

*Lifestyle management of hypertension:
International Society of Hypertension
position paper endorsed by the World
Hypertension League and European
Society of Hypertension*

Article

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Lifestyle management of hypertension: International Society of Hypertension position paper endorsed by the World Hypertension League and European Society of Hypertension

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Hypertension, defined as persistently elevated systolic blood pressure (SBP) >140 mmHg and/or diastolic blood pressure (DBP) at least 90 mmHg (International Society of Hypertension guidelines), affects over 1.5 billion people worldwide. Hypertension is associated with increased risk of cardiovascular disease (CVD) events (e.g. coronary heart disease, heart failure and stroke) and death. An international panel of experts convened by the International Society of Hypertension College of Experts compiled lifestyle management recommendations as first-line strategy to prevent and control hypertension in adulthood. We also recommend that lifestyle changes be continued even when blood pressure-lowering medications are prescribed. Specific recommendations based on literature evidence are summarized with advice to start these measures early in life, including maintaining a healthy body weight, increased levels of different types of physical activity, healthy eating and drinking, avoidance and cessation of smoking and alcohol use, management of stress and sleep levels. We also discuss the relevance of specific approaches including consumption of sodium, potassium, sugar, fibre, coffee, tea, intermittent fasting as well as integrated strategies to implement these recommendations using, for example, behaviour change-related technologies and digital tools.

Keywords: Blood pressure, hypertension, lifestyle, holistic approach, nutrition, diet, exercise, mindfulness, obesity, pollution

Abbreviations: ACE, angiotensin-converting enzyme; BP, blood pressure; CARDIA, Coronary Artery Risk Development in Young Adults; CI, confidence interval; CPAP, continuous positive airway pressure; CRF, cardiorespiratory fitness; CVD, cardiovascular disease; DALYS, disability-adjusted life years; DASH, Dietary Approaches to Stop Hypertension; DBP, diastolic blood pressure; ESADA, European Sleep Apnea Database; GLP-1, glucagon-like peptide-1; ISH, International Society of Hypertension; MBSR, mindfulness-based stress reduction; MET, one-metabolic equivalent increment; OSA, obstructive sleep apnoea; PRS, polygenic risk scores; RCT, randomized controlled trials; RPE, rating of perceived exertion; SBP, systolic blood pressure; SGLT2, sodium-glucose cotransporter-2; SSaSS, salt substitution and stroke study; WHO, World Health Organization

Table of Contents

I . INTRODUCTION	1
II. PROCESS OF WRITING	3
III. GENERAL RECOMMENDATION	3
IV. WEIGHT MANAGEMENT	3
V. PHYSICAL ACTIVITY	5
VI. GENERAL NUTRITION	7
A. Salt	8
B. Potassium	9
C. Sugar	9
D. Fibre	10

E. Alcohol	11
F. Non-alcoholic beverages	11
VII. INTERMITTENT FASTING	12
VIII. STRESS REDUCTION AND MINDFULNESS	12
IX. SLEEP	13
X. SMOKING	14
XI. POLLUTION EXPOSURE	15
XII. NONPRESCRIPTION MEDICINE, INCLUDING VITAMIN INTAKE	16
XIII. BEHAVIOURAL INTERVENTIONS, DIGITAL HEALTH, AND WEARABLES	16
XIV. HOLISTIC APPROACH TO BP PREVENTION ACROSS THE GLOBE	17
XV. OVERALL SUMMARY AND A GLIMPSE INTO THE FUTURE FOR ASSISTED LIFESTYLE INTERVENTIONS	18
ACKNOWLEDGMENTS	18
REFERENCES	18

I. INTRODUCTION

Hypertension is the single most significant modifiable risk factor for all-cause morbidity and mortality worldwide. Approximately 33% of the global population of 8 billion live with hypertension [1,2]. Lifestyle changes are the cornerstone of prevention and treatment of hypertension and both governments and industry are fundamental to endorse and implement these changes [3]. Moreover, lifestyle changes may not only reduce blood pressure (BP) and improve hypertension control but also improve cardiovascular and general health [4–11]. Patients also often view lifestyle changes as their highest priority area in their hypertension management [12]. Consequently, lifestyle modifications are the first-line recommendation in all major hypertension management guidelines to reduce the risk of CVD [13–17]. The importance of lifestyle changes in both treatment and prevention of hypertension is further emphasized in the ISH's Hypertension Zero Declaration [2] and the Society's public campaigns championing healthy lifestyle [18]. In addition to direct benefits, lifestyle modifications can also amplify the effect of the pharmacological treatment of hypertension, if needed [13]. Of importance, the well known lifestyle interventions such as change in diet, alcohol moderation, smoking cessation and aerobic exercise, have now been extended to less obvious strategies such as stress reduction, isometric exercise and reducing exposure to pollution [19–21]. Although lifestyle changes are effective, they remain difficult to implement and maintain long-term because many people live in environments that are not conducive to achieving and then sustaining a healthy lifestyle. Moreover, clinicians are often poorly trained in assisting patients to engage in healthy behaviours. This is further compounded by the paucity of information about lifestyle changes in people from different ethnic and cultural backgrounds and their health literacy and cultural beliefs concerning hypertension prevention and management. In this position paper, we discuss the updated evidence supporting the use of lifestyle interventions in patients with hypertension or people at risk of developing hypertension. Importantly, we also aim to

provide unique global perspectives on ways to implement these changes, taking into consideration different ethnicities and age groups, to aid healthcare providers and the scientific community to navigate and apply the current knowledge.

II. PROCESS OF WRITING

This position paper summarizes recommendations compiled by an international panel of experts convened by the International Society of Hypertension (ISH) College of Experts from 18 countries. Separate groups of experts were then assigned specific topics based on their area of expertise and developed draft recommendations by consensus based upon review of the literature and existing guidelines. Revisions were made by all authors and then submitted for external review internationally. Both World Hypertension League and the European Society of Hypertension reviewed and endorsed the recommendations.

III. GENERAL RECOMMENDATION

We recommend lifestyle goals (summarized in Table 1 and Fig. 1) to prevent or delay the onset of high BP and to reduce CVD risk. Lifestyle modifications should be the first line of antihypertensive treatment in grade 1 hypertension. However, if BP control is not achieved with lifestyle changes alone, we recommend a combination of lifestyle modifications and antihypertensive medications for diagnosed hypertension as the former enhance the effect of treatment [22]. Importantly,

pharmacotherapy should not be delayed when instituting lifestyle change in patients, for instance, in those with hypertension-mediated organ damage or at high level of risk for cardiovascular events [23]. Lastly, we recommend that healthcare providers and clinicians receive adequate training in working with patients to adopt lifestyle changes and take an active role in the implementation of these recommendations. These consensus recommendations are targeted towards primary care providers and public health programmers. In addition to the recommendations, we provide an international perspective to consider different resources, practices and cultural considerations for differing regions and implementation guidance at the individual and population level.

IV. WEIGHT MANAGEMENT

What is known and what is new?

Obesity is a worldwide epidemic with increasing prevalence in many countries. Over the past three decades, obesity prevalence in adults almost tripled worldwide, while in children and adolescents, the prevalence increased more than four-fold from 4 to 18% [24]. At the same time, globally, obesity-related deaths and disabilities have almost doubled. Obesity should be addressed with an extensive and encompassing methodology, including government and public health policies, clinical and self-directed approaches. Increased body mass measures have a well established causal effect on the development of hypertension, the risk of cardiovascular and kidney diseases as well

TABLE 1. Lifestyle modification for the management of hypertension recommended by the International Society of Hypertension

Recommendation	Action
Early start in life	Encourage healthy habits from childhood (governments/parents/carer) Increase awareness of healthy food types (parents/schools/governments)
Healthy eating	Increase consumption of vegetables, fresh fruits, fish, nuts, unsaturated fatty acids, low-fat dairy products and low consumption of red meat Reduce salt and sugar intake Increase dietary potassium intake Increase consumption of plant-based food and dietary fibre
Healthy drinking	Improve food labelling (governments/policy makers) Implement food taxation (governments/policy makers) Avoid or limit alcohol intake (if you do not drink, do not start) Avoid binge drinking Moderate consumption of unsweetened coffee and tea Consider hibiscus tea, pomegranate juice, beetroot juice and cocoa
Physical activity	Increase aerobic exercise and muscle strengthening exercise Increase incidental exercise Decrease prolonged periods of time seated Engage in regular physical exercise and avoid sedentary behaviour early in life Advocate for implementation of government and industry supported programs, environment, facilities, and infrastructure (governments/policy makers)
Healthy body weight	Ethnic-specific cut-offs for BMI and waist-to-height ratio or other indices Waist circumference and weight values to be monitored
Healthy stress levels and sleep	Improve sleep hygiene and increase quality sleep time Practice sleep stimulus control methods Promote mindfulness/stress reduction practice Consider meditation/yoga/breathing techniques Consider music therapy/self-soothing techniques/gratitude/acts of kindness
Reduce exposure to pollution and passive smoke	Promote smoking cessation Consider limiting time spent outdoors when pollution is high Consider ventilation systems with filtration
Use digital tools	Use apps to track calories, steps, and sleep patterns

Recommendations are aimed for individuals, unless stated otherwise.

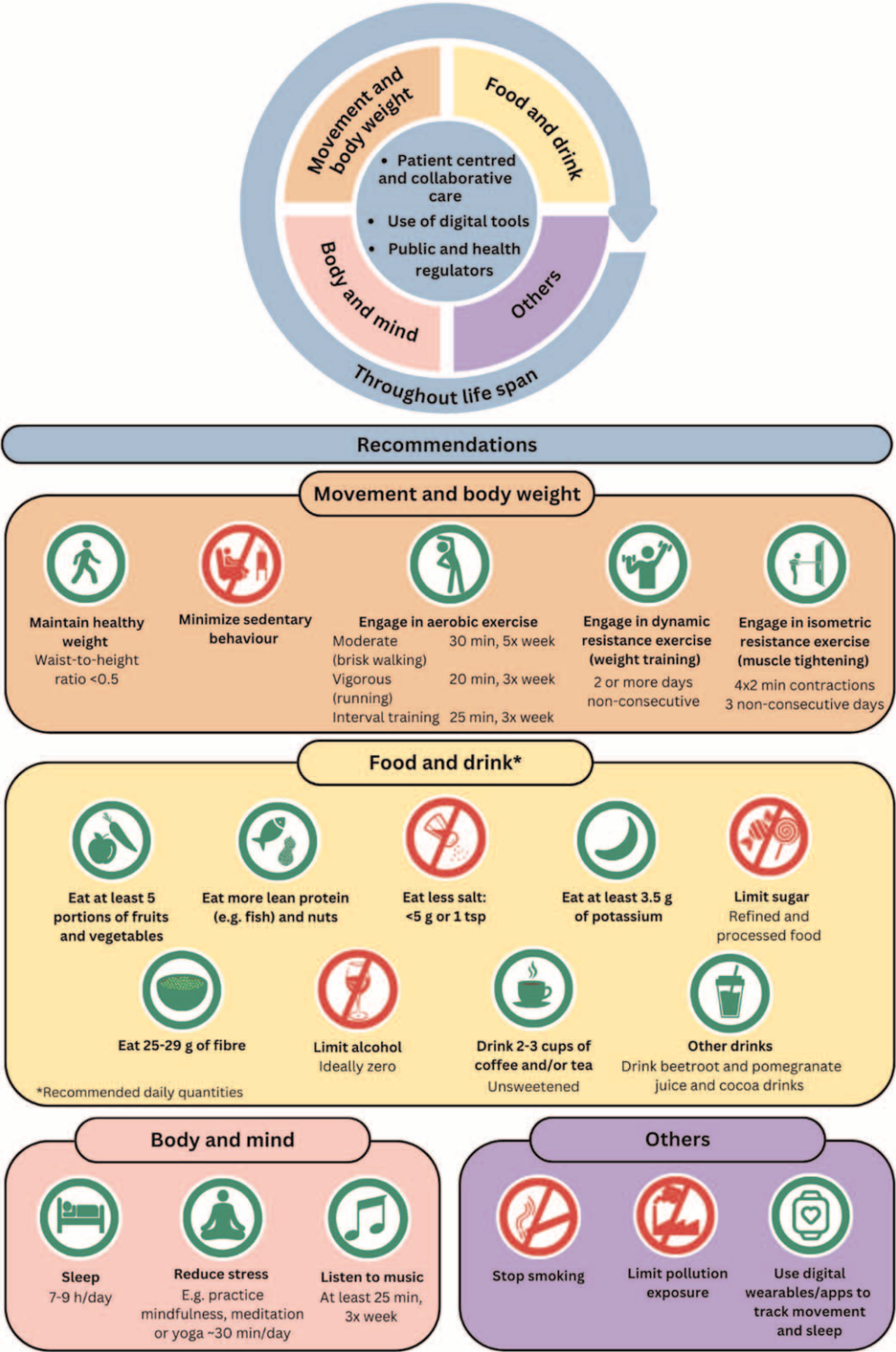


FIGURE 1 Lifestyle changes recommended in this position paper. In green are changes that should be encouraged, increased, and adhered to, for example, consumption of dietary fibre, improved sleep habits, increased exercise and adoption of mindfulness strategies. In red are changes to decrease or avoid, for example, smoking and alcohol use, sedentarism, consumption of refined sugars and salt and pollution exposure.

as cardiovascular mortality [25–27]. A meta-analysis of 57 prospective cohort studies, pooling 2.3 million individuals, reported a 1–2-fold increase in risk of developing hypertension with the increment of various obesity indices such as body mass index, waist circumference and waist-to-height ratio [28]. In addition, obesity is also associated with a higher risk of developing other cardiometabolic disturbances, including insulin resistance, type 2 diabetes, dyslipidaemia and metabolic syndrome, which can improve with weight loss [27]. The effect of obesity on BP is additive to other lifestyle components such as physical inactivity, diet, smoking and alcohol consumption [29].

All major hypertension guidelines recommend weight loss for the management of BP among those with overweight or obesity [30]. Furthermore, several large epidemiological studies have shown an association between body weight and BP over the life course [31–34]. Weight gain from early ages and in adult life, related to nutrition and/or lack of physical activity, is an important risk factor for hypertension development [3]. Weight loss has been recommended for patients with hypertension who are obese as an effective nonpharmacological treatment approach.

However, long-term results of weight loss programmes are disappointing as patients often regain most of the weight initially lost. A gradual and moderate weight loss approach, with support of dietitians, is more likely to be maintained over a longer period of time. The use of calorie-intake reduction, weight loss medication therapy or bariatric surgery for patients with significant obesity is an effective, longer lasting strategy for weight loss [35–37]. Furthermore, emerging treatment option for obesity include semaglutide and other glucagon-like peptide-1 (GLP-1) receptor agonist in effective management of obesity [38,39].

Weight reducing diets in overweight people with hypertension, leading to a modest weight loss (3–10% of baseline body weight), decreased/discontinued dosage requirements of antihypertensive medications and reduced SBP and DBP by ~3 mmHg resulting in lower BP levels even without reaching ideal weight [40]. Additionally, a meta-analysis of randomized controlled trials (RCTs) reported an average reduction of 1 mmHg SBP and DBP per kilogram of weight loss [41]. The BP-lowering effect of weight loss is most likely a result of a decrease in sympathetic nervous system activity and an improvement in insulin sensitivity [42] and occurs independent of salt restriction. Importantly, weight reduction following bariatric surgery reduced mortality and morbidity related to CVD [43].

International perspective

Currently, there is no global consensus on optimal weight reduction and preventive strategies and advice are not tailored at individual level considering bias, socioeconomic status and socio-cultural barriers and beliefs to weight loss in different parts of the world. Formalized, structured and/or supervised weight loss programs are not always possible or available. Furthermore, barriers to accessing appropriate exercise or diet guidance and professional advice are prevalent in some socio-economic and cultural backgrounds and the dissemination of many nonscience-based diets tends to be more harmful to some individuals with underlying conditions.

Recommendation

1. Weight loss should be incorporated into everyday life, that is, increasing incidental exercise, walking, or cycling more, or playing sports regularly, sitting for less hours, and so forth.
2. Emphasis should be placed upon early-life intervention and health education programs to sustain a healthy weight throughout life [44].
3. A modest weight loss that can be maintained over a long period of time is recommended for people with hypertension along with calorie-intake reduction.
4. Weight loss interventions should be based on cognitive-behavioural strategies (e.g. appetite awareness training, a self-monitoring strategy in which individuals learn to identify internal cues of moderate hunger and fullness and to use these cues to guide their eating behaviour) [45].
5. Abdominal obesity should be managed. Ethnic-specific cut-offs for body mass index and waist circumference should be used. Alternatively, a waist-to-height ratio less than 0.5 is recommended for all populations [46].
6. The chosen type of weight loss programmes should be tailored, considering individual baseline weight, age, sex, comorbidities and situational context with support from a dietitian.
7. Use of innovative approaches based on behavioural change technologies (e.g. apps, text messaging [47–49]) are encouraged for all people with overweight or obesity.

Implementation

Continual support for people with overweight or obesity is required to make a sustainable behavioural change and encourage healthy weight levels over the life course. The development of government-supported and industry-supported programmes, environment, facilities and infrastructure promoting healthy eating and weight reduction is paramount [50].

In the primary care of hypertension, healthcare providers should (whenever available) refer patients to appropriate weight loss programmes and prescribe individualized exercise and physical activity. Behavioural change techniques and coaching that include goal setting, monitoring and feedback should be considered to reduce overweight and maintain a healthy weight.

V. PHYSICAL ACTIVITY

What is known and what is new?

The CVD prevention and hypertension management guidelines [15,16,51–53] recommend exercise/physical activity across the lifespan, including childhood and adolescence, because of its demonstrated antihypertensive effects and also its favourable effects on other modifiable CVD risk factors. Guidelines follow current knowledge that regular engagement in physical activity leads to improved cardiorespiratory fitness (CRF) [54] and body habitus, as well as more favourable measures, of cardiometabolic health [55] and reduced inflammation [56], lowering the risk of

developing hypertension and CVD. Importantly, recent pooled-data highlights a dose–response relationship between CRF and relative risk of hypertension where a one-metabolic equivalent increment (MET) in CRF is associated with 8% hypertension risk reduction [57]. Recent data from the UK Biobank are consistent with these observations, with an adjusted inverse dose–response relationship for CVD risk observed between accelerometer-measured physical activity and stroke incidence, coronary artery disease and CVD and all-cause mortality, with no upper limit of activity (intensity or volume) detrimental in those with established hypertension [58].

Evidence from RCTs suggest that exercise training interventions may provide equivalent reductions in BP to that of antihypertensive medication [59]. When exercise is combined with pharmacotherapy, the magnitude of BP reduction is greater than with medication alone [60]. Meta-analysis of randomized studies has shown that, indeed, aerobic and dynamic resistance training interventions of varying designs reduce office and daytime ambulatory BP, with greater reductions occurring in those with established hypertension (up to 8.3/5.2 mmHg office BP reductions) [61–64]. Isometric resistance training also significantly reduces BP, which is supported by several meta-analyses [61,65–67] and is included as a nonpharmacological therapy in the American College of Cardiology/American Heart Association Task Force Guidelines on the Primary Prevention of Cardiovascular Disease [68].

Exercise of moderate-to-high intensity (including aerobic and dynamic resistance training; see Supplementary Box 1, <http://links.lww.com/HJH/C287>, for example is considered safe and currently recommended for hypertension management [20,69–71]. Adding a time-efficiency aspect, emerging data indicate that high-intensity interval training (a form of exercise training, which consists of repeated short and intense exercise), appropriately prescribed and supervised, can provide comparable BP reductions to that of moderate-intensity exercise training [72,73]. However, even modest intensity physical activity such as walking will have a BP-lowering effect [74,75], as does isometric resistance training of the upper or lower limb performed at ~30% of maximal voluntary contraction [20].

International perspective

Exercise as a therapy for hypertension is still not widely integrated into healthcare systems or insurance schemes worldwide. Indeed, socioeconomic and cultural barriers to structured exercise exist in many parts of the world [76], often combined with limited access to exercise facilities and lack of formalized physical activity guidance [77].

Neighbourhood safety, lack of community support, religious beliefs, financial position, employment, and social barriers may influence physical activity participation in certain groups with hypertension, especially ethnic minorities [78].

To overcome such barriers, individuals should be offered appropriate health education and be encouraged to use any means possible to be physically active, while creating facilities and programs that consider sensitive cultural needs [79,80]. Furthermore, healthcare practitioners

need to receive training to assist people to reduce sedentary behaviour and adopt lifestyle changes.

Recommendation

1. All individuals should be encouraged to be physically active for the prevention or management of hypertension and CVD [16,51,52,68,81].
2. Emphasis should be placed upon early-life intervention (from childhood) and health education programmes to sustain physical activity throughout the life-course (Table 1) [44].
3. Both aerobic and dynamic resistance exercise or their combinations can be used in prevention and management of hypertension and CVD (Box 1) [70].

Box 1 General physical activity recommendations for general health, wellness and cardiovascular disease prevention in (a) adults 18–64 years and (b) individuals aged at least 65 years [355]

(a) Physical activity recommendations in adults (18–64 years)

Adults should aim to be active on most days, preferably all days of the week. This should include either:

- 150–300 min of physical activity per week performed at a moderate intensity, or
- 75–150 min of physical activity per week performed at a vigorous intensity, or
- An equivalent combination of moderate and vigorous physical activities.

AND

- Include resistance (muscle strengthening) exercise on at least two nonconsecutive days each week, and
- Increase incidental physical activity.

(b) Physical activity recommendations in older adults (+65 years)

- Perform at least 30 min physical activity at a moderate intensity on most, preferably all days, of the week.
- If 30 min is difficult to achieve currently, it is recommended to commence at 10 min once or twice daily, which can be progressively increased.
- Try to incorporate different activities over the week, which may include:
 - **Aerobic (endurance) activities** to maintain and improve cardiovascular health such as brisk walking, swimming, and cycling.
 - **Dynamic resistance (muscle strengthening) activities** to maintain and improve muscle and bone strength such as weight-based, body weight, and resistance band exercises.
 - **Flexibility activities** to assist with ease of movement such as stretching, yoga and tai chi.
 - **Balancing activities** to improve balance and prevent falls such as semi-tandem, tandem and single leg standing.
- Try and reduce the amount of time spent sitting by breaking the time up as often and feasible as possible and increasing the amount of incidental exercise.

4. Adults should engage in 150–300 min of moderate-intensity or 75–150 min of vigorous exercise, or an equivalent combination, weekly (Box 1 and see Supplementary Box 1, <http://links.lww.com/HJH/C287> for examples) [82].
5. The chosen type of activity/exercise should be individually tailored, consider baseline fitness, comorbidities, pharmacological treatment, situational context and be progressive in nature (i.e. start slow and gradually build the amount/intensity of activity).

6. Emphasis should be on reducing daily sedentary time and increasing movement wherever possible [83].
7. Any type, amount and intensity of regular exercise/activity will be beneficial with minimum recommendations for maintenance of health and CVD prevention.

Implementation

Regular exercise should start early in life, and children and adolescents should aim to be active most days, decrease screen time and sedentary behaviours. This can be promoted and motivated by parents, guardians and educators and be implemented through the educational system's regular activities.

Healthcare professionals should encourage activities that are enjoyable to increase adherence and sustain life-long activity. Governments and town developers should plan environments conducive and safer to walk, cycle and exercise at all ages.

It is crucial to provide support and education to people at all life stages and levels of readiness for exercise to empower people to initiate/increase physical activity. Consumer resources and wearable devices are available at low cost, and may assist with and prompt increased incidental exercise [84].

Although exercise is considered generally very safe for those with high BP, first-time exercisers or those with complex medical history or chronic disease should be encouraged to seek guidance and approval to exercise from a doctor and or seek referral to/consult an exercise physiologist.

Behavioural change techniques and coaching that include goal setting, monitoring and feedback may increase physical activity uptake, sustainability and intervention effectiveness [85].

VI. GENERAL NUTRITION

Diet plays an important role in the healthy functioning of the body. Many different dietary patterns are hailed as useful for improving BP control, most without adequate evidence to substantiate their benefits. Studies show no age-related rise in BP or atherosclerosis in isolated non-Westernized tribal populations that live as hunter-gatherers and gardeners in remote regions in Brazil and Africa and consume diets low in fat and salt and high in fruit and fibre [86,87].

The Dietary Approaches to Stop Hypertension (DASH) low-sodium [88,89] and the Mediterranean diet [90] are most widely acknowledged for their benefits to reducing BP and improving cardiovascular health, albeit not always being viable because of food costs, availability and insecurity, incompatibility with a culturally and ethnically diverse society or food allergies. Broadly, both recommend low saturated fat, low salt, high fibre (from wholegrains, fruits and vegetables) and adequate lean protein. Both align with the World Health Organisation (WHO) guidance to reduce the risk of noncommunicable disease [91]. Such approaches are evidence-based and endorsed in ACC/AHA (USA), European and NICE (UK) guidelines [17,51,68,92].

The DASH diet, rich in calcium, potassium, magnesium and micronutrients, emphasizes consumption of fruits, vegetables, low-fat dairy products and reduced saturated fat and cholesterol, is recommended as an effective dietary intervention to reduce BP and maintain a healthy weight [15,93,94]. The adoption of the DASH diet was accompanied by substantial BP reduction in adults with and without hypertension with greater BP lowering among African Americans [88]. In addition, vegetarian diet has also been associated with lower BP in comparison to unrestricted omnivore diets (no salt, sugar or saturated and trans-fat restrictions) [95]. Vegetarian diets have a higher portion of glutamic acid and plant-based protein, which has blood-pressure-lowering effects [96–98]. Moreover, higher content of fibre, antioxidants and potassium as well as lower saturated fat and sodium content in healthy vegetarian diets can contribute to a lower body mass index and BP readings [99–101]. Vegetarian diet is thought to reduce BP via several mechanisms, including improving blood viscosity, vasodilation and insulin sensitivity; by altering the baroreceptors, renin–angiotensin and sympathetic nervous system; by its antioxidant and anti-inflammatory properties and by changing the colony and strain of gut microflora [102–107].

It is recommended that adopting the DASH diet or similar composition diet (Supplementary Box 2, <http://links.lww.com/HJH/C287>), where calorie-appropriate intake is considered depending on age, sex, build and levels of physical activity, is essential for maintenance of a healthy weight and should be considered a part of local hypertension management guidelines and introduced to people at risk of developing hypertension.

Long-term health benefits will depend on people's ability to make long-lasting dietary changes as long-term adherence/persistence to DASH and other diets remains a major concern. However, many population-level dietary recommendations are not followed, for example, 12% of Europeans and only 5% of the Australian population achieve the recommended five serves of fruit and vegetables per day (one serve is equivalent to 75–150 g depending on the type) [108–110]. There are also several barriers to adoption or adherence to these diets, which include lack of education (both with respect to health and/or food and nutrition) on a population level and by healthcare providers, food availability, including food insecurity or cost and difficulty or unwillingness to adhere because of cultural heritage, beliefs and behavioural norms [111]. To achieve long-term dietary change, healthcare professionals require specific training in effective behaviour change techniques to assist their patients in adopting new behaviours. There have been efforts to address some of these issues (e.g. reshaping of the UK medicine curriculum [112,113] and introduction of continuing professional development programmes for medical professionals [114]); however, the general public may be confused by (social) media reports delivering conflicting nutrition information [115,116], which leads to uncertainty and lack of trust in nutrition advice given by qualified professionals. Furthermore, many healthcare professions' training does not incorporate sufficient nutrition education and training in their undergraduate and professional development courses [114] and clinicians report a lack of time and confidence to deliver

comprehensive dietary and lifestyle advice during office visits [117]. These barriers need to be overcome if dietary approaches are to be implemented in preventive or primary healthcare and should start from an early age; methods such as targeted labelling, incentives (e.g. food vouchers) and policy change (e.g. taxation and fortification) are additional potential avenues but are widely debated by both the public and the food industry (Supplementary Box 3, <http://links.lww.com/HJH/C287>).

As nutrition research evolves, more complexity is realized. Looking beyond traditional macronutrients and micronutrients for health benefits might be important, for example, evidence suggests that cocoa flavanol intake could be beneficial for CVD risk prevention [118]. Further knowledge about how food components interact with each other could help maximize dietary approaches, for example, saturated fat in dairy may be less detrimental to CVD risk than in meat [119]. Evidence suggests that consumption of omega-3 fatty acids, found in fish, lowers BP, heart rate and triglycerides [120]. A recent dose–response meta-analysis suggests that the intake of 2–3 g/day of omega-3 fatty acids has optimal BP-lowering effects and more than 3 g/day might be beneficial for groups at higher risk of CVD [121]. An RCT showed independent additive effects of regular fish meals and weight reduction in people with overweight [122]. Unfortunately, consumption of omega-3 fatty acids is below recommended levels in a significant proportion of the population and fish meals should be considered as an alternative to daily meat meals as part of a healthy diet [123]. Emerging data suggest the gut microbiome is an important target for BP regulation [124,125], hence diets incorporating probiotics, prebiotics and/or postbiotics are essential. Indeed, this aligns with the relationship between fibre and BP control [126,127].

Personalized/precision nutrition approaches are also becoming widely appreciated [128], whether that be based on individuals' phenotype, genotype or even gut microbiota composition. Arguably they are harder to implement but may prove to be more effective. As the nutrition field advances, there needs to be a clear pathway for these recommendations to policy and clinical practice. We have included below separate sections for major nutritional components.

A. Salt

What is known and what is new?

High sodium consumption (over 2 g of sodium per day, equivalent to 5 g or one teaspoon of salt) resulting in increased BP is the leading dietary risk factor for death [129,130]. Sodium intake may influence BP by change of sodium–water homeostasis, vascular tone, immune cell homeostasis and tissue remodelling, but exact mechanisms remain subject of study [131]. The global mean sodium intake is estimated to be around 6 g/day [130], attributing to 1.89 million deaths and 44.87 million disability-adjusted life years (DALYs). Sodium is mainly consumed as salt, from processed foods or discretionary sodium added to food during preparation or at the table [129]. Meta-analysis of RCT examining sodium intake reduction to as low as 800 mg/day show a relatively linear decrease in BP

[132,133]. Moreover, the weighted average reduction in dietary sodium intake from 3646 to 2690 mg/day in RCT was associated with approximately 26% reduction in CVD [134]. Notably, the decrease in BP is greater in people from black African ancestry, people of older age, or with higher BP levels but is also observed in differing ethnicities, people with normal BP and in children [134].

A recent meta-analysis of prospective cohort studies investigating multiple nonconsecutive 24 h urine samples to assess sodium intake (the current recommended standard) reported a linear association between sodium intake [1846–5230 mg (equivalent to 4.6–13.8 g salt) per day] and cardiovascular events [135]. Unfortunately, these studies do not define a recommended lower level of sodium intake. However, many community-based salt reduction programs resulted in decreased salt intake, increased awareness and lowered BP [8,94,129,136–138].

International perspective

About 80% of the salt intake in high-income countries comes from manufactured and processed foods, whereas the majority of consumed salt in low-income and middle-income countries is added during food preparation or at the table [111,129,139]. Unfortunately, a shift in the future is to be expected as dietary patterns are transformed by increasing production and consumption of processed foods, rapid urbanization and changing lifestyles [140]. Studies consistently show that salt consumption in most parts of the world is greater than recommended and require urgent action [129,135]. Sub-Saharan Africa and the Caribbean have the lowest consumption of sodium worldwide, just under 3 g/day; whereas, East Asia has the highest intake, close to 8 g/day [130].

Recommendation

1. Intake of less than 2 g sodium (~5 g salt or one teaspoon) per day (Supplementary Box 2, <http://links.lww.com/HJH/C287>) [129].

Implementation

Globally reduce mean salt intake in the population by 30% by 2025, as proposed by the WHO and supported by the World Health Assembly. Approaches should be tailored considering local context, especially in regards to differences in food and lifestyle-related behaviours, including the source of salt intake [141]. Adopting the WHO SHAKE package at country level provides a unique opportunity for salt reduction at population and individual levels [141]. Importantly, salt substitution with low sodium and high potassium alternatives may provide an opportunity to reduce BP at the population level [142]. Translation of fundamental research into practice is crucial as mechanisms explaining how sodium increases BP is partly understood [131] and assessment of salt intake on the individual level is poorly associated with actual daily intake – complicating the interpretation of the epidemiological link between estimated salt intake and health outcomes [143]. To this end, academic organizations and learned societies called on scientists, clinicians, healthcare providers and other stakeholders to support global efforts to implement the

reduction in salt [129,144]. The global actors' policy priorities and funding allocation need to align, making salt reduction at the population level a key priority. Finally, industry and governments play a major role in shaping salt consumption at the population level. Governments should follow the expert advice and introduce legislation decreasing sodium in processed foods which, in turn should be endorsed by the food industry [129].

B. Potassium

What is known and what is new?

Enhancing potassium consumption lowers BP by direct effect on the vascular tone and stimulation of natriuresis [145]. The recent large randomized controlled salt substitution and stroke study (SSaSS) trial reported that increasing potassium intake as a sodium substitute – that is, replacing 25% sodium chloride with potassium chloride in salt – reduces the risk of stroke, CVD and death in patients with increased cardiovascular risk and low potassium intake and high sodium intake at baseline [142]. Potassium intake is reflective of healthy dietary patterns, as key sources of potassium include fruit, vegetables, nuts and legumes. The DASH diet recommends ~4.7 g potassium per day, possibly accounting for the BP-lowering effect reported. Dietary potassium, potassium supplements and implementation of salt substitutes reduce BP with little effect on the risk of hyperkalaemia in a healthy population [142,146].

Unlike comprehensive interventions that require implementing and long-term adherence to dietary or behavioural change, the use of salt substitutes containing potassium salt and/or adoption of potassium-rich diet may represent a more feasible intervention to implement [147]. Fruits, vegetables, legumes, fat-free or low-fat dairy foods and fish represent good natural dietary sources of potassium. Hyperkalaemia with higher potassium intake is a concern in certain populations such as people with advanced chronic kidney disease (excluded in the SSaSS trial), but the risk for developing clinically significant hyperkalaemia by increasing dietary potassium intake to recommended levels may be limited [148]. These individuals are at higher risk of adverse cardiovascular outcomes, and a modelling study demonstrates that there would be still a net benefit with consumption of a potassium salt in this setting [146].

International perspective

Estimates for potassium intake from cross-sectional studies show marked variations between countries ranging from 1.7 to 3.7 g per day, with highest consumption in Europe and lowest intakes in China [149] and indicate that most populations may benefit from increasing their daily potassium intake. BP reduction may depend on the population's background potassium and sodium consumption. Implementation of potassium salt substitutes in countries where the main source of salt consists of added salt during cooking at the household level (discretionary use) is expected to be more feasible and also more efficacious as compared with countries with usually very high potassium intake and low sodium consumption [147]. Importantly, a reduction of

sodium salt consumption is paramount, as stated above, and the use of low-sodium salt substitutes should be encouraged.

Recommendation

1. The European Food and Safety Administration and the WHO recommend potassium intake of over 3.5 g per day for adults, while an intake of more than 4.7 g per day for adults is recommended by the National Academies of Sciences and Medicine [129,134]. Foods high in potassium include cooked white cannellini beans (1200 mg/cup), unsalted boiled spinach (840 mg/cup), avocado (708 mg/cup) and bananas (450 mg per medium fruit).
2. Individual food and nutritional needs because of lower body composition between sex and larger differences between body mass within and between populations need to be considered.

Implementation

It is important to emphasize a healthy diet rich in vegetables and fruits, which will increase potassium intake. Potassium-enriched salts are also commercially available and could be used for cooking instead of regular salt in settings where most of the added salt happens at the household level. For other settings, partnership with the food industry is required to implement the substitution of sodium with potassium in food processing while preserving taste and other requirements. Pricing policies that promote the adoption of healthier diets will reduce sodium while increasing potassium intakes and should be a global priority.

C. Sugar

What is known and what is new?

Sugar is one of the most commonly used commodities globally. The term 'sugar' is vague, usually referring to refined sugar, it can be found in most packaged and processed foods and is commonly unrecognized by consumers because manufacturers use terms such as glucose or sucrose in nutritional labels. Recent studies have highlighted common side effects that accompany consistent consumption of sugar [150]. Cardiometabolic changes are well established as one of the biggest risk factors of long-term sugar consumption [151,152]. Regular consumption of foods and drinks high in fructose (fruit juices, sugar-sweetened beverages, dried fruits, sweetened yoghurt, honey) [151,153] and high glycaemic foods (white potato, white rice) predispose to the development of non-alcoholic fatty liver disease and is strongly associated with cardiometabolic changes related to increased adiposity such as hypertension, dyslipidaemia and insulin resistance [153,154]. Importantly, consumption of added sugar is strongly associated with increased risk of CVD mortality and contributes to the international overweight and obesity epidemic related to chronic illnesses [155].

Monosaccharides, such as fructose and glucose, have similar chemical structures; however, their metabolic pathways differ in both enterocytes and hepatocytes impacting overall health [151,153,156]. High glucose concentration as

well as high sodium in digested food are demonstrated to up-regulate the expression of sodium-glucose transporters. Therefore, the higher the consumption of sugar and salt, the more the gut absorbs glucose and sodium ('greedy gut') [152]. The same mechanism is observed in the kidney for re-absorption of glucose and sodium as for the expression of sodium-glucose cotransporter-2 (SGLT2) [152]. Recently, SGLT2 inhibitors have been highlighted not only in blood glucose lowering and body weight reducing effects but also for their renal and cardioprotective actions. It is recognized that SGLT2 inhibitors exert BP reduction possibly through renal sodium handling as well as body weight reduction [152,157].

International perspective

Diabetes linked with increased BP is rising, particularly in Asia. Furthermore, diabetes in Asians differs from that in Caucasians, commonly referred to as the "Asian phenotype" [158]. This phenotype is related to the impaired secretion of insulin by the pancreas and lower capacity of triglyceride storage by adipocytes [159].

Currently, 118 countries and jurisdictions (including Fiji, Germany, India, Mexico, Morocco, Nigeria, Qatar, South Africa, UK, Vietnam, some USA states) have introduced taxes on sugar-sweetened beverages, which have facilitated a decrease in sugar consumption, especially among those of lower socioeconomic backgrounds, and led to a trending decline in overweight and obesity rates, which could potentially contribute to lowering the BP levels [160–164].

Recommendation

1. Reduce or limit sugar intake both in raw form and in processed food, drinks and sweets (Supplementary Box 2, <http://links.lww.com/HJH/C287>).

Implementation

Governments should introduce a simpler food labelling system, such as 'traffic light' or 'health stars' to simplify the understanding of nutritional values by consumers. Meanwhile, it is important to educate the public on the importance of nutritional labels as sugar is present in most packaged food items, even the savoury type and especially low-fat products. This includes learning to read the nutrition label and recognize the terminology/wording for 'sugars'. There should be more emphasis on replacing fruit drinks for whole fruit (which contains fibre that can help in slowing the spike in blood sugar levels). Finally, everyone should be able to recognize that all forms of sugar are calorically equal, regardless of its origins, that is, table sugar, honey, jaggery or coconut sugar and that the main difference between refined sugar and other forms of sugar is the lack of essential micronutrients in the former.

D. Fibre

What is known and what is new?

Dietary fibres are carbohydrates with a minimum degree of polymerization that are neither digested nor absorbed in the small intestine [165]. Low intake of whole grains, fruits, nuts, seeds, vegetables and, thus, overall fibre are

some of the top dietary risks attributed to death globally, especially due to CVD [166]. A recent meta-analysis of 15 RCTs comparing fibre intake in 1064 patients and 988 controls showed an average 1.3 mmHg [95% confidence interval (CI): −2.5 to −0.04] decrease in SBP [127]. However, studies using around 135 million person-years of data found that high fibre intake reduced all-cause and cardiovascular mortality by 15–30% [127,166]. The effect of high fibre intake on CVD is mainly because of its high degree of fermentation by the gut microbiota, resulting in the release of short-chain fatty acids, which act on immune-dependent and independent mechanisms (e.g. vasodilation) [125,165,167]. A recent phase II RCT showed that short-chain fatty acids reduced SBP in untreated people with hypertension [167].

International perspective

Inadequate fibre consumption is a worldwide concern and often seen as a result of diets rich in processed foods and low in whole foods. The Westernized diet, characteristic of USA and Canada and the Westernization of traditional European diets (high in fat, refined grain and sugar and low on milk, fruits and vegetables) can be linked to an increase in CVD [168,169]. This contrast between low-fibre and high-fibre intake and its relationship with the incidence of CVD is also observed in Australia [168–170]. Interestingly, the high-fibre diet ingested by African hunter-gatherers seems to result in a protective gut microbiome and high production of short-chain fatty acids, preventing the development of chronic illnesses [171].

Lastly, undernutrition, present widely in sub-Saharan Africa and South Asia, also leads to an imbalance of the gut microbiota and morbidities, especially in childhood [168].

Recommendation

1. Fibre intake of 25–29 g/day conferred the greatest risk reduction, but dose–response data suggested more than 30 g/day results in additional benefits [127].

Implementation

Considering most people consume less than 20 g/day of fibre, community awareness programs, policies and tax incentives to promote an increase intake of foods high in fibre is recommended.

The consumption of high-fibre foods, particularly those high in fermentable fibre, should be increased, and substitutions be made instead of refined grain products (with low fibre content) to reach the recommended daily intake. For instance, lentils (16 g fibre per cup) and other beans such as edamame (9 g fibre per cup) are rich in fibre. Other foods high in fibre include brassica vegetables (kale, cabbage, cauliflower and broccoli – which contain an average of 5 g fibre per cup), blueberries (4 g fibre per cup), avocado (10 g fibre per cup) and whole grains (Supplementary Box 2, <http://links.lww.com/HJH/C287>) [172,173]. Importantly, people on diets with restrictive intake of carbohydrates (e.g. gluten-free diet) are at increased risk of poor fibre intake [174]. Lastly, the addition of high fibre flours (e.g.

beans, high amylose maize starch, green banana) in baked goods can also help increase fibre intake [172,173,175].

E. Alcohol

What is known and what is new?

The relationship between alcohol intake and high BP was established over a century ago [176], albeit clarity on the mechanisms involved is still lacking. Large-scale observational studies report that the incidence of hypertension increases with any amount of alcohol consumed by men and with more than two standard drinks per day by women [177]. Moreover, a systematic review and meta-analysis of 36 RCTs, described a significant reduction in BP associated with a decrease in alcohol consumption in people with hypertension who consumed three or more drinks per day in a dose-dependent manner [178]. The largest reduction in BP [SBP: -5.5 mmHg (95% CI: -6.7 to -4.3) and DBP: -4.0 mmHg (95% CI: -4.7 to -3.3)] was reported in individuals with a baseline consumption of six or more drinks per day. Additionally, both trial data and observational literature support the hypertensive effect of binge drinking [179,180].

Even though previous observational data have suggested a decrease in CVD with light drinking compared with abstainers, these analyses are limited by selection bias and confounding with other associated health behaviours [181]. Mendelian randomization studies do not support the purported protective effects of moderate alcohol consumption and suggest that the lowest risks for cardiovascular outcomes are in abstainers and that any amount of alcohol uniformly increases BP [181,182].

International perspective

Patterns of alcohol intake vary widely across the world, in terms of frequency, type of alcohol (e.g. beer, wine, spirits), and associated behaviours. Globally, the annual average consumption of pure alcohol per person is 6.18 l (the equivalent of 1 l of wine per week at 12% alcohol content) [183,184]. Consumption varies from particularly low across North Africa and Middle East (close to zero) to high in Europe (highest as 15 l per person annually or the equivalent of over two bottles of wine per week in the Czech Republic) [183,184]. Binge or heavy drinking (defined as consuming at least 60 g of pure alcohol or six standard alcoholic drinks on one occasion in the past 30 days) also varies internationally, and is more common in parts of Central and West Africa, as well as Russia [184]. Sex and gender differences in alcohol consumption (men drinking more) are wider when the average consumption in a country is low (e.g. in North Africa and Middle East) [184]. The type of alcoholic beverage consumed also varies, with spirits being predominant in countries such as India, Philippines and China, whereas beer is commonly consumed in Eastern Europe and wine in Western Europe [184].

Recommendation

1. Alcohol consumption should be zero for the best cardiovascular outcomes (Supplementary Box 2, <http://links.lww.com/HJH/C287>) [185]. However, the recommended daily upper limit for alcohol

consumption is two standard drinks for men and 1 for women (10 g alcohol/standard drink), while acknowledging that there is no safe limit for alcohol consumption to prevent hypertension and adverse cardiovascular outcomes.

2. Binge drinking should be avoided [185].

Implementation

Explicit assessment of alcohol intake should be a routine part of the history in the initial diagnosis and subsequent management of hypertension. The effect of counselling alone is uncertain, though data from several RCTs support the effect of brief interventions such as motivational interviews in reducing alcohol intake [186,187].

F. Non-alcoholic beverages

What is known and what is new?

Coffee and caffeinated drinks such as tea have beneficial effects on BP and overall cardiovascular health [188]. Drinking coffee shows a consistent nonlinear association with the reduced risk in cardiovascular mortality with the greatest benefits seen with three cups a day [189]. A recent study in the UK Biobank showed that those drinking two to three cups per day of coffee or tea 32 or 28%, respectively, have reduced risk of having a stroke and dementia when compared with nondrinkers [190]. It appears that a consumption greater than three cups daily is safe but the benefits (in terms of reduction in cardiovascular risk) were less apparent [189]. Interestingly, an inverse dose-response in coffee consumption has been associated with a 2% risk reduction of hypertension (relative risk = 0.98, 95% CI: 0.98–0.99) for each cup of coffee ingested daily [191]. In contrast, a Japanese study reported that consumption of more than two cups of coffee daily increased risk of CVD mortality in people with grade 2–3 hypertension [192]. This increased risk was not observed amongst green tea drinkers [192]. Increased coffee consumption has also been associated with increased risk of aortic valve stenosis in a Swedish prospective population study [193].

Emerging data also suggests the benefits of the consumption of nitrate compounds found in beetroot juice in lowering BP. A recent meta-analysis showed beetroot juice significantly decreases SBP but not DBP in patients with arterial hypertension [194]. Interestingly, the BP-lowering effects of beetroot juice in SBP and pulse pressure are enhanced when combined with grapefruit juice [195]. Other beverages that can be beneficial include karkade (hibiscus) tea, pomegranate juice and cocoa [196]. In contrast, the regular consumption of energy drinks, which have large amounts of caffeine [197], and liquorice root tea [198] promote a spike in SBP and heart rate leading to metabolic changes. The effects of prolonged consumption of energy drinks have not been determined but may be relevant to individuals at risk of CVD.

International perspective

Coffee and tea consumption varies between countries worldwide. However, studies suggest that coffee and tea intake may be accompanied by smoking, which may

revoke their benefits and could also be associated with BP differences in regions with high smoking rates.

Recommendation

1. Moderate regular coffee consumption (three to four cups per day) does not adversely affect BP and the cardiovascular system and can be moderately beneficial [81,199,200].
2. The addition of nitrate-rich beverages, such as beet-root juice, pomegranate juice and cocoa, may be considered.

Implementation

The moderate consumption of unsweetened coffee and tea can be assessed by healthcare professionals and recommended for potentially cardioprotective and/or BP-lowering effects.

VII. INTERMITTENT FASTING

What is known and what is new?

Fasting is the voluntary abstinence or restriction of food and/or drink consumption for short or long time periods. For over a thousand years, fasting has been used across the world for cultural, traditional, religious or health reasons. More recently, the use of short-term, intermittent fasting became a popular strategy for weight loss and improving metabolic health. This strategy gained interest with the recognition that periods of energy consumption have extended to predominately later into the evening, and this time factor, may contribute to chronic disease epidemics including obesity and hypertension [201,202].

There are various forms of intermittent fasting, including alternate day fasting where caloric intake varies on alternate days; daily time-restricted eating includes usual eating during an 8–12 h window and abstinence of food for the remaining time; and modified fasting where caloric intake is restricted on certain days (e.g. 5 : 2 diet) and *ad libitum* on other days of the week. Religious fasting, such as during the month of Ramadan, is a form of daily time-restricted fasting, where food and drink including medications are restricted from dawn until dusk.

It is postulated that intermittent fasting triggers a metabolic switch from hepatic glucose metabolism to adipose-derived ketone production during fasting periods.

Independent of weight loss, fasting lowers sympathetic nervous system activity, suppresses inflammation and improves glucose regulation [203]. Longer term clinical data on the safety and efficacy of intermittent fasting is emerging but is still limited. Several small studies demonstrate reduced BP among those who lose weight during intermittent fasting [204,205]. However, in a large clinical trial comparing intermittent fasting with reduced caloric intake versus a general calorie-restricted diet, both strategies were associated with reduced weight and BP, but there was no significant difference between the groups [206]. In a meta-analysis of 33 studies ($n = 3213$), fasting was associated with lower BP at end of Ramadan compared with before Ramadan, among those with diabetes, normotension and hypertension but no change in BP in those with chronic kidney

disease [207]. Further, there was no statistically significant increase in worsening renal outcomes associated with fasting [208,209]. However, these studies included generally low-risk or moderate-risk groups, and these results may not be extrapolated to high-risk patients including those with end-stage renal disease or severe or uncontrolled hypertension.

International perspective

Fasting is practiced around the globe across multiple regions, and the effects on health and BP can be affected by additional factors, including environmental climate, temperature, food availability, duration of daylight, culture and religious beliefs (e.g. Ramadan or Lent). Further, the timing of meals is heavily influenced by culture and region with some cultures typically consuming dinner after 10 pm. Finally, fasting is implemented for a variety of reasons and the patient-centred outcomes may extend beyond physical health, weight loss and BP. Therefore, the acceptability of intermittent fasting needs to consider the patient's personal preference. Lack of adequately powered longer term RCTs of fasting limit our understanding on the long-term effects of this strategy.

Recommendations

1. Intermittent fasting with calorie restrictions may be beneficial for weight loss and BP reduction but is not superior to general calorie-restricted diet. Therefore, use of intermittent fasting should depend on patient preferences, and other considerations.
2. Fasting, including during Ramadan or Lent, is generally safe in low-risk to moderate-risk populations.

Implementation

Patients with chronic conditions should seek guidance from their healthcare provider before initiating fasting.

VIII. STRESS REDUCTION AND MINDFULNESS

What is known and what is new?

Psychosocial stress is associated with an increased risk of hypertension and cardiovascular events [210,211]. Patients with psychological distress may develop a sudden increase in BP, which may usually normalize when the distress is relieved [212]. However, people who suffer from stress related to posttraumatic stress disorder have an increased risk of hypertension and history of poor medication adherence [213,214]. Importantly, post-COVID-19 pandemic-related stress, anxiety and depression may contribute to poorer BP control in patients with hypertension [215–218]. Stress is also associated with unhealthy habits such as poor diet or overeating and weight gain, heavy drinking and/or smoking and sedentary behaviours, increasing risk of hypertension and CVD [211].

Stress reduction can be achieved through a number of strategies, such as meditation, progressive muscle relaxation, yoga, deep breathing exercise and mindfulness [211]. In recent years, mindfulness practices, defined as the

awareness, including self-understanding, wisdom and compassion that comes from intentionally focusing on the present moment, have received interest as an alternative approach for the treatment of chronic diseases [219,220].

The mindfulness-based stress reduction (MBSR) program has been demonstrated to have a broad efficacy in improving physical and mental well being, as well as promoting office BP reduction [19,219]. A recent meta-analysis encompassing 12 studies reported that MBSR lessons of at least 8 weekly sessions of MBSR and over 30 min of meditation practices daily reduced stress, mood swings and SBP and DBP by 6.6 and 2.5 mmHg, respectively, in patients with hypertension [19].

A systematic review also reported that 45 min of yoga every day reduced SBP by 6.5 mmHg and DBP by 2.8 mmHg after a 12-week program [221].

Cognitive behavioural therapy helps individuals to identify and change negative thoughts and behaviours that contribute to stress. A meta-analysis of 15 RCTs revealed a reduction in SBP (8.7 mmHg), DBP (5.8 mmHg), cholesterol levels, depressive and anxiety symptoms and improved sleep [222]. Alternatively, other stress reduction techniques such as deep breathing, progressive muscle relaxation, self-soothing, gratitude and acts of kindness, learning problem-solving and goal-setting techniques can reduce stress levels and are better accepted than MBSR by some patients [7,210,223]. In addition, increasing physical activity, social connectedness and sleep hygiene (where warranted) can also be explored.

Overall, the best stress reduction interventions to reduce BP and hypertension are those that are maintainable and can be combined into daily life. A mixture of different approaches, such as MBSR, exercise and yoga, may be most effective and tolerated by individuals with hypertension. Importantly, it is vital to consult a healthcare professional to determine the best approach for each individual.

Another systematic review also reported that listening to music 1×/day to 3×/week showed a trend towards lower SBP and DBP by 10 and 6 mmHg, respectively [224]. Moreover, patients with hypertension who listened to 25 min of classical music over a 4-week period showed an average decrease of 6.6 mmHg in SBP [225].

International perspective

Evidence suggests that MBSR did not differ in reducing BP in patients between eastern and western countries [19]. Estimates for MBSR sessions from cross-sectional studies show a variation between countries ranging from 90 min/session in Iran [226] to 120 min/session in Spain over 8 weeks [227]. These studies indicate that patients with hypertension may benefit from practicing MBSR to reduce mental stress and BP, yet data on long-term BP effects and impact on future CVD risk are unknown.

Recommendation

1. Practice stress reductions techniques at least 3 h per week to reduce mental stress and BP [226,227].
2. Alternatively, practice activities such as yoga, meditation or tai chi at least 45 min a day [221].
3. Listen to music once a day to three times a week for at least 25 min [224].

Implementation

All healthcare professionals should assess psychological distress in all their patients with hypertension via visualized analogue scales or screening tools.

Patients with hypertension should seek to reduce stress and improve overall well being using mindfulness techniques. MBSR is a safe option and well accepted practice [19].

IX. SLEEP

What is known and what is new?

Sleep hygiene is important for vital function maintenance. Meanwhile, poor sleep hygiene, including that related to behavioural deviations, and sleep disruptions are associated with adverse health effects [228,229]. Inadequate sleep duration, both acute and chronic sleep deprivation and sleep disorders cause sympathetic hyperactivity resulting in blunted BP regulation, mainly absence or reversal of nocturnal BP dipping and/or nocturnal hypertension [229–231]. Sleep characteristics, including sleep quality, sleep duration and sleep efficiency, are associated with the risk of hypertension based on office BP [68,232]. According to the Coronary Artery Risk Development in Young Adults (CARDIA) study, shorter sleep duration and lower sleep efficiency were associated with higher office BP and BP increase over 5 years follow-up [233]. Prospective study from the UK Biobank showed that healthy sleep (sleep duration 7–8 h, no insomnia, no snoring, no excessive daytime sleepiness, early chronotype) is associated with lower incident hypertension risk [234] and, in hypertensive patients, it is inversely associated with the incidence of cardiometabolic morbidities, including coronary heart disease, diabetes mellitus and stroke [235]. Mendelian randomization analyses suggest potentially deleterious effects of more frequent daytime napping on cardiometabolic health (BP and abdominal obesity) [236]. A retrospective analysis from the UK Biobank suggested that frequent daytime napping is associated with a 12% increased risk of developing high BP [237]. Although both night-time sleep and daytime napping are associated with the decline in BP [238], and single cohort studies demonstrate negative association between daytime nap duration and hypertension presence [239], the evidence on long-term outcomes is lacking. Aligning sleep behaviours with endogenous circadian rhythms may be important in increasing SBP dipping, particularly if a part of the sleep period overlaps with the circadian peak in BP that occurs early in the evening [233,240,241].

Abnormal sleep patterns and obstructive sleep apnoea (OSA) are often associated with cardiometabolic risk factors, such as obesity, insulin resistance, impaired lipid and glucose metabolism, atherosclerosis and chronic inflammation [242–245]. Lifestyle factors and lifestyle interventions (in particular, weight reduction, diet, exercise training) significantly affect OSA severity metrics [246,247]. Salt intake might play a significant role in OSA severity in states associated with fluid retention and its redistribution in recumbent position (in particular, resistant hypertension and hyperaldosteronism) [248].

Sleep-disordered breathing, in particular, OSA, is considered a leading cause of secondary hypertension [249] and a risk factor for nocturnal and also resistant hypertension in up

to 85% of cases [17]. However, the antihypertensive effect of OSA treatment through the use of continuous positive airway pressure (CPAP) or other oral appliances is rather modest, comprising of a decrease in only 2–4 mmHg and depends on a number of other factors, including pretherapy BP, adherence and tolerance to the procedure (more than 6 h per night) and daytime sleepiness [250,251]. The effects of OSA treatment on metabolic disorders are partly inconsistent; it appears that some OSA patients' subgroups might benefit more than others from the European Sleep Apnea Database (ESADA) cohort demonstrates that some drug classes (beta-blockers in monotherapy or in combination) are associated with the better BP control in OSA patients depending on few clinical characteristics (age, gender, obesity) [253]. There is limited evidence on the association between habitual snoring at least four nights/week and around 1.5-fold increased incidence and prevalence of hypertension [254–256].

Recent studies suggest that irregular sleep timing [242,243], circadian disorders (including shift-work) [231], insomnia [257,258], parasomnias, periodic limb movements [258], central hypersomnias [259] and daytime napping [260] are associated with hypertension and other adverse cardiometabolic profiles. However, the evidence is scarce for any definite conclusions.

International perspective

Local, national and ethnic sleep-related traditions (e.g. polyphasic sleep, siesta, midnight sun patterns) as well as predisposing factors (i.e. ethnic/racial differences in sleep apnoea prevalence) should be considered [261–264]. Furthermore, sleep duration contributes to racial disparities in hypertension risk [265,266].

In general, data on daytime napping is limited; however, in countries with siesta tradition, it should not exceed 30 min daily.

In countries with midnight sun, the behavioural approaches are useful and should be recommended including the use of blackout blinds and sleep masks at night, as well as blue light blockers or sunglasses in the evening.

Mendelian randomization studies show an association between sleeping patterns and BP increase. Interestingly, genetic architecture of daytime napping is shared with BP traits [236], and genetically predicted short sleep duration is associated with risk of hypertension [232]. Meanwhile, data is less established related to genetically predicted longer sleep duration [232,267].

Recommendation

1. Weight control should be considered to improve sleep quality and management of sleep disorders (specially OSA).
2. Sleep hygiene should be addressed and implemented routinely [228,229,268,269]. Sleep duration of 7–9 h per night is recommended for adults [228,229,268,269].
3. Sleep hygiene approaches include appropriate sleep environment, regular sleep timing, sleep routine to prepare for sleep, avoidance of food intake, caffeine, alcohol and cigarettes close to bedtime and at night,

exposure to bright light and exercise during the day and their restriction before sleep.

4. Stimulus control and building a strong association between the bed/bedroom and sleep should be implemented: go to bed only when you are sleepy; establish a sleep routine; apply the 20 min rule (i.e. if you cannot get to sleep in 20 min, get up and go to another room and do nonstimulating activities).
5. Daytime napping for more than 30 min should not be routinely recommended given its effects on sleep pressure and nocturnal sleep and the lack of long-term outcome evidence.
6. Shift-workers might require an individualized approach to develop sleep–wakefulness patterns. Ambulatory BP monitoring should be considered for diagnosis and follow-up evaluations to assess nocturnal BP patterns.
7. Screening of sleep disorders (in particular, OSA [245]) should be performed in resistant hypertension, in patients with nocturnal hypertension and/or abnormal BP dipping. If applicable, treatment should be implemented, focusing on lifestyle changes and use of specific treatment (CPAP, oral appliances) to aid good sleeping habits.
8. People with known sleep disorders (snoring, OSA, insomnia, etc.) should undergo regular BP measurement. Ambulatory BP monitoring should be considered for BP nocturnal pattern evaluation.

Implementation

All healthcare professionals should be encouraged to evaluate sleep in a multimodal manner [270] (sleep satisfaction, timing, efficiency, duration, alertness, sleep disorders) in all patients with the involvement of partners/relatives, when needed. In case of abnormal sleep habits and/or sleep patterns, personalized counselling about the role of sleep in BP management and advice on modification of sleep habits should be implemented. Clear and achievable goals should be set in collaboration with the patient, potential challenges and barriers and motivating strategies should be discussed and strategies implemented. Patients experiencing persistent sleep difficulties (≥ 4 weeks) should be referred to a sleep specialist for further evaluation and management. Cognitive behavioural therapy for insomnia is beneficial for patients with hypertension, improving sleep quality and decreasing SBP and DBP by 8.6 and 5.6 mmHg, respectively [222].

At policy level, regulations of working schedules and educational programs in the sleep field must be promoted. Population-based awareness programs addressing healthy sleep must be supported.

X. SMOKING

What is known and what is new?

Smoking is not directly involved in the pathogenesis of essential hypertension but can cause acute increase of BP and is a known independent and strong risk factor for CVD. Smokers may have white coat and masked hypertension in the clinic, which is present on daytime ambulatory or home

BP monitoring [271]. Smoking affects the cardiovascular system structurally and functionally through interaction of hundreds of substances present in cigarettes and other tobacco products [272]. It serves as a major risk factor for atherosclerosis as evidenced from population studies [273,274].

Emerging data reveals the use of electronic cigarettes (e-cigarette) leads to acute increase of BP so we recommend against their use [275].

International perspective

Tobacco is the number one risk factor for global attributable deaths in men and the sixth in women [276]. Importantly, tobacco smoking, including second hand (passive) smoking, is within the top five risk factors for burden of disease worldwide, except sub-Saharan Africa [277].

Global prevalence of smoking has significant geographic, age and gender variation. Countries in Asia and Oceania have the highest prevalence of tobacco smoking among men while higher prevalence among females is seen in America and Western Europe [278].

Recommendation

- Smoking (including e-cigarettes) cessation is strongly recommended for its well established health benefits [274]. Importantly, strategies and approaches to avoid weight gain following smoking cessation should be implemented.
- Key target populations are those already exposed to smoking and those yet to be exposed, composed largely of young generations.
- Brief intervention counselling and motivational interviewing techniques in healthcare centres is shown to be very effective in tobacco cessation [279].

Implementation

Smoking cessation is the key factor in lowering risk impact on health. Enablers to quit smoking must be created through all levels, from the grass root healthcare settings to policy levels. At the healthcare centre level, brief intervention counselling for smoking and other tobacco use must be the standard practice care of any health service delivery model. Healthcare providers can be trained on simple modules of motivational interviewing techniques to reach the broader community. All patients must be assessed for smoking and on readiness to quit. Readiness to quit must be followed up till successful and continuing smoking cessation.

At policy level, population-based control programs can be administered. Policy regulations such as tobacco taxation, and separation of smoking and nonsmoking areas at public places must be sustained and expanded. Population-based awareness and advocacy programs addressing harmful effects of smoking must be advocated.

XI. POLLUTION EXPOSURE

What is known and what is new?

Each year, air pollution causes millions of premature deaths globally and is recognized as a modifiable risk factor for

CVD, respiratory disease and cancer [280–283]. Air pollution is a complex mixture of thousands of gaseous and particulate matter components, with the primary source of urban air pollution coming from motor vehicle exhausts [284].

The main components of air pollution associated with detrimental health effects are airborne particulate matter (especially fine particulate with an aerodynamic diameter $\leq 2.5 \mu\text{m}$) and gaseous substances including ozone, nitrogen dioxide, sulphur dioxide and carbon monoxide [285].

Inhaled ultrafine particulate matter rapidly passes from the lungs into the systemic circulation [286] where acute BP-raising effects are exerted through vasoconstriction of conduit arteries, systemic inflammation, endothelial dysfunction and autonomic nervous system imbalance [21,282,287–289].

Epidemiological studies show a strong correlation between chronic exposure to elements of air pollution and increased BP, as well as incident hypertension and organ damage associated with hypertension [290–296]. Human experimental data also demonstrate a dose–response relationship between air pollution and SBP [297].

Reducing population exposure to air pollution would significantly improve health, reduce mortality and lower health system expenditure [298,299].

Emerging data also indicates that noise pollution may correlate with increased risk of hypertension [300], and reducing exposure to noise could help control BP. More data with adjustment for confounding variables is needed to evaluate this relationship more comprehensively [301].

International perspective

Air quality has improved in high-income countries but has worsened in most low-income and middle-income countries alongside increased urbanization and growing economies [283].

Inequity regarding increased air pollution exposure is experienced by the poor and disempowered, and women and children in low-economic settings where polluting fuels have to be used for cooking, heating and lighting [302].

Policy makers internationally should follow the WHO Global Air Quality Guidelines and Good Practice Statements to inform evidence-based legislation and policies to improve air quality and mitigate population exposure to air pollution [283]. Use public transport, or travel by walking or cycling in preference to car or motorcycle to reduce pollution at population level should be encouraged. These guidelines should be used by healthcare professionals and medical societies to raise awareness and advise the public and patients [283,302]. Individuals can reduce exposure to air pollution by modifying the location, timing and type of outdoor activity [303–305].

Recommendations

1. Exercise in parks and gardens away from busy roadways.
2. Limit time spent outdoors during highly polluted periods.
3. Avoid inefficient burning of biomass for domestic heating.

4. Consider using ventilation systems with filtration for homes in high pollution areas.

Implementation

The greatest benefits to health can be achieved through national level policies that reduce air pollution at the point of source. These policies are specific to context and place, and must be applied across sectors, including transport, urban planning, industry, agriculture, energy and power generation [302].

Governments must play a major role by engaging widely at regional, national and international levels to advise policy options to prevent air pollution exposure and attain the greatest health benefits for communities [283,302].

XII. NONPRESCRIPTION MEDICINE AND NUTRITIONAL SUPPLEMENTS

What is known and what is new?

Nonprescription medicines include an array of medications ranging from anti-inflammatories to herbal medications. Herein, we focused on specific nutrient supplements and their relationship with BP, excluding herbal medicines and nonnutritional foods.

Although the link between vitamin intake and hypertension is indicated, evidence-based research studies are scarce. Studies suggested multivitamin supplementation decreases high-BP-related mortality [306,307]. However, these effects were not observed in middle-aged and older women [308]. Several studies support the role played by hydrosoluble vitamins in maintaining both SBP and DBP in various populations [309–311] and a meta-analysis of 29 RCTs revealed that vitamin C supplementation reduced SBP and DBP [312]. Circulating micronutrient status is associated with adiposity, SBP and cardiometabolic risk factors, such as vitamin D with obesity parameters in women [313–315], imposing alternative pathways for the vitamin D role in sustaining optimal BP. Importantly, vitamin D metabolism is dependent on the appropriate intake of calcium and magnesium. It was reported that daily magnesium supplementation lowers BP [316]. Furthermore, a meta-regression also indicated a decrease in BP with long-term potassium supplementation of 3500 mg/day, especially in subjects not taking antihypertensives [317]. Similarly, chlorogenic acids, compounds contained in apples, coffee, tomatoes and other vegetables, might decrease BP, also indirectly through effects on adiposity, then hypoglycaemic, hypouricemic effects and similar [318].

Additionally, intake of functional foods based on onion and garlic, rich in flavanols, as well as catechin-containing green tea with angiotensin-converting enzyme (ACE) inhibitory properties, can be a food-based nutritional strategy to aid BP management and CVD prevention [319].

Importantly, extensive research on the impact of these supplementations is needed and evidence of effects on BP is still minimal.

International perspective

Lack of adequately powered RCTs does not allow streamlined and efficient implementation process for the

suggested intake of dietary supplements and nonprescription medications for BP optimization.

Micronutrient and bioactive intake personalization based on individual characteristics, mainly related to ethnicity and genetic background, is an envisaged strategy for the future of hypertension primary prevention.

Recommendation

1. Vitamins, minerals and micronutrients intake should be obtained as part of a healthy, nutrient-rich and balanced diet.
2. Health supplements are not a replacement for a balanced diet and are not recommended as a treatment to hypertension.

Implementation

Considering interpersonal variability in response to a nutrient intake, preexisting health concerns and dietary preferences, a personalized approach to nutrition and supplementation is essential for the nutritional management of hypertension, while focusing on higher intake of certain micronutrients (as part of a normal diet) and functional foods.

XIII. BEHAVIOURAL INTERVENTIONS, DIGITAL HEALTH AND WEARABLES

What is known and what is new?

Behavioural intervention is an effective approach for the treatment of CVD [320,321]. A recent large-scale systematic review [322], including 150 RCTs identified the clinically supervised Patient Self-Management Support, frequently based on Wagner's Chronic Care Model, the most dominant intervention method [323]. Importantly, many chronic illnesses share the same modifiable behavioural lifestyle risk factors.

Behavioural habits known to modulate cardiovascular risk and BP control include oral hygiene practices as both dental care access and periodontitis are independently linked to hypertension and target organ damage [324]. This relationship is independent of socioeconomic status [325].

The use of digital technologies for prevention and/or treatment of health conditions, termed digital health interventions, is an emerging and promising approach to support people with hypertension [326,327].

Recent studies showed clinically modest but significant effects of digital interventions to reduce SBP, improve medication adherence and lifestyle behavioural changes longer term [328,329].

Use of personalization through digital health (e.g. electronic health records or mobile devices to support health practices) interventions can, especially aid in the delivery of timely precision medicine strategies and be delivered directly by the healthcare practitioner and/or digitally, through the utilization of end-user manual inputs and/or digital system-derived inputs [330].

Providing a dynamic and adaptive hypertensive digital intervention into the future that can adjust its responses when the individual changes (in real-time) should better facilitate behaviour change outcomes [331–333]. Additionally,

antihypertensive digital interventions that can minimize the amount of manual end-user input may also increase patient adherence [328,330,332], especially through the consistent use of validated digital wearables to support and maintain lifestyle modifications. There is, however, a need to ensure that any digital BP device has its accuracy validated, and we caution the use of unvalidated devices.

Importantly, digital health approaches should not supplant the role of clinicians but are tools that clinicians and patients can use to improve the efficiency and effectiveness of clinical objectives. Digital health platforms enable the efficient exchange of clinical data and clinician–patient interactions to achieve effective lifestyle modifications.

International perspective

Behaviour change techniques are included in the medical curriculum for healthcare providers in many countries [334–336] reinforcing the importance of chronic illnesses burden control through effective population-based intervention programs.

The US Centres for Disease Control and Prevention websites contain a large compendium of local and global studies centred on the benefits of behavioural interventions upon modifiable risk factors for chronic disease prevalence and outcomes.

Recommendations

1. Healthcare providers should be trained in behaviour change approaches and digital support technologies to facilitate action and control hypertension [337].
2. Behavioural changes should be comprehensive and encompass oral hygiene improvement and maintenance to prevent periodontitis and improve BP control [324].
3. Continuous use of validated digital wearables and digital tools to improve symptom awareness, introduce potential reminders (e.g. for medication intake or promote increased incidental exercise), aid behaviour change (e.g. goal setting, decision support, self-monitoring, which may integrate wearable data, reminder and alert systems, digital support conversational agents to improve user adherence, motivation and self-efficacy, intervention personalization programming) should be encouraged.
4. Introduce communication technologies to facilitate professional guidance and input or social support (digital spaces to communicate with healthcare providers or peers, SMS messaging) as a standalone digital intervention or integrated into a multicomponent intervention [326].

Implementation

The use of digital health interventions that incorporate behaviour change models and include personalization is the most promising but reliable RCTs showing efficacy are lacking.

Data collection, either using manual or system-based input, can be achieved through smart programming and/or artificial intelligence software to continually personalize digital intervention (e.g. offering relevant tailored content, sending reminders about targets or encouragement if

motivation is declining, alerting the patient (and practitioner) if symptoms are deteriorating), in real-time.

Lastly, the greater inclusion of wearables and integration of smarter programming and artificial intelligence into such interventions is predicted to strengthen behaviour change and improve health outcomes [332,338].

XIV. HOLISTIC APPROACH TO BLOOD PRESSURE PREVENTION ACROSS THE GLOBE

What is known and what is new?

Urbanization is considered a major contributor to the rising prevalence of hypertension worldwide. Higher prevalence of hypertension in urban areas includes socioeconomic and lifestyle changes that increase the diagnosis of overweight, obesity, smoking and diabetes [339]. Other mechanisms are thought to involve psychosocial stress in urban residents resulting from financial stress, redefinition of cultural identity and movement away from traditional coping mechanisms, including social support from extended family. These changes have been reported in different epidemiological studies worldwide [340].

Another study shows the relationship between education and lifestyle and healthcare changes [341]. The risk factors for unhealthy nutritional habits, stress, poor working conditions or inadequate access to education and appropriate medical services are present in a large part of the urban population [342], but evidence is lacking in the rural population. Furthermore, these risk factors are more pronounced between gender differences.

There is undeniable need to change our current standards of lifestyle approach, such as an integral treatment of hypertension patients. Implementation of HEARTS-WHO model based on the premise that a person needs wholesome care and not just the treatment for the disease should be implemented. The holistic and wholesome patient care approach takes into consideration all patient aspects, including physical, emotional, social, economic, cultural and religious/spiritual needs to provide the most appropriate advice and treatment [343].

International perspective

Worldwide, factors are similar, although inequalities in health access between populations have been described [344–346], and several studies have shown that diagnosis, control and treatment of hypertension are worse in rural areas than in urban areas [347]. Indigenous populations worldwide experience a huge gap in access to general health and health treatments and mortality [348], including CVD and diabetes [347,349,350]. This divide is observed as a combination of racial discrimination, poverty, lifestyle and access to services due to geographical isolation [348,351]. Socioeconomic differences between poorer and richer countries and populations also impact general health access and CVD morbidity and mortality [346,351].

Recommendation

1. Implementation of holistic care for patients, as evidenced in Bhutan, Latin America, Canada and some

countries in Africa, to provide the optimal treatment and management of BP [23,352–354].

2. Improve access to BP diagnosis, treatment and management in regional and rural areas.

Implementation

Healthcare providers should consider a holistic, multicomponent approach addressing many lifestyle changes simultaneously. Comprehensive hypertension lifestyle changes such as exercise, education, nutritional habits and policies should be targeted for both rural and urban areas.

Exercise and nutritional habits according to culture, ethnicity and religion should be considered in populations around the world to achieve equitable population outreach.

Differential work must be done on vulnerable populations to adequately identify the necessary policies to be implemented and maintained.

XV. OVERALL SUMMARY AND A GLIMPSE INTO THE FUTURE FOR ASSISTED LIFESTYLE INTERVENTIONS

In this position paper, we presented recommendations that support a healthy lifestyle to manage and maintain a healthy BP – more exercise, less salt, sugar and saturated fats in diet, more fruit, vegetables, and dietary fibre, no smoking and alcohol. We also presented emerging evidence for newer modalities and discussed the potential for increased use of digital technologies to help improve lifestyle behaviours in a personalized way. Indeed, there are many mobile phone ‘apps’ that can be used to help track movement, diet and habits. These technologies can be used for weight management, assessment of meal composition and calorie content by scanning meals using phones. They can also be used to help individuals meditate and reduce stress. These apps are international in scope and reach and will play an important part in encouraging interventions. These technologies will probably integrate the next ‘more efficient’ step in precision lifestyle interventions alongside assessment by genetic components using polygenic risk scores (Supplementary Box 4, <http://links.lww.com/HJH/C287>), which has the potential to be used to calculate risk of CVD and allow early personalized approaches to treatment. Importantly, data to support many of these revolutionary interventions are scarce and larger studies are essential to assess efficacy.

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Conflicts of interest

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REFERENCES

1. NCD Risk Factor Collaboration. Worldwide trends in hypertension prevalence and progress in treatment and control from 1990 to 2019: a pooled analysis of 1201 population-representative studies with 104 million participants. *Lancet* 2021; 398:957–980.
2. Tomaszewski M, Itoh H. ISH2022KYOTO Hypertension Zero Declaration. *Hypertens Res* 2023; 46:1–2.
3. Cheema KM, Dicks E, Pearson J, Samani NJ. Long-term trends in the epidemiology of cardiovascular diseases in the UK: insights from the British Heart Foundation statistical compendium. *Cardiovasc Res* 2022; 118:2267–2280.
4. Eljovich F, Weinberger MH, Anderson CA, Appel LJ, Burszty N, Cook NR, *et al.*, American Heart Association Professional and Public Education Committee of the Council on Hypertension; Council on Functional Genomics and Translational Biology; and Stroke Council. Salt sensitivity of blood pressure: a scientific statement from the American Heart Association. *Hypertension* 2016; 68:e7–e46.
5. Appel LJ, Brands MW, Daniels SR, Karanja N, Elmer PJ, Sacks FM, *et al.*, American Heart Association. Dietary approaches to prevent and treat hypertension: a scientific statement from the American Heart Association. *Hypertension* 2006; 47:296–308.
6. Cushman WC, Cutler JA, Hanna E, Bingham SF, Follmann D, Harford T, *et al.* Prevention and Treatment of Hypertension Study (PATHS): effects of an alcohol treatment program on blood pressure. *Arch Intern Med* 1998; 158:1197–1207.
7. Grossman E, Grossman A, Schein MH, Zimlichman R, Gavish B. Breathing-control lowers blood pressure. *J Hum Hypertens* 2001; 15:263–269.
8. He FJ, Li J, Macgregor GA. Effect of longer term modest salt reduction on blood pressure: Cochrane systematic review and meta-analysis of randomised trials. *BMJ* 2013; 346:f1325.
9. Whelton SP, Chin A, Xin X, He J. Effect of aerobic exercise on blood pressure: a meta-analysis of randomized, controlled trials. *Ann Intern Med* 2002; 136:493–503.
10. Saneei P, Salehi-Abargouei A, Esmailzadeh A, Azadbakht L. Influence of Dietary Approaches to Stop Hypertension (DASH) diet on blood pressure: a systematic review and meta-analysis on randomized controlled trials. *Nutr Metab Cardiovasc Dis* 2014; 24:1253–1261.

11. Svetkey LP, Simons-Morton D, Vollmer WM, Appel LJ, Conlin PR, Ryan DH, *et al.* Effects of dietary patterns on blood pressure: subgroup analysis of the Dietary Approaches to Stop Hypertension (DASH) randomized clinical trial. *Arch Intern Med* 1999; 159:285–293.
12. Khan N, Bacon SL, Khan S, Perlmutter S, Gerlinsky C, Dermer M, *et al.*, Hypertension Canada Priority Setting Partnership Group. Hypertension management research priorities from patients, caregivers, and healthcare providers: a report from the Hypertension Canada Priority Setting Partnership Group. *J Clin Hypertens (Greenwich)* 2017; 19:1063–1069.
13. Carey RM, Wright JT Jr, Taler SJ, Whelton PK. Guideline-driven management of hypertension: an evidence-based update. *Circ Res* 2021; 128:827–846.
14. Leung AA, Daskalopoulou SS, Dasgupta K, McBrien K, Butalia S, Zarnke KB, *et al.*, Hypertension Canada. Hypertension Canada's 2017 Guidelines for diagnosis, risk assessment, prevention, and treatment of hypertension in adults. *Can J Cardiol* 2017; 33:557–576.
15. Unger T, Borghi C, Charchar F, Khan NA, Poulter NR, Prabhakaran D, *et al.* 2020 International Society of Hypertension Global Hypertension Practice Guidelines. *Hypertension* 2020; 75:1334–1357.
16. Whelton PK, Carey RM, Aronow WS, Casey DE Jr, Collins KJ, Dennison Himmelfarb C, *et al.* 2017 ACC/AHA/AAPA/ABC/ACPM/AGS/APhA/ASH/ASPC/NMA/PCNA Guideline for the prevention, detection, evaluation, and management of high blood pressure in adults: executive summary: a report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. *Circulation* 2018; 138:e426–e483.
17. Williams B, Mancia G, Spiering W, Agabiti Rosei E, Azizi M, Burnier M, *et al.* 2018 ESC/ESH Guidelines for the management of arterial hypertension. *Eur Heart J* 2018; 39:3021–3104.
18. International Society of Hypertension. 8 simple rules for living with hypertension. Available at: <https://www.youtube.com/watch?v=9YU9Xhhjin4>. YouTube2021.
19. Lee EKP, Yeung NCY, Xu Z, Zhang D, Yu CP, Wong SYS. Effect and acceptability of mindfulness-based stress reduction program on patients with elevated blood pressure or hypertension: a meta-analysis of randomized controlled trials. *Hypertension* 2020; 76:1992–2001.
20. Hanssen H, Boardman H, Deiseroth A, Moholdt T, Simonenko M, Krankel N, *et al.* Personalized exercise prescription in the prevention and treatment of arterial hypertension: a Consensus Document from the European Association of Preventive Cardiology (EAPC) and the ESC Council on Hypertension. *Eur J Prev Cardiol* 2022; 29:205–215.
21. Parsanathan R, Palanichamy R. Air pollution impairs endothelial function and blood pressure. *Hypertens Res* 2022; 45:380–381.
22. Silvestri D, Goutos D, Lloren A, Zhou S, Zhou G, Farietta T, *et al.* Factors associated with disparities in hospital readmission rates among US adults dually eligible for medicare and medicaid. *JAMA Health Forum* 2022; 3:e214611.
23. Campbell NRC, Ordunez P, Giraldo G, Rodriguez Morales YA, Lombardi C, Khan T, *et al.* WHO HEARTS: a global program to reduce cardiovascular disease burden: experience implementing in the Americas and opportunities in Canada. *Can J Cardiol* 2021; 37:744–755.
24. World Health Organisation. Obesity and overweight. 2021.
25. Kaess BM, Jozwiak J, Mastej M, Lukas W, Grzeszczak W, Windak A, *et al.* Association between anthropometric obesity measures and coronary artery disease: a cross-sectional survey of 16,657 subjects from 444 Polish cities. *Heart* 2010; 96:131–135.
26. Xu X, Eales JM, Jiang X, Sanderson E, Drzal M, Saluja S, *et al.* Contributions of obesity to kidney health and disease: insights from Mendelian randomization and the human kidney transcriptomics. *Cardiovasc Res* 2022; 118:3151–3161.
27. Kim MS, Kim WJ, Khara AV, Kim JY, Yon DK, Lee SW, *et al.* Association between adiposity and cardiovascular outcomes: an umbrella review and meta-analysis of observational and Mendelian randomization studies. *Eur Heart J* 2021; 42:3388–3403.
28. Jayedi A, Rashidy-Pour A, Khorshidi M, Shab-Bidar S. Body mass index, abdominal adiposity, weight gain and risk of developing hypertension: a systematic review and dose-response meta-analysis of more than 2.3 million participants. *Obes Rev* 2018; 19:654–667.
29. Appel LJ, Moore TJ, Obarzanek E, Vollmer WM, Svetkey LP, Sacks FM, *et al.* A clinical trial of the effects of dietary patterns on blood pressure. DASH Collaborative Research Group. *N Engl J Med* 1997; 336:1117–1124.
30. Bakris G, Ali W, Parati G. ACC/AHA versus ESC/ESH on hypertension guidelines: JACC guideline comparison. *J Am Coll Cardiol* 2019; 73:3018–3026.
31. Mertens IL, Van Gaal LF. Overweight, obesity, and blood pressure: the effects of modest weight reduction. *Obesity research* 2000; 8:270–278.
32. Azegami T, Uchida K, Arima F, Sato Y, Awazu M, Inokuchi M, *et al.* Association of childhood anthropometric measurements and laboratory parameters with high blood pressure in young adults. *Hypertens Res* 2021; 44:711–719.
33. Vallée A, Perrine A-L, Deschamps V, Blacher J, Olié V. Relationship between dynamic changes in body weight and blood pressure: the ESTEBAN Survey. *Am J Hypertens* 2019; 32:1003–1012.
34. Islam SMS, Mainuddin A, Islam MS, Karim MA, Mou SZ, Arefin S, *et al.* Prevalence of risk factors for hypertension: a cross-sectional study in an urban area of Bangladesh. *Global Cardiol Sci Pract* 2015; 2015:43.
35. Wang L, Lin M, Yu J, Fan Z, Zhang S, Lin Y, *et al.* The impact of bariatric surgery versus non-surgical treatment on blood pressure: systematic review and meta-analysis. *Obes Surg* 2021; 31:4970–4984.
36. Fernstrom JD, Courcoulas AP, Houck PR, Fernstrom MH. Long-term changes in blood pressure in extremely obese patients who have undergone bariatric surgery. *Arch Surg* 2006; 141:276–283.
37. Schiavon CA, Ikeoka D, Santucci EV, Santos RN, Damiani LP, Bueno PT, *et al.* Effects of bariatric surgery versus medical therapy on the 24-hour ambulatory blood pressure and the prevalence of resistant hypertension. *Hypertension* 2019; 73:571–577.
38. le Roux CW, Astrup A, Fujioka K, Greenway F, Lau DCW, Van Gaal L, *et al.* 3 years of liraglutide versus placebo for type 2 diabetes risk reduction and weight management in individuals with prediabetes: a randomised, double-blind trial. *Lancet* 2017; 389:1399–1409.
39. Wilding JPH, Batterham RL, Calanna S, Davies M, Van Gaal LF, Lingway I, *et al.*, STEP 1 Study Group. Once-weekly semaglutide in adults with overweight or obesity. *N Engl J Med* 2021; 384:989–1002.
40. Mulrow CD, Chiquette E, Angel L, Cornell J, Summerbell CD, Anagnostis BB, *et al.* Dieting to reduce body weight for controlling hypertension in adults. *Cochrane Database Syst Rev* 1998; 2: CD000484.
41. Neter JE, Stam BE, Kok FJ, Grobbee DE, Geleijnse JM. Influence of weight reduction on blood pressure. *Hypertension* 2003; 42:878–884.
42. Biaggioni I. Should we target the sympathetic nervous system in the treatment of obesity-associated hypertension? *Hypertension* 2008; 51:168–171.
43. Doumouras AG, Wong JA, Paterson JM, Lee Y, Sivapathasundaram B, Tarride JE, *et al.* bariatric surgery and cardiovascular outcomes in patients with obesity and cardiovascular disease: a population-based retrospective cohort study. *Circulation* 2021; 143:1468–1480.
44. Olsen MH, Angell SY, Asma S, Boutouyrie P, Burger D, Chirinos JA, *et al.* A call to action and a lifecourse strategy to address the global burden of raised blood pressure on current and future generations: the Lancet Commission on hypertension. *Lancet* 2016; 388:2665–2712.
45. Blumenthal JA, Babyak MA, Hinderliter A, Watkins LL, Craighead L, Lin P-H, *et al.* Effects of the DASH diet alone and in combination with exercise and weight loss on blood pressure and cardiovascular biomarkers in men and women with high blood pressure: the ENCORE study. *Arch Intern Med* 2010; 170:126–135.
46. Katzmarzyk PT, Bray GA, Greenway FL, Johnson WD, Newton RL Jr, Ravussin E, *et al.* Ethnic-specific BMI and waist circumference thresholds. *Obesity (Silver Spring)* 2011; 19:1272–1278.
47. Khan NA, Stergiou GS, Omboni S, Kario K, Renna N, Chapman N, *et al.* Virtual management of hypertension: lessons from the COVID-19 pandemic-International Society of Hypertension position paper endorsed by the World Hypertension League and European Society of Hypertension. *J Hypertens* 2022; 40:1435–1448.
48. Islam SMS, Farmer AJ, Bobrow K, Maddison R, Whittaker R, Pfaeffli Dale LA, *et al.* Mobile phone text-messaging interventions aimed to prevent cardiovascular diseases (Text2PreventCVD): systematic review and individual patient data meta-analysis. *Open Heart* 2019; 6: e001017.
49. Islam SMS, Talukder A, Awal MA, Siddiqui MMU, Ahamad MM, Ahammed B, *et al.* Machine learning approaches for predicting hypertension and its associated factors using population-level data from three South Asian countries. *Front Cardiovasc Med* 2022; 9:839379.

50. Semlitsch T, Krenn C, Jeitler K, Berghold A, Horvath K, Siebenhofer A. Long-term effects of weight-reducing diets in people with hypertension. *Cochrane Database Syst Rev* 2021; 3:CD008274.
51. Williams B, Mancia G, Spiering W, Agabiti Rosei E, Azizi M, Burnier M, et al., List of authors/Task Force members. 2018 Practice Guidelines for the management of arterial hypertension of the European Society of Hypertension and the European Society of Cardiology: ESH/ESC Task Force for the Management of Arterial Hypertension. *J Hypertens* 2018; 36:2284–2309.
52. Rabi DM, McBrien KA, Sapir-Pichhadze R, Nakhla M, Ahmed SB, Dumanski SM, et al. Hypertension Canada's 2020 Comprehensive Guidelines for the prevention, diagnosis, risk assessment, and treatment of hypertension in adults and children. *Can J Cardiol* 2020; 36:596–624.
53. Barone Gibbs B, Hivert MF, Jerome GJ, Kraus WE, Rosenkranz SK, Schorr EN, et al., American Heart Association Council on Lifestyle and Cardiometabolic Health; Council on Cardiovascular and Stroke Nursing; and Council on Clinical Cardiology. Physical activity as a critical component of first-line treatment for elevated blood pressure or cholesterol: who, what, and how?: a scientific statement from the American Heart Association. *Hypertension* 2021; 78:e26–e37.
54. Denham J, Brown NJ, Tomaszewski M, Williams B, O'Brien BJ, Charchar FJ. Aortic augmentation index in endurance athletes: a role for cardiorespiratory fitness. *Eur J Appl Physiol* 2016; 116:1537–1544.
55. Rao P, Belanger MJ, Robbins JM. Exercise, physical activity, and cardiometabolic health: insights into the prevention and treatment of cardiometabolic diseases. *Cardiol Rev* 2022; 30:167–178.
56. Tomaszewski M, Charchar FJ, Przybycin M, Crawford L, Wallace AM, Gosek K, et al. Strikingly low circulating CRP concentrations in ultramarathon runners independent of markers of adiposity: how low can you go? *Arterioscler Thromb Vasc Biol* 2003; 23:1640–1644.
57. Cheng C, Zhang D, Chen S, Duan G. The association of cardiorespiratory fitness and the risk of hypertension: a systematic review and dose-response meta-analysis. *J Hum Hypertens* 2021; 36:744–752.
58. Del Pozo Cruz B, Ahmadi M, Inan-Eroglu E, Huang BH, Stamatakis E. Prospective associations of accelerometer-assessed physical activity with mortality and incidence of cardiovascular disease among adults with Hypertension: the UK Biobank Study. *J Am Heart Assoc* 2022; 11: e023290.
59. Naci H, Salcher-Konrad M, Dias S, Blum MR, Sahoo SA, Nunan D, et al. How does exercise treatment compare with antihypertensive medications? A network meta-analysis of 391 randomised controlled trials assessing exercise and medication effects on systolic blood pressure. *Br J Sports Med* 2019; 53:859–869.
60. Pescatello LS, Wu Y, Gao S, Livingston J, Sheppard BB, Chen MH. Do the combined blood pressure effects of exercise and antihypertensive medications add up to the sum of their parts? A systematic meta-review. *BMJ Open Sport Exerc Med* 2021; 7:e000895.
61. Cornelissen VA, Smart NA. Exercise training for blood pressure: a systematic review and meta-analysis. *J Am Heart Assoc* 2013; 2: e004473.
62. Cornelissen VA, Buys R, Smart NA. Endurance exercise beneficially affects ambulatory blood pressure: a systematic review and meta-analysis. *J Hypertens* 2013; 31:639–648.
63. de Sousa EC, Abrahim O, Ferreira ALL, Rodrigues RP, Alves EAC, Vieira RP. Resistance training alone reduces systolic and diastolic blood pressure in prehypertensive and hypertensive individuals: meta-analysis. *Hypertens Res* 2017; 40:927–931.
64. MacDonald HV, Johnson BT, Huedo-Medina TB, Livingston J, Forsyth KC, Kraemer WJ, et al. Dynamic resistance training as stand-alone antihypertensive lifestyle therapy: a meta-analysis. *J Am Heart Assoc* 2016; 5:e003231.
65. Carlson DJ, Dieberg G, Hess NC, Millar PJ, Smart NA. Isometric exercise training for blood pressure management: a systematic review and meta-analysis. *Mayo Clin Proc* 2014; 89:327–334.
66. Lopez-Valenciano A, Ruiz-Perez I, Ayala F, Sanchez-Meca J, Vera-Garcia FJ. Updated systematic review and meta-analysis on the role of isometric resistance training for resting blood pressure management in adults. *J Hypertens* 2019; 37:1320–1333.
67. Inder JD, Carlson DJ, Dieberg G, McFarlane JR, Hess NC, Smart NA. Isometric exercise training for blood pressure management: a systematic review and meta-analysis to optimize benefit. *Hypertens Res* 2016; 39:88–94.
68. Arnett DK, Blumenthal RS, Albert MA, Buroker AB, Goldberger ZD, Hahn EJ, et al. 2019 ACC/AHA Guideline on the primary prevention of cardiovascular disease: a report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. *J Am Coll Cardiol* 2019; 74:e177–e232.
69. Pescatello LS, Franklin BA, Fagard R, Farquhar WB, Kelley GA, Ray CA, American College of Sports Medicine. American College of Sports Medicine position stand. Exercise and hypertension. *Med Sci Sports Exerc* 2004; 36:533–553.
70. Vanhees L, Geladas N, Hansen D, Koudi E, Niebauer J, Reiner Z, et al. Importance of characteristics and modalities of physical activity and exercise in the management of cardiovascular health in individuals with cardiovascular risk factors: recommendations from the EACPR. Part II. *Eur J Prev Cardiol* 2012; 19:1005–1033.
71. Pescatello LS, MacDonald HV, Ash GI, Lamberti LM, Farquhar WB, Arena R, Johnson BT. Assessing the existing professional exercise recommendations for hypertension: a review and recommendations for future research priorities. *Mayo Clin Proc* 2015; 90:801–812.
72. Leal JM, Galliano LM, Del Vecchio FB. Effectiveness of high-intensity interval training versus moderate-intensity continuous training in hypertensive patients: a systematic review and meta-analysis. *Curr Hypertens Rep* 2020; 22:26.
73. Costa EC, Hay JL, Kehler DS, Boreskie KF, Arora RC, Umpierre D, et al. Effects of high-intensity interval training versus moderate-intensity continuous training on blood pressure in adults with pre to established hypertension: a systematic review and meta-analysis of randomized trials. *Sports Med* 2018; 48:2127–2142.
74. Oja P, Kelly P, Murtagh EM, Murphy MH, Foster C, Titze S. Effects of frequency, intensity, duration and volume of walking interventions on CVD risk factors: a systematic review and meta-regression analysis of randomised controlled trials among inactive healthy adults. *Br J Sports Med* 2018; 52:769–775.
75. Hanson S, Jones A. Is there evidence that walking groups have health benefits? A systematic review and meta-analysis. *Br J Sports Med* 2015; 49:710–715.
76. Jaeschke L, Steinbrecher A, Luzak A, Puggina A, Aleksovskaya K, Buck C, et al., DEDIPAC consortium. Socio-cultural determinants of physical activity across the life course: a 'Determinants of Diet and Physical Activity' (DEDIPAC) umbrella systematic literature review. *Int J Behav Nutr Phys Act* 2017; 14:173.
77. Sanchis-Gomar F, Lavie CJ, Marin J, Perez-Quilis C, Eijssvogels TMH, O'Keefe JH, et al. Exercise effects on cardiovascular disease: from basic aspects to clinical evidence. *Cardiovasc Res* 2022; 118:2253–2266.
78. Beune EJ, Haafkens JA, Agyemang C, Bindels PJ. Inhibitors and enablers of physical activity in multiethnic hypertensive patients: qualitative study. *J Hum Hypertens* 2010; 24:280–290.
79. Samara A, Nistrup A, Al-Rammah TY, Aro AR. Lack of facilities rather than sociocultural factors as the primary barrier to physical activity among female Saudi university students. *Int J Womens Health* 2015; 7:279–286.
80. Joseph RP, Ainsworth BE, Keller C, Dodgson JE. Barriers to physical activity among African American women: an integrative review of the literature. *Women Health* 2015; 55:679–699.
81. Visseren FLJ, Mach F, Smulders YM, Carballo D, Koskinas KC, Back M, et al., ESC National Cardiac Societies, ESC Scientific Document Group. 2021 ESC Guidelines on cardiovascular disease prevention in clinical practice. *Eur Heart J* 2021; 42:3227–3337.
82. Bull FC, Al-Ansari SS, Biddle S, Borodulin K, Buman MP, Cardon G, et al. World Health Organization 2020 guidelines on physical activity and sedentary behaviour. *Br J Sports Med* 2020; 54:1451–1462.
83. Dempsey PC, Larsen RN, Dunstan DW, Owen N, Kingwell BA. Sitting Less and Moving More: Implications for Hypertension. *Hypertension* 2018; 72:1037–1046.
84. American Heart Association. Getting active. 2022. Available at: <https://www.heart.org/en/healthy-living/fitness/getting-active>. [Accessed 15 February 2023]
85. Howlett N, Trivedi D, Troop NA, Chater AM. Are physical activity interventions for healthy inactive adults effective in promoting behavior change and maintenance, and which behavior change techniques are effective? A systematic review and meta-analysis. *Transl Behav Med* 2019; 9:147–157.
86. Mueller NT, Noya-Alarcon O, Contreras M, Appel LJ, Dominguez-Bello MG. Association of Age With Blood Pressure Across the Lifespan

- in Isolated Yanomami and Yekwana villages. *JAMA Cardiol* 2018; 3:1247–1249.
87. Carvalho JJ, Baruzzi RG, Howard PF, Poulter N, Alpers MP, Franco LJ, *et al.* Blood pressure in four remote populations in the INTERSALT Study. *Hypertension* 1989; 14:238–246.
 88. Filippou CD, Tsioufis CP, Thomopoulos CG, Mihos CC, Dimitriadis KS, Sotiropoulou LI, *et al.* Dietary Approaches to Stop Hypertension (DASH) Diet and Blood Pressure Reduction in Adults with and without Hypertension: A Systematic Review and Meta-Analysis of Randomized Controlled Trials. *Adv Nutr* 2020; 11:1150–1160.
 89. Juraschek SP, Miller ER 3rd, Weaver CM, Appel LJ. Effects of Sodium Reduction and the DASH Diet in Relation to Baseline Blood Pressure. *J Am Coll Cardiol* 2017; 70:2841–2848.
 90. Widmer RJ, Flammer AJ, Lerman LO, Lerman A. The Mediterranean diet, its components, and cardiovascular disease. *Am J Med* 2015; 128:229–238.
 91. Nishida C, Uauy R, Kumanyika S, Shetty P. The joint WHO/FAO expert consultation on diet, nutrition and the prevention of chronic diseases: process, product and policy implications. *Public Health Nutr* 2004; 7:245–250.
 92. National Institute for Health and Care Excellence. Prevention of cardiovascular disease. Manchester, UK: National Institute for Health and Care Excellence. 2014. p. 38.
 93. Savica V, Bellinghieri G, Kopple JD. The effect of nutrition on blood pressure. *Annu Rev Nutr* 2010; 30:365–401.
 94. Sacks FM, Svetkey LP, Vollmer WM, Appel LJ, Bray GA, Harsha D, *et al.*, DASH-Sodium Collaborative Research Group. Effects on blood pressure of reduced dietary sodium and the Dietary Approaches to Stop Hypertension (DASH) diet. DASH-Sodium Collaborative Research Group. *N Engl J Med* 2001; 344:3–10.
 95. Yokoyama Y, Nishimura K, Barnard ND, Takegami M, Watanabe M, Sekikawa A, *et al.* Vegetarian diets and blood pressure: a meta-analysis. *JAMA Intern Med* 2014; 174:577–587.
 96. Gorska-Warzewicz H, Laskowski W, Kulykovets O, Kudlinska-Chylak A, Czechtoko M, Rejman K. Food Products as Sources of Protein and Amino Acids-The Case of Poland. *Nutrients* 2018; 10:1977.
 97. Poppitt SD. Milk proteins and human health. *Milk Proteins: Elsevier*. Massachusetts, USA: Academic Press; 2020. pp. 651–669.
 98. Dasinger JH, Fehrenbach DJ, Abais-Battad JM. Dietary Protein: Mechanisms Influencing Hypertension and Renal Disease. *Curr Hypertens Rep* 2020; 22:13.
 99. Soeters PB. Editorial: vegan diets: what is the benefit? *Curr Opin Clin Nutr Metab Care* 2020; 23:151–153.
 100. Willett W. Lessons from dietary studies in Adventists and questions for the future. *Am J Clin Nutr* 2003; 78 (3 Suppl):539S–543S.
 101. Alexander S, Ostfeld RJ, Allen K, Williams KA. A plant-based diet and hypertension. *J Geriatr Cardiol* 2017; 14:327.
 102. Suter PM, Sierro C, Vetter W. Nutritional factors in the control of blood pressure and hypertension. *Nutr Clin Care* 2002; 5:9–19.
 103. Baradaran A, Nasri H, Rafieian-Kopaei M. Oxidative stress and hypertension: possibility of hypertension therapy with antioxidants. *J Res Med Sci* 2014; 19:358.
 104. Upadhyay S, Dixit M. Role of Polyphenols and Other Phytochemicals on Molecular Signaling. *Oxid Med Cell Longev* 2015; 2015:504253.
 105. Zhou MS, Wang A, Yu H. Link between insulin resistance and hypertension: What is the evidence from evolutionary biology? *Diabetol Metab Syndr* 2014; 6:12.
 106. Ernst E, Pietsch L, Matrai A, Eisenberg J. Blood rheology in vegetarians. *Br J Nutr* 1986; 56:555–560.
 107. Marques FZ, Nelson E, Chu PY, Horlock D, Fiedler A, Ziemann M, *et al.* High-Fiber Diet and Acetate Supplementation Change the Gut Microbiota and Prevent the Development of Hypertension and Heart Failure in Hypertensive Mice. *Circulation* 2017; 135:964–977.
 108. European Statistics. Daily consumption of fruit and vegetables by sex, age and educational attainment level. 2022.
 109. Australian Institute of Health and Welfare. Australia's health 2018. Canberra: Australian Govt. Pub. Service; 2018.
 110. Miller V, Mente A, Dehghan M, Rangarajan S, Zhang X, Swaminathan S, *et al.*, Prospective Urban Rural Epidemiology (PURE) study investigators. Fruit, vegetable, and legume intake, and cardiovascular disease and deaths in 18 countries (PURE): a prospective cohort study. *Lancet* 2017; 390:2037–2049.
 111. Campbell NRC, Schutte AE, Varghese CV, Ordunez P, Zhang XH, Khan T, *et al.* Sao Paulo call to action for the prevention and control of high blood pressure: 2020. *J Clin Hypertens (Greenwich)* 2019; 21:1744–1752.
 112. Martin S, Sturgiss E, Douglas K, Ball L. Hidden curriculum within nutrition education in medical schools. *BMJ Nutr Prev Health* 2020; 3:18–23.
 113. Blunt SB, Kafatos A. Clinical Nutrition Education of Doctors and Medical Students: Solving the Catch 22. *Adv Nutr* 2019; 10:345–350.
 114. DiMaria-Ghalili RA, Mirtallo JM, Tobin BW, Hark L, Van Horn L, Palmer CA. Challenges and opportunities for nutrition education and training in the healthcare professions: intraprofessional and interprofessional call to action. *Am J Clin Nutr* 2014; 99:1184S–1193S.
 115. Vijaykumar S, McNeill A, Simpson J. Associations between conflicting nutrition information, nutrition confusion and backlash among consumers in the UK. *Public Health Nutr* 2021; 24:914–923.
 116. Nagler RH. Adverse outcomes associated with media exposure to contradictory nutrition messages. *J Health Commun* 2014; 19:24–40.
 117. Hills S, Terry D, Gazula S, Browning C. Practice nurses' communication with people living with type 2 diabetes: a scoping review. *Patient Educ Couns* 2022; 105:2664–2670.
 118. Sesso HD, Manson JE, Aragaki AK, Rist PM, Johnson LG, Friedenberg G, *et al.*, COSMOS Research Group. Effect of cocoa flavanol supplementation for the prevention of cardiovascular disease events: the COcoa Supplement and Multivitamin Outcomes Study (COSMOS) randomized clinical trial. *Am J Clin Nutr* 2022; 115:1490–1500.
 119. de Oliveira Otto MC, Mozaffarian D, Kromhout D, Bertoni AG, Sibley CT, Jacobs DR Jr, Nettleton JA, *et al.* Dietary intake of saturated fat by food source and incident cardiovascular disease: the Multi-Ethnic Study of Atherosclerosis. *Am J Clin Nutr* 2012; 96:397–404.
 120. Nestel PJ, Beilin LJ, Clifton PM, Watts GF, Mori TA. Practical Guidance for Food Consumption to Prevent Cardiovascular Disease. *Heart Lung Circ* 2021; 30:163–179.
 121. Zhang X, Ritonja JA, Zhou N, Chen BE, Li X. Omega-3 Polyunsaturated Fatty Acids Intake and Blood Pressure: A Dose-Response Meta-Analysis of Randomized Controlled Trials. *J Am Heart Assoc* 2022; 11:e025071.
 122. Bao DQ, Mori TA, Burke V, Puddey IB, Beilin LJ. Effects of dietary fish and weight reduction on ambulatory blood pressure in overweight hypertensives. *Hypertension* 1998; 32:710–717.
 123. Jayedi A, Shab-Bidar S. Fish Consumption and the Risk of Chronic Disease: An Umbrella Review of Meta-Analyses of Prospective Cohort Studies. *Adv Nutr* 2020; 11:1123–1133.
 124. Muralitharan RR, Jama HA, Xie L, Peh A, Snelson M, Marques FZ. Microbial Peer Pressure: The Role of the Gut Microbiota in Hypertension and Its Complications. *Hypertension* 2020; 76:1674–1687.
 125. O'Donnell JA, Zheng T, Meric G, Marques FZ. The gut microbiome and hypertension. *Nat Rev Nephrol* 2023; 19:153–167.
 126. Aljuraiban GS, Griep LM, Chan Q, Daviglus ML, Stamler J, Van Horn L, *et al.* Total, insoluble and soluble dietary fibre intake in relation to blood pressure: the INTERMAP Study. *Br J Nutr* 2015; 114:1480–1486.
 127. Reynolds A, Mann J, Cummings J, Winter N, Mete E, Te Morenga L. Carbohydrate quality and human health: a series of systematic reviews and meta-analyses. *Lancet* 2019; 393:434–445.
 128. Ordoas JM, Ferguson LR, Tai ES, Mathers JC. Personalised nutrition and health. *BMJ* 2018; 361:; bmj.k2173.
 129. Campbell NRC, Whelton PK, Ories M, Wainford RD, Cappuccio FP, Ide N, *et al.* 2022 World Hypertension League, Resolve To Save Lives and International Society of Hypertension dietary sodium (salt) global call to action. *J Hum Hypertens* 2022; 37:428–437.
 130. GBD Diet Collaborators. Health effects of dietary risks in 195 countries, 1990–2017: a systematic analysis for the Global Burden of Disease Study 2017. *Lancet* 2019; 393:1958–1972.
 131. Wenstedt EFE, Olde Engberink RHG, Vogt L. Sodium Handling by the Blood Vessel Wall: Critical for Hypertension Development. *Hypertension* 2018; 71:990–996.
 132. Filippini T, Malavolti M, Whelton PK, Naska A, Orsini N, Vinceti M. Blood Pressure Effects of Sodium Reduction: Dose-Response Meta-Analysis of Experimental Studies. *Circulation* 2021; 143:1542–1567.
 133. Huang L, Trieu K, Yoshimura S, Neal B, Woodward M, Campbell NRC, *et al.* Effect of dose and duration of reduction in dietary sodium on blood pressure levels: systematic review and meta-analysis of randomised trials. *BMJ* 2020; 368:m315.
 134. National Academies of Sciences, Engineering, and Medicine. *Dietary Reference Intakes for sodium and potassium*. Washington, DC: The National Academies Press. 2019.

135. Ma Y, He FJ, Sun Q, Yuan C, Kieneker LM, Curhan GC, et al. 24-Hour Urinary Sodium and Potassium Excretion and Cardiovascular Risk. *N Engl J Med* 2022; 386:252–263.
136. He FJ, MacGregor GA. Reducing population salt intake worldwide: from evidence to implementation. *Prog Cardiovasc Dis* 2010; 52:363–382.
137. Do HT, Santos JA, Trieu K, Petersen K, Le MB, Lai DT, et al. Effectiveness of a Communication for Behavioral Impact (COMBI) Intervention to Reduce Salt Intake in a Vietnamese Province Based on Estimations From Spot Urine Samples. *J Clin Hypertens (Greenwich)* 2016; 18:1135–1142.
138. Land MA, Wu JH, Selwyn A, Crino M, Woodward M, Chalmers J, et al. Effects of a community-based salt reduction program in a regional Australian population. *BMC Public Health* 2016; 16:388.
139. Brown IJ, Tzoulaki I, Candeias V, Elliott P. Salt intakes around the world: implications for public health. *Int J Epidemiol* 2009; 38:791–813.
140. Popkin BM. Global nutrition dynamics: the world is shifting rapidly toward a diet linked with noncommunicable diseases. *Am J Clin Nutr* 2006; 84:289–298.
141. LINKS. Sodium Reduction Framework. 2022.
142. Neal B, Wu Y, Feng X, Zhang R, Zhang Y, Shi J, et al. Effect of Salt Substitution on Cardiovascular Events and Death. *N Engl J Med* 2021; 385:1067–1077.
143. Olde Engberink RHG, van den Hoek TC, van Noordenne ND, van den Born BH, Peters-Sengers H, Vogt L. Use of a Single Baseline Versus Multiyear 24-Hour Urine Collection for Estimation of Long-Term Sodium Intake and Associated Cardiovascular and Renal Risk. *Circulation* 2017; 136:917–926.
144. Jones ESW, Lee HY, Khan N, Charchar FJ, Williams B, Chia YC, et al. Reduction of salt intake: time for more action. *J Hypertens* 2022; 40:2130–2132.
145. Welling PA. WNKs on the Fly. *J Am Soc Nephrol* 2018; 29:1347–1349.
146. Marklund M, Singh G, Greer R, Cudhea F, Matsushita K, Michal R, et al. Estimated population wide benefits and risks in China of lowering sodium through potassium enriched salt substitution: modelling study. *BMJ* 2020; 369:m824.
147. Ruzicka M, Hiremath S, Steiner S, Helis E, Szczotka A, Baker P, Fodor G. What is the feasibility of implementing effective sodium reduction strategies to treat hypertension in primary care settings? A systematic review. *J Hypertens* 2014; 32:1388–1394.
148. Gritter M, Wouda RD, Yeung SMH, Wieers MLA, Geurts F, de Ridder MAJ, et al., on behalf of K consortium. Effects of Short-Term Potassium Chloride Supplementation in Patients with CKD. *J Am Soc Nephrol* 2022; 33:1779–1789.
149. van Mierlo LA, Greyling A, Zock PL, Kok FJ, Geleijnse JM. Suboptimal potassium intake and potential impact on population blood pressure. *Arch Intern Med* 2010; 170:1501–1502.
150. Jeemon P, Severin T, Amodeo C, Balabanova D, Campbell NRC, Gaita D, et al. World Heart Federation Roadmap for Hypertension - A 2021 Update. *Glob Heart* 2021; 16:63.
151. Hannou SA, Haslam DE, McKeown NM, Herman MA. Fructose metabolism and metabolic disease. *J Clin Invest* 2018; 128:545–555.
152. Itoh H, Tanaka M. 'Greedy Organs Hypothesis' for sugar and salt in the pathophysiology of noncommunicable diseases in relation to sodium-glucose co-transporters in the intestines and the kidney. *Metabol Open* 2022; 13:100169.
153. Mortera RR, Bains Y, Gugliucci A. Fructose at the crossroads of the metabolic syndrome and obesity epidemics. *Front Biosci (Landmark Ed)* 2019; 24:186–211.
154. Nakagawa T, Johnson RJ, Andres-Hernando A, Roncal-Jimenez C, Sanchez-Lozada LG, Tolan DR, et al. Fructose Production and Metabolism in the Kidney. *J Am Soc Nephrol* 2020; 31:898–906.
155. Yang Q, Zhang Z, Gregg EW, Flanders WD, Merritt R, Hu FB. Added sugar intake and cardiovascular diseases mortality among US adults. *JAMA Intern Med* 2014; 174:516–524.
156. Ferraris RP, Choe JY, Patel CR. Intestinal Absorption of Fructose. *Annu Rev Nutr* 2018; 38:41–67.
157. Savarese G, Butler J, Lund LH, Bhatt DL, Anker SD. Cardiovascular effects of noninsulin glucose-lowering agents: a comprehensive review of trial evidence and potential cardioprotective mechanisms. *Cardiovasc Res* 2022; 118:2231–2252.
158. Chan JC, Yeung R, Luk A. The Asian diabetes phenotypes: challenges and opportunities. *Diabetes Res Clin Pract* 2014; 105:135–139.
159. Spracklen CN, Horikoshi M, Kim YJ, Lin K, Bragg F, Moon S, et al. Identification of type 2 diabetes loci in 433,540 East Asian individuals. *Nature* 2020; 582:240–245.
160. Acton RB, Vanderlee L, Adams J, Kirkpatrick SI, Pedraza LS, Sacks G, et al. Tax awareness and perceived cost of sugar-sweetened beverages in four countries between 2017 and 2019: findings from the international food policy study. *Int J Behav Nutr Phys Act* 2022; 19:38.
161. von Philipsborn P, Huizinga O, Leibinger A, Rubin D, Burns J, Emmert-Fees K, et al. Interim Evaluation of Germany's Sugar Reduction Strategy for Soft Drinks: Commitments versus Actual Trends in Sugar Content and Sugar Sales from Soft Drinks. *Ann Nutr Metab* 2023; [Epub ahead of print].
162. Salgado Hernandez JC, Ng SW, Colchero MA. Changes in sugar-sweetened beverage purchases across the price distribution after the implementation of a tax in Mexico: a before-and-after analysis. *BMC Public Health* 2023; 23:265.
163. Silver LD, Padon AA, Li L, Simard BJ, Greenfield TK. Changes in sugar-sweetened beverage consumption in the first two years (2018–2020) of San Francisco's tax: a prospective longitudinal study. *PLOS Glob Public Health* 2023; 3:e0001219.
164. Nguyen DT, Hoang MV, Dao S, Do PH, Nguyen QD, Jewell J, et al. Estimating the health impacts of sugar-sweetened beverage tax for informing policy decisions about the obesity burden in Vietnam. *PLoS One* 2023; 18:e0274928.
165. Gill SK, Rossi M, Bajka B, Whelan K. Dietary fibre in gastrointestinal health and disease. *Nat Rev Gastroenterol Hepatol* 2021; 18:101–116.
166. Collaborators GBDD. Health effects of dietary risks in 195 countries, 1990–2017: a systematic analysis for the Global Burden of Disease Study 2017. *Lancet* 2019; 393:1958–1972.
167. Jama HA, Rhys-Jones D, Nakai M, Yao CK, Climie RE, Sata Y, et al. Prebiotic intervention with HAMSAB in untreated essential hypertensive patients assessed in a phase II randomized trial. *Nat Cardiovasc Res* 2023; 2:35–43.
168. Wilson AS, Koller KR, Ramaboli MC, Nesengani LT, Ocvirk S, Chen C, et al. Diet and the Human Gut Microbiome: An International Review. *Dig Dis Sci* 2020; 65:723–740.
169. Stephen AM, Champ MM, Cloran SJ, Fleith M, van Lieshout L, Mejbörn H, Burley VJ. Dietary fibre in Europe: current state of knowledge on definitions, sources, recommendations, intakes and relationships to health. *Nutr Res Rev* 2017; 30:149–190.
170. Fayet-Moore F, Cassettari T, Tuck K, McConnell A, Petocz P. Dietary Fibre Intake in Australia. Paper II: Comparative Examination of Food Sources of Fibre among High and Low Fibre Consumers. *Nutrients* 2018; 10:1223.
171. Rampelli S, Schnorr SL, Consolandi C, Turroni S, Severgnini M, Peano C, et al. Metagenome sequencing of the Hadza hunter-gatherer gut microbiota. *Curr Biol* 2015; 25:1682–1693.
172. Magee E. 6 Foods and Tips for More Fiber.. In: Chang L, editor. *WebMD*. 2010.
173. Li F, Hullar MA, Schwarz Y, Lampe JW. Human gut bacterial communities are altered by addition of cruciferous vegetables to a controlled fruit- and vegetable-free diet. *J Nutr* 2009; 139:1685–1691.
174. McKeown NM, Fahey GC Jr, Slavin J, van der Kamp JW. Fibre intake for optimal health: how can healthcare professionals support people to reach dietary recommendations? *BMJ* 2022; 378:e054370.
175. Belghith-Fendri L, Chaari F, Kallel F, Zouari-Ellouzi S, Ghorbel R, Besbes S, et al. Pea and Broad Bean Pods as a Natural Source of Dietary Fiber: The Impact on Texture and Sensory Properties of Cake. *J Food Sci* 2016; 81:C2360–C2366.
176. Lian C. L'alcoolisme, cause d'hypertension arterielle. *Bull Acad Natl Med Paris* 1915; 7:525–528.
177. Roerecke M, Tobe SW, Kaczorowski J, Bacon SL, Vafaei A, Hasan OSM, et al. Sex-Specific Associations Between Alcohol Consumption and Incidence of Hypertension: A Systematic Review and Meta-Analysis of Cohort Studies. *J Am Heart Assoc* 2018; 7:e008202.
178. Roerecke M, Kaczorowski J, Tobe SW, Gmel G, Hasan OSM, Rehm J. The effect of a reduction in alcohol consumption on blood pressure: a systematic review and meta-analysis. *Lancet Public Health* 2017; 2:e108–e120.
179. Seppa K, Sillanaukee P. Binge drinking and ambulatory blood pressure. *Hypertension* 1999; 33:79–82.
180. Piano MR, Burke L, Kang M, Phillips SA. Effects of Repeated Binge Drinking on Blood Pressure Levels and Other Cardiovascular Health

- Metrics in Young Adults: National Health and Nutrition Examination Survey, 2011–2014. *J Am Heart Assoc* 2018; 7:e008733.
181. Biddinger KJ, Emdin CA, Haas ME, Wang M, Hindy G, Ellinor PT, *et al.* Association of Habitual Alcohol Intake With Risk of Cardiovascular Disease. *JAMA Netw Open* 2022; 5:e223849.
 182. Holmes MV, Dale CE, Zuccolo L, Silverwood RJ, Guo Y, Ye Z, *et al.*, InterAct Consortium. Association between alcohol and cardiovascular disease: Mendelian randomisation analysis based on individual participant data. *BMJ* 2014; 349:g4164.
 183. Global Health Workforce Statistics WHO. *Global status report on alcohol and health* 2018. Geneva: WHO; 2018.
 184. Ritchie H, Roser M. Alcohol Consumption. 2018. Available at: <https://ourworldindata.org/alcohol-consumption>. [Accessed 11 March 2023]
 185. Arora M, ElSayed A, Beger B, Naidoo P, Shilton T, Jain N, *et al.* The Impact of Alcohol Consumption on Cardiovascular Health: Myths and Measures. *Glob Heart* 2022; 17:45.
 186. Staton CA, Vissoci JRN, El-Gabri D, Adewumi K, Concepcion T, Elliott SA, *et al.* Patient-level interventions to reduce alcohol-related harms in low- and middle-income countries: a systematic review and meta-summary. *PLoS Med* 2022; 19:e1003961.
 187. Platt L, Melendez-Torres GJ, O'Donnell A, Bradley J, Newbury-Birch D, Kaner E, Ashton C. How effective are brief interventions in reducing alcohol consumption: do the setting, practitioner group and content matter? Findings from a systematic review and meta-regression analysis. *BMJ Open* 2016; 6:e011473.
 188. Terentes-Prinzios D, Vlachopoulos C. Coffee and cardiovascular health: looking through the steaming cup. *Cardiovasc Res* 2022; 118:e51–e53.
 189. Poole R, Kennedy OJ, Roderick P, Fallowfield JA, Hayes PC, Parkes J. Coffee consumption and health: umbrella review of meta-analyses of multiple health outcomes. *BMJ* 2017; 359:j5024.
 190. Zhang Y, Yang H, Li S, Li WD, Wang Y. Consumption of coffee and tea and risk of developing stroke, dementia, and poststroke dementia: a cohort study in the UK Biobank. *PLoS Med* 2021; 18:e1003830.
 191. Xie C, Cui L, Zhu J, Wang K, Sun N, Sun C. Coffee consumption and risk of hypertension: a systematic review and dose-response meta-analysis of cohort studies. *J Hum Hypertens* 2018; 32:83–93.
 192. Teramoto M, Yamagishi K, Muraki I, Takamashi A, Iso H. Coffee and Green Tea Consumption and Cardiovascular Disease Mortality Among People With and Without Hypertension. *J Am Heart Assoc* 2023; 12:e026477.
 193. Larsson SC, Wolk A, Hakansson N, Back M. Coffee consumption and risk of aortic valve stenosis: a prospective study. *Nutr Metab Cardiovasc Dis* 2018; 28:803–807.
 194. Benjamim CJR, Porto AA, Valenti VE, Sobrinho A, Garner DM, Gualano B, Bueno Júnior CR. Nitrate Derived From Beetroot Juice Lowers Blood Pressure in Patients With Arterial Hypertension: A Systematic Review and Meta-Analysis. *Front Nutr* 2022; 9:823039.
 195. O'Gallagher K, Borg Cardona S, Hill C, Al-Saedi A, Shahed F, Floyd CN, *et al.* Grapefruit juice enhances the systolic blood pressure-lowering effects of dietary nitrate-containing beetroot juice. *Br J Clin Pharmacol* 2021; 87:577–587.
 196. Cicero AFG, Grassi D, Tocci G, Galletti F, Borghi C, Ferri C. Nutrients and Nutraceuticals for the Management of High Normal Blood Pressure: An Evidence-Based Consensus Document. *High Blood Press Cardiovasc Prev* 2019; 26:9–25.
 197. Basrai M, Schweinlin A, Menzel J, Mielke H, Weikert C, Dusemund B, *et al.* Energy Drinks Induce Acute Cardiovascular and Metabolic Changes Pointing to Potential Risks for Young Adults: A Randomized Controlled Trial. *J Nutr* 2019; 149:441–450.
 198. Allcock E, Cowdery J. Hypertension induced by liquorice tea. *BMJ Case Rep* 2015; 2015:E581–E583.
 199. Cosentino F, Grant PJ, Aboyans V, Bailey CJ, Ceriello A, Delgado V, *et al.* 2019 ESC Guidelines on diabetes, prediabetes, and cardiovascular diseases developed in collaboration with the EASD. *Eur Heart J* 2020; 41:255–323.
 200. Mancia Chairperson G, Kreutz Co-Chair R, Brunstrom M, Burnier M, Grassi G, Januszewicz A, *et al.*, Authors/Task Force Members. 2023 ESH Guidelines for the management of arterial hypertension The Task Force for the management of arterial hypertension of the European Society of Hypertension Endorsed by the European Renal Association (ERA) and the International Society of Hypertension (ISH). *J Hypertens* 2023; [Epub ahead of print].
 201. Berg C, Lappas G, Wolk A, Strandhagen E, Toren K, Rosengren A, *et al.* Eating patterns and portion size associated with obesity in a Swedish population. *Appetite* 2009; 52:21–26.
 202. Davis R, Rogers M, Coates AM, Leung GKW, Bonham MP. The Impact of Meal Timing on Risk of Weight Gain and Development of Obesity: a Review of the Current Evidence and Opportunities for Dietary Intervention. *Curr Diab Rep* 2022; 22:147–155.
 203. Mager DE, Wan R, Brown M, Cheng A, Wareski P, Abernethy DR, Mattson MP. Caloric restriction and intermittent fasting alter spectral measures of heart rate and blood pressure variability in rats. *FASEB J* 2006; 20:631–637.
 204. St-Onge MP, Ard J, Baskin ML, Chiuve SE, Johnson HM, Kris-Etherton P, Varady K, American Heart Association Obesity Committee of the Council on Lifestyle and Cardiometabolic Health; Council on Cardiovascular Disease in the Young; Council on Clinical Cardiology; and Stroke Council. Meal Timing and Frequency: Implications for Cardiovascular Disease Prevention: A Scientific Statement From the American Heart Association. *Circulation* 2017; 135:e96–e121.
 205. Adafer R, Messaadi W, Meddahi M, Patey A, Haderbache A, Bayen S, *et al.* Food Timing, Circadian Rhythm and Chrononutrition: A Systematic Review of Time-Restricted Eating's Effects on Human Health. *Nutrients* 2020;12.
 206. Liu D, Huang Y, Huang C, Yang S, Wei X, Zhang P, *et al.* Calorie Restriction with or without Time-Restricted Eating in Weight Loss. *N Engl J Med* 2022; 386:1495–1504.
 207. Hammoud S, Kurdi M, van den Bemt BJF. Impact of Fasting on Cardiovascular Outcomes in Patients With Hypertension. *J Cardiovasc Pharmacol* 2021; 78:481–495.
 208. Baynouna AlKetbi L, Nagelkerke N, AlZarouni A, Al Kuwaiti M, Al Ghafl M, Al Qahtani S, *et al.* Ramadan fasting outcome among high-risk patients. *BMC Nephrol* 2022; 23:304.
 209. AlAbdan NA, Almohammed OA, Altukhaimeh MS, Farooqui MA, Abdalla MI, Al Otaibi HQ, *et al.* Fasting during Ramadan and acute kidney injury (AKI): a retrospective, propensity matched cohort study. *BMC Nephrol* 2022; 23:54.
 210. Albus C, Waller C, Fritzsche K, Gunold H, Haass M, Hamann B, *et al.* Significance of psychosocial factors in cardiology: update 2018: Position paper of the German Cardiac Society. *Clin Res Cardiol* 2019; 108:1175–1196.
 211. Liu MY, Li N, Li WA, Khan H. Association between psychosocial stress and hypertension: a systematic review and meta-analysis. *Neurol Res* 2017; 39:573–580.
 212. Liu M-Y, Li N, Li WA, Khan H. Association between psychosocial stress and hypertension: a systematic review and meta-analysis. *Neurol Res* 2017; 39:573–580.
 213. Howard JT, Sosnov JA, Janak JC, Gundlapalli AV, Pettey WB, Walker LE, Stewart JJ. Associations of Initial Injury Severity and Posttraumatic Stress Disorder Diagnoses With Long-Term Hypertension Risk After Combat Injury. *Hypertension* 2018; 71:824–832.
 214. Persu A, Petit G, Georges C, de Timary P. Hypertension, a Posttraumatic Stress Disorder? Time to Widen Our Perspective. *Hypertension* 2018; 71:811–812.
 215. Celik M, Yilmaz Y, Karagoz A, Kahyaoglu M, Cakmak EO, Kup A, *et al.* Anxiety Disorder Associated with the COVID-19 Pandemic Causes Deterioration of Blood Pressure Control in Primary Hypertensive Patients. *Medeni Med J* 2021; 36:83–90.
 216. Rubio-Guerra AF, Rodriguez-Lopez L, Vargas-Ayala G, Huerta-Ramirez S, Serna DC, Lozano-Nuevo JJ. Depression increases the risk for uncontrolled hypertension. *Exp Clin Cardiol* 2013; 18:10–12.
 217. Task Force for the management of C-otESoC: Baigent C, Windecker S, Andreini D, Arbelo E, Barbato E, *et al.* European Society of Cardiology guidance for the diagnosis and management of cardiovascular disease during the COVID-19 pandemic: part 1-epidemiology, pathophysiology, and diagnosis. *Cardiovasc Res* 2022; 118:1385–1412.
 218. Task Force for the management of C-otESoC. ESC guidance for the diagnosis and management of cardiovascular disease during the COVID-19 pandemic: part 2-care pathways, treatment, and follow-up. *Cardiovasc Res* 2022; 118:1618–1666.
 219. Niazi AK, Niazi SK. Mindfulness-based stress reduction: a nonpharmacological approach for chronic illnesses. *N Am J Med Sci* 2011; 3:20–23.
 220. Kabat-Zinn J. *Wherever you go, there you are: mindfulness meditation in everyday life*. New York: Hachette Books; 2009.

221. Nalbant G, Hassanein ZM, Lewis S, Chattopadhyay K. Content, Structure, and Delivery Characteristics of Yoga Interventions for Managing Hypertension: A Systematic Review and Meta-Analysis of Randomized Controlled Trials. *Front Public Health* 2022; 10:846231.
222. Li Y, Buys N, Li Z, Li L, Song Q, Sun J. The efficacy of cognitive behavioral therapy-based interventions on patients with hypertension: a systematic review and meta-analysis. *Prev Med Rep* 2021; 23:101477.
223. Perciavalle V, Blandini M, Fecarotta P, Buscemi A, Di Corrado D, Bertolo L, et al. The role of deep breathing on stress. *Neurol Sci* 2017; 38:451–458.
224. Kuhlmann AY, Etnel JR, Roos-Hesselink JW, Jeekel J, Bogers AJ, Takkenberg JJ. Systematic review and meta-analysis of music interventions in hypertension treatment: a quest for answers. *BMC Cardiovasc Disord* 2016; 16:69.
225. do Amaral MA, Neto MG, de Queiroz JG, Martins-Filho PR, Saquetto MB, Oliveira Carvalho V. Effect of music therapy on blood pressure of individuals with hypertension: A systematic review and Meta-analysis. *Int J Cardiol* 2016; 214:461–464.
226. Nejati S, Zahiroddin A, Afrookhteh G, Rahmani S, Hoveida S. Effect of Group Mindfulness-Based Stress-Reduction Program and Conscious Yoga on Lifestyle, Coping Strategies, and Systolic and Diastolic Blood Pressures in Patients with Hypertension. *J Tebran Heart Cent* 2015; 10:140–148.
227. Ponte Marquez PH, Feliu-Soler A, Sole-Villa MJ, Matas-Pericas L, Filella-Agullo D, Ruiz-Herrerias M, et al. Benefits of mindfulness meditation in reducing blood pressure and stress in patients with arterial hypertension. *J Hum Hypertens* 2019; 33:237–247.
228. Hirshkowitz M, Whiton K, Albert SM, Alessi C, Bruni O, DonCarlos L, et al. National Sleep Foundation's sleep time duration recommendations: methodology and results summary. *Sleep Health* 2015; 1:40–43.
229. Ohayon M, Wickwire EM, Hirshkowitz M, Albert SM, Avidan A, Daly FJ, et al. National Sleep Foundation's sleep quality recommendations: first report. *Sleep Health* 2017; 3:6–19.
230. Johnson KA, Gordon CJ, Chapman JL, Hoyos CM, Marshall NS, Miller CB, Grunstein RR. The association of insomnia disorder characterised by objective short sleep duration with hypertension, diabetes and body mass index: a systematic review and meta-analysis. *Sleep Med Rev* 2021; 59:101456.
231. Manohar S, Thongprayoon C, Cheungpasitporn W, Mao MA, Herrmann SM. Associations of rotational shift work and night shift status with hypertension: a systematic review and meta-analysis. *J Hypertens* 2017; 35:1929–1937.
232. Ai S, Zhang J, Zhao G, Wang N, Li G, So HC, et al. Causal associations of short and long sleep durations with 12 cardiovascular diseases: linear and nonlinear Mendelian randomization analyses in UK Biobank. *Eur Heart J* 2021; 42:3349–3357.
233. Knutson KL, Van Cauter E, Rathouz PJ, Yan LL, Hulley SB, Liu K, Lauderdale DS. Association between sleep and blood pressure in midlife: the CARDIA sleep study. *Arch Intern Med* 2009; 169:1055–1061.
234. Li ZH, Huang QM, Gao X, Chung VCH, Zhang PD, Shen D, et al. Healthy Sleep Associated With Lower Risk of Hypertension Regardless of Genetic Risk: A Population-Based Cohort Study. *Front Cardiovasc Med* 2021; 8:769130.
235. He L, Ma T, Li J, Luo Y, Zhang G, Cheng X, Bai Y. Adherence to a healthy sleep pattern and incidence of cardiometabolic multimorbidity among hypertensive patients: a prospective study of UK Biobank. *Sleep* 2022; 45:zsac141.
236. Dashti HS, Daghlasi I, Lane JM, Huang Y, Udder MS, Wang H, et al. Genetic determinants of daytime napping and effects on cardiometabolic health. *Nat Commun* 2021; 12:900.
237. Yang MJ, Zhang Z, Wang YJ, Li JC, Guo QL, Chen X, Wang E. Association of nap frequency with hypertension or ischemic stroke supported by prospective cohort data and mendelian randomization in predominantly middle-aged European subjects. *Hypertension* 2022; 79:1962–1970.
238. Bursztyjn M, Mekler J, Wachtel N, Ben-Ishay D. Siesta and ambulatory blood pressure monitoring. Comparability of the afternoon nap and night sleep. *Am J Hypertens* 1994; 7:217–221.
239. Huang M, Yang Y, Huang Z, Yuan H, Lu Y. The association of nighttime sleep duration and daytime napping duration with hypertension in Chinese rural areas: a population-based study. *J Hum Hypertens* 2021; 35:896–902.
240. Vgontzas AN, Liao D, Bixler EO, Chrousos GP, Vela-Bueno A. Insomnia with objective short sleep duration is associated with a high risk for hypertension. *Sleep* 2009; 32:491–497.
241. Thomas SJ, Booth JN 3rd, Jaeger BC, Hubbard D, Sakhuja S, Abdalla M, et al. Association of Sleep Characteristics With Nocturnal Hypertension and Nondipping Blood Pressure in the CARDIA Study. *J Am Heart Assoc* 2020; 9:e015062.
242. Fritz J, Phillips AJK, Hunt LC, Imam A, Reid KJ, Perreira KM, et al. Cross-sectional and prospective associations between sleep regularity and metabolic health in the Hispanic Community Health Study/Study of Latinos. *Sleep* 2021; 44:zsaa218.
243. Chen J, Patel SR, Redline S, Durazo-Arvizu R, Garside DB, Reid KJ, et al. Weekly sleep trajectories and their associations with obesity and hypertension in the Hispanic/Latino population. *Sleep* 2018; 41:zsy150.
244. Garbarino S, Lanteri P, Bragazzi NL, Magnavita N, Scoditti E. Role of sleep deprivation in immune-related disease risk and outcomes. *Commun Biol* 2021; 4:1304.
245. Randerath W, Bassetti CL, Bonsignore MR, Farre R, Ferini-Strambi L, Grote L, et al. Challenges and perspectives in obstructive sleep apnoea: report by an ad hoc working group of the Sleep Disordered Breathing Group of the European Respiratory Society and the European Sleep Research Society. *Eur Respir J* 2018; 52:1702616.
246. Carneiro-Barrera A, Diaz-Roman A, Guillen-Riquelme A, Buella-Casal G. Weight loss and lifestyle interventions for obstructive sleep apnoea in adults: systematic review and meta-analysis. *Obes Rev* 2019; 20:750–762.
247. Stelmach-Mardas M, Brajer-Luftmann B, Kusnierczak M, Batura-Gabryel H, Piorunek T, Mardas M. Body Mass Index Reduction and Selected Cardiometabolic Risk Factors in Obstructive Sleep Apnea: Meta-Analysis. *J Clin Med* 2021; 10:1485.
248. White LH, Bradley TD, Logan AG. Pathogenesis of obstructive sleep apnoea in hypertensive patients: role of fluid retention and nocturnal rostral fluid shift. *J Hum Hypertens* 2015; 29:342–350.
249. Nieto FJ, Young TB, Lind BK, Shahar E, Samet JM, Redline S, et al. Association of sleep-disordered breathing, sleep apnea, and hypertension in a large community-based study. Sleep Heart Health Study. *JAMA* 2000; 283:1829–1836.
250. Pengo MF, Soranna D, Giontella A, Perger E, Mattaliano P, Schwarz EI, et al. Obstructive sleep apnoea treatment and blood pressure: which phenotypes predict a response? A systematic review and meta-analysis. *Eur Respir J* 2020; 55:1901945.
251. Pedrosa RP, Drager LF, Gonzaga CC, Sousa MG, de Paula LK, Amaro AC, et al. Obstructive sleep apnea: the most common secondary cause of hypertension associated with resistant hypertension. *Hypertension* 2011; 58:811–817.
252. Cattazzo F, Pengo MF, Giontella A, Soranna D, Bilo G, Zambon A, et al. Effect of Continuous Positive Airway Pressure on Glucose and Lipid Profiles in Patients With Obstructive Sleep Apnoea: A Systematic Review and Meta-Analysis of Randomized Controlled Trials. *Arch Bronconeumol* 2023; 59:370–376.
253. Svedmyr S, Hedner J, Bonsignore MR, Lombardi C, Parati G, Ludka O, et al., European Sleep Apnea Database (ESADA) study group (full collaborator list in E-supplement). Hypertension treatment in patients with sleep apnea from the European Sleep Apnea Database (ESADA) cohort - towards precision medicine. *J Sleep Res* 2022; 32:e13811.
254. Kim J, Yi H, Shin KR, Kim JH, Jung KH, Shin C. Snoring as an independent risk factor for hypertension in the nonobese population: the Korean Health and Genome Study. *Am J Hypertens* 2007; 20:819–824.
255. Niu Y, Sui X, He Y, Xi H, Zhu R, Xu H, et al. Association between self-reported snoring and hypertension: a systematic review and meta-analysis. *Sleep Med* 2021; 88:140–148.
256. Han B, Chen WZ, Li YC, Chen J, Zeng ZQ. Sleep and hypertension. *Sleep Breath* 2020; 24:351–356.
257. Li X, Sotres-Alvarez D, Gallo LC, Ramos AR, Aviles-Santa L, Perreira KM, et al. Associations of Sleep-disordered Breathing and Insomnia with Incident Hypertension and Diabetes. The Hispanic Community Health Study/Study of Latinos. *Am J Respir Crit Care Med* 2021; 203:356–365.
258. Maiolino G, Bisogni V, Soranna D, Pengo MF, Pucci G, Vettor R, et al., Sleep Disorders Working Group of the Italian Society of Hypertension. Effects of insomnia and restless legs syndrome on sleep arterial

- blood pressure: a systematic review and meta-analysis. *Sleep Med Rev* 2021; 59:101497.
259. Mohammadi S, Moosaie F, Saghaizadeh A, Mahmoudi M, Rezaei N. Metabolic profile in patients with narcolepsy: a systematic review and meta-analysis. *Sleep Med* 2021; 81:268–284.
 260. Cheungpasitporn W, Thongprayoon C, Srivali N, Vijayvargiya P, Andersen CA, Kittanamongkolchai W, *et al.* The effects of napping on the risk of hypertension: a systematic review and meta-analysis. *J Evid Based Med* 2016; 9:205–212.
 261. Kingsbury JH, Buxton OM, Emmons KM. Sleep and its Relationship to Racial and Ethnic Disparities in Cardiovascular Disease. *Curr Cardiovasc Risk Rep* 2013; 7: 10.1007/s12170-013-0330-0.
 262. Johnson DA, Jackson CL, Williams NJ, Alcantara C. Are sleep patterns influenced by race/ethnicity - a marker of relative advantage or disadvantage? Evidence to date. *Nat Sci Sleep* 2019; 11:79–95.
 263. Dudley KA, Patel SR. Disparities and genetic risk factors in obstructive sleep apnea. *Sleep Med* 2016; 18:96–102.
 264. Alkhazna A, Bhat A, Ladesich J, Barthel B, Bohnam AJ. Severity of obstructive sleep apnea between black and white patients. *Hosp Pract (1995)* 2011; 39:82–86.
 265. Pandey A, Williams N, Donat M, Ceide M, Brimah P, Ogedegbe G, *et al.* Linking sleep to hypertension: greater risk for blacks. *Int J Hypertens* 2013; 2013:436502.
 266. Lieu SJ, Curhan GC, Schernhammer ES, Forman JP. Rotating night shift work and disparate hypertension risk in African-Americans. *J Hypertens* 2012; 30:61–66.
 267. van Oort S, Beulens JWJ, van Ballegooijen AJ, Grobbee DE, Larsson SC. Association of Cardiovascular Risk Factors and Lifestyle Behaviors With Hypertension: A Mendelian Randomization Study. *Hypertension* 2020; 76:1971–1979.
 268. Watson NF, Badr MS, Belenky G, Bliwise DL, Buxton OM, Buysse D, *et al.* Recommended Amount of Sleep for a Healthy Adult: A Joint Consensus Statement of the American Academy of Sleep Medicine and Sleep Research Society. *Sleep* 2015; 38:843–844.
 269. Consensus Conference P, Watson NF, Badr MS, Belenky G, Bliwise DL, Buxton OM, *et al.* Joint Consensus Statement of the American Academy of Sleep Medicine and Sleep Research Society on the Recommended Amount of Sleep for a Healthy Adult: Methodology and Discussion. *J Clin Sleep Med* 2015; 11:931–952.
 270. Buysse DJ. Sleep health: can we define it? Does it matter? *Sleep* 2014; 37:9–17.
 271. Zhang DY, Huang JF, Kang YY, Dou Y, Su YL, Zhang LJ, *et al.* The prevalence of masked hypertension in relation to cigarette smoking in a Chinese male population. *J Hypertens* 2020; 38:1056–1063.
 272. Rahman MM, Laher I. Structural and functional alteration of blood vessels caused by cigarette smoking: an overview of molecular mechanisms. *Curr Vasc Pharmacol* 2007; 5:276–292.
 273. Howard G, Wagenknecht LE, Burke GL, Diez-Roux A, Evans GW, McGovern P, *et al.* Cigarette smoking and progression of atherosclerosis: the Atherosclerosis Risk in Communities (ARIC) Study. *JAMA* 1998; 279:119–124.
 274. Rigotti NA, Kruse GR, Livingstone-Banks J, Hartmann-Boyce J. Treatment of Tobacco Smoking: A Review. *JAMA* 2022; 327:566–577.
 275. Martinez-Morata I, Sanchez TR, Shimbo D, Navas-Acien A. Electronic Cigarette Use and Blood Pressure Endpoints: a Systematic Review. *Curr Hypertens Rep* 2020; 23:2.
 276. Collaborators GBD. Global burden of 87 risk factors in 204 countries and territories, 1990–2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet* 2020; 396:1223–1249.
 277. Lim SS, Vos T, Flaxman AD, Danaei G, Shibuya K, Adair-Rohani H, *et al.* A comparative risk assessment of burden of disease and injury attributable to 67 risk factors and risk factor clusters in 21 regions, 1990–2010: a systematic analysis for the Global Burden of Disease Study 2010. *Lancet* 2012; 380:2224–2260.
 278. Collaborators GBD. Spatial, temporal, and demographic patterns in prevalence of smoking tobacco use and attributable disease burden in 204 countries and territories, 1990–2019: a systematic analysis from the Global Burden of Disease Study 2019. *Lancet* 2021; 397:2337–2360.
 279. Utap MS, Tan C, Su AT. Effectiveness of a brief intervention for smoking cessation using the 5A model with self-help materials and using self-help materials alone: a randomised controlled trial. *Malays Fam Physician* 2019; 14:2–9.
 280. Lelieveld J, Evans JS, Fnais M, Giannadaki D, Pozzer A. The contribution of outdoor air pollution sources to premature mortality on a global scale. *Nature* 2015; 525:367–371.
 281. Hamanaka RB, Mutlu GM. Particulate Matter Air Pollution: Effects on the Cardiovascular System. *Front Endocrinol (Lausanne)* 2018; 9:680.
 282. Brook RD, Franklin B, Cascio W, Hong Y, Howard G, Lipsett M, *et al.* Air Pollution and Cardiovascular Disease: A Statement for Healthcare Professionals from the Expert Panel on Population and Prevention Science of the American Heart Association. *Circulation* 2004; 109:2655–2671.
 283. WHO global air quality guidelines. Particulate matter (PM_{2.5} and PM₁₀), ozone, nitrogen dioxide, sulfur dioxide and carbon monoxide. Geneva: World Health Organization; 2021. Licence: CC BY-NC-SA 3.0 IGO.
 284. HEI. Understanding the health effects of ambient ultrafine particles. HEI Review Panel on Ultrafine Particles. Boston (MA) Health Effects Institute (HEI Perspectives 3. 2013. Available at: <https://www.health-effects.org/system/files/Perspectives3.pdf>. [Accessed 29 April 2022]
 285. Newby DE, Mannucci PM, Tell GS, Baccarelli AA, Brook RD, Donaldson K, *et al.*, ESC Working Group on Thrombosis, European Association for Cardiovascular Prevention and Rehabilitation, ESC Heart Failure Association. Expert position paper on air pollution and cardiovascular disease. *Eur Heart J* 2015; 36:83–93b.
 286. Nemmar A, Hoet PH, Vanquickenborne B, Dinsdale D, Thomeer M, Hoylaerts MF, *et al.* Passage of inhaled particles into the blood circulation in humans. *Circulation* 2002; 105:411–414.
 287. Sanidas E, Papadopoulos DP, Grassos H, Velliou M, Tsioufis K, Barbetseas J, Papademetriou V. Air pollution and arterial hypertension. A new risk factor is in the air. *J Am Soc Hypertens* 2017; 11:709–715.
 288. Ohlwein S, Kappeler R, Kutlar Joss M, Kunzli N, Hoffmann B. Health effects of ultrafine particles: a systematic literature review update of epidemiological evidence. *Int J Public Health* 2019; 64:547–559.
 289. Brook RD, Brook JR, Urch B, Vincent R, Rajagopalan S, Silverman F. Inhalation of fine particulate air pollution and ozone causes acute arterial vasoconstriction in healthy adults. *Circulation* 2002; 105:1534–1536.
 290. Zanobetti A, Canner MJ, Stone PH, Schwartz J, Sher D, Eagan-Bengston E, *et al.* Ambient Pollution and Blood Pressure in Cardiac Rehabilitation Patients. *Circulation* 2004; 110:2184–2189.
 291. Chen H, Burnett RT, Kwong JC, Villeneuve PJ, Goldberg MS, Brook RD, *et al.* Spatial association between ambient fine particulate matter and incident hypertension. *Circulation* 2014; 129:562–569.
 292. Adar SD, Sheppard L, Vedral S, Polak JF, Sampson PD, Diez Roux AV, *et al.* Fine particulate air pollution and the progression of carotid intima-medial thickness: a prospective cohort study from the multi-ethnic study of atherosclerosis and air pollution. *PLoS Med* 2013; 10:e1001430.
 293. Hee VCV, Adar SD, Szpiro AA, Barr RG, Bluemke DA, Roux AVD, *et al.* Exposure to traffic and left ventricular mass and function. *Am J Respir Crit Care Med* 2009; 179:827–834.
 294. Yang B-Y, Qian Z, Howard SW, Vaughn MG, Fan S-J, Liu K-K, Dong GH. Global association between ambient air pollution and blood pressure: a systematic review and meta-analysis. *Environ Pollut* 2018; 235:576–588.
 295. Qin P, Luo X, Zeng Y, Zhang Y, Li Y, Wu Y, *et al.* Long-term association of ambient air pollution and hypertension in adults and in children: a systematic review and meta-analysis. *Sci Total Environ* 2021; 796:148620.
 296. Ibaldo-Mulli A, Stieber J, Wichmann HE, Koenig W, Peters A. Effects of air pollution on blood pressure: a population-based approach. *Am J Public Health* 2001; 91:571–577.
 297. Hudda N, Eliasziw M, Hersey SO, Reisner E, Brook RD, Zamore W, *et al.* Effect of Reducing Ambient Traffic-Related Air Pollution on Blood Pressure. *Hypertension* 2021; 77:823–832.
 298. Pascal M, Corso M, Chancel O, Declercq C, Badaloni C, Cesaroni G, *et al.*, Aphekom group. Assessing the public health impacts of urban air pollution in 25 European cities: results of the Aphekom project. *Sci Total Environ* 2013; 449:390–400.
 299. Munzel T, Hahad O, Sorensen M, Lelieveld J, Duerr GD, Nieuwenhuijsen M, Daiber A. Environmental risk factors and cardiovascular diseases: a comprehensive expert review. *Cardiovasc Res* 2022; 118:2880–2902.

300. Fu W, Wang C, Zou L, Liu Q, Gan Y, Yan S, et al. Association between exposure to noise and risk of hypertension: a meta-analysis of observational epidemiological studies. *J Hypertens* 2017; 35:2358–2366.
301. Kawada T. Noise exposure and hypertension. *J Hypertens* 2018; 36:2478.
302. World Health Organization Sixty-Eighth World Health Assembly 68.18 Health and the Environment: Addressing the Health Impact of Air Pollution. 2015.
303. Personal interventions and risk communication on air pollution. Geneva: World Health Organization; 2020. Licence: CC BY-NC-SA 3.0 IGO.
304. Johan de Hartog J, Boogaard H, Nijland H, Hoek G. Do the health benefits of cycling outweigh the risks? *Environ Health Perspect* 2010; 118:1109–1116.
305. Sharman JE. Clinicians prescribing exercise: is air pollution a hazard? *Med J Aust* 2005; 182:606–607.
306. Mark SD, Wang W, Fraumeni JF Jr, Li JY, Taylor PR, Wang GQ, et al. Lowered risks of hypertension and cerebrovascular disease after vitamin/mineral supplementation: the Linxian Nutrition Intervention Trial. *Am J Epidemiol* 1996; 143:658–664.
307. Wang C, Li Y, Zhu K, Dong YM, Sun CH. Effects of supplementation with multivitamin and mineral on blood pressure and C-reactive protein in obese Chinese women with increased cardiovascular disease risk. *Asia Pac J Clin Nutr* 2009; 18:121–130.
308. Rautiainen S, Wang L, Lee IM, Manson JE, Gaziano JM, Buring JE, Sesso HD. Multivitamin use and the risk of hypertension in a prospective cohort study of women. *J Hypertens* 2016; 34:1513–1519.
309. Chen J, He J, Hamm L, Batuman V, Whelton PK. Serum antioxidant vitamins and blood pressure in the United States population. *Hypertension* 2002; 40:810–816.
310. Zhang Y, Liu M, Zhou C, Zhang Z, He P, Li Q, et al. Inverse association between dietary vitamin A intake and new-onset hypertension. *Clin Nutr* 2021; 40:2868–2875.
311. Ran L, Zhao W, Tan X, Wang H, Mizuno K, Takagi K, et al. Association between Serum Vitamin C and the blood pressure: a systematic review and meta-analysis of observational studies. *Cardiovasc Ther* 2020; 2020:4940673.
312. Juraschek SP, Guallar E, Appel LJ, Miller ER. Effects of vitamin C supplementation on blood pressure: a meta-analysis of randomized controlled trials. *Am J Clin Nutr* 2012; 95:1079–1088.
313. Poss J, Mahfoud F, Ukena C, Esler MD, Schlaich M, Hering D, et al. Association of vitamin D status and blood pressure response after renal denervation. *Clin Res Cardiol* 2014; 103:41–47.
314. Qasemi R, Ghavamzadeh S, Faghfour AH, Valizadeh N, Mohammadi A, Sayyadi H. The effect of vitamin D supplementation on flow-mediated dilatation, oxidized LDL and intracellular adhesion molecule 1 on type 2 diabetic patients with hypertension: a randomized, placebo-controlled, double-blind trial. *Diabetes Metab Syndr* 2021; 15:102200.
315. Pantovic A, Zec M, Zekovic M, Obrenovic R, Stankovic S, Glibetic M. Vitamin D is inversely related to obesity: cross-sectional study in a small cohort of serbian adults. *J Am Coll Nutr* 2019; 38:405–414.
316. Zhang X, Li Y, Del Gobbo LC, Rosanoff A, Wang J, Zhang W, et al. Effects of magnesium supplementation on blood pressure: a meta-analysis of randomized double-blind placebo-controlled trials. *Hypertension* 2016; 68:324–333.
317. Filippini T, Violi F, D'Amico R, Vinceti M. The effect of potassium supplementation on blood pressure in hypertensive subjects: a systematic review and meta-analysis. *Int J Cardiol* 2017; 230:127–135.
318. Li L, Su C, Chen X, Wang Q, Jiao W, Luo H, et al. Chlorogenic acids in cardiovascular disease: a review of dietary consumption, pharmacology, and pharmacokinetics. *J Agric Food Chem* 2020; 68:6464–6484.
319. Sosnowska B, Penson P, Banach M. The role of nutraceuticals in the prevention of cardiovascular disease. *Cardiovasc Diagn Ther* 2017; 7 (Suppl 1):S21–S31.
320. Neupane D, McLachlan CS, Mishra SR, Olsen MH, Perry HB, Karki A, Kallestrup P. Effectiveness of a lifestyle intervention led by female community health volunteers versus usual care in blood pressure reduction (COBIN): an open-label, cluster-randomised trial. *Lancet Glob Health* 2018; 6:e66–e73.
321. Jafar TH, Gandhi M, de Silva HA, Jehan I, Naheed A, Finkelstein EA, et al., COBRA-BPS Study Group. A Community-based intervention for managing hypertension in rural South Asia. *N Engl J Med* 2020; 382:717–726.
322. Reynolds R, Dennis S, Hasan I, Slewa J, Chen W, Tian D, et al. A systematic review of chronic disease management interventions in primary care. *BMC Fam Pract* 2018; 19:11.
323. Wagner EH, Austin BT, Davis C, Hindmarsh M, Schaefer J, Bonomi A. Improving chronic illness care: translating evidence into action. *Health Aff (Millwood)* 2001; 20:64–78.
324. Munoz Aguilera E, Suvan J, Buti J, Czesnikiewicz-Guzik M, Barbosa Ribeiro A, Orlandi M, et al. Periodontitis is associated with hypertension: a systematic review and meta-analysis. *Cardiovasc Res* 2020; 116:28–39.
325. Del Pinto R, Monaco A, Ortu E, Czesnikiewicz-Guzik M, Munoz Aguilera E, Giannoni M, et al. Access to dental care and blood pressure profiles in adults with high socioeconomic status. *J Periodontol* 2022; 93:1060–1071.
326. Baderol Allam FN, Ab Hamid MR, Buhari SS, Md Noor H. Web-based dietary and physical activity intervention programs for patients with hypertension: scoping review. *J Med Internet Res* 2021; 23:e22465.
327. Yatabe J, Yatabe MS, Ichihara A. The current state and future of internet technology-based hypertension management in Japan. *Hypertens Res* 2021; 44:276–285.
328. Kassavou A, Wang M, Mirzaei V, Shpendi S, Hasan R. The association between smartphone app-based self-monitoring of hypertension-related behaviors and reductions in high blood pressure: systematic review and meta-analysis. *JMIR Mhealth Uhealth* 2022; 10:e34767.
329. Stogios N, Kaur B, Huszti E, Vasanthan J, Nolan RP. Advancing digital health interventions as a clinically applied science for blood pressure reduction: a systematic review and meta-analysis. *Can J Cardiol* 2020; 36:764–774.
330. Tong HL, Quiroz JC, Kocaballi AB, Fat SCM, Dao KP, Gehringer H, et al. Personalized mobile technologies for lifestyle behavior change: a systematic review, meta-analysis, and meta-regression. *Prev Med* 2021; 148:106532.
331. Nahum-Shani I, Smith SN, Spring BJ, Collins LM, Witkiewitz K, Tewari A, et al. Just-in-Time Adaptive Interventions (JITIs) in Mobile Health: Key Components and Design Principles for Ongoing Health Behavior Support. *Ann Behav Med* 2018; 52:446–462.
332. Kang HS, Exworthy M. Wearing the Future-Wearables to Empower Users to Take Greater Responsibility for Their Health and Care: Scoping Review. *JMIR Mhealth Uhealth* 2022; 10:e35684.
333. Wongvibulsin S, Martin SS, Saria S, Zeger SL, Murphy SA. An Individualized, Data-Driven Digital Approach for Precision Behavior Change. *Am J Lifestyle Med* 2020; 14:289–293.
334. Sun W, Li Y, Hu Y, Rao X, Xu X, Browning CJ, et al. Perspectives on the Training of Chinese Primary Healthcare Physicians to Reduce Chronic Illnesses and Their Burden. *Front Public Health* 2019; 7:168.
335. Rao X, Lai J, Wu H, Li Y, Xu X, Browning CJ, et al. The Development of a Competency Assessment Standard for General Practitioners in China. *Front Public Health* 2020; 8:23.
336. Dragomir AI, Julien CA, Bacon SL, Boucher VG, Lavoie KL, Canadian Network for Health Behavior Change and Promotion (CAN-Change). Training physicians in behavioural change counseling: a systematic review. *Patient Educ Couns* 2019; 102:12–24.
337. Ren Y, Yang H, Browning C, Thomas S, Liu M. Therapeutic effects of motivational interviewing on blood pressure control: a meta-analysis of randomized controlled trials. *Int J Cardiol* 2014; 172:509–511.
338. Awad A, Trenfield SJ, Pollard TD, Ong JJ, Elbadawi M, McCoubrey LE, et al. Connected healthcare: improving patient care using digital health technologies. *Adv Drug Deliv Rev* 2021; 178:113958.
339. Schutte AE, Jafar TH, Poulter NR, Damasceno A, Khan NA, Nilsson PM, et al. Addressing global disparities in blood pressure control: perspectives of the International Society of Hypertension. *Cardiovasc Res* 2023; 119:381–409.
340. Sani RN, Connelly PJ, Toft M, Rowa-Dewar N, Delles C, Gasevic D, Karaye KM. Rural-urban difference in the prevalence of hypertension in West Africa: a systematic review and meta-analysis. *J Hum Hypertens* 2022.
341. Chaves G, Brites N, Munzinger J, Uhlmann L, Gonzalez G, Oviedo G, et al. Education to a Healthy Lifestyle Improves Symptoms and Cardiovascular Risk Factors - AsuRiesgo Study. *Arq Bras Cardiol* 2015; 104:347–355.
342. Dastan I, Erem A, Cetinkaya V. Urban and rural differences in hypertension risk factors in Turkey. *Anatol J Cardiol* 2017; 18:39–47.

343. Ventegodt S, Kandel I, Ervin D, Merrick J. Concepts of holistic care. In: Rubin IL, Merrick J, Greydanus DE, Patel DR, editors. *Health care for people with intellectual and developmental disabilities across the lifespan*. Springer: Cham; 2016. pp. 1935–1941.
344. Lopez-Jaramillo P, Joseph P, Lopez-Lopez JP, Lanas F, Avezum A, Diaz R, *et al*. Risk factors, cardiovascular disease, and mortality in South America: a PURE substudy. *Eur Heart J* 2022; 43:2841–2851.
345. Torlasco C, Faini A, Makil E, Bilo G, Pengo M, Beaney T, *et al*. Nationwide hypertension screening in Italy: data from May Measurements Month 2017–Europe. *Eur Heart J Suppl* 2019; 21 (Suppl d):D66–D70.
346. Salazar MR, Garcia Vazquez F, Espeche WG, Marquez D, Becerra P, Martinez Marissi E, *et al*. May Measurement Month 2019: an analysis of blood pressure screening results from Argentina. *Eur Heart J Suppl* 2021; 23:B12–B14.
347. Wang Z, Knight S, Wilson A, Rowley KG, Best JD, McDermott R, *et al*. Blood pressure and hypertension for Australian Aboriginal and Torres Strait Islander people. *Eur J Cardiovasc Prev Rehabil* 2006; 13:438–443.
348. Davy C, Harfield S, McArthur A, Munn Z, Brown A. Access to primary healthcare services for Indigenous peoples: a framework synthesis. *Int J Equity Health* 2016; 15:163.
349. Okpechi IG, Hariramani VK, Sultana N, Ghimire A, Zaidi D, Muneer S, *et al*. The impact of community-based nonpharmacological interventions on cardiovascular and kidney disease outcomes in remote dwelling Indigenous communities: a scoping review protocol. *PLoS One* 2022; 17:e0269839.
350. Rawal S, Johnson BR, Young HN, Gaye B, Sattler ELP. Association of Life's Simple 7 and ideal cardiovascular health in American Indians/Alaska Natives. *Open Heart* 2023; 10:e002222.
351. Marmot M. The health gap: the challenge of an unequal world. *Lancet* 2015; 386:2442–2444.
352. DiPette DJ, Goughnour K, Zuniga E, Skeete J, Ridley E, Angell S, *et al*. Standardized treatment to improve hypertension control in primary healthcare: the HEARTS in the Americas Initiative. *J Clin Hypertens (Greenwich)* 2020; 22:2285–2295.
353. Husain MJ, Allaire BT, Hutchinson B, Ketgudee L, Srisuthisak S, Yueyay K, *et al*. Assessing costs of a hypertension management program: an application of the HEARTS costing tool in a program planning workshop in Thailand. *J Clin Hypertens (Greenwich)* 2020; 22:111–117.
354. Boudreaux C, Barango P, Adler A, Kabore P, McLaughlin A, Mohamed MOS, *et al*. Addressing severe chronic NCDs across Africa: measuring demand for the Package of Essential Noncommunicable Disease Interventions-Plus (PEN-Plus). *Health Policy Plan* 2022; 37:452–460.
355. Australian Government Department of Health and Aged Care. Exercise and physical activity for adults (18 to 64 years). 2021.
356. Sharman JE, Smart NA, Coombes JS, Stowasser M. Exercise and sport science australia position stand update on exercise and hypertension. *J Hum Hypertens* 2019; 33:837–843.
357. Pescatello L. What's new in the ACSM pronouncement on exercise and hypertension? *Am Coll Sports Med* 2019; Available at: [https://www.acsm.org/blog-detail/acsm-blog/2019/06/11/new-acsm-pronouncement-exercise-hypertension#:~:text=\(13\)%20that%20shows%20resistance%20exercise,hypertension%20\(3%2D5\)](https://www.acsm.org/blog-detail/acsm-blog/2019/06/11/new-acsm-pronouncement-exercise-hypertension#:~:text=(13)%20that%20shows%20resistance%20exercise,hypertension%20(3%2D5).). [Accessed 27 March 2023].
358. Mozaffarian D. Dietary and policy priorities for cardiovascular disease, diabetes, and obesity: a comprehensive review. *Circulation* 2016; 133:187–225.
359. Lichtenstein AH, Appel LJ, Vadiveloo M, Hu FB, Kris-Etherton PM, Rebholz CM, *et al*. 2021 dietary guidance to improve cardiovascular health: a scientific statement from the American Heart Association. *Circulation* 2021; 144:e472–e487.
360. Wang W, Liu Y, Li Y, Luo B, Lin Z, Chen K, *et al*. Dietary patterns and cardiometabolic health: clinical evidence and mechanism. *MedComm* 20202023; 4:e212.
361. World Health Organisation. Draft recommendation for the prevention and management of obesity over the life course, including potential targets. 2021.
362. Leeson P. Chronic Hypertension in Pregnancy Project and the Control of Hypertension in Pregnancy Study: impact of blood pressure control in pregnancy on maternal and fetal outcomes. *Cardiovasc Res* 2022; 118:e98–e100.
363. World Health Organisation. WHO child growth standards: length/height-for-age, weight-for-age, weight-for-length, weight -for-height and body mass index-for-age: methods and development.: World Health Organisation; 2006.
364. de Onis M, Onyango AW, Borghi E, Siyam A, Nishida C, Siekmann J. Development of a WHO growth reference for school-aged children and adolescents. *Bull World Health Organ* 2007; 85:660–667.
365. World Health Organization. Guideline: Protecting, Promoting and Supporting Breastfeeding in Facilities Providing Maternity and Newborn Services. Guideline: Protecting, Promoting and Supporting Breastfeeding in Facilities Providing Maternity and Newborn Services. Geneva 2017.
366. World Health Organization Regional Office for Europe. Digital food environments: factsheet. 2021.
367. Raine KD, Atkey K, Olstad DL, Ferdinands AR, Beaulieu D, Buhler S, *et al*. Healthy food procurement and nutrition standards in public facilities: evidence synthesis and consensus policy recommendations. *Health Promot Chronic Dis Prev Can* 2018; 38:6–17.
368. World Health Organization. Tackling NCDs: 'best buys' and other recommended interventions for the prevention and control of non-communicable diseases. 2017.
369. Khera AV, Emdin CA, Drake I, Natarajan P, Bick AG, Cook NR, *et al*. Genetic risk, adherence to a healthy lifestyle, and coronary disease. *N Engl J Med* 2016; 375:2349–2358.
370. Sun L, Pennells L, Kaptoge S, Nelson CP, Ritchie SC, Abraham G, *et al*. Polygenic risk scores in cardiovascular risk prediction: a cohort study and modelling analyses. *PLoS Med* 2021; 18:e1003498.
371. Vaura F, Kauko A, Suvila K, Havulinna AS, Mars N, Salomaa V, *et al*. Polygenic risk scores predict hypertension onset and cardiovascular risk. *Hypertension* 2021; 77:1119–1127.
372. Ye Y, Chen X, Han J, Jiang W, Natarajan P, Zhao H. Interactions between enhanced polygenic risk scores and lifestyle for cardiovascular disease, diabetes, and lipid levels. *Circ Genom Precis Med* 2021; 14:e003128.
373. Rutten-Jacobs LC, Larsson SC, Malik R, Rannikmae K, consortium M, International Stroke Genetics C, *et al*. Genetic risk, incident stroke, and the benefits of adhering to a healthy lifestyle: cohort study of 306 473 UK Biobank participants. *BMJ* 2018; 363:k4168.
374. Chiuve SE, Cook NR, Shay CM, Rexrode KM, Albert CM, Manson JE, *et al*. Lifestyle-based prediction model for the prevention of CVD: the Healthy Heart Score. *J Am Heart Assoc* 2014; 3:e000954.