Does silodosin offer better results than tamsulosin as medical expulsive treatment after shock wave lithotripsy for single distal ureteric stones?

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Abstract

Introduction: Different antagonists of α-adrenergic receptors (α-blockers) have been used as medical expulsive treatment (MET) after extracorporeal shock wave lithotripsy (ESWL).

Aim: To retrospectively evaluate the expulsion rate of fragments after extracorporeal shock wave lithotripsy performed for single ureteral stones followed by different medical expulsive treatments.

Material and methods: We retrospectively analyzed stone expulsion rates of 190 patients treated by shock wave lithotripsy (SWL) for single, 5 to 10 mm, symptomatic and uncomplicated distal ureteric stones, treated with tamsulosin 0.4 mg, silodosin 8 mg or silodosin 4 mg as MET. Beside the stone-free rate after 4 weeks of treatment, we also investigated the pain intensity using the visual analogue scale (VAS), adverse events induced by the medication, safety of drug administration and the reasons for possible early treatment discontinuation.

Results: Silodosin 8 mg and tamsulosin 0.4 mg have similar results in terms of stone-free rate. For silodosin 4 mg the stone-free rate was significantly lower than for the previous two drugs. In patients treated with silodosin 4 mg the VAS was significantly higher than in patients treated with silodosin 8 mg or tamsulosin 0.4 mg, for all the follow-up visits.

Conclusions: Alpha-blocker treatment after ESWL with silodosin 8 mg offers a similar stone-free rate compared with tamsulosin 0.4 mg, being well tolerated. A lower dose of silodosin (4 mg) has significantly poor results, irrespective of ureteric stone size, with more frequent renal colic and severe pain.

Key words: extracorporeal shock wave lithotripsy, expulsion rate, α-blockers.
of the patients for distal ureteric stones of 5 to 10 mm diameter (or even higher if the follow-up period is prolonged) but we need to consider the possible complications that may occur, from bothersome symptoms to urinary infection or renal dysfunction [7]. In fact, “watchful waiting” is not an option for such patients, various MET schemes, SWL, URS, or diverse combinations being recommended [8–10].

Tamsulosin is the most commonly used α-blocker in MET. Within the last years, silodosin – an α-blocker – has also been used instead of tamsulosin as MET. However, there are few studies comparing these substances for MET, and none comparing their role as MET after SWL.

Aim

The aim of this study was to retrospectively evaluate the expulsion rate of the ureteric stone fragments after SWL in patients with a single distal ureteric stone, which underwent SWL and one of the following adjuvant therapies: (A) tamsulosin 0.4 mg once daily, (B) silodosin 8 mg once daily, or (C) silodosin 4 mg once daily, for a treatment period of up to 4 weeks. All patients received the NSAID lornoxicam as an additional analgesic. Beside the major aspect evaluated, i.e. success rate up to 4 weeks of treatment, we also investigated the following: pain intensity (using the visual analogue scale), adverse events induced by the medication, safety of drug administration and the reasons for possible early treatment discontinuation.

Material and methods

This retrospective study was performed in a university, tertiary stone center and evaluated the records of 190 adult patients with single distal radiopaque ureteric stones (on Kidney Ureter Bladder X-ray study (KUB)) treated by ESWL (single SWL session, 4000 shock waves applied at a rate of 1/s) by the same experienced urologist. The indication for SWL was represented by recurrent renal colic non-responsive to medical treatment with NSAID. Only the record files of the patients who met the inclusion criteria (normal renal function – as determined by serum creatinine levels – and single distal ureteric stone, measuring 5 to 10 mm diameter in maximum size) were evaluated. We excluded from the evaluation patients with concomitant or previous β-blocker (antagonist of β-adrenergic receptors) treatment, pre-existing treatment with an α-blocker for a prostatic disease, allergy/intolerance to α-blockers or NSAID, single kidney (surgical, congenital, or functional), JJ stent inserted prior to SWL, bilateral ureteric stones or confirmed infection of the urinary tract.

This retrospective study obtained the review board approval, and met all the local and national ethical requirements applying to the study type, being conducted in accordance with the ethical standards laid down in the Declaration of Helsinki. As the α-blocker treatment is an off-the label treatment for MET, informed consent for all the patients was obtained.

For statistical analysis, we defined three study groups, according to the medical expulsive treatment administered after the SWL sessions, as follows: group A – patients who received tamsulosin 0.4 mg/daily, group B – patients who received silodosin 8 mg/daily, and group C – patients treated with silodosin 4 mg/daily. The local protocol involved the treatment of all the patients (additional analgesic treatment) with lornoxicam 8 mg twice a day during the first week and then once a day until stone fragments’ elimination or up to 28 days. Each patient received information concerning the daily water intake and had to fill out a questionnaire regarding: pain intensity on the visual analogue scale, adverse events. As there is no consensus regarding the scheme for MET, the choice of α-blocker type and dose was made randomly by the urologist at the time of SWL. The patients were followed up weekly (weeks 1–3 for adverse events and pain evaluation, the 4th evaluation for evaluation of the stone-free status).

Therapeutic success was defined as complete lack of fragments detected on unenhanced CT at 4 weeks after the procedure – as it appeared on patients’ record files.

The stone-free rate after 4 weeks in each group was used to compare the effectiveness of the three medical treatments (either of them was adjuvant for stone expulsion after the initial SWL). Paired comparisons were made of the stone-free rates between the groups, e.g. A-B, A-C, and B-C. For each comparison the χ² test was applied and 0.05 was considered the statistical significance threshold.

To evaluate the effects of repeated measures and size, a generalized estimating equations (GEE) approach was applied by means of a logistic regression
for a repeated measures model. The GEE-based multivariate response profile approach is an analogue to repeated measures analysis of variance (repeated measures ANOVA and MANOVA) and multivariate analysis of covariance (MANCOVA). These more conventional longitudinal approaches require Gaussian response variables. In contrast, the GEE approach is well suited for skewed binary response variables (e.g. via a logarithmic link). In this model repeated measures of the response variable (success or failure of stone expulsion) in each subject were managed as clusters, while stone size was included as a factor.

Results

Patient age varied between 21 and 65, while the men/women ratio was 1.8 for the entire study. The groups were similar regarding the number of patients, sex distribution, age, stone size and degree of distension induced by the obstruction ($p > 0.05$ for each aspect; $\chi^2$ test) (Table I).

The stone-free rates for the three groups are presented in Table II.

One can see that silodosin 8 mg ensured a similar success rate in comparison with tamsulosin 0.4 mg ($p = 0.9083$), while silodosin 4 mg is significantly less effective than tamsulosin and silodosin 8 mg.

The results of the logistic regression model, transformed into odds ratios, are presented in Table III.

Regarding the renal colic episodes in the three groups, we found that patients in groups A and B reported less pain and required only sporadic administration of additional doses of pain killers (tramadol chloride) (Tables IV and V). One can see that in group C, the VAS was significantly higher than in groups A and B for all the follow-up visits.

In groups A and B, the visual analogue scale (VAS) for pain due to renal colic was seldom evaluated at

| Group | Stone size | Total | % |
|-------|------------|-------|---|
|       | 5 mm       | 6 mm  | 7 mm | 8 mm | 9 mm | 10 mm |
| A (tamsulosin 0.4 mg/day) | 17 | 19 | 8 | 8 | 4 | 5 | 61 | 32.1 |
| B (silodosin 8 mg/day) | 20 | 19 | 9 | 7 | 5 | 6 | 66 | 34.7 |
| C (silodosin 4 mg/day) | 21 | 16 | 9 | 7 | 5 | 5 | 63 | 33.2 |
| Total | 58 | 54 | 26 | 22 | 14 | 16 | 190 | 100.0 |

| Group | Stone size | Total % success | A vs. B vs. |
|-------|------------|-----------------|-------------|
|       | 5 mm | 6 mm | 7 mm | 8 mm | 9 mm | 10 mm | A vs. | B vs. |
| A | 100.0 | 100.0 | 100.0 | 87.5 | 75.0 | 60.0 | 93.4 | – | 0.9083 |
| B | 100.0 | 100.0 | 100.0 | 85.7 | 80.0 | 66.7 | 93.9 | 0.9083 | – |
| C | 100.0 | 93.8 | 66.7 | 57.1 | 60.0 | 40.0 | 81.0 | 0.0381 | 0.0253 |

| Test variables | Logistic regression model |
|----------------|--------------------------|
| A (tamsulosin 0.4 mg/day) vs. C (silodosin 4 mg/day) | B (silodosin 8 mg/day) vs. C (silodosin 4 mg/day) |
| OR (95% CI) | P-value | OR (95% CI) | P-value |
| Stone-free status | 2.3 (1.73–4.56) | 0.0004 | 2.41 (1.78–4.92) | < 0.0001 |
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Table IV. Required additional pain treatment in study groups

| Group | Patients who required and received additional doses of pain killers (tramadol chloride) after SWL |
|-------|--------------------------------------------------------------------------------------------------|
| A     | 2 patients                                                                                       |
|       | 1 patient – 100 mg/1 day                                                                         |
|       | 1 patient – 2 × 100 mg/1 day                                                                     |
| B     | 4 patients                                                                                       |
|       | 1 patient 100 mg/1 day                                                                           |
|       | 1 patient 2 × 50 mg/1 day                                                                         |
|       | 2 patients 2 days, 50 mg/1 day                                                                    |
| C     | 9 patients                                                                                       |
|       | 3 patients 1 day, 100 mg/1 day                                                                    |
|       | 2 patients 2 days, 100 mg/1 day                                                                    |
|       | 2 patients 2 days, 50 mg/1 day                                                                    |
|       | 2 patients 1 day, 50 mg/1 day                                                                    |

Table V. Mean visual analogue scale (VAS) values by study groups at 1, 2, 3 and 4 weeks follow-up

| Variable | Mean VAS Week 1 | Mean VAS Week 2 | Mean VAS Week 3 | Mean VAS Week 4 |
|----------|-----------------|-----------------|-----------------|-----------------|
| Group A  (tamsulosin 0.4 mg) | 6.2             | 4.1             | 4.2             | 3.2             |
| Group B  (silodosin 8 mg) | 5.9             | 4.3             | 4.1             | 3.1             |
| Group C  (silodosin 4 mg) | 7.5             | 5.1             | 4.8             | 4.1             |
| A vs. B (p-value)           | 0.031           | 0.25            | 0.32            | 0.27            |
| A vs. C (p-value)           | 0.038           | 0.026           | 0.047           | 0.039           |
| B vs. C (p-value)           | 0.026           | 0.032           | 0.041           | 0.037           |

8 points (4 patients in each group) or 9 points (1 patient in group A).

One patient from each of groups A and B came back to the emergency room suffering from pain, but both had symptoms relieved by pain killers. VAS of 8 was reported in 6 patients and VAS of 9 was reported in 3 patients.

From the 22 cases of failure reported after 4 weeks of treatment, in 19 cases another ESWL procedure was performed, in 1 case semirigid ureteroscopy was done at the patient’s choice, and in two symptomatic cases (from group C), without significant ureterohydronephrosis, the expulsion therapy with silodosin 8 mg was continued at the patient’s preference, stone-free status being reached in both cases after an additional 2 and 3 weeks respectively.

Discussion

To our knowledge this retrospective study appears to be the first aiming to compare the effects of tamsulosin 0.4 mg with silodosin 4 mg and 8 mg as medical expulsive treatment after SWL. A recent meta-analysis showed that all the previous published studies compared tamsulosin 0.4 mg with silodosin 8 mg, revealing higher expulsion rates for silodosin 8 mg, none of them evaluating the effect of lower dose silodosin [11]. Although controversies exist, α-blocker therapy is recommended by the European Association of Urology Guidelines as medical expulsive treatment, especially for distal ureteric stones, facilitating the stone passage due to the relaxation of the smooth muscle of the ureter [12].

α1A-, α1B- and α1D-ARs are the three types of α1 adrenoceptors, the most frequent being the α1D subtype, followed by α1A and α1B [13].

Tamsulosin is a uroselective α-blocker owing to its selectivity as an antagonist of α1A receptors. Tamsulosin 0.4 mg once daily was reported to increase the rate of ureteric stone expulsion [7, 14–18]. The existing literature is abundant in studies that recommend α-blockers in the scheme of expulsion treatment, motivated by the fact that the ureter contracted by stimulation of α1-adrenergic receptors may im-
pede elimination of the stone and stone fragments, while the use of α-blockers may allow an increase of urine flow by ureteric muscle relaxation, with the respective washout effect, but also by an increase of intra-ureteric pressure gradient around the stone, which may help stone expulsion [19]. There are data suggesting that similar results can be obtained with doxazosin or terazosin, by a class effect [12].

The affinity of silodosin for α1A receptors is 150 times higher than for α1D in comparison with tamsulosin, which is 100-fold higher; thus, we could, at least theoretically, expect better results [20]. As silodosin 4 mg can be successfully used for treatment of lower urinary tract symptoms, we decided to use it in order to evaluate its efficacy as MET.

Adrenergic receptors α1A and 1D are more frequent in the ureter than 1B, and α1D has the highest density in the distal ureter [21]. However, the literature regarding the impact of specific 1A vs. 1D antagonist is not very conclusive. Comparison between naftopidil (selective α1D antagonist) and silodosin 8 mg, having an affinity for α1A, which is 56 times higher than for α1D, revealed that inhibition of α1A receptors is more important for expulsion facilitation than that of α1D [22, 23]. Alpha 1D receptors are better represented than 1A in each ureteric region, so they could play the key role in stone expulsion [24].

Our results reveal similar expulsion rates for tamsulosin 0.4 mg and silodosin 8 mg. Meanwhile, it seems that silodosin 4 mg has inferior results in terms of stone-free rate, the stone-free odds ratio being 2.3 and 2.41 for tamsulosin 0.4 mg and silodosin 8 mg compared to silodosin 4 mg, respectively. Moreover, the pain score (VAS) was significantly higher in patients treated with silodosin 4 mg compared with the other two groups.

The strong points of this study are the homogeneity of the groups, the inclusion criteria, the single person performing the ESWL procedure (thus, biases related to the technique being avoided) and the simple follow-up schedule. Also, we acknowledge the relatively small size of the groups, the retrospective nature of the study and the lack of stone characterization by density (as only KUB was performed before the procedure), further larger studies being necessary.

Conclusions

α-Blocker treatment after ESWL with silodosin 8 mg offers a similar stone-free rate compared with tamsulosin 0.4 mg, being well tolerated. A lower dose of silodosin (4 mg) has significantly poor results, irrespective of ureteric stone size, with more frequent renal colic and severe pain.

Conflict of interest

The authors declare no conflict of interest.

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Received: 3.11.2019, accepted: 12.12.2019.