereas Group B received silodosin (8mg) daily, for a maximum period of 6 weeks. All the patients were prescribed the 50 mg diclofenac tablet on demand for pain relief. The patients were advised that on experiencing an episode of unbearable ureteric colic, they should Table 1: Demographic data of the two study groups. immediately report to us. The patients were followed up weekly or 3 times weekly with X-rays of the abdomen and the pelvis and ultrasonography. The patients were instructed to record the time and date of the stone passage. The follow up continued until the stone spontaneously passed, as reported by the patient, or for a maximum period of 6 weeks. The primary endpoint was the stone expulsion rate and the secondary endpoints were the stone expulsion time. The stone expulsion rate was defined as the percentage of patients that spontaneously pass their stones within the follow up period (i.e.6 weeks), whereas the stone expulsion time was defined as the number of days from the random allocation to the stone expulsion.

Data Analysis: The statistical analysis was performed by using the Student's t-test to compare continuous variables between the two groups, and the Chi-square test was used for categorical variables. A p value of < 0.05 was considered to be statistically significant.

III Results

Four patients in Group A and six patients in Group B were lost to follow-up, with 86 patients remaining for per-protocol analyses. No significant differences were found between the groups with respect to age, stone size, or stone location (Table 1). Spontaneous stone expulsion was observed in 26 of 44 patients (59%) in Group A and in 34 of 42 patients (80%) in Group B (P=0.027). The stone expulsion rate was significantly higher in Group B than in Group A. There was also a significant difference between the groups with regards to the mean stone expulsion time (p=0.01). The mean expulsion time was 19.5 ± 7.5 days in Group A vs. 12.5 ± 3.5 days in Group B (Table 2). In table (1) we notice that despite the random allocation of the patients into the two treatment groups, the difference in stone size, stone location, sex of the patients and laterality was not significant; meaning that the difference in these variables is negligible, and there was no bias in patients' randomization.

NO. of patients	Group A (n=44)	Group B (n=42)	P value	
	(Tamsulosin)	(Silodosin)		
Sex:			0.47	
Male	25	27		
Female	19	15		
Mean age± SD(years)	37±11	35±10	0.22	
Mean stone size±SD (mm)	6.9±1.9	7.0±2.1	0.51	
Stone location:			0.49	
Left	23	25		
Right	21	17		
Stone position:			0.19	
Lower ureter	18	23		
Mid ureter	26	19		
None of the differences are statistically significant.				

Table 2. Results according to treatment				
Endpoint	Group A (n=44)	Group B (n=42)	P value	
	(Tamsulosin)	(Silodosin)		
Primary end point: Stone expulsion rate	26/44 (59%)	34/42 (80%)	0.027	
Secondary end point: Time to stone expulsion (days).	19.5±7.5	12.5± 3.5	0.01	
Both differences are statistically significant				

Table 2: Results according to treatment.

IV Discussion

Ureteral colic, which is mainly due to stones, represents 1 to 2% of the emergency room admissions [28]. With the introduction of effective medical therapeutic agents in the market, there has been a significant improvement in the medical management of the ureteral calculi [5, 8, 12]. Several studies findings indicate that alpha blockers facilitate ureteral stone passage while nifedipine may provide a marginal benefit [6]. These have demonstrated that this approach may facilitate and accelerate the spontaneous passage of ureteral stones [2, 5, 15]. Similar findings have been reported by Hollingsworth and associates, who recently performed a metaanalysis of studies involving alpha blockers or nifedipine in patients with ureteral stones [29]. The likelihood of a ureteral stone passage is dependent on several factors, which include the stone size and the location and the ureteral conditions [12-14]. Ibrahim AI et al. has demonstrated that stone passage rates between 71-98% for the distal ureteral stones which are less than 5 mm and from 25-53% for those which are between 5 and 10 mm [30]. The role of adrenergic receptors in the human ureter was first described in 1970 [31]. It was shown later, that the alphaadrenergic receptors were classified into three different subtypes of α_{1A} , α_{1B} and α_{1D} , of which the distribution in the human ureter was $\alpha_{1D} > \alpha_{1A} > \alpha_{1B}$ [22]. It was also shown that the alpha-adrenergic receptor agonists had a stimulatory effect on the ureteral smooth muscle, whereas the beta-adrenergic receptor agonists had an inhibitory effect [32]. The alphaadrenergic receptor agonists prevent the uncoordinated muscle activity which is seen in renal colic, while maintaining ureteral peristalsis, which might facilitate a spontaneous stone passage [33]. The alpha blockers mainly produce relaxation of the distal human ureter by reducing the ureteric smooth muscle tone rather than completely ablating its activity [33]. Two meta-analyses provided a high level of evidence for the clinical benefit of the alpha blockers in the patients with distal ureteral calculi, in which the patients who were given alpha blockers had 52% and 44% greater likelihoods of stone passage than those who were not given such treatment [29, 34]. The treatment effect on the expulsion rate was partially lost, as the sizes of the stones decreased, because of the high spontaneous expulsion rate of the small stones (4). By way of example only, De Sioet al., Wang et al., and Yilmazet al. reported better stone expulsion rates (81%, 79%, AND 90%, respectively) in patients who received 0.4 mg tamsulosin daily than in controls (54%, 53%, AND 58%, respectively) [35-37]. Although most of the studies used tamsulosin, which is a selective α_{1A}/α_{1D} adrenergic receptor antagonist, the efficacies of the other alpha blockers such as doxazosin, terazosin, alfuzosin and naftopidil were also indicated [36, 38, 39]. Wang et al., Yilmazet al., and Agrawalet al. demonstrated the efficacy of a1adrenoceptor antagonists in the management of lower ureteral stones regardless of the type of alpha-blocker used [36, 37, 40]. Many studies have been published on α 1-adrenoceptors in the human ureter since the first report in 1970, Malinet al. first described the presence of α - and β -adrenoceptors through the entire length of the human ureter and the physiological response (increased tone and frequency of contractions) of the ureter when exposed to a adrenoceptor agonists [31]. In 2005, Sigalaet al. found that α_{1D} - and α_{1A} -adrenoceptors were expressed in significantly larger amounts than α_{1B} -adrenoceptors in the human ureter, and these authors also demonstrated that the distal ureter expressed a greater amount of g1-adrenoceptor mRNA than the proximal and middle ureter [41]. Itohet al. reported that α_{1D} -adrenoceptor mRNA is more highly expressed than α_{1A} -adrenoceptor mRNA in each region of the ureter [33]. According to their results, a α_{1D} -adrenoceptor blocker can be expected to be more effective for the expulsion of ureteral stones than a α_{1A} -adrenoceptor blocker 41]. [22, However, Tomiyamaet al. reported that, in the hamster ureter, ureteral contraction was mediated mainly by α_{1A} adrenoceptors, even though a1Dadrenoceptors were more prevalent [42]. Recently, it was found that α_{1A} adrenoceptors is the main participant in phenylephrine-induced ureteral contraction in the human isolated ureter [43]. Our results indicate that a α_{1A} -adrenoceptor blocker is more effective than a α_{1D} adrenoceptor blocker with respect to stone expulsion rate and the time to stone expulsion suggesting more clinical usefulness of α_{1A} -adrenoceptor blockers. Silodosin was approved for BPH by the US Food and Drug Administration in October 2008 [44]. Silodosin is a highly selective α_{1A} -adrenoceptor antagonist, which has 56-fold affinity for α_{1A} -over α_{1D} -adrenoceptors [33]. Our study has compared the efficacy between tamsulosin and silodosin and our results are also very encouraging with stone expulsion rate of (80%) in group B who received silodosin (8mg) compared to (59%) of group A who received tamsulosin (0.4 mg) which was a significant difference (P value=0.027). Regarding the incidence of the retrograde ejaculation,

which is the most common side effect of silodosin (which has been stated to be very common among other side effects) [45-49], there has been a consensus among many urologists, that its occurrence should be considered as a sign of the efficacy, rather than an adverse effect of the treatment [45]. Silodosin appears to relax the smooth muscles of the genital tract and the lower urinary tract enough to induce a retrograde ejaculation [46]. This was reflected in the finding that the patients who had the greatest relief from the lower urinary tract symptoms had a higher likelihood of the retrograde ejaculation (46). This observation suggests that the retrograde ejaculation is actually an indirect indicator of the relaxation of the smooth musculature that induced by silodosin [48]. The advantage of the medical expulsive therapy is important, because the risks which are related to a surgical intervention are not trivial [50]. Studies have reported the overall complication rates after ureteroscopic lithotripsies to be 11-22%, with major complications such as ureteral perforations, avulsions and strictures occurring during 4-6% of the procedures [50]. Urinomas and sub capsular bleeds have been reported in16-33% of the patients who are treated with shock wave lithotripsy (ESWL) (51). Therefore the medical expulsive therapy should be offered as a cost-effective treatment for the patients with distal ureteral calculi, who are amenable to a waiting management. Limitations encountered during study were: (1) Relatively small sample size, and (2) The cost of silodosin was much higher than any available alpha blocker.

V Conclusions

From this study we identified that: (1). Silodosin (as an example of a selective α_{1A} -adrenoceptor antagonist) was more effective than tamsulosin (as an example of a α_{1D} and α_{1A} -adrenoceptor antagonist) with respect to stone expulsion rate for ureteral stones and the time to stone expulsion, despite the abundance of α_{1D} -adrenoceptors in human ureter. (2). A conservative approach should be considered as an option in the management of the uncomplicated, small, distal ureteral calculi.

VI Recommendations

We recommend the use of silodosin in the medical expulsive therapy for ureteric stones, since it is clinically superior to tamsulosin in this type of therapy. Further studies on medical expulsive therapy for ureteric stones, are required to determine the superiority of α_{1A} adrenoceptor antagonist (silodosin) versus α_{1D}/α_{1A} adrenoceptor antagonist (tamsulosin). These studies should include larger sample size.

References:

1. Shah J, Whitfield HN. Urolithiasis through the ages. BJU International 2002; 89(8): 801–810.

2. Resim S, Ekerbicer H, Ciftci A. Effect of tamsulosin on the number and intensity of ureteral

colic in patients with lower ureteral calculus. Int. J Urol. 2005; 12: 615–20.

3. Phillips E, Kieley S, Johnson EB, et al. Emergency room management of ureteral calculi: current practices. J Endourol 2009 Jun; 23(6):1021-4.

4. Seitz C, Liatsikos E, Porpiglia F, et al. Medical Therapy to Facilitate the Passage of Stones: What Is the Evidence? EurUrol 2009 Sep; 56(3):455-71.

5. Porpiglia F, Ghignone G, Fiori C, Fontana D, Scarpa RM. Nifedipine versus tamsulosin for the management of lower ureteral stones. J Urol. 2004; 172:568-71

6. Preminger GM, Tiselius HG, Assimos DG, et al. American Urological Association Education and Research, Inc; European Association of Urology. 2007 Guideline for the management of ureteral calculi.EurUrol 2007 Dec; 52(6):1610-31.

7. Miller OF, Kane CJ. Time to stone passage for observed ureteral calculi: a guide for patient education. J Urol 1999 Sep; 162(3 Pt 1):688-90; discussion 690-1.

8. Liatsikos EN, Katsakiori PF, Assimakopoulos K, et al. Doxazosin for the management of distalureteral stones. J Endourol 2007 May; 21(5):538-41.

9. "Drugs.com, Watson Announces Silodosin NDA Accepted for Filing by FDA for the Treatment of Benign Prostatic Hyperplasia". Retrieved 2008-02-13.

10. Ferre RM, Wasielewski JN, Strout TD, et al. Tamsulosin for ureteral stones in the emergency department: a Randomized controlled trial. Ann Emerg Med 2009 Sep; 54(3):432-9.

11. Hermanns T, Sauermann P, Rufibach K, et al. Is there a role for tamsulosin in the treatment of distal ureteral stones of 7 mm or less? Results of a randomized, double-blind, placebo-controlled trial.EurUrol 2009 Sep;56(3):407-12.

12. Vincendeau S, Bellissant E, Houlgatte A, et al; Tamsulosin Study Group. Tamsulosin hydrochloride vs placebo for management of distal ureteral stones: a multicentric, randomized, doubleblind trial. Arch Intern Med 2010 Dec 13; 170(22):2021-7.

13. Ochoa-Gómez R, Prieto-Díaz-Chávez E, Trujillo-Hernández B, et al. Tamsulosin does not have greater efficacy than conventional treatment for distal ureteral stone expulsion in Mexican patients. Urol Res 2011 Dec; 39(6)491-5.

14. Yencilek F, Erturhan S, Canguven O, et al. Does tamsulosin change the management of proximally located ureteral stones? Urol Res 2010 Jun; 38(3):195-9.

15. Gravina GL, Costa AM, Ronchi P, et al. Tamsulosin treatment increases clinical success rate of single extracorporeal shock wave lithotripsy of renal stones. Urology 2005 Jul; 66(1):24-8. 16. Naja V, Agarwal MM, Mandal AK, et al. Tamsulosin facilitates earlier clearance of stone fragments and reduces pain after shockwave lithotripsy for renal calculi; results from an open-label randomized study. Urology 2008 Nov; 72(5):1006-11.

17. Bhagat SK, Chacko NK, Kekre NS, et al. Is there a role for tamsulosin in shock wave lithotripsy for renal and ureteral calculi? J Urol 2007 Jun; 177(6):2185-8.

18. John TT, Razdan S. Adjunctive tamsulosin improves stone free rate after ureteroscopic lithotripsy of large renal and ureteric calculi: a prospective randomized study. Urology 2010 May; 75(5):1040-2.

19. Wen CC, Nakada SY. Treatment selection and outcomes: renal calculi. UrolClin North Am 2007 Aug; 34(3):409-19.

20. McCune DF, Edelmann SE, Olges JR et al. Regulation of the cellular localization and signaling properties of the alpha (1B)- and alpha (1D)adrenoceptors by agonists and inverse agonists. Mol. Pharmacol. 2006; 57: 659

21. Hieble JP, Bylund DB, Clarke DE et al. International union of pharmacology. X. Recommendation for nomenclature of alpha 1adrenoceptors: consensus update. Pharmacol. Rev.1995; 47: 267

22. Itoh Y, Kojima Y, Yasui T, Okada A, Tozawa K, Kohri K. Examination of alpha 1 adrenoceptor subtypes in the human ureter. Int. J. Urol.2007; 14: 749–53.

23. Kawabe K, Yoshida M, and Homma Y et al. Silodosin, a new alpha1A-adrenoceptor-selective antagonist for treating benign prostatic hyperplasia: results of a phase III randomized, placebo-controlled, double-blind study in Japanese men. BJU Int. 2006; 98:1019-24.

24. Anon. KMD 3213: KAD 3213, Silodosin. Drugs R D. 2004; 5:50-1.

25. Eri LM, Tveter KJ. A-blockade in the treatment of symptomatic benign prostatic hyperplasia. J Urol. 1995; 154:923-34.

26. Chapple CR, Wyndaele JJ, Nordling J et al et al. Tamsulosin, the first prostate-selective α 1A-adrenoceptor antagonist: a meta-analysis of two randomized, placebo-controlled, multicenter studies in patients with benign prostatic obstruction (symptomatic BPH). Eur Urol. 1996; 29:155-67

27. Faure C., Pimoule C., Vallancien G., Langer S.Z., Graham D. Identification of $\alpha 1$ -adrenoceptor subtypes present in the human prostate. Life Sci 1994; 54(21):1595-1605.

28. Galvin DJ, Pearle MS. The contemporary management of renal and ureteric calculi. BJU Int 2006 Dec; 98(6):1283-8.

29. Hollingsworth JM, Rogers MA, Kaufman SR et al. Medical therapy to facilitate urinary stone passage: a meta-analysis. Lancet.2006; 368: 1171-79.

30. Ibrahim AI, Shetty SD, Awad RM, Patel KP. Prognostic factors in the conservative treatment of ureteric stones.Br. J. Urol. 1991; 67: 358-61.

31. Malin JM Jr, Deane RF, Boyarsky S. Characterisation of adrenergic receptors in human ureter.Br. J. Urol. 1970; 42: 171-74.

32. Weiss RM, Bassett AL, Hoffman BF. Adrenergic innervation of the ureter. Invest. Urol. 1978; 16: 123–7.

33. Tzortzis V, Mamoulakis C, Rioja J, Gravas S, Michel MC, de la Rosette JJ. Medical expulsive therapy for distal ureteral stones.Drugs.2009; 69: 677-92.

34. Parsons JK, Hergan LA, Sakamoto K, Lakin C. Efficacy of alpha-blockers for the treatment of ureteral stones. J. Urol. 2007; 177: 983-87.

35. De Sio M, Autorino R, Di Lorenzo G et al. Medical expulsive treatment of distal ureteral stones using tamsulosin: a single-center experience. J. Endourol.2006; 20: 12–16.

36. Wang CJ, Huang SW, Chang CH. Efficacy of an alpha 1 blocker in expulsive therapy of lower ureteral stones. J. Endourol.2008; 22: 41–6.

37. Yilmaz E, Batislam E, Basar MM, Tuglu D, Ferhat M, Basar H. The comparison and efficacy of 3 different alpha1-adrenergic blockers for distal ureteral stones.J.Urol. 2005; 173: 2010–12.

38. Pedro RN, Hinck B, Hendlin K, Feia K, Canales BK, Monga M. Alfuzosin stone expulsion therapy for distal ureteral calculi: a double-blind, placebo controlled study. J. Urol. 2008; 179: 2244-47.

39. Sun X, He L, Ge W, Lv J. Efficacy of selective alpha1D-blocker naftopidil as medical expulsive therapy for distal ureteral stones. J. Urol. 2009; 181: 1716-20.

40. Agrawal M, Gupta M, Gupta A, Agrawal A, Sarkarl A, Lavania P. Prospective randomized trial comparing efficacy of alfuzoisin and tamsulosin in management of lower ureteral stones.Urology2009; 73: 706–9.

41. Sigala S, Dellabella M, Milanese G et al. Evidence for the presence of alpha 1 adrenoceptor subtypes in the human ureter. Neurourol. Urodyn.2005; 24: 142–8.

42. Tomiyama Y, Kobayashi K, Tadachi M et al. Expressions and mechanical functions of alpha 1 adrenoceptor subtypes in hamster ureter. Eur. J. Pharmco.2007; 573: 201–5.

43. Sasaki S, Tomiyama Y, Kobayashi S, Kojima Y, Kubota Y, Kohri K. Characterization of α (1)-adrenoceptor subtypes mediating contraction in

human isolated ureters. Urology 2011; 77: 762 e13-17.

44. Cantrell MA, Bream-Rouwenhorst HR, Steffensmeier A. et al. Intraoperative floppy iris syndrome associated with alpha1-adrenergic receptor antagonists. Ann Pharmacother. 2008; 42:558-63.

45. Kobayashi K, Masumori N, Kato R, et al. Orgasm is preserved regardless of ejaculatory dysfunction with selective alpha1A-blocker administration. Int J Impot Res 2009; 21:306–10

46. Kobayashi K, Masumori N, Hisasue S, et al. Inhibition of seminal emission is the main cause of anejaculation induced by a new highly selectivea1Ablocker in normal volunteers. J Sex Med 2008; 5:2185–90.

47. Morganroth J, Lepor H, Hill LA, Volinn W, Hoel G. Effects of the selective alpha (1A)adrenoceptor antagonist silodosin on ECGs of healthy men in a randomized, double-blind, placebo- and moxifloxacin-controlled study. ClinPharmacolTher 2010; 87: 609–13. 48. Tatemichi S, Kiguchi S, Kobayashi M, Yamazaki Y, Shibata N, Uruno T. Cardiovascular effects of the selective alpha1A-adrenoceptor antagonist silodosin (KMD-3213), a drug for the treatment of voiding dysfunction. Arzneimittelforschung 2006; 56:682–7.

49. Guimaraes S, Moura D. Vascular adrenoceptors: an update. Pharmacol Rev 2001; 53:319–56.

50. Picozzi SC, Ricci C, Gaeta M, et al. Urgent ureteroscopy as first-line treatment for ureteral stones: a meta-analysis of 681 patients. Urol Res 2012 Oct; 40(5):581-6.

51. Lotan Y, Gettman MT, Roehrborn CG, Cadeddu JA, Pearle MS. Management of ureteral calculi: a cost comparison and decision making analysis. J. Urol. 2002; 167:1621-29.