Pumpkin seed oil (Cucurbita pepo) versus tamsulosin for benign prostatic hyperplasia symptom relief: a single-blind randomized clinical trial

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Abstract

Background: Benign prostatic hyperplasia (BPH) is very common in aging men. We aimed to compare the effects of tamsulosin and pumpkin (Cucurbita pepo) seed oil on BPH symptoms.

Methods: This single-blind randomized clinical trial included patients with BPH aged ≥ 50 years referred to the Urology Clinic of Shahid Beheshti Hospital, Hamadan, Iran, from August 23, 2019 to February 19, 2020. Patients were randomized into two groups. One group received 0.4 mg tamsulosin every night at bedtime and the other received 360 mg pumpkin seed oil twice a day. Patients' age, weight, height, and body mass index (BMI) were recorded. The International Prostate Symptom Score (IPSS) was filled out by the patients at baseline and then 1 month and 3 months after the initiation of treatment. The BPH-associated quality of life (QoL), serum prostate-specific antigen, prostate and postvoid residual volume, and maximum urine flow were also assessed at baseline and 3 months later. Drug side effects were also noted.

Results: Of the 73 patients included in this study with a mean age of 63.59 ± 7.04 years, 34 were in the tamsulosin group and 39 in the pumpkin seed oil group. Patients were comparable with respect to age, weight, height, BMI, and baseline principal variables in both groups. Also, there was no significant difference between groups in terms of principal variables at any time point. However, there was a significant decrease in IPSS and a significant improvement in QoL in both groups. Although the decrease in IPSS from baseline to 1 month and 3 months was significantly higher in the tamsulosin group compared to the pumpkin group (P = 0.048 and P = 0.020, respectively), the decrease in IPSS from 1 to 3 months was similar (P = 0.728). None of the patients in the pumpkin group experienced drug side effects, while dizziness (5.9%), headache (2.9%), retrograde ejaculation (2.9%), and erythema with pruritus occurred in the tamsulosin group.

Conclusions: Pumpkin (Cucurbita pepo) seed oil relieved BPH symptoms with no side effects, but was not as effective as tamsulosin. Further studies are required to confirm the role of pumpkin seed oil as an option for the treatment of BPH symptoms.

Trial registration: Iranian Registry of Clinical Trials, IRCT20120215009014N340. Registered 19.02.2020. Retrospectively registered, https://en.irct.ir/trial/45335.

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Introduction

Benign prostatic hyperplasia (BPH) is a common age-dependent chronic disease that results from the progressive enlargement of the prostate gland due to the non-malignant proliferation of epithelial prostate cells and smooth muscle cells [1]. A recent systematic review and meta-analysis including data from 25 countries reported a lifetime prevalence of 26.2% for BPH [2]. Patients with BPH become symptomatic when the tissue overgrowth around the urethra constricts its opening leading to lower urinary tract symptoms (LUTS), including incomplete urination, frequency, urgency, nocturia, and decreased urine flow [3]. The prevalence of BPH-associated LUTS increases with age and it has been reported that approximately 80% of men experience BPH-associated LUTS by 70 years of age [4]. However, not every man with BPH symptoms seeks medical attention; most often, BPH patients only seek medical care when BPH-associated LUTS become bothersome or intolerable [1].

There is a variety of treatment strategies for BPH symptoms depending on symptom severity, together with patient discomfort and preference. These treatment strategies include lifestyle alterations, medical therapy, and surgical treatment [5]. Alpha-blockers such as tamsulosin are excellent first-line options of medical therapy [6]. Recently, there has been an increasing tendency towards the use of herbal medicines for different medical conditions. Pumpkin (Cucurbita) seeds are traditionally known around the world for their remedial effects on urinary tract complications, such as nocturia, urinary frequency, and stress urinary incontinence [7]. Pumpkins belong to the Cucurbitaceae family which includes various species such as Cucurbita pepo, Cucurbita moschata, and Cucurbita maxima [8]. Cucurbita pepo seed oil consists of high amounts of free fatty acids serving as a natural source of vitamins, proteins, trace elements, and polyunsaturated fatty acids, such as omega 3, 6, and 9 [9]. It is the phytosterol content of Cucurbita pepo seed oil that appears to interfere with the function of dihydrotestosterone produced by 5α-reductase which plays a major role in the process of BPH [10]. The phytosterol content varies among different Cucurbita species, including Cucurbita pepo and the previously mentioned Cucurbita moschata and Cucurbita maxima [11, 12]. However, only Cucurbita pepo is available in the Iranian market. Thus, in the current study we aimed to compare the effects of Cucurbita pepo with tamsulosin for the treatment of BPH symptoms.

Methods

Participants

This single-blind randomized clinical trial included patients aged ≥ 50 years with the clinical diagnosis of BPH by an expert urologist based on history, digital rectal examination (DRE), and paraclinical tests including serum prostate-specific antigen (PSA), who had been referred to the Urology Clinic of Shahid Beheshti Hospital, Hamadan, Iran, from August 23, 2019 to February 19, 2020. Exclusion criteria were PSA > 10 ng/ml, indications of surgical treatment (patients who were planned to receive surgical treatment for BPH due to absolute or relative indications, including renal insufficiency caused by benign prostatic obstruction, intractable urinary retention, recurrent cystitis, failure of medical therapy, bladder calculi, and persistent hematuria due to prostatic bleeding), previous surgical treatment of BPH, symptom exacerbation during the study period mandating surgical intervention, change in the diagnosis during the study period, any morbidities interfering with the course of treatment, having taken BPH-related medications within the past 6 months, and the development of drug side effects leading to its discontinuation. In patients with PSA > 4 ng/ml and < 10 ng/ml, in order to rule out prostatic cancer, DRE was done and free/total PSA ratio was measured. Also, in suspicious cases, prostatic biopsy was performed. The sample size was calculated as at least 35 patients in each group using a level of significance of 5%, power of 90%, and non-inferiority margin of 50%. The study was approved by the Institutional Review Board of Hamadan University of Medical Sciences (IR.UMSHA.REC.1398.429) and it complies with the statements of the Declaration of Helsinki. Written informed consent was obtained from all the participants. The study has also been retrospectively registered at the Iranian Registry of Clinical Trials (IRCT) under the registration number: IRCT20120215009014N340.

Study design

Initially, 80 patients were eligible to enter the study. Patients were randomized into two equal groups using random generated numbers by the Random Allocation software. Seven patients were lost to follow-up. Details of patient enrollment are demonstrated in Fig. 1. Patients’ age, weight, and height were recorded. Body mass index (BMI) was calculated for each patient by dividing their weight (kg) by the square of their height (m). Patients in the tamsulosin group received 0.4 mg tamsulosin capsules (Farabi Pharmaceutical Co., Iran)
every night at bedtime, while patients in the pumpkin group received 360 mg pumpkin (*Cucurbita pepo*) seed oil capsules (Tehran Darou Pharmaceutical Co., Iran) containing 1% phytosterol, twice a day. The reason for choosing this dose and type of pumpkin (*Cucurbita pepo*) was that it was the only form and dose available in the Iranian market. In previous studies, Vahlensieck et al. and Friedrich et al. used 1000 mg of pumpkin [13, 14]; however, with the available dose per capsule in our study, taking 3 capsules (1080 mg) could have reduced patient compliance. Thus, patients in the pumpkin group received 2 capsules a day (720 mg). Patients in both groups continued taking their medications till the last round of evaluations at 3 months. Patients were asked to fill out the International Prostate Symptom Score (IPSS) for the assessment of BPH symptom severity, before treatment and then 1 and 3 months later. The BPH-associated quality of life (QoL) scale was also used once before treatment and then at 3 months. The QoL scale evaluates how the patient would feel if he were to spend the rest of his life with his current urinary condition. It is scored from 0 to 6 with 0 indicating the highest and 6 the lowest QoL. Before treatment and at 3 months, all patients underwent ultrasonography for the measurement of postvoid residual (PVR) and prostate volume. Uroflowmetry was also performed for all the participants at the same time points and serum PSA was measured as well. Drug side effects, including dizziness, headache, retrograde ejaculation, and erythema with pruritus during the study period were noted. The laboratory personnel and the individuals in charge of ultrasonography and uroflowmetry were blinded to the grouping of the patients.
Data analysis
We used the Statistical Package for the Social Sciences (SPSS) software (version 25.0, Armonk, NY: IBM Corp.) for data analysis. Mean, standard deviation, frequency, and percentages were used to describe the variables. Based on the results of Kolmogorov–Smirnov normality test, independent t test and Mann–Whitney test were used to compare quantitative variables between groups. Friedman test was used to assess the significance of changes in qualitative variables across different time points. Pairwise comparisons were made using paired t-test and Wilcoxon test. P values ≤ 0.05 were regarded as statistically significant.

Results
Of the 73 patients included in this study with a mean age of 63.59 ± 7.04 years, 39 (53.4%) were in the pumpkin group and 34 (46.6%) in the tamsulosin group. Participants’ general characteristics are shown in Table 1. Participants in both groups were comparable regarding age, weight, height, and BMI. There was no significant difference between groups in terms of baseline IPSS, IPSS at one month, and IPSS at three months (Table 2). However, there was a significant change in IPSS from baseline to the end of three months in both groups (P < 0.001). The reduction in IPSS from baseline to 1 month and 3 months was significantly higher in the tamsulosin group compared to the pumpkin group (−3.30 ± 3.15 vs. −1.74 ± 3.02, P = 0.048 and −5.33 ± 3.64 vs. −3.19 ± 3.63, P = 0.020, respectively). Pairwise comparisons (Additional file 1: Table S1) also showed significant decrease in IPSS between any two time points in both groups. Moreover, although the QoL scores were comparable between groups at baseline and at 3 months, these scores significantly decreased between these time points in both groups (Table 2). Serum PSA was also similar in both groups at baseline and at 3 months. There was a slight increase in serum PSA between these time points in both groups; nevertheless, this was not statistically significant. Prostate volume was comparable between groups at baseline and at 3 months with no significant increase in any of the groups. The same was true for PVR and maximum urine flow.

Furthermore, none of the patients in the pumpkin group experienced drug side effects, while in the tamsulosin group, dizziness occurred in 2 patients (5.9%), headache in 1 (2.9%), retrograde ejaculation in 1 (2.9%), and erythema with pruritus in 1 (2.9%).

Discussion
In the current study, we found that both tamsulosin and pumpkin (Cucurbita pepo) seed oil significantly reduced BPH symptoms assessed by IPSS. However, the decrease in IPSS from baseline to 1 month and 3 months was significantly higher in the tamsulosin group compared to the pumpkin group, while the decrease from 1 to 3 months was similar. Moreover, no patients in the pumpkin group experienced drug side effects, while dizziness (5.9%), headache (2.9%), retrograde ejaculation (2.9%), and erythema with pruritus occurred in the tamsulosin group.

With a global prevalence of 20–62% in men over 50 years, BPH is considered a very common condition [15]. BPH causes LUTS which markedly affect the quality of life in many patients [16, 17]. Multiple strategies have been proposed and used for the relief of BPH-related symptoms. Alpha-blockers such as tamsulosin are regarded as the first-line options. By selectively blocking α1A-adrenergic receptors leading to the relaxation of the prostate smooth muscles, tamsulosin is reported to improve dysuria and other BPH symptoms [18]. Although tamsulosin is generally preferred due to its lower side effects compared to other alpha-blockers, it can still cause some complications and unwanted reactions, including dizziness, headache, and retrograde ejaculation [19]. We also observed some of these side effects in patients taking tamsulosin.

The effects of tamsulosin and pumpkin seed oil on BPH symptoms have separately been investigated in multiple studies, yet none have compared them with regard to BPH symptom relief. In vitro and in vivo experiments have shown promising results for pumpkin seeds. In their study on Sprague–Dawley rats, Gossell-Williams et al. observed a reduction of prostate size with pumpkin

Table 1 General characteristics of the study population

| Variables          | Total (n = 73) | Tamsulosin (n = 34) | Pumpkin (n = 39) | P value* |
|--------------------|---------------|---------------------|------------------|---------|
| Age (years) mean ± SD | 63.59 ± 7.04  | 62.71 ± 6.63       | 64.36 ± 7.38     | 0.320   |
| Height (cm) mean ± SD | 169.49 ± 6.74 | 169.12 ± 6.49      | 169.82 ± 7.01    | 0.916†  |
| Weight (kg) mean ± SD | 73.04 ± 14.21 | 71.27 ± 12.11      | 74.54 ± 12.25    | 0.261   |
| BMI (kg/m²) mean ± SD | 25.38 ± 3.64  | 24.89 ± 3.68       | 25.78 ± 3.61     | 0.309   |

n, number; SD, standard deviation; BMI, body mass index
† Analyzed by independent t-test
‡ Analyzed by Mann–Whitney test
### Table 2: Comparison of IPSS, QoL, serum PSA, maximum urine flow, PVR, and prostate volume between groups at different time points

| Variables                                | Tamsulosin (n = 34) | Pumpkin (n = 39) | P-value* |
|------------------------------------------|---------------------|------------------|----------|
| IPSS at 1 month                          | 7.35 ± 4.53 (5.77–9.93) | 9.33 ± 5.57 (7.53–11.14) | 0.174 |
| IPSS at 3 months                         | 5.65 ± 4.25 (4.09–7.20) | 7.46 ± 5.84 (5.51–9.41) | 0.231 |
| IPSS change from baseline to 1 month     | <0.001†             | <0.001†          |          |
| IPSS change from baseline to 3 months    | −3.30 ± 3.15 (−4.42–−2.19) | −1.74 ± 3.02 (−2.72–−0.77) | 0.048 |
| IPSS change from baseline to 3 months    | −5.33 ± 3.64 (−6.69–−3.97) | −3.19 ± 3.63 (−4.40–−1.98) | 0.020 |
| IPSS change from 1 to 3 months           | −1.84 ± 2.59 (−2.79–−0.89) | −1.46 ± 3.81 (−2.73–−0.19) | 0.728 |
| Baseline QoL score                       | 2.35 ± 1.41 (1.86–2.85) | 2.41 ± 1.33 (1.98–2.84) | 0.767 |
| QoL score at 3 months                    | 1.38 ± 0.99 (1.04–1.73) | 1.67 ± 1.06 (1.32–2.01) | 0.148 |
| Serum PSA change (ng/ml)                 | 0.48 ± 1.41 (0.11–1.07) | 0.13 ± 1.36 (−0.47–0.73) | 0.396 |
| Serum PSA change (ng/ml)                 | 50.93 ± 22.74 (42.87–58.99) | 53.53 ± 23.53 (45.79–61.26) | 0.511 |
| Prostate volume at 3 months (ml)         | 55.00 ± 21.22 (46.21–63.75) | 58.32 ± 23.78 (48.85–69.77) | 0.629 |
| PVR at 3 months (ml)                     | 56.43 ± 81.63 (19.27–93.59) | 51.61 ± 33.54 (34.93–68.29) | 0.080 |
| PVR change (ml)                          | 0.723‡              | 0.087‡            |          |
| Maximum urine flow change (ml/sec)       | −9.56 ± 94.93 (−56.76–37.65) | −7.67 ± 21.98 (−19.84–4.50) | 0.563 |
| Maximum urine flow at 3 months (ml/sec)  | 12.27 ± 5.62 (9.83–14.69) | 11.18 ± 6.09 (8.61–13.75) | 0.344 |
| P-value                                  | 0.698‡              | 0.091†            |          |
| Maximum urine flow change (ml/sec)       | 0.11 ± 3.09 (−1.42–1.64) | 1.82 ± 4.39 (0.12–3.77) | 0.328 |

**SD**, standard deviation; CI, confidence interval; IPSS, International Prostate Symptom Score; QoL, quality of life; PSA, prostate-specific antigen; PVR, postvoid residual
* Analyzed by Mann–Whitney test
† Analyzed by Friedman test
‡ Analyzed by Wilcoxon test
§ Analyzed by paired t-test
¶ Analyzed by independent t-test

Seed oil [20]. This effect has been confirmed by other in vitro and animal studies [21–23]. In line with our findings, Vahlensieck et al. demonstrated a clinically relevant reduction of IPSS compared to placebo after 12 months of taking pumpkin seed extract [13]. They used 500 mg *Cucurbita pepo* seed oil extract capsules twice a day. Nevertheless, the period of treatment was much longer in their study reflecting the long-term effects of pumpkin seeds compared to the 3-month period of treatment in our study. In a study on 100 patients by Shirvan et al., pumpkin seed oil (360 mg *Cucurbita pepo* twice a day) was compared with prazosin for the treatment of BPH symptoms [24]. They found pumpkin seed oil to be safe and effective but not as effective as prazosin. Their findings were consistent with ours. We also found a significant reduction in IPSS with both tamsulosin and pumpkin seed oil from baseline 3 months; nonetheless, the reduction in IPSS from baseline to the end of 1 month and 3 months was significantly higher with tamsulosin. Moreover, they further evaluated patients after 6 months of treatment, while we only assessed the results after 3 months. The results of a large study by Friederich et al. on 2245 BPH patients taking pumpkin seed extract (1–2 500 mg *Cucurbita pepo* capsules) for 3 months have also confirmed the effectiveness of this herbal medications showing a 41.4% decrease in IPSS and 46.1% improvement in their QoL [14]. Dihydrotestosterone, converted from testosterone by 5α-reductase, is responsible for the
overgrowth of the prostate gland which is characteristic of BPH. [25]. Targeting this pathway has been the mainstay of medical treatment in BPH. Pumpkin seeds appear to affect BPH through the same pathway [10]. Anti-inflammatory properties of pumpkin seeds have been proposed as another mechanism for the effectiveness of pumpkin seed oil in BPH [26]. BPH is associated with the inflammation of the prostate gland and overexpression of cytokines, leukotriene, inducible nitric oxide synthase, NF-κB, and cyclooxygenase-2 is linked with prostatitis [27]. Another proposed mechanism is the diuretic effects of pumpkin seed due to its fatty acids content [28].

Of note, IPSS is a subjective tool to assess the improvement of BPH symptoms. An objective method to evaluate BPH improvement is maximum urine flow and we found no significant change in this variable in any of the groups. One reason for this can be that we had included patients with > 15 ml/s maximum urine flow at baseline only because they complained of BPH symptoms and this could have interfered with the results.

Also, we did not observe significant reductions in PSA and prostate volume with pumpkin seed oil. The results of the study by Hong et al. were similar, as they found no significant reduction in PSA and prostate volume after 3 months in patients receiving 320 mg/day pumpkin seed oil. They did not specify the Cucurbita species used in their study (29). As the effect of pumpkin seed oil has been attributed to its phytosterol content, higher doses of pumpkin seed oil potentially include higher amounts of phytosterol leading to increased effects. Nevertheless, patients’ adherence to medications would be of concern with higher doses.

The strength of the current study was the homogeneity of patients taking tamsulosin or pumpkin seed oil regarding the potential effectors, as well as baseline characteristics, including symptom severity, QoL, serum PSA, maximum urine flow, and prostate and PVR volumes. This study was not without limitations. First, it has been conducted on a limited number of patients. The study did not reach the minimal sample size of 35; 6 patients were lost to follow-up in the tamsulosin group and 34 patients remained in this group. Further studies with a larger sample size are required to confirm our findings. Second, we only evaluated the efficacy of pumpkin seeds for a 3-month period of treatment and its long-term effects need to be determined in future studies. Also, patients’ adherence to treatment, which we failed to assess, is another factor that could have influenced the results. Moreover, we included patients with IPSS < 8 who were mildly symptomatic. Although the initial design of the study was to only include patients with moderate to severe BPH symptoms, we also included mildly symptomatic patients due to the persistent complaint of these patients of BPH symptoms causing them discomfort, as well as the limited recruitment time.

Conclusions
Both pumpkin seed oil and tamsulosin significantly reduced BPH symptoms; however, due to the higher reduction in IPSS scores from baseline to 1 month and 3 months, tamsulosin was more effective. The advantage of pumpkin seed was its lower side effects of pumpkin seeds. Further studies are required to confirm the role of pumpkin seed oil as an option for the treatment of BPH symptoms.

Abbreviations
BMI: Body mass index; BPH: Benign prostatic hyperplasia; DRE: Digital rectal examination; IPSS: International Prostate Symptom Score; IRCT: Iranian Registry of Clinical Trials; LUTS: Lower urinary tract symptoms; PSA: Prostate-specific antigen; PVR: Postvoid residual; QoL: Quality of life.

Supplementary Information
The online version contains supplementary material available at https://doi.org/10.1186/s12894-021-00910-8.

Additional file 1. Pairwise comparisons of IPSS scores at different time points in each group.

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Authors’ contributions
Conceptualization and study validation: MA. Study supervision: MB and FS. Implementation: NZ. Data analysis and interpretation: MB and FS. Writing and reviewing: NZ. Editing and revision: GZ. All the authors have read and approved the manuscript.

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Availability of data and materials
The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Declarations
Ethics approval and consent to participate
The study was approved by the Institutional Review Board of Hamadan University of Medical Sciences (IR.UUMSHA.REC.1398.429) and it complies with the statements of the Declaration of Helsinki. Written informed consent was obtained from all the participants. The study has also been retrospectively registered at the Iranian Registry of Clinical Trials (IRCT) with the registration number: IRCT20120215009014N340 and is accessible at the following website: www.irct.ir.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.
1. Egan KB. The epidemiology of benign prostatic hyperplasia associated with lower urinary tract symptoms: prevalence and incident rates. Urol Clin. 2016;43(3):289–97.

2. Lee SWH, Chan EMC, Lai YK. The global burden of lower urinary tract symptoms suggestive of benign prostatic hyperplasia: a systematic review and meta-analysis. Sci Rep. 2017;7(1):1–10.

3. McVary KT, Roehrborn CG, Avins AL, Barry MJ, Bruskewitz RC, Donnell RF, et al. Update on AUA guideline on the management of benign prostatic hyperplasia. J Urol. 2011;185(5):1793–803.

4. Litman HJ, McKinlay JB. The future magnitude of urological symptoms in the USA: projections using the Boston Area Community Health survey. BJU Int. 2007;100(4):820–5.

5. Nickel JC, Aaron L, Barkin J, Etterman D, Nachabé M, Zorn KC. Canadian Urological Association guideline on male lower urinary tract symptoms/benign prostatic hyperplasia (MLUTS/BPH): 2018 update. Can Urol Assoc J. 2018;12(10):303.

6. Rendon RA, Mason RJ, Marzouk R, Finelli A, Saad F, So A, et al. Canadian Urological Association recommendations on prostate cancer screening and early diagnosis. Can Urol Assoc J. 2017;11(10):298.

7. Medjakovic S, Hoberg S, Arjmand-Weelkart K, Bucar F, Jungbauer A. Pumpkin seed extract: Cell growth inhibition of hyperplastic and cancer cells, independent of steroid hormone receptors. Fitoterapia. 2016;10:150–6.

8. Lee YK, Chung WI, Ezura H. Efficient plant regeneration via organogenesis in winter squash (Cucurbita maxima Duch.). Plant Sci. 2003;164(3):413–8.

9. Ratnam N, Naijjibullah M, Ibrahim MD. A review on Cucurbita pepo. Int J Pharm Phytochem Res. 2017;9:1190–4.

10. Perez Gutierrez RM. Review of Cucurbita pepo (pumpkin) its phytochemistry and pharmacology. Med cham. 2016;6(1):1012–21.

11. Martínez Aguilar Y, Martínez Yero O, Córdova López J, Valdivié Navarro M, Estarrón EM. Fitosteroides y escualeno como hipocolesterolémicos en cinco variedades de semillas de Cucurbita maxima y Cucurbita moschata (calabaza). Revista Cubana de Plantas Medicinales. 2011;16(1):72–81.

12. Elias MG, Hassan KD, Odeh S, Mohiaddin SR. Study of growth, yield and phytosterol of squash (Cucurbita pepo) and medical pumpkin (Cucurbita pepo) on their hybrid. Iraqi J Agric Sci. 2020;52(2):675–84.

13. Vah lensiek W, Theurer C, Pfister E, Patz B, Banik N, Engelmann U. Effects of pumpkin seed in men with lower urinary tract symptoms due to benign prostatic hyperplasia in the one-year, randomized, placebo-controlled GRANU study. Urol Int. 2015;94(3):286–95.

14. Friederich M, Theurer C, Schiebel-Schlosser G. Prosta Fink Forte capsules in the treatment of benign prostatic hyperplasia. Multicentric surveillance study in 2245 patients. Res Complement Natural Class Med. 2000;7(4):200–4.

15. Venoba ED. Prevalence of benign prostatic hyperplasia and prostate cancer in Africans and Africans in the diaspora. J West Afr College Surg. 2016;6(4):1.

16. Bushman W. Etiology, epidemiology, and natural history. Urol Clin. 2009;36(4):403–15.

17. Woodard TJ, Manigault KR, McBurnows NN, Wray TL, Woodard LM. Management of benign prostatic hyperplasia in older adults. The Consultant Pharmacist®. 2016;31(8):412–24.

18. Zhou Z, Cui Y, Wu J, Ding R, Cai T, Gao Z. Meta-analysis of the efficacy and safety of combination of tamsulosin plus dutasteride compared with tamsulosin monotherapy in treating benign prostatic hyperplasia. BMC Urol. 2019;19(1):1–12.

19. Sun Y-Y, Liu Z-j, Zhang Z-s, Xu C-L, Ji J-T, Wu Y-Y, et al. Long-term efficacy and safety of tamsulosin hydrochloride for the treatment of lower urinary tract symptoms associated with benign prostatic hyperplasia: data from China. Chin Med Journal. 2011;124(1):56–60.

20. Gossell-Williams M, Davis A, O’Connor N. Inhibition of testosterone-induced hyperplasia of the prostate of sprague-dawley rats by pumpkin seed oil. J Med Food. 2006;9(2):284–6.

21. Abdel-Rahman MK. Effect of pumpkin seed (Cucurbita pepo L) diets on benign prostatic hyperplasia (BPH): chemical and morphometric evaluation in rats. World J Chem. 2006;1(1):33–40.

22. Tsai Y-S, Tong Y-C, Cheng J-T, Lee C-H, Yang F-S, Lee H-Y. Pumpkin seed oil and phytosterol-F can block testosterone/prazosin-induced prostate growth in rats. Urol Int. 2006;77(3):269–74.

23. Schleich S, Papaioannou M, Bainahmad A, Matusch R. Extracts from Pygurum africanum and other ethnochemical species with antiandrogenic activity. Planta Med. 2006;72(9):807–13.

24. Shirvan MK, Mahboob MRD, Masuminia M, Mohammadi S. Pumpkin seed oil (prostafit) or prazosin? Which one is better in the treatment of symptomatic benign prostatic hyperplasia? JPMJ J Pak Med Assoc. 2011;64:683–5.

25. Banerjee PP, Banerjee S, Brown TR, Zirkin BR. Androgen action in prostate function and disease. Am J Clin Experim Urol. 2018;6(2):62.

26. Fahim AT, Abd-El Fattah AA, Agha AM, Gad MZ. Effect of pumpkin-seed oil on the level of free radical scavengers induced during adjuvant-arthritis in rats. Pharmaco Res. 1995;31(1):73–9.

27. Sciarra A, Di Silviero F, Saliccia S, Gomez AMA, Gentilucci A, Gentile V. Inflammation and chronic prostatic diseases: evidence for a link? Eur Urol. 2007;52(4):964–72.

28. Ramak P, Mahboubi M. The beneficial effects of pumpkin (Cucurbita pepo L) seed oil for health condition of men. Food Rev Int. 2019;35(2):166–76.

29. Hong H, Kim C-S, Maeng S. Effects of pumpkin seed oil and saw palmetto on prostate cancer in Koreans. Jpn J Cancer Res. 2000;91(4):323–7.

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