

Review Article

Clinical antihypertensive efficacy and safety of Iran plants: a systematic review

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ABSTRACT

Background: Antihypertensive plants are one of the means of hypertension control.

Objective: To examine the clinical antihypertensive efficacy and safety of the plants found in Iran. **Methods:** PUBMED, MEDLINE, SCOPUS, EMBASE, SCIENCECIRECT, PROQUEST, OVID, EBSCO, GOOGLE, and GOOGLE SCHOLAR were searched. The PRISMA guideline was observed. The search terms were Iran, Iranian, plant, herb, antihypertensive, hypertension and randomized controlled trial (RCT). English-language articles published until the end of 2022 were included. In-vitro and animal studies, editorials, and reviews were excluded. The methodological quality of the RCTs was evaluated using the JADAD scale. **Results:** Two hundred and eight studies were found. Only 74 of them were eligible. For *Berberis vulgaris* (5 studies), *Nigella sativa* (10 studies), *Allium sativum* (12 studies), *Hibiscus sabdariffa* (11 studies), *Beta vulgaris* L (15 studies), *Solanum lycopersicum* (5 studies), *Cinnamomum verum* (9 studies), *Rhus coriaria* (1 study), *Phyllanthus emblica* (1 study), *Olea europaea* (4 studies), and *Vaccinium arctostaphylos* (3 studies) were found. Most RCTs had high methodological quality and reported efficacy and no side effects.

Conclusion: While most trials demonstrate antihypertensive efficacy and safety, there are more evidence regarding *Hibiscus sabdariffa*, *Olea europaea*, *Vaccinium arctostaphylos* and *Allium sativum* versus the other plants.

Abbreviations: 2hPPG, 2 hour post-prandial plasma glucose; ABPM, ambulatory blood pressure monitoring; ALP, alkaline phosphatase; ALT, alanine aminotransferase; AST, aspartate aminotransferase; BID, twice a day; BL, base line; BMI, body mass index; BP, blood pressure; BS, blood sugar; BUN, blood urea nitrogen; CBC, complete blood count; CK-MB, creatine kinase-myoglobin binding; Cr, creatinine; CVLT-II, California Verbal Learning Test Second Edition; DBP, diastolic blood pressure; EC, enteric coated; eGFR, estimated glomerular filtration rate; EHTN, essential hypertension; ET, extract; FBS, fasting blood sugar; FID, 4 times a day; HAM, healthy adolescent male; HbA1c, glycated hemoglobin; HDL, high-density lipoprotein; HDL-C, high density lipoproteins-cholesterol; HR, heart rate; HS, *hibiscus sabdariffa*; hs-CRP, high-Sensitivity C-Reactive Protein; LDL, low-density lipoprotein; ICAM-1, intercellular adhesion molecule-1; LDL-C, low density lipoproteins-cholesterol; HTN, hypertension; HCL, hypercholesterolemia; LV, left ventricular; MDA, malondialdehyde; MetS, metabolic syndrome; N/A, not applicable; NS, *nigella sativa*; NS, Not Significant; Po, placebo; RDB, randomized double blind; PAH, Pulmonary arterial hypertension; Pre-DM, pre diabetes mellitus; s-ICAM-1, soluble intercellular adhesion molecule-1; sVCAM-1, soluble vascular cell adhesion molecule-1; T2DM, type 2 diabetic mellitus; TID, 3 times a day; SBP, systolic blood pressure; NAFLD, non-alcoholic fatty liver disease; NHV, Normal healthy volunteer; OLE, olive leaf extract; RCT, randomized controlled trial; SBP, systolic blood pressure; SGOT, serum glutamic oxaloacetic transaminase; SGPT, serum glutamic pyruvic transaminase; STAI, State-Trait Anxiety Inventory; TC, total cholesterol; TDS, three times a day; TE, tomato extract; TG, triglycerides; TNF- α , tumor necrosis factor alpha; Tsp, tea spoonful; UHTN, uncontrolled hypertension.

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1. Introduction

Hypertension is a prevalent condition categorized into two types: essential hypertension and secondary hypertension. Essential and secondary hypertension respectively constitute 90-95 % and 5-10 % of the hypertensive cases [1, 2]. Uncontrolled hypertension leads to cardiovascular complications such as stroke, myocardial infarction, nephropathy, and retinopathy. Hypertension is the leading cause of morbidity and mortality. It is usually defined as SBP (systolic blood pressure) ≥ 140 mm Hg and DBP (diastolic BP) ≥ 90 mm Hg [1, 3]. Normal BP is defined as SBP < 120 mm Hg and DBP < 80 mm Hg. Every 20/10 mm Hg increase of BP above 115/75 mm Hg is associated with a doubling of the risk of cardiovascular diseases (CVD). The target of 120/80 mm Hg is more beneficial than the previous goal of 140/90 mm Hg in hypertension control [4]. There are around 1.4 billion hypertensive adults in the world, but the BP of less than 14 % is controlled with antihypertensive pharmacotherapy to a SBP/DBP $< 140/90$ mm Hg [5, 6]. One of the main causes of inadequate BP control is limited efficacy and safety profiles of the conventional antihypertensive drugs, which necessitates alteration of drug regimen or combination therapy [7, 8]. More than one-half of the patients need two or more antihypertensive drugs to achieve the goal BP [9]. Antihypertensive plants can be used as alternative or complementary therapies to conventional antihypertensive drugs [10]. Some plants found in Iran may have antihypertensive effects. This review aims to evaluate the clinical antihypertensive efficacy and safety of the plants existing in Iran.

2. Materials and Methods

2.1. Method of search and assessment of trial quality

This systematic review was conducted in accordance with the Preferred Items for Systematic Reviews and Meta-analyses (PRISMA) guidelines. The terms Iran, Iranian, plant, herb, antihypertensive, hypertension, randomized controlled trial (RCT) were searched in the databases PubMed, Medline, Scopus, Embase, ScienceDirect, ProQuest, Ovid, Ebsco, Google, and Google Scholar. References from relevant articles were searched manually. Also, the methodological quality of the trials was evaluated by the JADAD scale as described previously [11].

2.2. Eligibility of the articles

RCTs in the English language studying the antihypertensive efficacy and safety of the plants found in Iran were eligible. The articles published until the end of the year 2022 were included. Reviews, editorials and animal and in vitro studies were excluded.

2.3. Selection of articles

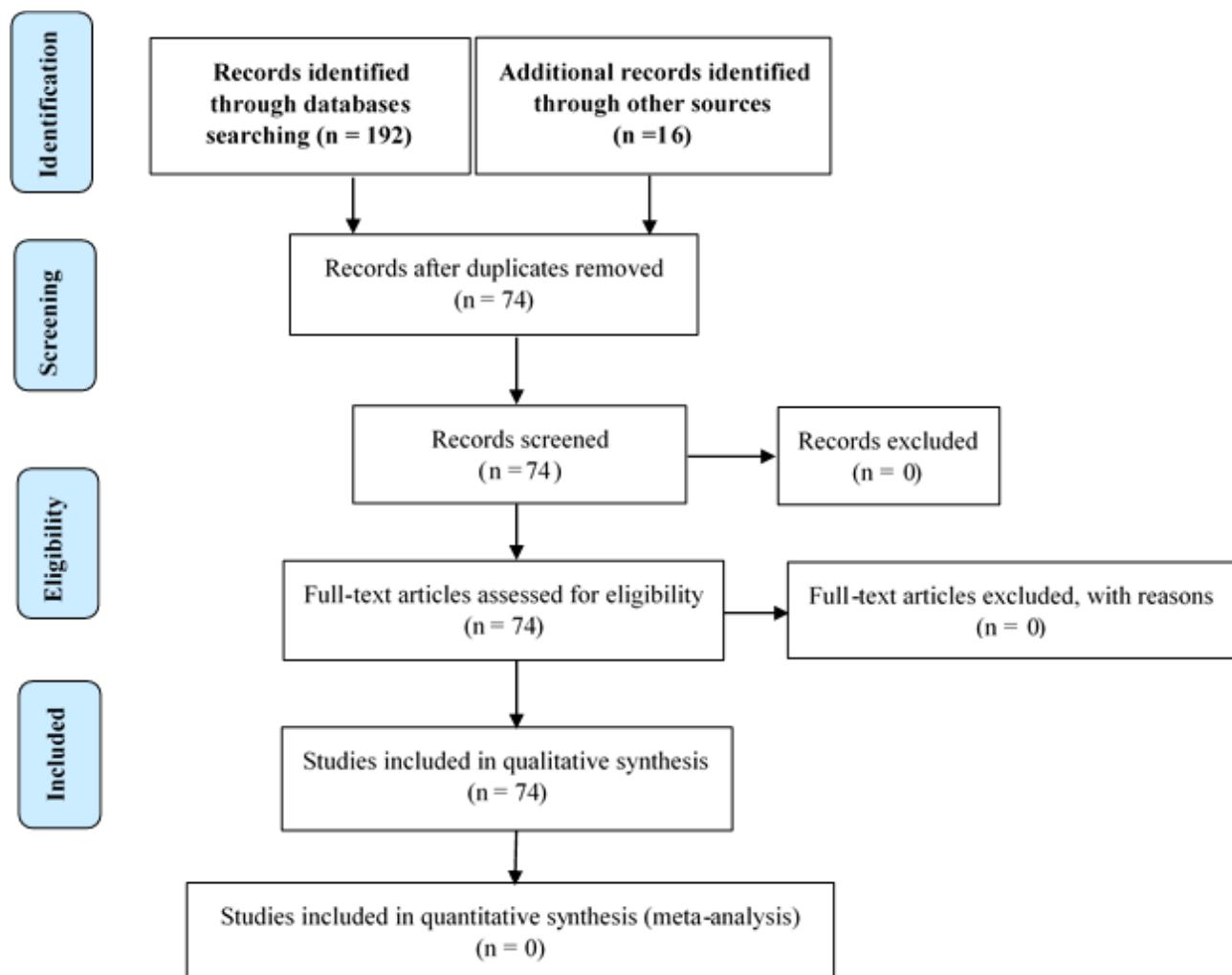
The authors independently screened the retrieved articles and resolved differences through discussion. The titles, abstracts and full texts of the retrieved articles were assessed.

2.4. Collation of data

The process of data collation is depicted in the PRISMA flow diagram (Fig. 1).

3. Results

The results were summarized and presented in the Table.

**Fig. 1.** The PRISMA flow diagram**Table 1.** Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran

Ref	JADAD score (Out of 5)	*Level of evidence/Stu- dy design/Participants/Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
<i>Berberis vulgaris</i>					
12	> 3	I/RCT/57 (19 <i>B. vulgaris</i> 1Tsp, 19 2Tsp /19 Po)/T2DM	Group1- n = 19, daily consumption of 1Tsp processed <i>B. vulgaris</i> in apple vinegar Group 2- n = 19, daily consumption of 2Tsp processed <i>B. vulgaris</i> in apple vinegar Group3- n = 19 with no change in their diet (Po) for 4 weeks	BP and inflammatory markers	NS* SBP/ DBP change compared with Po and BL No adverse effect reported

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/Stu dy design/Participants/Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
13	> 3	I/RCT/80 (40 <i>B. vulgaris</i> and 40 Po)/ NAFLD	Case group- n = 40 Two capsules (750 mg) including <i>B. vulgaris</i> extract every day for 3 months Control group n = 40 Po every day for 3 months	BP and weight	↓ SBP/ DBP compared with Po No adverse effect reported
14	> 3	I/RCT/101 (51 <i>B. vulgaris</i> 50 Po)/MetS and Aged between 18-65 years	Case group n = 51 receiving a capsule of barberry 600 mg/daily for 6 weeks Control group n = 50 receiving a Po capsule for 6 week s	Anthropometric measurements, BP and FBS	↓ SBP/ DBP compared with Po No adverse effect reported
15	> 3	I/RCT/(23) <i>B. vulgaris</i> and 23 Po)/T2DM	Barberry juice group n = 23 who consumed 200 ml of barberry juice daily for 8 weeks Control group n = 23 with no intervention for 8 weeks	BP and biochemical markers	↓ SBP/ DBP compared with BL and ↓ DBP compared with Po No adverse effect reported
16	> 3	I/RCT/80 (40 <i>B. vulgaris</i> and 40 Po)/T2DM and Age 20-65 years	Group1- n = 40, 1000mg dry extract (157.3 mg berberine per day) for 6 weeks Group2- n = 40, daily consumption of Po for 6 weeks	Blood glucose and Lipid profile levels	NS* SBP/ DBP change compared with Po and BL No adverse effect reported
17	> 3	I/RCT/70 (35 NS and 35 Po)/NHV	Group1- n = 35 received 2.5 ml NS oil BID for 3 months Group2- n = 35 received similarly 2.5 ml mineral oil BID for 3 months	SBP/DSB, BMI and blood levels of SGOT, ALT, ALP, Cr and BUN	↓ SBP/ DBP compared with Po and BL No adverse effect reported
18	> 3	I/RCT/20 (10 NS and 10 Po)/stage 1 HTN	NS group n = 10 receiving 1000 mg of powdered NS BID for 50 days Control group n = 10 receiving the same doses of Po for 50 days	SBP/DBP, lipid profile, BS, some anthropometric indicators	↓ SBP/DBP compared with Po. ↓ SBP compared with BL No adverse effect reported
19	< 3	I/NRCT/114/(57 NS and 57 Po)/T2DM	NS group n = 57 receiving 2 g of powdered NS daily for 1 year Control group n= 57 receiving the same doses of Po for 1 year	Lipid levels, BP and HR	↓ SBP/ DBP compared with BL and ↓ DBP compared with Po No adverse effect reported
21	> 3	I/RCT 119/ (36 NS, 39 NS and 33 Po)/Mild HTN	Group1- n = 36, 200 mg NS extract Group2- n = 39, 400 mg NS extract Group3- n = 33 Po for /8 weeks	HTN	↓ SBP, DBP in both groups compared with Po and BL No adverse effect reported

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/Stu dy design/Participants/Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
22	> 3	I/RCT/73(39 NS and 34 Po)/HCL	Group1- n = 39, 2 g NS powder Group2- n = 34 Po/ 6 weeks	BMI, waist-hip ratio, BP, FBS, serum lipids, and serum ALT and Cr	NS* SBP/DBP change compared with Po and BL No adverse effect reported
23	> 3	I/RCT/39 (19 NS and 20 Po)/obese Men	Group1- n = 19, 1.5 g NS powder Group2- n = 20 Po/ 12 weeks	BW, waist circumference, and BP, serum free testosterone, FBS, TG, HDL-Cholesterol, UA, Cr, SGOT and SGPT, adiponectin, and hs-CRP	↓ SBP compared with BL No adverse effect reported
24	> 3	I/RCT /40 (20 NS and 20 Po)/HAM	Group1- n=20 1g NS powder Group2 n= 20 Po /9 weeks	TC, TG and HDL cholesterol, VLDL, LDL cholesterol, CK-MB; AST, ALT, ALP, Total protein, Albumin, Bilirubin, Cr and BUN	NS* SBP/DBP change compared with Po and BL No adverse effect reported
25	> 3	I/RCT /80 (40 NS and 40 Po)/MetS	Group1- n= 40 NS powder 500mg Control group n= 40 Po /8 weeks Aspirin 150mg once a day was given in both groups.	FBG, PPBG & HbA1C at the beginning of the trial, the once every two weeks during the trial. Venous blood was also collected from each subject before and after the trial	↓ SBP, DBP compared with BL and control group No adverse effect reported
26	> 3	I/RCT /48 (24 NS and 24 Po)/HAM	Group1- n = 24 500 mg NS powder Group2- n = 24 Po /4 weeks	Cognition with CVLT-II, mood with Bond-Lader scale and anxiety with STAI	NS* SBP/DBP change compared with Po and BL No adverse effect reported
<i>Allium sativum</i>					
27	< 3	I/RCT /20 (In 1 group)/EHTN	Group1- n= 20 Garlic pearls 250mg/day for 2 months	Lipids and lipoprotein subfractions, plasma-ox-LDL, plasma and urinary concentration of 8-iso-PGF _{2α} and the TOS	↓ SBP/DBP compared with BL No adverse effect reported

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/ Study design/ Participants/ Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
28	> 3	I/RCT /62 (30 garlic 32 Po)/normo-lipidaemic volunteers	Group1- n = 30 Garlic powder (10.8 mg alliin) Group2- n = 32 Po/ 12 weeks	Serum lipids, BP and arterial stiffness	NS* SBP/DBP change compared with Po and BL No adverse effect ***reported
29	> 3	I/RCT /47 (24 garlic 23 Po)/mild HTN	Group1- n = 24 Garlic powder 600 mg Group2- n = 23 Po/ 12 weeks	BP and plasma lipids	↓ Supine DBP compared with BL No adverse effect reported
30	> 3	I/RCT /42 (garlic 23 and 19 Po)/normotensives mild HCL men	Group1- n = 23 Garlic powder 600 mg Group2- n = 19 Po/ 12 weeks	TC, LDL, HDL cholesterol	↓ SBP/DBP compared with Po No adverse effect reported
31	> 3	I/RCT /84 (garlic 30/18/16 and 20 Po)/mild & moderate HTN	One tablet containing 300 mg garlic powder BID, n = 30 or continued to receive a Po of identical appearance, n = 20 Some patients were randomly switched to the open-label branch and received either 2400 mg Allicor daily (2 tablets FID, n = 18 or 900 mg Kwai 1 tablet containing 300 mg TID, n=16/Po 12 weeks	TC, LDL, HDL cholesterol	↓ SBP in 480/960 compared with Po No adverse effect reported
32	> 3	I/RCT/189 (seven groups; A:300, B: 600, C: 900, D: 1200, E: 1500 mg garlic, F: 100 mg atenolol and G: Po n = 27/EHTN	Group A- n = 27, Garlic 300 mg Group B- n = 27, Garlic 600 mg Group C- n = 27, Garlic 900 mg Group D- n = 27, Garlic 1200 mg Group E- n = 27, Garlic 1500 mg Group F- n= 27, Atenolol 100 mg Group G- n = 27 Po/ 24 weeks	BP readings recorded at weeks 0, 12 and 24	↓ SBP, DBP compared with Po
33	> 3	I/RCT /50 (garlic 25 and 25 Po)/UHTN	Group1- n = 25 Garlic extract 960 mg Group2- n = 25 Po/ 12 weeks	SBP and DBP at baseline, 4, 8 and 12 weeks, and change over time	↓ SBP compared with Po No adverse effect reported
34	> 3	I/RCT /37 (garlic 16 and 23 Po)/pre HTN	Group1- n = 16, Garlic powder 600 mg Group2- n = 23 Po/ 12 weeks	BP were recorded at visits 1 and 2. Then all participants were further instructed to self-measure their BP at home during the 2-week interval between clinic visits 2 & 3	↓ SBP, DBP compared with Po No adverse effect reported

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/ Study design/ Participants/Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
35	> 3	I/RCT /88 (garlic 50 and 38 Po)/UHTN	Group1- n = 50 Aged- garlic extract 1.2 g Group2- n = 38 Po/ 12 weeks	BP, and secondary outcome measures of central- hemodynamics and other cardiovascular markers, including cholesterol, homocysteine, platelet function, and inflammatory markers	↓SBP, DBP compared with Po No adverse effect reported
36	> 3	I/RCT /49 (garlic 23 and 26 Po)/UHTN	Group1- n = 23 Aged- garlic extract 1.2 g Group2- n = 26 Po/ 12 weeks	BP, pulse wave velocity and arterial stiffness, inflammatory markers, and gut microbiota	↓SBP, DBP compared with Po No adverse effect reported
37	> 3	I/RCT /98 (garlic 23 and 26 Po)/ NAFLD	Group1- n = 47 Garlic tablet 400 mg (EC-coated tablet containing 1.5 mg Allicin) BID Group2- n = 51 Po (EC- coated tablet containing 400mg microcrystalline cellulose) BID/ 15 weeks	BP and hs-CRP	↓SBP, DBP compared with Po No adverse effect reported
<i>Hibiscus sabdariffa</i>					
38	> 3	I/RCT /54 (HS 31 and 23 Po)/moderate EHTN	Group1- n = 31 10g HS (decocotion) Group2- n = 23 Po/ 2 weeks	SBP and DBP were measured before and 15 days after the intervention	↓SBP, DBP compared with BL and control groups No adverse effect reported
39	> 3	I/RCT/ 7 5 (39 HS 36 Po)/ T2DM with mild HTN	Group1- n = 39 10 g HS (decocotion) Group2- n = 36 Po/ 4 weeks	tolerability, diastolic reduction > or = 10 mm Hg and, in the experimental group, urinary electrolytes modification	↓SBP, DBP compared with BL and Po groups No adverse effect reported
40	> 3	I/RCT /171 (were in HS and Po)/stage I or II HTN	Group1- n = 86 HS equivalent to 250 mg anthocyanin Control group n = 85 (10mg Lisinopril)/ 4 weeks	Effectiveness (DBP reduction ≥10 mmHg), Safety (absence of pathological modifications in the biochemical tests of hepatic & renal function), Tolerability (absence of intense side effects), effect on serum electrolytes, and ACE activity	↓SBP, DBP compared with BL and Po group No adverse effect reported

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/Stu dy design/Participants/Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
41	> 3	I/RCT /53 (27 HS 26 Po)/ T2DM with mild HTN	Group1- n = 27 HS 2 g (decoction) Group2- n = 26 Po (black tea)/ 4 weeks	BP was measured on days 0, 15 and 30 of the study	↓SBP compared with BL and Po group No adverse effect reported
42	< 3	I/RCT /20 (10 HS and 10 captopril)/mild HTN	Group1- n = 10 HS extract 1000 mg (250 mg anthocyanin) Control group n = 10, 25 mg Captopril/ 6 weeks	ABPM	↓SBP, DBP compared with BL No adverse effect reported
43	> 3	I/RCT /41 (20 HS 21 Po)/diabetic nephropathy	Group1- n = 20 HS extract 850mg Group2- n = 21 Po/ 8 weeks	BP and Urinary albumin concentration	↓SBP compared with BL No adverse effect reported
44	> 3	I/RCT /65 (35 HS 30 Po)/ pre and mild HTN	Group1- n = 35 HS 3.75g (decoction)/ Group2- n = 30 Po/ 6 weeks	A standardized method was used to measure BP at baseline and weekly intervals	↓SBP compared with Po group No adverse effect reported
45	> 3	I/RCT/36 (19 HS 17 Po)/T2DM with mild HTN	HSE-treated group n = 19 HS extract 900 mg Control group n = 17 Po/ 12 weeks	BMI, body fat, waist-to-hip ratio, and FFA	↓SBP compared with BL and Po group No adverse effect reported
46	> 3	I/RCT/50 (25 HS 25 Po)/mild to moderate HTN	Group1- n = 25 HS 9 g (decoction)/ Group2- n = 25 Po/ 4 weeks	BP, serum, and urine electrolytes were measured at baseline, weekly during treatment and 1 week after withdrawal of treatment	↓SBP, DBP compared with BL and Po group No adverse effect reported
47	> 3	I/RCT/35 (18 HS 17 Po)/MetS	Group1- n = 18 HS extract 500 mg Group2- n = 17 Po/ 4 weeks	SBP and DBP and BMI FBS, Insulin, lipoproteins, TG, hs-CRP, and MDA were determined pre- and post-intervention	↓SBP compared with Po group No adverse effect reported
48	> 3	I/RCT/33 (17 HS 16 Po)/healthy adult	Group1- n = 17 HS extract 450 mg	48	>3

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/Stu dy design/Participants/Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
<i>Beta vulgaris</i>					
49	N/A	47 intervention (n = 650) and 43 control (n = 598) groups.	2-56 days, 70-500 ml/d Beetroot juice supplemented	SBP, DBP	↓ SBP, DBP in nitrate-rich compared with control groups No adverse effect reported
50	> 3	I/RCT/ 12 (5 male, 7 female)/healthy older adults	3 h after ingestion 140 ml of nitrate-rich/ 140 ml nitrate-depleted beetroot juice	BP, blood coagulation, vascular inflammation markers, plasma nitrate and nitrite before, and 3 h and 6 h after ingestion	↓ SBP, DBP in nitrate-rich compared with BL No adverse effect reported
51	3	I/RCT/24 in either group/mild HTN	Group1- n = 24 250 ml Raw beetroot juice Group2- n = 24 250 g Cooked beetroot juice/ 2 weeks	SBP, DBP, FMD and TNF- α	↓ SBP, DBP compared with BL in both groups No adverse effect reported
52	> 3	I/RCT/40 (20 nitrate-rich and 20 nitrate-depleted beetroot juice)/ HTN pregnant women	Group1- n = 20 beetroot juice 70 ml Group2- n = 20 nitrate- depleted beetroot juice 70ml/ 8 day	BP, cardiovascular function and utero- placental blood flow	No overall reduction in BP in the nitrate- treated group; however there was a highly significant correlation between changes in plasma nitrite concentrations and changes in DBP in the nitrate-treated arm only
53	> 3	I/RCT/18 cross-over/untreated HTN	Group-1 n = 18 consumed randomly, a nitrate-rich (8.1 mmol- BRJ nitrate) and a nitrate-depleted (BRJ placebo) BRJ	Participants performed submaximal isometric handgrip with beat-by-beat monitoring of hemodynamics and cBRs. AMBP assessment followed.	Office/ambulatory BP were lower following BRJnitrate vs BRJpo No adverse effect reported
54	< 3	II/RCT/30 (10 in each group)/ healthy non-smoking men and women, aged 55–70 years, with a BMI between 25 and 40 kg/m ²	Group1- n = 10 beetroot juice Group2- n = 10 r isometric handgrip exercise Group3- n=10 control/7 days	Clinic and 24-h ABP, peripheral arterial function quantified by pulse wave velocity and arterial volume distensibility were assessed before and after intervention.	No change in SBP and DBP No adverse effect reported

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/ Study design/ Participants/ Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
55	> 3	I/RCT/14/ non-hypertensive obese individuals	Fourteen were randomly assigned to 3 experimental sessions: 1) Beetroot juice with exercise (BJE, 200 ml with \approx 800 mg nitrate and 40 minutes of moderate-intensity aerobic exercise at an intensity of 50% of the heart rate reserve), 2) fruit soda with exercise (FSE, 200ml of a low-nitrate drink and the same exercise session) and 3) control (CON, 200ml of water, an insignificant nitrate drink without exercise).	Subjects were instructed to shower after each experimental session. ABMP device was fitted on their non-dominant arm. They were fitted with the ABPM device ~60 minutes after the experimental sessions and had it removed on the following day. The device was programmed to measure BP every 15 minutes while the subject was awake and every 30 minutes during their periods of sleep	NS* changes were observed for ambulatory DBP No adverse effect reported
56	< 3	II/RCT/21 (10 & 11) subjects completing the study	56	< 3	II/RCT/21 (10 & 11) subjects completing the study
51	< 3	II/RCT/24 twelve raw beet juice and 12 cooked beet/ HTN	24 hypertensive subjects aged 25-68 years	51	< 3
57	< 3	I/RCT/ 15 / patients with PAH	Group 1- n=15 The patients received nitrate-rich beetroot juice (~16 mmol nitrate per day) and Po in 2 treatment periods of 7 days each.	The primary outcome: change in peak oxygen consumption (VO ₂ peak) and VO ₂ at the anaerobic threshold. The secondary outcome: changes in; the 6-minute walking test, WHO-functional class, right and left ventricular function, right and left atrial/ventricular dimensions, systolic pulmonary artery pressure, exhaled NO, systemic BP, N-terminal pro-brain natriuretic peptide, biochemical variables involved in the NO system, a range of standard variables obtained from the ergo-spirometry PAH	SBP and DBP did not differ between interventions No adverse effect reported

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/ Study design/ Participants/ Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
58	> 3	I/RCT/64 NHV	Group1- n = 32 receive daily dietary supplementation with dietary nitrate (250 ml daily, as beetroot juice) Group2- n = 32 Po (250 ml daily, as nitrate-free beetroot juice)/ 4 weeks	ABP, BP, and HR	Robust BP lowering No adverse effect reported
59	> 3	I/RCT/ 47 middle-aged and older participants	Group1- n = 16 Combined intervention (high-nitrate beetroot juice and folic acid) Group2- n = 16 Single intervention (high-nitrate beetroot juice and Po) Group3- n = 15 control (nitrate-depleted beetroot juice and Po)/ 60 days	Clinic and 24-h ambulatory BP and measurements of compliance in plasma (nitrate and folate concentrations) and saliva (nitrate and nitrite) were obtained at baseline, 30 d, and 60 d	↓ BP No adverse effect reported
60	> 3	I/RCT/ 87/ patients with/at risk of T2DM	Group1- n = 27 Doxazocin + Po juice Group2- n = 16 Doxazocin + Active beetroot juice Group3- n = 20 Spironolactone + Po juice Group4- n = 24 Spironolactone + Active beetroot juice/3 and 6 months	Haemodynamic parameters, Echocardiographic morphological Parameters, and Echocardiographic Systo-Diastolic function	BP did not differ between the juices, or between the drugs. However, 6 months' dietary nitrate decreased LV volumes ~5% No adverse effect reported
61	> 3	I/RCT/ 87/ UHTN	Group1- n = 20 (13 + 7) 7-d, double-blind, randomized, Po - controlled, cross-over trial to assess the effect of dietary nitrate. Subjects were tested on three separate occasions – baseline (day 1), midpoint (day 8) and endpoint (day 15) – before and after each intervention period	On all 3 testing days (days 1, 8 and 15) in an identical manner and at the same time of day, non-fasting blood was drawn and subjects were fitted with an ABPM for 24 h	It is noteworthy that BP values decreased after Po, as well as after NO ₃ – No adverse effect reported
62	> 3	I/RCT/27/HTN men and women	Group1 – n = 27 The effect of 1-week intake of nitrate-rich beetroot juice was compared with 1-week intake of nitrate-depleted beetroot juice (Po)	The primary outcome was BP assessed by measuring home BP during the intervention and 24-h AMBP on day 7 of the intervention. Other outcomes included nitrate metabolism assessed by measuring nitrate and nitrite in plasma, saliva, and urine	An increase in dietary nitrate intake may not be an effective short-term approach to further lower BP in treated hypertensive subjects No adverse effect reported

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/ Study design/ Participants/ Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
<i>Solanumly copersicum</i>					
63	< 3	I/RCT/31/grade-1 HTN	Group1-n=31 Po period/ 4 weeks The same group (Group1) n =31 TE 250mg/ 8 weeks	SBP, DBP	↓SBP, DBP in TE group compared with BL No adverse effect reported
64	> 3	I/RCT/25 pre-HTN 15 TE, 11 dark chocolate, 10 Po	Group1- n= 11 Fifty grams daily dose of dark chocolate with 70% cocoa containing 750mg polyphenols Group2- n= 15 Were allocated one tomato extract capsule containing 15mg lycopene per day, and Control group n= 10 received 1 placebo capsule daily/ Over 8 weeks followed by a 4- week washout period.	Median BP, weight, and abdominal circumference	No change in SBP and DBP No adverse effect reported
65	> 3	I/RCT /50 (26 TE, 24 Po)/uncontrolled HTN	Group1- n= 26 TE 250mg (15 mg lycopene) Group2- n=24 Po/ 6 weeks	Plasma concentrations of lycopene, nitrite and nitrate	↓SBP, DBP in TE group compared with BL No adverse effect reported
66	> 3	I/RCT/126 (41 lycopen 6 mg; 37 lycopen 15 mg; 38 Po)/NHV	Group1- n= 41 Lycopen 6 mg Group2- n= 37 Lycopen 15 mg Group3- n= 38 Po/ 8 weeks	hs-CRP, SBP, sICAM-1 and sVCAM-1 β- carotene and LDL	↓SBP in 15 mg Lycopen group with BL No adverse effect reported
67	> 3	I/RCT /24 in either group/Mild HTN	Group1- n= 24 Synthetic lycopene 15mg Group2- n=24 Po/ 8 weeks	BP	↓SBP compared with control groups No adverse effect reported

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/ Study design/ Participants/ Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
<i>Cinnamomum verum</i>					
68	> 3	I/RCT /22 (12 in cinnamon 10 in Po)/Pre-DM	Group1- n = 12 Cinnamon extract 250 mg TID Group2- n = 10 Po/ 12 Weeks	FBG, SBP, and Body composition	↓ SBP No change in DBP No adverse effect reported
69	> 3	I/RCT /58 (30 in cinnamon 28 in Po)/T2DM	Group1- n = 30 Cinnamon 2g daily Group2- n = 28 Po/ 12 Weeks	FBG, SBP, DBP, HbA1c	↓ SBP, DBP compared with Po No adverse effect reported
70	> 3	I/RCT/59 (29 in cinnamon 30 in Po)/T2DM	Group1- n = 29 Cinnamon extract 400 mg TID Group2- n = 30 Po/ 12 Weeks	BP, HbA1c, FBG, lipid profile, physical examination, and Blood and Urine chemistry	↓ SBP ↓DBP compared with Po and BL No adverse effect reported
71	> 3	I/RCT /37 T2DM 19 in cinnamon 18 in Po	Group1- n = 19; 2 capsule each contain 500 mg Cinnamon powder TID (3g daily) Group2- n = 18 Po/ 8 weeks	Weight, height, body fat mass, SBP and DBP	No change in SBP and DBP No adverse effect reported
72	> 3	I/RCT/ 135 (63 in cinnamon 72 in Po)/Hyperglycemic individual	Group1- n = 63 Cinnamon extract 250 mg BID Group2- n=72 Po/ 8 Weeks	FBG, SBP and DBP, serum lipids, and Fructosamine.	No change in SBP and DBP No adverse effect reported
73	> 3	I/RCT /79 (40 cinnamon/39 Po)/T2DM	Group1- n = 40 Cinnamon 3g daily + Black tea Control group n =39 Po + Black tea/ 8 Weeks	ICAM-1, SBP, DBP and Anthropometric measures	No change in SBP and DBP No adverse effect reported
74	> 3	I/RCT /99 (49 cinnamon/50 Po)/T2DM	Group1- n = 49 Cinnamon extract 500 mg TID Group2- n = 50 Po/ 8 Weeks	Glucose, TG, HDL-C levels, TG/HDL-C ratio, BP, and eGFR	↓ SBP ↓DBP compared with Po and BL No adverse effect reported
75	> 3	I/RCT /116 (58 cinnamon/58 Po /MetS	Group1- n = 58 Cinnamon powder 3g daily Control group n = 58 Po/ 16 Weeks	Body composition, BP and Metabolic parameters	↓ SBP ↓DBP compared with Po and BL No adverse effect reported
76	> 3	I/RCT /36 (18 in cinnamon 18 in Po)/RA women	Group1- n = 18 Cinnamon capsule 500 mg FID Control group n = 18 Po/ 8 Weeks	FBS, lipid profile, liver enzymes, serum levels of CRP, TNF- α , ESR, BP, and Clinical symptoms	↓ SBP ↓DBP compared with Po and BL No adverse effect reported

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/ Study design/ Participants/ Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
<i>Rhus coriaria</i>					
77	> 3	I/RCT /80/HTN	Group1- n = 40 received <i>R. Coriaria</i> capsules (500mg BID) and captopril (25mg daily) Group2- n = 40 Po capsules (500mg starch BID) and captopril (25mg daily)/ 8 weeks	BP and BMI	↓ SBP ↓DBP compared with Po and BL No adverse effect reported
<i>Olea europaea</i>					
78	> 3	I/RCT/64 (32 in OLE 32 Po)/mild to moderate HTN	Group1- n = 32 OLE 500mg Group2- n = 32 Po/ 8 Weeks	Risk factors of atherosclerosis and co-morbid medical conditions, SBP, DBP, and HR	↓SBP in OLE group compared with Po and BL. ↓DBP compared with Po No adverse effect reported
79	< 3	I/RCT /40 monozygotic twins/ borderline HTN	Group1- n = 10; 500 Olive leaf extract Control group1- n = 10; No medication but advice on how HTN be ameliorated by an adequate lifestyle Group2- n = 10; 500 Olive leaf extract Control group2- n = 10; No medication but advice on how HTN be ameliorated by an adequate lifestyle/ 8 Weeks	Body weight, HR, BP, glucose and lipids	↓SBP, DBP in 1000 mg OLE group compared with BL No adverse effect reported
80	> 3	I/RCT /148 stage-1 HTN/ 72 olive leaf extract 76 captopril	Group1- n = 72; 500 mg Olive leaf extract BID Control group n = 76 /12.5 mg captopril BID/ 8 Weeks	SBP/DBP	↓SBP, DBP in 1000 mg OLE group and 25 mg Captopril compared with BL No adverse effect reported
81	< 3	II/Non-controlled, non-randomized pilot study/663/T2DM and pre T2DM with Grade 1 HTN	100 mg/d of Oleuropein and 20 mg/d of Hydroxytyrosol BID	SBP/DBP	↓SBP, DBP in TE group compared with BL No adverse effect reported
<i>Vaccinium arctostaphylos</i>					
82	> 3	I/RCT /100 50 <i>Vaccinium arctostaphylos</i> /50 Po/ HTN, T2DM with hyperlipidemia	Group1- n = 50 <i>Vaccinium arctostaphylos</i> leaf extract 350 mg TID Control group n = 50 Po/ 2 Months	BP, FG, 2hPPG, HbA1c, TC, LDL-C, TG, HDL-C, SGOT, SGPT and Cr	↓SBP, DBP in leaf extract group compared with Po and BL No adverse effect reported
83	> 3	I/RCT /39 (in one group)/ T2DM with HTN	Group1- n = 39; 1, 2, 3 and 4 weeks Decoction of 7 g <i>V. arctostaphylos</i> oral solution	FBS and BP	↓SBP, DBP compared with BL Not reported any harmful effect

Table 1. Summary of the clinical trials conducted on hypertensive patients to evaluate antihypertensive efficacy and safety of plants found in Iran (Continued)

Ref	JADAD score (Out of 5)	*Level of evidence/ Study design/ Participants/ Inclusion criteria	Intervention/Control group	Outcome measure	Results / Safety
84	> 3	I/RCT/100 (50 <i>Vaccinium arctostaphylos</i> /50 Po)/HTN	Extract group n = 50 One extract capsule TID alongside the standard anti-hypertensive treatments for 3 months Po group n = 50 One Po capsule TID alongside the standard anti-hypertensive treatments for 3 months	SBP, DBP, BMI, and waist circumference CBC, blood levels of AST, ALT, ALP, BUN, and Cr	↓ SBP, ↓ DBP compared with Po No adverse effect reported
<i>Phyllanthus emblica</i>					
85	> 3	I/RCT/81 (<i>Ph.. emblica</i> group n = 41 and Po group n = 40)/uncontrolled HTN	EO and Po groups took 500 mg extract and Po respectively TDS after meal with standard anti-hypertensives for 8 weeks	SBP, DBP	↓ SBP/DBP compared with Po. No adverse effect reported

4. Discussion

Hypertension is the leading cause of cardiovascular morbidity and mortality. It causes 9.4 million deaths annually in the world. The global prevalence of hypertension will increase 30 % by the year 2025. Effective treatment of hypertension is a major development in medicine. Development of numerous antihypertensive drugs have extended life expectancy and reduced complications of hypertension. Medicinal plants are considered as one of the modalities for the treatment of hypertension [5]. In this systematic review, the antihypertensive efficacy and safety of the plants found in Iran as evaluated in randomized controlled trials were examined. Most clinical trials as cited in the Table had Jadad scores larger than 3, showing high methodological quality. Virtually all the medicinal plants listed in the Table were able to reduce SBP and DBP. *Hibiscus sabdariffa* was able to reduce SBP and DBP in all studies. Although, the number of studies on *Olea europaea*, *Rhus coriaria*, and

Vaccinium arctostaphylos was small, but they showed antihypertensive efficacy. The results of studies of other plants were not as consistent as studies of these plants; i.e., the results of some studies showed antihypertensive efficacy, while some other studies did not show efficacy in lowering BP. Among these studies, most studies on *Allium sativum* exhibited antihypertensive efficacy. The studies on *Cinnamomum verum* did not show consistent results. One study concluded that it was able to lower only SBP. *Beta vulgaris* and *Solanum lycopersicum* had conflicting effects on hypertension. No adverse effect was reported in virtually all clinical trials.

Systemic reflexes, namely the baroreceptor neural reflex and the renin-angiotensin-aldosterone hormonal response, control the BP. The reflexes regulate BP via action on the hemodynamic factors. As such, BP equals cardiac output (CO) multiplied by peripheral vascular resistance (PVR). In turn, CO and PVR can be subdivided into their determinants (Fig. 2) [86, 87]:

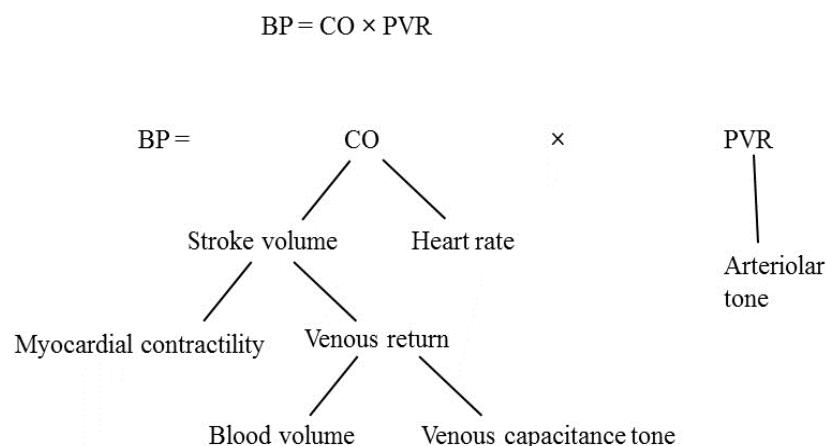


Fig. 2. A schema on the hemodynamic factors involved in blood pressure regulation and hypertension.

BP is a function of peripheral vascular resistance and cardiac output. Hypertension can be depicted in terms of its cardiovascular hemodynamics (Fig. 2). Increased vascular resistance or cardiac output raise BP and cause hypertension. Therefore, antihypertensive plants lower BP by decreasing vascular resistance or/and cardiac output through various mechanisms [86, 87]. Delineation of the mechanisms and sites of antihypertensive action of plants is important for optimal use of plants, for understanding the pathophysiology of hypertension and for development of new drugs [86, 87].

This systematic review suggests that limited clinical trials have been conducted on the antihypertensive efficacy and safety of most plants. There are more clinical evidence regarding the antihypertensive efficacy and safety of *Hibiscus sabdariffa*, *Olea europaea*, *vaccinium arctostaphylos* and *Allium sativum* versus the other plants. These plants seem to have efficacy and safety for the treatment of hypertension.

5. Conclusion

Most clinical trials evaluating the antihypertensive effects of the plants had high methodological quality. Most clinical trials demonstrated antihypertensive efficacy and no adverse effect was reported for virtually all plants. Most clinical trials show promising antihypertensive efficacy and safety of the plants. There are more clinical evidence denoting antihypertensive efficacy and safety of *Hibiscus sabdariffa*, *Olea europaea*, *vaccinium arctostaphylos* and *Allium sativum* compared with the other plants.

Author contributions

SK conceived the title of study. HFH, BF and SK searched for the clinical trials, and wrote the manuscript.

Conflicts of interest

The authors declare that there is no conflict of interest.

References

1. Afzal M. Recent updates on novel therapeutic targets of cardiovascular diseases. *Mol. Cell. Biochem.* 2021; 476(1): 145-155. doi: 10.1007/s11010-020-03891-8.
2. Oparil S, Acelajado MC, Bakris GL, Berlowitz DR, Cifkova R, Dominiczak AF, Grassi G, Jordan J, Poulter NR, Rodgers A and Whelton PK. Hypertension. *Nat. Rev. Dis. Primers.* 2018; 4: 18014. doi: 10.1038/nrdp.2018.14.
3. Taler SJ. Initial treatment of hypertension. *N. Engl. J. Med.* 2018; 378(7): 636-644. doi: 10.1056/NEJMcp1613481.
4. Volpe M, Gallo G, Battistoni A and Tocci G. Highlights of ESC/ESH 2018 guidelines on the management of hypertension: what every doctor should know. *High Blood Press. Cardiovasc. Prev.* 2019; 26: 1-8. doi: 10.1007/s40292-018-00297-y.
5. Al-Makki A, DiPette D, Whelton PK, Murad MH, Mustafa RA, Acharya S, Beheiry HM, Champagne B, Connell K, Cooney MT, Ezeigwe N, Gaziano TA, Gidio A, Lopez-Jaramillo P, Khan UI, Kumarapeli V, Moran AE, Silwimba MM, Rayner B, Sukonthasan A, Yu J, Saraffzadegan N, Reddy KS and Khan T. Hypertension Pharmacological Treatment in Adults: A World Health Organization Guideline Executive Summary. *Hypertension* 2022; 79: 293-301. doi: 10.1161/HYPERTENSIONAHA.121.18192.
6. Bosch A and Schmieder RE. Novel approaches to management of hypertension. *Curr. Opin. Nephrol. Hypertens* 2021; 30(1): 54-62. doi: 10.1097/MNH.0000000000000668.
7. Gorostidi M and de la Sierra A. Combination therapies for hypertension - why we need to look beyond RAS blockers. *Expert Rev. Clin. Pharmacol.* 2018; 11(9): 841-853. doi: 10.1080/17512433.2018.1509705.
8. Guerrero-Garcia C and Rubio-Guerra AF. Combination therapy in the treatment of hypertension. *Drugs Context* 2018; 7: 212531. doi: 10.7573/dic.212531.
9. Chobanian AV. Guidelines for the management of Hypertension. *Med. Clin. North Am.* 2017; 101(1): 219-227. doi: 10.1016/j.mcna.2016.08.016.
10. Verma T, Sinha M, Bansal N, Yadav SR, Shah K and Chauhan NS. Plants used as antihypertensive. *Nat. Prod. Bioprospect.* 2021; 11(2): 155-184. doi: 10.1007/s13659-020-00281-x.
11. Jadad AR, Moore RA, Carroll D, Jenkinson C, Reynolds DJ, Gavaghan DJ and McQuay HJ. Assessing the quality of reports of randomized clinical trials: is blinding necessary? *Control Clin. Trials.* 1996; 17(1): 1-12. doi: 10.1016/0197-2456(95)00134-4.
12. Golzarand M, Ebrahimi Mamaghani M, Arefhosseini SR and Aliasgarzadeh A. Short term-effect of processed berberis consumption vulgaris on cardiovascular risk factors in type II diabetes patients with metabolic syndrome. *Med. J. Tabriz Uni. Med. Sciences Health Services* 2009; 31(2): 89-94.
13. SalehZadeh H, Iloun kashkooli R, Najafi SS, Hosseini Asl MK, Hamed A and Kalateh Sadati A. The Effect of *Berberis vulgaris* extract on blood pressure and weight of the patients suffered from non-alcoholic fatty liver

disease. *J. Res. Dev. Nurs. Midw.* 2013; 10(Supplementary): 21-7.

14. Zilaee M, Safarian M, Kermany T, Emamian M, Mobarhan MG and Ferns GAA. Effect of barberry treatment on blood pressure in patients with metabolic syndrome. *J. Natural. Products* 2015; 8: 59-63.

15. Lazavi F, Mirmiran P, Sohrab G, Nikpayam O, Angoorani P and Hedayati M. The barberry juice effects on metabolic factors and oxidative stress in patients with type 2 diabetes: a randomized clinical trial. *Complement. Ther. Clin. Pract.* 2018; 31: 170-174. doi: 10.1016/j.ctcp.2018.01.009.

16. Tahmasebi L, Zakerkish M, Golafkhrabadi F and Namjoyan F. Randomised clinical trial of *Berberis vulgaris* root extract on glycemic and lipid parameters in type 2 diabetes mellitus patients. *European J. Integrative. Medicine* 2019; 32: 1000998. doi: 10.1016/j.eujim.2019.100998.

17. Fallah Huseini H, Amini M, Mohtashami R, Ghamarchehre ME, Sadeghi Z, Kianbakht S and Fallah Huseini A. Blood pressure lowering effect of *Nigella sativa* L. seed oil in healthy volunteers: a randomized, double-blind, placebo-controlled clinical trial. *Phytother. Res.* 2013; 27(12): 1849-53. doi: 10.1002/ptr.4944.

18. Saumi R and Bukhari A. Phytotherapy *Nigella sativa* lowers blood pressure in patients with stage 1 hypertension. *J. Hypertension* 2015; 33(pe43). doi: 10.1097/01.hjh.0000469884.66111.bc.

19. Badar A, Kaatabi H, Bamosa A, Al-Elq A, Abou-Hozaifa B, Lebda F, Alkhadra A and Al-Almaie S. Effect of *Nigella sativa* supplementation over a one-year period on lipid levels, blood pressure and heart rate in type-2 diabetic patients receiving oral hypoglycemic agents: nonrandomized clinical trial. *Ann. Saudi. Med.* 2017; 37(1): 56-63. doi: 10.5144/0256-4947.2017.56.

20. Rizka A, Setiati S, Lydia A and Dewiasty E. Effect of *Nigella sativa* seed extract for hypertension in elderly: a double-blind, randomized controlled trial. *Acta Med. Indones* 2017; 49(4): 307-313.

21. Roghani Dehkordi F and Kamkhah AF. Antihypertensive effect of *Nigella sativa* seed extract in patients with mild hypertension. *Fundam. Clin. Pharmacol.* 2008; 22(4): 447-52. doi: 10.1111/j.1472-8206.2008.00607.x.

22. Qidwai W, Hamza HB, Qureshi R and Gilani AH. Effectiveness, safety, and tolerability of powdered *Nigella sativa* (kalonji) seed in capsules on serum lipid levels, blood sugar, blood pressure, and body weight in adults: results of a randomized, double-blind controlled trial. *J. Altern. Complement. Med.* 2009; 15(6): 639-44. doi: 10.1089/acm.2008.0367.

23. Datau EA, Wardhana, Surachmanto EE, Pandelaki K, Langi JA and Fias. Efficacy of *Nigella sativa* on serum free testosterone and metabolic disturbances in central obese male. *Acta Med. Indones* 2010; 42(3): 130-4.

24. Bin Sayeed MS, Asaduzzaman M, Morshed H, Hossain MM, Kadir MF and Rahman MR. The effect of *Nigella sativa* Linn. seed on memory, attention and cognition in healthy human volunteers. *J. Ethnopharmacol.* 2013; 148(3): 780-6. doi: 10.1016/j.jep.2013.05.004.

25. Najmi A, Nasiruddin M, Khan RA and Haque SF. Indigenous herbal product *Nigella*

sativa proved effective as an antihypertensive in metabolic syndrome. *Asian J. Pharm. Clin. Res.* 2013; 6(1): 61-64.

26. Bin Sayeed MS, Shams T, Fahim Hossain S, Rahman MR, Mostofa A, Fahim Kadir M, Mahmood S and Asaduzzaman Md. *Nigella sativa* L. seeds modulate mood, anxiety and cognition in healthy adolescent males. *J. Ethnopharmacol.* 2014; 152(1): 156-62. doi: 10.1016/j.jep.2013.12.050.

27. Dhawan V and Jain S. Effect of garlic supplementation on oxidized low density lipoproteins and lipid peroxidation in patients of essential hypertension. *Mol. Cell. Biochem.* 2004; 266: 109-15. doi: 10.1023/ b:mcbi.0000049146.89059.53.

28. Turner B, Molgaard C, and Markmann P. Effect of garlic (*Allium sativum*) powder tablets on serum lipids, blood pressure and arterial stiffness in normo-lipidaemic volunteers; a randomised, double-blind, placebo-controlled trial. *Br. J. Nutr.* 2004; 92(4): 701-706. doi: 10.1079/bjn20041255.

29. Auer W, Eiber A, Hertkorn E, Hoehfeld E, Koehrle U, Lorenz A, Mader F, Merx W, Otto G, Schmid-Otto B and et al. Hypertension and hyperlipidaemia: garlic helps in mild cases. *Br. J. Clin. Pract. Suppl.* 1990; 69: 3-6.

30. Sobenin IA, Andrianova IV, Demidova ON, Gorchakova T and Orekhov AN. Lipid-lowering effects of time-released garlic powder tablets in double-blinded placebo-controlled randomized study. *J. Atheroscler. Thromb.* 2008; 15(6): 334-8. doi: 10.5551/jat.e550.

31. Sobenin IA, Andrianova IV, Fomchenkov IV, Gorchakova TV and Orekhov AN. Time-released garlic powder tablets lower systolic and diastolic blood pressure in men with mild

and moderate arterial hypertension. *Hypertens. Res.* 2009; 32: 433-7. doi: 10.1038/hr.2009.36.

32. Ashraf R, Khan RA, Ashraf I and Qureshi AA. Effects of *Allium sativum* (garlic) on systolic and diastolic blood pressure in patients with essential hypertension. *Pak. J. Pharm. Sci.* 2013; 26(5): 859-63.

33. Ried K, Frank OR and Stocks NP. Aged garlic extract lowers blood pressure in patients with treated but uncontrolled hypertension: a randomised controlled trial. *Maturitas* 2010; 67(2): 144-50. doi: 10.1016/j.maturitas.2010.06.001.

34. Nakasone Y, Nakamura Y, Yamamoto T and Yamaguchi H. Effect of a traditional Japanese garlic preparation on blood pressure in prehypertensive and mildly hypertensive adults. *Exp. Ther. Med.* 2013; 5(2): 399-405. doi: 10.3892/etm.2012.819.

35. Ried K, Travica N and Sali A. The effect of aged garlic extract on blood pressure and other cardiovascular risk factors in uncontrolled hypertensives: the AGE at heart trial. *Integr. Blood Press Control* 2016; 9: 9-21. doi: 10.2147/IBPC.S93335.

36. Ried K, Travica N and Sali A. The Effect of kyolic aged garlic extract on gut microbiota, inflammation, and cardiovascular markers in hypertensives: the GarGIC trial. *Front. Nutr.* 2018; 5: 122. doi: 10.3389/fnut.2018.00122.

37. Soleimani D, Parisa Moosavian S, Zolfaghari H and Paknahad Z. Effect of garlic powder supplementation on blood pressure and hs-C-reactive protein among nonalcoholic fatty liver disease patients: a randomized, double-blind, placebo-controlled trial. *Food Sci. Nutr.* 2021; 9(7): 3556-3562. doi: 10.1002/fsn3.2307.

38. Haji Faraji M and Haji Tarkhani A. The effect of sour tea (*Hibiscus sabdariffa*) on essential hypertension. *J. Ethnopharmacol.* 1999; 65(3): 231-6. doi: 10.1016/s0378-8741(98)00157-3.

39. Herrera-Arellano A, Flores-Romero S, Chavez-Soto MA and Tortoriello J. Effectiveness and tolerability of a standardized extract from *Hibiscus sabdariffa* in patients with mild to moderate hypertension: a controlled and randomized clinical trial. *Phytomedicine* 2004; 11(5): 375-82. doi: 10.1016/j.phymed.2004.04.001.

40. Herrera-Arellano A, Miranda-Sanchez J, Avila-Castro P, Herrera-Alvarez S, Jimenez-Ferrer JE, Zamilpa A, Roman-Ramos R, Ponce-Monter H and Tortoriello J. Clinical effects produced by a standardized herbal medicinal product of *Hibiscus sabdariffa* on patients with hypertension. A randomized, double-blind, lisinopril-controlled clinical trial. *Planta Med.* 2007; 73(1): 6-12. doi: 10.1055/s-2006-957065.

41. Mozaffari-Khosravi H, Jalali-Khanabadi BA, Afkhami-Ardekani M, Fatehi F and Noori-Shadkam M. The effects of sour tea (*Hibiscus sabdariffa*) on hypertension in patients with type II diabetes. *J. Hum. Hypertens.* 2009; 23: 48-54. doi: 10.1038/jhh.2008.100.

42. Soleimani AR, Akbari H, Soleimani S, Beladi Mousavi SS and Tamadon MR. Effect of sour tea (Lipicom) pill versus captopril on the treatment of hypertension. *J. Renal. Inj. Prev.* 2015; 4(3): 73-9. doi: 10.12861/jrip.2015.15.

43. Najjarzade A, Hemayati R, Zavar Reza J, Fallahzade H, Mozaffari-Khosravi H, Taghizadeh M and Esmaeili A. Effects of *Hibiscus sabdariffa* on albuminuria and hypertension in patients with diabetic nephropathy. *The J. Toloo-e-behdash.* 2016; 14: 107-118.

44. McKay DL, Chen CY, Saltzman E and Blumberg JB. *Hibiscus sabdariffa* L. tea (tisane) lowers blood pressure in prehypertensive and mildly hypertensive adults. *J. Nutr.* 2010; 140(2): 298-303. doi: 10.3945/jn.109.115097.

45. Chang HC, Peng CH, Yeh DM, Kao ES and Wang CJ. *Hibiscus sabdariffa* extract inhibits obesity and fat accumulation, and improves liver steatosis in humans. *Food Funct.* 5(4): 734-9. doi: 10.1039/c3fo60495k.

46. Nwachukwu DC, Aneke E, Nwachukwu NZ, Obika LF, Nwagha UI and Eze AA. Effect of *Hibiscus sabdariffa* on blood pressure and electrolyte profile of mild to moderate hypertensive Nigerians: a comparative study with hydrochlorothiazide. *Niger. J. Clin. Pract.* 2015; 18(6): 762-70. doi: 10.4103/1119-3077.163278.

47. Asgary S, Soltani R, Zolghadr M, Keshvari M and Sarrafzadegan N. Evaluation of the effects of roselle (*Hibiscus sabdariffa* L.) on oxidative stress and serum levels of lipids, insulin and hs-CRP in adult patients with metabolic syndrome: a double-blind placebo-controlled clinical trial. *J. Complement. Integr. Med.* 2016; 13(2): 175-80. doi: 10.1515/jcim-2015-0030.

48. Kafeshani M, Entezari MH, Karimian J, Pourmasoumi M, Maracy MR, Amini MR and Hadi A. A comparative study of the effect of green tea and sour tea on blood pressure and lipid profile in healthy adult men. *ARYA Atheroscler.* 2017; 13(3): 109-116.

49. Bahadoran Z, Mirmiran P, Kabir A, Azizi F and Ghasemi A. (2017). The Nitrate-independent blood pressure-lowering effect of beetroot juice: a systematic review and meta-analysis. *Adv. Nutr.* 2017; 8(6): 830-838. doi: 10.1093/advances/nmy004.

50. Raubenheimer K, Hickey D, Leveritt M, Fassett R, Ortiz de Zevallos Munoz J, Allen JD, Briskey D, Parker TJ, Kerr G, Peake JM, Pechenik NM and Neubauer O. Acute effects of nitrate-rich

beetroot juice on blood pressure, hemostasis and vascular inflammation markers in healthy older adults: a randomized, placebo-controlled crossover study. *Nutrients* 2017; 9(11): doi: 10.3390/nu9111270.

51. Asgary S, Afshani MR, Rafieian-Kopaei M and Keshvari M. Clinical effects of consumption of raw beet juice and beet cooked on improving blood pressure, FMD and inflammatory cytokines TNF- α level of blood pressure on hypertensive patients volunteer. *J. Shahrekord Univ. Med. Sci.* 2017; 19(2): 148-157.

52. Ormesher L, Myers JE, Chmiel C, Wareing M, Greenwood SL, Tropea T, Lundberg JO, Weitzberg E, Nihlen C, Sibley CP, Johnstone ED, and Cottrell EC. Effects of dietary nitrate supplementation, from beetroot juice, on blood pressure in hypertensive pregnant women: a randomised, double-blind, placebo-controlled feasibility trial. *Nitric Oxide* 2018; 80: 37-44. doi: 10.1016/j.niox.2018.08.004.

53. Zafeiridis A, Triantafyllou A, Papadopoulos S, Koletsos N, Touplikioti P, Zafeiridis AS, Gkaliagkousi E, Dipla K and Douma S. Dietary nitrate improves muscle microvascular reactivity and lowers blood pressure at rest and during isometric exercise in untreated hypertensives. *Microcirculation* 2019; 26(3): e12525. doi: 10.1111/micc.12525.

54. Lara J, Ogbonmwan I, Oggioni C, Zheng D, Qadir O, Ashor A, Brandt K, Mathers JC and Siervo M. Effects of handgrip exercise or inorganic nitrate supplementation on 24-h ambulatory blood pressure and peripheral arterial function in overweight and obese middle age and older adults: a pilot RCT. *Maturitas.* 2015; 82(2): 228-35. doi: 10.1016/j.maturitas.2015.07.028.

55. de Lima Bezerra AD, Costa EC, Pacheco DA, Souza DC, Farias-Junior LF, Ritti-Dia RM, Grigolo GB, de Bittencourt Junior PIH, Krause M, and Fayh APT. Effect of acute dietary nitrate supplementation on the post-exercise ambulatory blood pressure in obese males: a randomized, controlled, crossover trial. *J. Sports Sci. Med.* 2019; 18(1): 118-127.

56. Jajja A, Suttyarjoko A, Lara J, Rennie K, Brandt K, Qadir O and Siervo M. Beetroot supplementation lowers daily systolic blood pressure in older, overweight subjects. *Nutr. Res.* 2014; 34(10): 868-75. doi: 10.1016/j.nutres.2014.09.007.

57. Henrohn D, Bjorkstrand K, Lundberg JO, Granstam SO, Baron T, Ingimarsdottir IJ, Hedenstrom H, Malinovschi A, Wernroth ML, Jansson M, Hedeland M and Wikstrom G. Effects of oral supplementation with nitrate-rich beetroot juice in patients with pulmonary arterial hypertension-results from BEET-PAH, an exploratory randomized, double-blind, placebo-controlled, crossover study. *J. Card. Fail.* 2018; 24(10): 640-653. doi: 10.1016/j.cardfail.2018.09.010.

58. Kapil V, Khambata RS, Robertson A, Caulfield MJ Ahluwalia A. Dietary nitrate provides sustained blood pressure lowering in hypertensive patients: a randomized, phase 2, double-blind, placebo-controlled study. *Hypertension* 2015; 65(2): 320-7. doi: 10.1161/HYPERTENSIONAHA.114.04675.

59. Siervo M, Shannon O, Kandhari N, Prabhakar M, Fostier W, Kochl C, Rogathi J, Temu G, Stephan BCM, Gray WK, Haule I, Paddick SM, Mmbaga BT and Walker R. Nitrate-rich beetroot juice reduces blood pressure in tanzanian adults with elevated blood pressure: a double-blind randomized controlled feasibility trial. *J. Nutr.* 2020; 150(9): 2460-2468. doi: 10.1093/jn/nxa170.

60. Faconti L, Mills CE, Govoni V, Gu H, Morant S, Jiang B, Cruickshank JK and Webb AJ. Cardiac effects of 6 months' dietary nitrate and spironolactone in patients with hypertension and with/at risk of type 2 diabetes, in the factorial design, double-blind, randomized controlled VaSera trial. *Br. J. Clin. Pharmacol.* 2019; 85(1): 169-180. doi: 10.1111/bcp.13783.

61. Kerley CP, Dolan E, James PE and Cormican L. Dietary nitrate lowers ambulatory blood pressure in treated, uncontrolled hypertension: a 7-d, double-blind, randomised, placebo-controlled, cross-over trial. *Br. J. Nutr.* 2018; 119(6): 658-663. doi: 10.1017/S0007114518000144.

62. Bondonno CP, Liu AH, Croft KD, Ward NC, Shinde S, Moodley Y, Lundberg JO, Puddey IB, Woodman RJ and Hodgson JM. Absence of an effect of high nitrate intake from beetroot juice on blood pressure in treated hypertensive individuals: a randomized controlled trial. *Am. J. Clin. Nutr.* 2015; 102(2): 368-75. doi: 10.3945/ajcn.114.101188.

63. Engelhard YN, Gazer B and Paran E. Natural antioxidants from tomato extract reduce blood pressure in patients with grade-1 hypertension: a double-blind, placebo-controlled pilot study. *Am. Heart. J.* 2006; 151(1): 100. doi: 10.1016/j.ahj.2005.05.008.

64. Ried K, Frank OR and Stocks NP. Dark chocolate or tomato extract for prehypertension: a randomised controlled trial. *BMC Complement Altern. Med.* 2009; 9: 22. doi: 10.1186/1472-6882-9-22.

65. Paran E, Novack V, Engelhard YN and Hazan-Halevy I. The effects of natural antioxidants from tomato extract in treated but uncontrolled hypertensive patients. *Cardiovasc. Drugs Ther.* 2009; 23: 145-51. doi: 10.1007/s10557-008-6155-2.

66. Kim JY, Paik JK, Kim OY, Park HW, Lee JH, Jang Y and Lee JH. Effects of lycopene supplementation on oxidative stress and markers of endothelial function in healthy men. *Atherosclerosis* 2011; 215(1): 189-95. doi: 10.1016/j.atherosclerosis.2010.11.036.

67. Wolak T, Sharoni Y, Levy J, Linnewiel-Hermoni K, Stepensky D and Paran E. Effect of tomato nutrient complex on blood pressure: a double blind, randomized dose(-)response study. *Nutrients* 2019; 11(5): 950. doi: 10.3390/nu11050950.

68. Ziegenfuss TN, Hofheins JE, Mendel RW, Landis J and Anderson RA. Effects of a water-soluble cinnamon extract on body composition and features of the metabolic syndrome in pre-diabetic men and women. *J. Int. Soc. Sports Nutr.* 2006; 3(2): 45-53. doi: 10.1186/1550-2783-3-2-45.

69. Akilen R, Tsiami A, Devendra D and Robinson N. Glycated haemoglobin and blood pressure-lowering effect of cinnamon in multi-ethnic Type 2 diabetic patients in the UK: a randomized, placebo-controlled, double-blind clinical trial. *Diabet. Med.* 2010; 27(10): 1159-67. doi: 10.1111/j.1464-5491.2010.03079.x.

70. Wainstein J, Stern N, Heller S and Boaz M. Dietary cinnamon supplementation and changes in systolic blood pressure in subjects with type 2 diabetes. *J. Med. Food* 2011; 14(12): 1505-10. doi: 10.1089/jmf.2010.0300.

71. Vafa MR, Mohammadi F, Shidfar F, Sormaghi MS, Heidari I, Golestan B and Amiri FS. Effects of cinnamon consumption on glycemic status, lipid profile and body composition in type 2 diabetic patients. *Int. J. Prev. Med.* 2012; 3(8): 531-6.

72. Anderson RA, Zhan Z, Luo R, Guo X, Guo Q, Zhou J, Kong J, Davis PA and Stoecker BJ. Cinnamon extract lowers glucose, insulin and cholesterol in people with elevated serum glucose. *J. Tradit. Complement. Med.* 2016; 6(4), 332-336. doi: 10.1016/j.jtcme.2015.03.005.

73. Azimi P, Ghiasvand R, Feizi A, Hosseinzadeh J, Bahreynian M, Hariri M and Khosravi-Boroujeni H. Effect of cinnamon, cardamom, saffron and ginger consumption on blood pressure and a marker of endothelial function in patients with type 2 diabetes mellitus: A randomized controlled clinical trial. *Blood Press.* 2016; 25(3): 133-40. doi: 10.3109/08037051.2015.1111020.

74. Sengsuk C, Sanguanwong S, Tangvarasittichai O and Tangvarasittichai S. Effect of cinnamon supplementation on glucose, lipids levels, glomerular filtration rate, and blood pressure of subjects with type 2 diabetes mellitus. *Diabetol. Int.* 2016; 7: 124-132. doi: 10.1007/s13340-015-0218-y.

75. Gupta Jain S, Puri S, Misra A, Gulati S and Mani K. Effect of oral cinnamon intervention on metabolic profile and body composition of Asian Indians with metabolic syndrome: a randomized double -blind control trial. *Lipids Health Dis.* 2017; 16: 113. doi: 10.1186/s12944-017-0504-8.

76. Shishehbor F, Rezaeyan Safar M, Rajaei E and Haghizadeh MH. Cinnamon consumption improves clinical symptoms and inflammatory markers in women with rheumatoid arthritis. *J. Am. Coll. Nutr.* 2018; 37(8): 685-690. doi: 10.1080/ 07315724.2018.1460733.

77. Ardalani HR, Hassanpour Moghadam M, Rahimi R, Soltani J, Mozayanimonfared A, Moradie M and Azizi A. Sumac as a novel adjunctive treatment in hypertension: a randomized, double-blind, placebo-controlled clinical trial *RSC Adv.* 2016; 6(14): 11507-11512. doi: 10.1039/C5RA22840A.

78. Saberi, M., Kazemisaleh D and Bolurian V. Effect of Olive Leaf on Mild to Moderate Hypertension Resistant to Normal Treatments. *J. Med. Plants* 2008; 7(27): 52-59.

79. Perrinjaquet-Moccetti T, Busjahn A, Schmidlin C, Schmidt A, Bradl B and Aydogan C. Food supplementation with an olive (*Olea europaea* L.) leaf extract reduces blood pressure in borderline hypertensive monozygotic twins. *Phytother. Res.* 2008; 22(9): 1239-42. doi: 10.1002/ptr.2455.

80. Susalit E, Agus N, Effendi I, Tjandrawinata RR, Nofiarny D, Perrinjaquet-Moccetti T and Verbruggen M. Olive (*Olea europaea*) leaf extract effective in patients with stage-1 hypertension: comparison with Captopril. *Phytomedicine* 2011; 18(4): 251-8. doi: 10.1016/j.phymed.2010.08.016.

81. Hermans MP, Lempereur P, Salembier JP, Maes N, Albert A, Jansen O and Pincemail J. Supplementation effect of a combination of Olive (*Olea europaea* L.) leaf and fruit extracts in the clinical management of hypertension and metabolic syndrome. *Antioxidants* 2020; 9(9): doi: 10.3390/antiox9090872.

82. Mohtashami R, Fallah Huseini H, Nabati F, Hajiaghaei R Kianbakht S. Effects of standardized hydro-alcoholic extract of *Vaccinium arctostaphylos* leaf on hypertension and biochemical parameters in hypertensive hyperlipidemic type 2 diabetic patients: a randomized, double-blind and placebo-controlled clinical trial. *Avicenna J. Phytomed.* 2019; 9(1): 44-53.

83. Zolfaghari F, Pourzadi N, Sahbaei F, Zolfaghari F, Kazemi SS and Davari AR. The effect of blueberry solution on blood pressure and fasting blood sugar in patients with non-insulin dependent diabetes mellitus: a double-blind clinical trial. *Feyz* 2015; 19(4): 278-283.

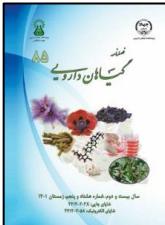
84. Kianbakht S and Hashem-Dabaghian F. Antihypertensive efficacy and safety of *Vaccinium arctostaphylos* berry extract in overweight/obese hypertensive patients: a randomized, double-blind and placebo-controlled clinical trial. *Complement. Ther. Med.* 2019; 44: 296-300. doi: 10.1016/j.ctim.2019.05.010.

85. Ghaffari S, Navabzadeh M, Ziae M, Ghobadi A, Ghods R and Hashem-Dabaghian F. A randomized, triple-blind, placebo-controlled, add-on clinical trial to evaluate the efficacy of *Emblica officinalis* in uncontrolled hypertension. *Evid. Based Complement. Alternat. Med.* 2020; 2020: 8592869. doi: 10.1155/2020/8592869.

86. Carey RM, Moran AE and Whelton PK. Treatment of hypertension: a review. *JAMA*. 2022; 328(18): 1849-1861. doi: 10.1001/jama.2022.19590.

87. Nadar SK and Lip GYH. *Hypertension*. Third ed., New York; Oxford University Press; 2023.

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مقاله مروی

اثربخشی و ایمنی بالینی ضد پرفشاری خون گیاهان ایران: یک مرور نظام مند

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اطلاعات مقاله	چکیده
گل و ازگان:	مقدمه: استفاده از گیاهان ضدپرفشاری خون یکی از راههای کنترل پرفشاری خون است. هدف: بررسی اثربخشی و ایمنی بالینی ضدپرفشاری خون گیاهانی که در ایران یافت می شوند. روش بررسی: SCOPUS, MEDLINE, PUBMED, Google Scholar, Google, EBSCO, OVID, PROQUEST, SCIENCE DIRECT, EMBASE, جستجو قرار گرفت و از راهنمای PRISMA تبعیت شد. کلمات مورد جستجو عبارت بودند از herb, plant, Iranian, hypertension, antihypertensive, randomized controlled trial. مقالات زبان انگلیسی که تا آخر ۲۰۲۲ چاپ شده استفاده شد. مطالعات برون تنی و حیوانی، مقالات سردبیر و مروری استفاده نشد. کیفیت روش شناختی مطالعات با استفاده از مقیاس JADAD ارزیابی شد. نتایج: دویست و هشت مطالعه یافت شد. فقط ۷۴ مطالعه واجد شرایط بود.
ایران	برای زرشک (<i>Berberis vulgaris</i>) (۵ مطالعه)، سیاهدانه (<i>Nigella sativa</i>) (۱۰ مطالعه)، سیر (<i>Allium sativum</i>) (۱۲ مطالعه)، چای ترش (<i>Hibiscus sabdariffa</i>) (۱۱ مطالعه)، چغندر (<i>Beta vulgaris</i>) (۱۵ مطالعه)، گوجه فرنگی (<i>Rhus coriaria</i>) (۱ مطالعه)، آمله (<i>Phyllanthus emblica</i>) (۱ مطالعه)، زیتون (<i>Olea europaea</i>) (۴ مطالعه) و قره قاط (۳ مطالعه) یافت شد. اکثر مطالعات کیفیت روش شناختی بالا داشتند و اثربخشی بدون عوارض جانبی گزارش کردند. نتیجه گیری: در حالی که، اکثر مطالعات اثربخشی و ایمنی ضدپرفشاری خون نشان می دهند، شواهد بیشتری در رابطه با چای ترش، زیتون، قره قاط و سیر در مقایسه با سایر گیاهان وجود دارد.
پرفشاری خون	
فشار خون	
گیاه	
طب مکمل	
طب جایگزین	

مخفف ها: 2hPPG، گلوكز پلاسمای 2 ساعت بعد غذا؛ ABPM، آپیش فشار خون در حالت حرکت؛ ALP، آلكالین فسفاتاز؛ ALT، آلتین آمینوتانسferاز؛ AST، آسپارتات آمینوتانسferاز؛ BID، دو بار در روز؛ BL، شروع مطالعه؛ BMI، باشخص توده بدنی؛ BP، فشار خون؛ BS، قند خون؛ CBC، شمارش کامل خون؛ CK-MB، اتصال کراتین کیناز- میوگلوبین؛ Cr، کرآتینین؛ CVLT-II، چاپ دوم آزمایش یادگیری کلامیفیرینا؛ DBP، فشار خون دیاستولی؛ EC، باز شونده در روده؛ eGFR، سرعت تخلیقی فیلتر اسیون گلومرولی؛ EHTN، برفشار خون ضروری؛ FBS، قند خون ناشتا؛ FID، نوجوان مذکر سالم؛ HAM، روزی ۴ بار؛ HbA1c، HbA1c؛ HR، سرعت ضربان قلب؛ ICAM-1، لیپوپروتین با چگالی بالا؛ ICAM-C، HDL-C، لیپوپروتین-کلسیترول با چگالی بالا؛ IC-LDL، لیپوپروتین-کلسیترول با چگالی پایین؛ IC-LV، لیپوپروتین با چگالی پایین؛ IC-MetS، مولکول جیسیندگی بین سلولی ۱-۳-C، HCL، برفشار خون؛ HCTN، برفشار خون بالا؛ LV، بطن چپ؛ MDA، مالون دی الید؛ N/A، سندروم متابولیک؛ NS، غیر معنیدار؛ NS، شامل نیشیدن؛ NS، سیاهدهن؛ NS، دارونا؛ PAH، تصادفی شده دوسوپربر؛ PAH، برفشار خون شریان ریوی؛ Pre-DM، پیش دیابت شریین؛ RDB، دارو؛ RCT، کارآزمایی کنترل و تصادفی شده؛ SBP، فشار خون سیستولی؛ S-VCAM-1، مولکول جیسیندگی بین سلولی ۱-۱ محلول؛ T2DM، دیابت شیرین نوع ۲؛ TID، ۳ بار در روز؛ TSP، فشار خون سیستولی؛ NAFLD، بیماری کبد چرب غیرالکلی؛ NHV، داوطلب سالم طبیعی؛ OLE، عصاره برگ زیتون؛ RCT، کارآزمایی کنترل و تصادفی شده؛ SGOT، گلوتامیک اکسالواستیک ترانسامیانز سرم؛ STAI، برسننامه اضطراب حالت-صفت؛ TCD، کلسترول تام؛ TDS، سه بار در روز؛ TE، عصاره گوجه فرنگی؛ TG، تری گلیسریدها-0؛ TNF- α ، عامل نکروز قاسیچ چایخوری؛ UHTN، برفشار خون کنترل شده.

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