Alpha blockers for treatment of ureteric stones: systematic review and meta-analysis

John M Hollingsworth,¹ Benjamin K Canales,² Mary A M Rogers,³ Shyam Sukumar,⁴ Phyllis Yan,¹ Gretchen M Kuntz,⁵ Philipp Dahm⁴

¹Department of Urology, University of Michigan, 2800 Plymouth Rd, Building 16, 1st Floor, Ann Arbor, MI 48109, USA

²Department of Urology, University of Florida, 1600 SW Archer Rd, Gainesville, FL 32610, USA

³Department of Internal Medicine, Division of General Medicine, University of Michigan, 2800 Plymouth Rd, Building 16, 4th Floor, Ann Arbor, MI 48109, USA

⁴Minneapolis Veterans Administration Health Care System and Department of Urology, University of Minnesota, Mayo Memorial Building, 420 Delaware St SE, MMC 394, Minneapolis, MN 55455, USA

⁵Borland Library, University of Florida, 653-1 W 8th St, Jacksonville, FL 32209, USA

Correspondence to: J M Hollingsworth kinks@med.umich.edu

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ABSTRACT

OBJECTIVE

To investigate the efficacy and safety of alpha blockers in the treatment of patients with ureteric stones.

DESIGN

Systematic review and meta-analysis.

DATA SOURCES

Cochrane Central Register of Controlled Trials, Web of Science, Embase, LILACS, and Medline databases and scientific meeting abstracts to July 2016.

REVIEW METHODS

Randomized controlled trials of alpha blockers compared with placebo or control for treatment of ureteric stones were eligible. Two team members independently extracted data from each included study. The primary outcome was the proportion of patients who passed their stone. Secondary outcomes were the time to passage; the number of pain episodes; and the proportions of patients who underwent surgery, required admission to hospital, and experienced an adverse event. Pooled risk ratios and 95% confidence intervals were calculated for the primary outcome with profile likelihood random effects models. Cochrane Collaboration's tool for assessing risk of bias and the GRADE approach were used to evaluate the quality of evidence and summarize conclusions.

RESULTS

55 randomized controlled trials were included. There was moderate quality evidence that alpha blockers facilitate passage of ureteric stones (risk ratio 1.49, 95% confidence interval 1.39 to 1.61). Based on a priori subgroup analysis, there seemed to be no benefit to treatment with alpha blocker among patients with smaller ureteric stones (1.19, 1.00 to 1.48). Patients with larger stones treated with an alpha blocker,

however, had a 57% higher risk of stone passage compared with controls (1.57, 1.17 to 2.27). The effect of alpha blockers was independent of stone location (1.48 (1.05 to 2.10) for upper or middle stones; 1.49 (1.38 to 1.63) for lower stones). Compared with controls, patients who received alpha blockers had significantly shorter times to stone passage (mean difference -3.79 days, -4.45 to -3.14; moderate quality evidence), fewer episodes of pain (-0.74 episodes, -1.28 to -0.21; low quality evidence), lower risks of surgical intervention (risk ratio 0.44, 0.37 to 0.52; moderate quality evidence), and lower risks of admission to hospital (0.37, 0.22 to 0.64; moderate quality evidence). The risk of a serious adverse event was similar between treatment and control groups (1.49, 0.24 to 9.35; low quality evidence).

CONCLUSIONS

Alpha blockers seem efficacious in the treatment of patients with ureteric stones who are amenable to conservative management. The greatest benefit might be among those with larger stones. These results support current guideline recommendations advocating a role for alpha blockers in patients with ureteric stones.

SYSTEMATIC REVIEW REGISTRATION

PROSPERO registration No CRD42015024169.

Introduction

Contemporary practice guidelines from leading professional societies recognize the off-label use of alpha adrenergic antagonists (or alpha blockers) as an initial treatment option for patients with newly diagnosed, uncomplicated ureteric stones <10 mm in size, whose symptoms are controlled.³⁻⁹ This endorsement is based on several systematic reviews and meta-analyses of numerous randomized controlled trials, which, in aggregate, showed a higher risk of stone passage among patients treated with alpha blockers (their use has been termed medical expulsive therapy) compared with controls.³⁻⁹ Consequently, such treatment has become part of the routine management algorithm for ureteric colic.

Even ardent proponents of medical expulsive therapy concede that many supporting data come from small, single centre, low quality studies, and a large confirmatory trial has been recommended. This prompted a recent multicentre randomized controlled trial in the United Kingdom that involved over 1100 patients with ureteric stones.¹⁰ The trial showed this treatment to be no more efficacious than placebo at decreasing four week rates of intervention for stone clearance. In light of these results, the investigators concluded that medical expulsive therapy "should not

WHAT IS ALREADY KNOWN ON THIS TOPIC

Several small randomized controlled trials have suggested a role of alpha blockers in promoting passage of ureteric stones

Consequently, contemporary practice guidelines recommend their use as medical expulsive therapy

A recent large methodologically rigorous trial has raised questions about the efficacy of this treatment

WHAT THIS STUDY ADDS

Medical expulsive therapy is beneficial for multiple health outcomes such as passage of ureteric stone and need for surgical interventions; adverse effects associated with alpha blocker use were relatively infrequent and not severe

Medical expulsive therapy with alpha blockers seems of particular benefit in patients with larger ureteric stones

be offered to patients with ureteric colic managed expectantly, giving providers of health care an opportunity to reallocate resources elsewhere.”

Despite the rigor of this trial, concerns have been raised about the choice of primary endpoint and the possibility that other important data might have been overlooked (for example, the high background rate of spontaneous stone passage).^{11–14} Further, while intervention rates were similar between the treatment and placebo groups for smaller and upper/middle ureteric stones, results were consistent with a clinically important effect only in patients with larger, lower calculi.¹⁰ To help reconcile these issues, we conducted a systematic review, identifying all randomized controlled trials examining alpha blockers for treatment of ureteric stones. We then pooled data to derive estimates of the effect of alpha blockers on stone passage, including a priori subgroup analyses to assess the impact that stone size and location have on efficacy.

Methods

Data sources and searches

Following the PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) statement,¹⁵ we prospectively registered our review on PROSPERO (CRD42015024169). We established inclusion criteria before beginning our search. We considered all randomized controlled trials in any language that looked at alpha blockers compared with placebo or control for treatment of ureteric stones. Controls were defined a priori as patients who had not received any additional treatment to facilitate stone passage (such as antispasmodics, antimuscarinics) with the exception of corticosteroids and only if they were applied equally to both treatment arms. We included only those trials in which alpha blockers were used as the main treatment. Thus, we excluded trials in which alpha blockers were examined as an adjuvant to surgery (for example, after shockwave lithotripsy or ureteroscopy).

A Cochrane Collaboration systematic review identified all randomized controlled trials on alpha blockers for ureteric stones published up to 9 July 2012.⁶ This review served as the foundation for our search. One member of our study team, a trained medical librarian, performed an updated electronic search of the Cochrane Central Register of Controlled Trials (via Wiley), Web of Science, Embase, LILACS, and Medline (via PubMed) databases up to 10 July 2016. We used boolean logic to incorporate various terms and synonyms for concepts in each of three distinct filters: an anatomic filter for ureteric stones, a treatment filter for alpha blockers, and a publication type filter for randomized controlled trials. When possible, we used controlled vocabulary (such as MeSH in PubMed, Emtree in Embase) and key words. Although we tailored the precise strategy to accommodate each database's features, as an example appendix 1 shows the strategy we used in PubMed. Other search strategies are available on request.

In addition, we scanned the reference lists of other published narrative and systematic reviews to identify potential additional studies not retrieved by our electronic

search. In an effort to find unpublished studies, we also hand searched abstracts from the annual meetings of the World Congress of Endourology and SWL, the European Association of Urology, and the American Urological Association to 10 July 2016. To identify ongoing trials, we used the WHO International Clinical Trials Registry Platform (ICTRP) and ClinicalTrials.gov search portals.

Study selection

We used the reference manager software Endnote to identify and remove duplicate records. Next, we imported references into Covidence (www.covidence.org), and two study team members independently scanned each title and abstract. For studies that advanced beyond this stage, two study team members then performed independent full text reviews. We mapped publications relating to the same trial to unique studies. When we found multiple publications from a research group, we contacted the corresponding author to determine whether their reports were from the same study population, and we removed duplicates. We had non-English language articles translated before review. Once we identified all potentially relevant articles, two study team members met to achieve consensus. When necessary, we used third party adjudication to settle disputes.

Data extraction and risk of bias assessment

For each study selected for inclusion, two team members independently extracted data using a pilot tested standardized form. We resolved inconsistencies between the two through discussion, with a third team member serving as the arbitrator. We collected information on study characteristics (study year, country of origin, publication type, use of placebo, maximum length of follow-up, and imaging use), patients' characteristics (age, sex, and stone size and location), and data on outcomes. Our primary outcome was the proportion of patients who passed their stone. Our secondary outcomes were the time to passage, the number of episodes of pain, and the proportions of patients who underwent surgical intervention, required admission to hospital, and experienced a serious adverse event (as defined by the study authors). We also examined specific adverse events including dizziness, headache, fatigue, malaise, insomnia, hypotension, palpitations, collapse, retrograde/abnormal ejaculation, sexual dysfunction, dyspepsia, diarrhea, nausea, vomiting, constipation, flatulence, abdominal pain, nasal congestion, cough, arthralgia, and rash.

To assess the risk of bias of the selected studies, we used the Cochrane Collaboration's tool on an outcome specific basis.¹⁶ Relying only on the information presented in the study report and making no assumptions, two team members evaluated each trial across four domains. These domains included sequence generation/allocation concealment (selection bias), blinding of participants and personnel (performance bias), blinding of outcome assessors (detection bias), and completeness of follow-up (attrition bias). For each domain, individual team members judged whether the risk of bias in a given study was “low,” “high,” or “unclear.” Any disagreements were referred to a third team member.

Data synthesis and primary analysis

We conducted two tailed statistical tests and set the probability of type I error at 0.05. We performed all calculations using Stata/MP, version 14.1 (StataCorp, College Station, TX). In preliminary analyses, we pooled the proportion of patients who passed stones in each study group (treatment, control) using the random effects profile likelihood method and the Freeman-Tukey double arcsine transformation to stabilize the variance.¹⁷ Our main estimates of effect were pooled risk ratios with 95% confidence intervals. Specifically, we compared the proportion of people taking an alpha blocker who passed their stones (numerator) with the proportion not taking an alpha blocker (controls) who passed their stones (denominator). For these comparisons, we fitted random effects models and calculated 95% confidence intervals using two different estimators—the profile likelihood and restricted maximum likelihood methods.¹⁸

When the number of pooled studies was small (fewer than 10), we used the more conservative Hartung-Knapp-Sidik-Jonkman method to calculate confidence intervals,^{19–21} which is based on a *t* distribution. For studies that involved different alpha blockers (for example, tamsulosin and terazosin) or different doses of the same drug (for example, tamsulosin 0.2 mg and 0.4 mg), we collapsed the separate treatment arms into one for our main analysis. We computed pooled risk differences to generate number needed to treat (calculated as the inverse of the pooled risk difference) with 95% confidence intervals from the profile likelihood method. To determine whether an early extreme result for treatment with an alpha blocker deviated from the results of later studies (the Proteus phenomenon), we used the approach described by Ioannidis and Trikalinos.²² The prediction interval was calculated for the primary outcome (stone passage); this interval reflects the expected future benefits of treatment to patients.²³

Based on a priori decisions, we also performed subgroup analyses, stratifying by stone size and location. In some trials, the investigators reported outcomes in patients who had smaller versus larger ureteric stones; we used these size thresholds, ranging from 5 mm to 8 mm, to stratify the results. For our analyses of stone location, we collapsed upper and middle ureteric stones into one category.

To assess statistical heterogeneity, we calculated τ^2 , which represents variance between studies (τ is the estimated standard deviation of the effects across studies).²⁴ We also calculated the I^2 statistic, which estimates the proportion of variability in the meta-analysis caused by differences between studies rather than sampling error.²⁵ Prior empirical work suggests that τ^2 performs better than I^2 as precision increases.²⁶

Sensitivity analyses

To better understand the sources of statistical heterogeneity between studies, as well as test the robustness of our findings, we then performed a series of sensitivity analyses.

First, we repeated our analysis, excluding those studies in which corticosteroids were co-administered. Second, we restricted our analysis to only those studies

published as full length, peer reviewed research articles. Third, we pooled data only from those studies that had placebo controls. Fourth, to evaluate the impact that baseline risk had on our estimates, we conducted meta-regression using the restricted maximum likelihood method for variance between studies with the Knapp-Hartung modification to calculate P values and confidence intervals. We also used meta-regression to assess differences in effect related to length of follow-up and sex composition. Fifth, we performed sensitivity analyses based on variation between studies in the risk of bias. For each of the four domains described above, a study was awarded a point if the risk of bias was judged low. As such, a study could be awarded a maximum of four points. We categorized each study by its point totals as being of low (three or four points), moderate (two points), or high risk of bias (zero or one point). Finally, to examine the effect of individual studies on our summary estimates, we conducted an influence analysis, in which we recalculated pooled estimates omitting one study at a time.

Secondary analyses

In addition to our primary analysis, we derived summary estimates of effect for our secondary outcomes. When we used continuous scales of measurement (time to stone passage, number of pain episodes), we summarized our findings with the pooled mean difference. For dichotomous outcomes (need for surgical intervention, subsequent admission to hospital, occurrence of an adverse event), we again used pooled risk ratios.

Assessing for publication bias

We explored the possibility of publication bias visually, checking for asymmetry in contoured funnel plots of treatment effect against its standard error, and with formal statistical procedures. Specifically, we used Harbord's test for dichotomous endpoints and Egger's test for continuous endpoints to examine possible small study effects.²⁷

Patient involvement

No patients were involved in setting the research question or the outcome measures, nor were they involved in developing plans for design or implementation of our study. We did not ask patients for advice on interpretation or writing up of our results. We have no plans to disseminate our results to patient participants from the pooled studies.

Rating quality of evidence for each pooled analysis

Finally, we rated our confidence in the estimates of effect for each outcome according to the GRADE approach, taking into account study limitations (risk of bias), inconsistency, imprecision, indirectness and publication bias.²⁸ For each comparison, two team members independently rated the quality of evidence for each outcome as “high,” “moderate,” “low,” or “very low” using GRADEpro GDT (<http://grade.pro.org>). We resolved discrepancies by consensus, and if needed, with arbitration by a third team member.

Results

Search strategy results

From our electronic search, we identified 286 studies. We found an additional 20 records by hand searching meeting abstracts and from reference lists of other review articles. When we combined these with the 29 references from the 2014 Cochrane review,⁶ there were 443 potentially relevant studies in total. Of these, we excluded 355 during the initial screening phase based on the title and abstract. For the remaining 88 studies, we performed a full text screening and eliminated 33 studies because they included duplicate populations, had an observational design, or their treatment groups received additional treatment to facilitate stone passage. This yielded 55 unique studies (involving 5990 randomized patients) that examined the effect of alpha blockers on ureteric stones (fig 1) and met our inclusion criteria.^{10 29-82}

Descriptive statistics

Table A in appendix 2 shows all the studies included in our systematic review, most of which were conducted in Europe and Asia. Thirty nine studies reported mean age in treatment and control groups,^{10 29-32 34-36 39 40 43-50 52-56 58 61-65 67 70 72 75-79 82} which was 40.7 (SD 6.9) and 40.4 (SD 6.1), respectively. The percentage of women varied from 0% to 59.6%. Average stone size was reported by treatment group in 41 studies,^{10 29-32 34-36 39 40 43-56 58 60-67 69-72 75 76 78 79 82} with a mean of 5.7 (SD 1.2) mm in the treatment group and 5.7 (SD 1.1) mm in the control group. While most studies were limited to patients with lower ureteric stones, 11 studies included stones located in the upper and middle ureter.^{10 38 40 49 59 60 63 68 69 77 79}

Table B in appendix 2 shows the interventions and follow-up, as well as the primary and secondary outcomes and recorded adverse events, from all included studies. Most studies had two arms, but 10 had three arms,^{43 46 47 52 63 65 68 70 71 78} and two had four arms.^{34 38} Although tamsulosin was the most common intervention (40 studies),^{10 29 30 32-35 38-40 43 45-49 51-63 65-69 71 73-75 77 78 81} additional alpha blockers were evaluated, including alfuzosin (six),^{44 47 52 63 72 76} doxazosin (four),^{37 41 42 55}

naftopidil (three),^{50 65 70} silodosin (six),^{64 71 78 79 80 82} and terazosin (four).^{31 34 36 46} The duration of follow-up varied across studies from seven to 42 days, with 28 days being the most common (37),^{10 30 31 36 37 40-45 47 51 53-56 58-69 71 72 74 76-79 81} or until stone passage. Placebo controls were used in 14 studies.^{10 44 47 51 53 56-58 73-75 79 81 82} In three studies, corticosteroids were given to the intervention and/or control groups.^{30 65 68}

The baseline rate of stone passage without treatment with an alpha blocker varied across countries (fig A in appendix 3), at 7% in Thailand, 14% in Sri Lanka, 80% in the UK, and 82% in Australia.

Figure 2 summarizes the assessment of risk of bias for individual studies. Fifty two studies had at least one domain judged as unclear risk of bias, nine of which had at least one domain considered at high risk.^{32 43 55 60 63 69 75 76 82} Only three had all bias domains judged as low risk.^{10 53 79} Eight studies were explicit about the reporting of an appropriate method of allocation concealment,^{10 44 53 58 65 78 79 81} and only six studies reported blinding of outcome assessors.^{10 29 53 75 79 81}

Effect of alpha blockers on passage of ureteric stones

The random effects pooled risk ratio (and 95% confidence interval estimated with the profile likelihood method) was 1.49 (1.39 to 1.61), indicating a 49% higher risk of stone passage associated with treatment with an alpha blocker (fig 3). The 95% confidence interval obtained with the restricted maximum likelihood method was similar (1.39 to 1.59). The quality of evidence for stone passage was rated “moderate” according to the GRADE approach (table 1). The pooled risk difference was 0.27 (0.22 to 0.31). In other words, four patients would need treatment for one patient to realize benefit from alpha blockers. The prediction interval for the pooled risk ratio was 1.12 to 1.86. The pooled percentages for stone passage in the intervention and control groups were 75.8% (72.1% to 79.2%) and 48.4% (43.5% to 53.1%), respectively. When the follow-up time was held constant at 28 days, these pooled percentages were 75.8% (71.4% to 80.0%) and 48.2% (42.0% to 54.4%), respectively.

In a cumulative meta-analysis, where we added studies one at a time (based on the publication date) and summarized the pooled risk ratio of ureteric stone passage as each new study was added, we observed a fairly stable pattern since 2006 (fig B in appendix 3). We saw no significant Proteus effect ($P=0.32$).

To determine if the effect size varied by the type of alpha blocker used, we performed meta-regression. There were no significant differences between the risk ratios for tamsulosin compared with alfuzosin ($P=0.96$), doxazosin ($P=0.91$), naftopidil ($P=0.31$), silodosin ($P=0.37$), or terazosin ($P=0.88$).

Not all studies reported on the use of imaging during follow-up. When we restricted analysis to studies in which computed tomography was used,^{29 44 51 52 58 60 65 67 6 9 72 79 81} the pooled risk ratio for stone passage was 1.64 (95% confidence interval 1.31 to 2.16). When we restricted analysis to studies reporting any imaging

Fig 1 | PRISMA diagram of trials investigating efficacy of treatment with alpha blocker in patients with ureteric stones

<div><div><div></div></div>Low risk of bias</div> <div><div><div></div></div>Unclear risk of bias</div> <div><div><div></div></div>High risk of bias</div>										
	Sequence generation	Allocation concealment	Blinding of participants and personnel	Blinding of outcome assessors	Incomplete outcome data					
Study										
Abdel-Meguid 2010 ⁵¹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Abhishek 2015 ⁷³	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Agrawal 2009 ⁴⁷	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Ahmad 2015 ⁷⁴	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Ahmad 2010 ⁵²	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Al-Ansari 2010 ⁵³	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Alizadeh 2014 ⁶⁶	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Aravinthan 2012 ⁵⁷	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Ayubov 2007 ³⁷	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Bajwa 2013 ⁶²	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Bak 2007 ³⁸	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Balci 2014 ⁶⁷	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Berger 2015 ⁷⁵	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Desai 2014 ⁶⁸	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
De Sio 2006 ³⁵	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
El-Gamal 2012 ⁵⁸	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
El Said 2015 ⁷⁶	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Ertuhan 2007 ³⁹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Eryildirim 2015 ⁷⁷	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Ferre 2009 ⁴⁸	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Furyk 2015 ⁸¹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Georgescu 2015 ⁷⁸	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Ibrahim 2013 ⁶³	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Itoh 2013 ⁶⁴	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Kaneko 2010 ⁵⁴	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Kang 2009 ⁴⁹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Kim 2007 ⁴⁰	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Kumar 2013 ⁶⁵	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					

<div><div><div></div></div>Low risk of bias</div> <div><div><div></div></div>Unclear risk of bias</div> <div><div><div></div></div>High risk of bias</div>										
	Sequence generation	Allocation concealment	Blinding of participants and personnel	Blinding of outcome assessors	Incomplete outcome data					
Study										
Kupeli 2004 ²⁹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Lee 2014 ⁶⁹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Liatsikos 2007 ⁴¹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Lojanapiwat 2008 ⁴³	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Lojanapiwat 2012 ⁵⁹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Lv 2014 ⁷⁰	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Mohseni 2006 ³⁶	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Mukhtarov 2007 ⁴²	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Ochoa-Gomez 2011 ⁵⁶	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Park 2012 ⁶⁰	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Pedro 2008 ⁴⁴	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Pickard 2015 ¹⁰	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Porpiglia 2004 ³⁰	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Rahim 2012 ⁶¹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Rathi 2014 ⁷¹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Resim 2005 ³²	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Sameer 2014 ⁷²	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Sayed 2008 ⁴⁵	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Sun 2009 ⁵⁰	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Sur 2015 ⁷⁹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Taghavi 2005 ³³	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Tekin 2004 ³¹	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Wang 2008 ⁴⁶	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Wang 2016 ⁸²	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Yilmaz 2005 ³⁴	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Yuksel 2015 ⁸⁰	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>					
Zehri 2010 ⁵⁵	<div></div>	<div></div>	<div></div>	<div></div>	<div></div>

Fig 3 | Risk ratios for passage of ureteric stones in randomized controlled trials on efficacy of treatment with alpha blockers, stratified by baseline risk

Table 1 | Summary of findings and assessment of quality of evidence for outcomes

Outcomes	No of participants (studies)	Quality of evidence (GRADE)*	Anticipated absolute effects	
			Risk with no alpha blockers	Risk difference with alpha blocker†
Stone passage (follow-up 1-6 weeks)	5701 (55)	Moderate (1,2)	545/1000	267 more per 1000 (213 more to 333 more)
Surgical intervention (follow-up 1-6 weeks)	3758 (32)	Moderate (1,3)	289/1000	162 fewer per 1000 (182 fewer to 139 fewer)
Admission to hospital	1007 (8)	Moderate (1)	168/1000	1.06 fewer per 1000 (131 fewer to 61 fewer)
No of pain episodes	1235 (13)	Low (1,4)	Mean No of pain episodes 2.3	SMD 0.74 SD lower (1.28 lower to 0.21 lower)
Time to stone passage	2862 (24)	Moderate (1,2)	Mean time to stone passage 13.3 days	MD 3.79 days lower (4.45 lower to 3.14 lower)
Serious adverse events	1205 (4)	Low (5,6)	5/1000	2 more per 1000 (4 fewer to 42 more)

SMD=standardized mean difference; MD=mean difference.

*High=very confident that true effect lies close to that of estimate of effect; moderate=moderately confident in effect estimate. True effect is likely to be close to estimate of effect, but there is a possibility that it is substantially different; low=confidence in effect estimate is limited. True effect might be substantially different from estimate of effect; very low quality=very little confidence in effect estimate. True effect is likely to be substantially different from estimate of effect. Numbers in parentheses are: 1=most studies rated as unclear risk of bias for allocation concealment (selection bias), blinding of patients and personnel (performance bias), and blinding of outcome assessors (detection bias); 2=not downgraded for unexplained heterogeneity ($I^2=76\%$) as unlikely to affect clinical decision-making; 3=not downgraded for publication bias as effect size unchanged in sensitivity analysis based on large studies alone; 4=downgraded for substantial unexplained heterogeneity ($I^2=94\%$); 5=wide confidence interval consistent with possibility of substantially increased risk; 6=adverse events addressed in only small proportion of studies.

†Showing increase or decrease in numbers of patients. Risk in intervention group (and 95% confidence interval) based on assumed risk in comparison group and relative effect of intervention (and its 95% CI).

Fig 4 | Risk ratios for passage of ureteric stones in randomized controlled trials on efficacy of treatment with alpha blockers, stratified by stone size

Effect of alpha blockers on secondary outcomes

As displayed in figure E in appendix 3, compared with controls, patients who received alpha blockers had a significantly lower risk of surgical intervention (32 studies, 3758 patients: pooled risk ratio 0.44, 95% confidence interval 0.37 to 0.52; $\tau^2=0.07$; $I^2=39\%$; moderate quality evidence).^{10 30 32-36 39 41 43-46 48 52 55 56 60 63-65 67-70 74 76-78 80-82}

When we restricted analysis to full length articles,^{10 30 32 34-36 39 41 43-46 48 52 55 56 63-65 67 69 70 74 76 77 78 80-82} the pooled risk ratio for surgery with alpha blockers was 0.46 (0.38 to 0.53; $\tau^2=0.07$; $I^2=38\%$). In meta-regression, the results for surgical intervention were similar to those for stone passage—that is, there was no significant association between maximum follow-up days and surgery ($P=0.70$) and no association between sex and surgery ($P=0.42$). The results remained significant regardless of the baseline risk (fig E, appendix 3). From the Hartung-Knapp-Sidik-Jonkman method, the pooled risk ratio for surgery in the category of low baseline risk was 0.46 (0.37 to 0.57; $\tau^2<0.001$; $I^2=0\%$).^{43 55 60 65 67 76 77} From the profile likelihood method, the pooled risk ratio for surgery in the middle risk category was 0.39 (0.32 to 0.47; $\tau^2<0.001$; $I^2=0\%$),^{30 33-35 39 41 45 46 52 63 64 69 74 78 82} and the pooled risk ratio for surgery in the high risk category was 0.53 (0.33 to 0.78; $\tau^2=0.18$; $I^2=53\%$).^{10 32 36 44 48 56 68 70 80 81}

Twenty four studies reported data regarding time to stone passage (means and standard deviations).^{10 34-36 40 44-46 52-54 56 61 62 64-66 69 70 71 72 78 80 82}

Figure F in appendix 3 shows that treatment with an alpha blocker was associated with an overall shorter time to stone passage (2862 patients: pooled mean difference -3.79 days, 95% confidence interval -4.45 to -3.14 ; $\tau^2=1.42$, $I^2=74\%$; moderate quality evidence). Mean time to stone passage (unweighted) was 8.8 and 13.3 days in the treatment and control groups, respectively. Figure F in appendix 3 shows that there was also a significant difference in the mean number of episodes of pain, favoring treatment with an alpha blocker (13 studies, 1235 patients: -0.74 episodes, -1.28 to -0.21 ; $\tau^2=0.78$; $I^2=94\%$; low quality evidence).^{32 34 45 46 52 53 65 70-72 77 80 82} Figure H in appendix 3 shows that patients treated with an alpha blocker needed admission to hospital less often than controls (8 studies, 1007 patients: pooled risk ratio 0.37, 0.22 to 0.64; $\tau^2=0.23$; $I^2=39\%$; moderate quality evidence).^{35 39 52 72 74 76 78 81} Five studies reported no patients needing admission in either the treatment or control groups and were excluded.^{30 37 46 50 82}

Adverse events were uncommon among treatment and control groups (fig I, appendix 3). Men receiving alpha blockers were more likely than male controls to experience abnormal ejaculation (pooled risk ratio 4.09, 95% confidence interval 1.73 to 12.04). The risk of a serious adverse event, however, was similar between the two groups (1.49, 0.24 to 9.35; low quality evidence).

Risk of publication bias

Inspection of the funnel plots showed asymmetry (fig J, appendix 3), indicating evidence of a small study effect for stone passage (Harbord's test: $P<0.01$), surgical intervention (Harbord's test: $P<0.01$), and pain (Egger's test: $P=0.01$), but not for time to stone passage (Egger's test: $P=0.40$) or admission to hospital (Harbord's test: $P=0.48$). Thus, we calculated the pooled risk ratio only for large studies (sample size ≥ 100).^{10 34 38 47 51 57 63-65 68-70 73 75 77-79 81 82} This yielded a pooled risk ratio of 1.39 (95% confidence interval 1.26 to 1.58) for stone passage and 0.46 (0.33 to 0.60) for surgery.

Discussion

Statement of principal findings

The pooled results of the randomized controlled trials suggest that alpha blockers help facilitate the passage of larger ureteric stones regardless of their location. Given the low risk profile of these drugs and their wide therapeutic window, our findings suggest that clinicians who manage patients with ureteric colic should consider prescribing a course of an alpha blocker, unless it is medically contraindicated.

Our findings on the effectiveness of medical expulsive therapy as it relates to the size of ureteric stones have face validity. Data from several observational studies suggest that nearly all smaller ureteric stones (that is, <5 mm) will pass without difficulty.^{83 84} The expected benefit of medical expulsive therapy (for augmenting stone passage) is therefore probably minimal in this subgroup. Along these lines, our findings corroborate

Fig 5 | Risk ratios for passage of ureteric stones in randomized controlled trials on efficacy of treatment with alpha blockers, stratified by stone location

results from another high profile randomized controlled trial reported by Furyk and colleagues,⁸¹ which was published shortly after the Pickard study.¹⁰ This double blind, placebo controlled, multicenter trial from Australia found no benefit overall of treatment with an alpha blocker for patients with lower ureteric calculi ≤ 10 mm in terms of spontaneous passage; however, in a pre-specified subgroup analysis of large stones (5-10 mm), use of tamsulosin was associated with significantly higher rates of passage.

Regarding location of ureteric stones, our findings contrast with those from a rigorous randomized controlled trial conducted by Sur and colleagues.⁷⁹ Despite the fact that use of sildosin achieved a significantly greater rate of passage of lower ureteric stones than placebo, this trial showed no benefit to treatment for stones in the upper and middle ureter (because of wide confidence intervals around the effect estimates), but there was a consistent trend towards benefit. The effects of alpha blockers on passage are thought to result from relaxation of ureteric smooth muscle mediated by binding of the drug to alpha adrenergic receptors in the region of the stone. Yet, while alpha adrenergic receptors are concentrated in the lower ureter,⁸⁵ studies have shown that they are present along the entire length of the human ureter.^{86,87} Thus, one could anticipate that alpha blockers exert their effect throughout the ureter, as we found in our pooled analysis.

Strengths and weaknesses of study

The strengths of our study include the comprehensive nature of our literature search (which included studies irrespective of language and publication status); the thoroughness of our study selection, data abstraction, and risk of bias assessment (carried out in duplicate by two independent members of our study team); our pre-defined analytic plan; and our use of the GRADE approach for assessing the quality of evidence on an outcome specific basis.

Despite these strengths, several limitations merit discussion. The first relates to clinical (not statistical) heterogeneity between pooled studies, in view of the variation in the types of alpha blockers given, the inconsistent use of post-treatment imaging, and differential follow-up. We found, however, that the pooled relative risk of passage of ureteric stones was independent of these clinical differences. Another limitation pertains to the overall methodological rigor of the pooled studies. Only a handful concealed allocation adequately, and just six studies reported blinding of outcome assessment. Given that the inclusion of less rigorous studies could lead to misestimation of the true intervention effect, we conducted separate sensitivity analyses, in which we recalculated our summary estimates excluding data from non-placebo controlled studies and those in which the risk of bias was judged high. While our summary estimates were attenuated, the benefit of alpha blockers persisted. Differences in stone passage by type of alpha blocker were not found through meta-regression, but this could be because of insufficient power. Finally, our results could be affected

by publication bias, whereby smaller studies with negative results might never have been published. To account for this possibility, we repeated our models pooling data from only larger studies and found our results to be robust.

Strengths and weaknesses in relation to other studies

Several previous systematic reviews and meta-analyses have examined the role of alpha blockers for treatment of ureteric stones, most notably the initial study by Hollingsworth and colleagues.³ They identified nine randomized controlled trials that yielded a pooled risk ratio of 1.54 (95% confidence interval 1.29 to 1.85), favoring medical expulsive therapy. The recent Cochrane review published by Campshroer and colleagues included 32 randomized controlled trials and found a 48% higher risk of stone passage with treatment with alpha blockers than control (risk ratio 1.48, 1.33 to 1.64).⁶ Similar to our findings, the benefit of alpha blockers was attenuated when the analysis was limited to only four placebo controlled trials (1.22, 0.99 to 1.55) but consistent with a potentially large effect.

Since the date of the Cochrane group's last search, we identified 24 additional randomized controlled trials on alpha blockers for the treatment of ureteric stones. We are unaware of any recent high quality systematic reviews that include the randomized controlled trial from Pickard and colleagues, findings from which have called the concept of medical expulsive therapy into question.¹⁰ Although the Pickard study was large and methodologically rigorous in its design, its results need to be placed into the context of the entire body of evidence as in a systematic review like ours. We can explain the discrepancies in findings largely due to the high rate of spontaneous stone passage in the control arm of the trial perhaps because of the large proportion of patients with smaller stones.

A second important difference between the Pickard study and most other randomized controlled trials on medical expulsive therapy relates to how stone passage was defined. Imaging evidence has been the standard assessment, yet investigators in the Pickard study chose instead "absence of need for additional interventions to assist stone passage at four weeks after randomisation."¹⁰ Compelling arguments can be made that this endpoint is highly relevant to surgeons, but the degree to which intervention rates accurately approximate spontaneous stone passage is uncertain. While additional imaging was obtained when "clinically indicated" (such as for continued pain, development of infection), just over half of participants were reimaged, raising the possibility of silent obstruction and late secondary complications.⁸⁸ Further, defining passage by the absence of intervention, rightly or wrongly, does not reflect routine clinical practice in many countries outside the UK, nor does it align with contemporary practice guidelines from the European Association of Urology, under which reimaging is recommended to monitor the position of ureteric stones and assess for hydronephrosis.

Implications for clinicians and policy makers

Findings from our study emphasize the role of systematic reviews to examine focused clinical questions. The recent publication of the Pickard study brought into question the effectiveness of alpha blockers in patients with ureteric colic, leading to calls from the urologic community to reformulate treatment guidelines and even abandon medical expulsive therapy altogether. Our findings suggest that this would be an over-reaction as large subgroups of patients could be spared from stone surgery and its attendant risks with a trial of an alpha blocker.

Further, our study highlights the challenges of trials with undifferentiated cohorts that often lack the sample size necessary to tease out important clinical nuances. Therefore, while not all patients seen in the emergency department for ureteric colic will be helped by alpha blockers, our meta-analysis, which draws power from the pooling of data across multiple studies, suggests that those with stones ≥ 5 mm in size could.

Such a size based approach to the use of alpha blockers requires patients with suspected ureteric stones to undergo radiologic testing. Under current guidelines on urolithiasis from the European Association of Urology, immediate imaging is indicated in the evaluation of all patients who present with acute flank pain.² In many emergency departments, however, particularly for patients with known histories of urinary stone disease, imaging is often deferred until the time of primary or specialty care follow-up, which could delay initiation of expulsive therapy. Moreover, renal ultrasonography is often used as the primary diagnostic tool, from which stone size cannot always be accurately assessed. While a non-contrast computed tomogram should be obtained subsequently to confirm a stone diagnosis,² many times it is not. Clearly, there are some clinical challenges for the implementation of our findings.

Unanswered questions and future research

Our findings on international differences in the baseline rates of stone passage suggest that patient related factors could modify the effects of expulsive therapy. To better assess their role, investigators could consider including variables like patient age, sex, and race/ethnicity in the design of a large international trial. Future studies should also examine patients' values and preferences concerning acceptable risk of retained ureteric stones versus the potential inconvenience, radiation exposure, and the direct and indirect costs of repeat imaging.

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Appendix 1: PubMed search strategy
Appendix 2: Supplementary tables A-F
Appendix 3: Supplementary figures