

1
2
3
4 Preprint - Preprints and early-stage research may not have been peer reviewed yet.
5

6 **Risk factors for hypertension in Vietnam**

7 **Truong Dinh Cam^{1*}, Do Thanh Huyen², Ho Thi Le¹, Pham Toan Trung¹, Ta Anh Hoang¹**

8 ¹Cardiology - Rheumatology - Endocrinology Department, Military Hospital 175, Ho Chi Minh City,
9 Vietnam

10 ²General internal medicine Department, Military Hospital 121, Can Tho City, Vietnam

11
12 ***Corresponding Author:** truongcam1967@gmail.com
13

14 **Abstract.** Hypertension (or high blood pressure) affects many adults (mostly people over 65) in
15 Vietnam. Quantifying risk factors associated with hypertension may help to inform prevention
16 efforts, such as hypertension awareness, treatment, and control in low- income countries. A cross-
17 sectional study was conducted in a study area Vietnam. The study was conducted at Military
18 Hospital 121 consisting of 166 patients, who were randomly selected. Engaging in less than 150
19 minutes per week of moderate-to- vigorous physical activity, eating fruits and vegetables fewer than
20 five times per day, being overweight or obese, having diabetes, and having chronic kidney disease
21 were all independently associated with an increased risk of hypertension. When these factors were
22 combined into a risk score, there was a linear increase in the predicted risk of hypertension with each
23 additional risk factor. The predicted prevalence of hypertension for those with such risk factors was
24 75% in women and 25% in men with mean aged 60 years. Being overweight or obese, consuming
25 fruits and vegetables less often, being inactive, and having diabetes contributed to the largest attributable
26 fractions for hypertension in the Vietnam population. Physical activity, diet, body mass index, the
27 presence of diabetes, and the presence of chronic kidney disease were strong risk factors for
28 hypertension. Many of these risk factors are modifiable and highlight targets for future prevention
29 strategies.
30

31 **Key words:** hypertension risk, prevalence, survey, cross-sectional, lifestyle, exercise, diet, obesity,
32 diabetes, chronic kidney disease, cholesterol

INTRODUCTION

The global epidemic of hypertension (HPN) is largely uncontrolled and hypertension remains the leading cause of non-communicable disease deaths worldwide (Burnier and Egan, 2019; Ramli et al., 2012). High blood pressure (BP) is estimated to account for 6% of deaths worldwide and is the most common treatable risk factor for cardiovascular disease (Wolf-Maier et al., 2004). High BP, also known as “silent killer,” is associated with approximately 9.4 million cases of death a year worldwide (Lim et al., 2012). Traditionally, raised blood pressure has been estimated based on the definition of systolic blood pressure ≥ 140 mmHg and/or diastolic ≥ 90 mmHg. With this definition, the global prevalence of raised blood pressure in adults aged 18 years and above was around 22% in 2014 (World Health Organization, 2014).

The increase in prevalence of raised BP is associated with population ageing and the rise in occurrence of behavioral risk factors, such as tobacco use, unhealthy diet, harmful use of alcohol, physical inactivity, overweight, and persistent stress (Margolis et al., 2019; Trudel et al., 2019; Chu et al., 2018, 2019). Also, there are six psychosocial factors that may induce HNT in a person including occupational stress, personality, sleep quality, mental health, housing instability, and social support/ isolation (Lelong et al., 2019; Cuffee et al., 2014). Concurrently, high BP as a physiological risk factor contributes to a high burden of other non-communicable diseases, for example, strokes and chronic kidney diseases and ocular illness (Naghavi et al., 2017; Gordon et al., 2002; Johnson et al., 2002). The majority of deaths related to BP mainly occurs in middle-aged and elderly people. There is an estimated 80% of the population aged 60 years or over will be affected by HNT (World Health Organization, 2013). However, the elderly can improve their quality of life through good blood pressure control. HNT is known as a result from the aging and atherosclerosis of arterial walls, when the artery wall decreases elasticity, becomes sclerotic with accumulation of lipids in the blood, arteriosclerosis and vascular stenosis will be formatted (Rubio-Ruiz et al., 2014; Sun, 2015). Regarding to the age, the risk of HNT in the lifetime in males is higher than that in females (Van et al., 2019).

During the last 30 years, hypertension treatment has improved dramatically, contributing to a decrease in the incidence of mortality due to stroke and coronary heart disease. However, accumulating evidence suggests that healthy lifestyle factors (such as a healthy diet and increased physical activity) contribute to lowering blood pressure and that managing these risks can offset, at least to some extent, genetic predisposition towards hypertension and the development of subsequent cardiovascular sequelae (Leung et al., 2019). A crucial step in this process is therefore the explicit recommendations given to medical professionals and the practical consequences of these treatment strategies in particular health systems.

However, control rates of hypertension is currently inappropriate and the majority of the hypertensive patients will require two or more antihypertensive agents to reach target BP goals. Antihypertensive medicines can effectively reduce blood pressure and the risk of associated diseases (Ettehad et al., 2016; Olsen et al., 2016). As clinical trials have shown the benefits of pharmacological treatment for patients with low to moderate blood pressure, clinical guidelines have evolved to

81 recommend lower blood pressure thresholds for initiating treatment (Zhou et al.,
82 2019). Beta-blockers have been used for more than 40 years to treat hypertension.
83 Data from clinical trials that used these agents to manage BP have demonstrated
84 reductions in cardiovascular mortality and this has resulted in recommendations of
85 β -blockers as first- or second-line antihypertensive agents (Stafylas and Sarafidis,
86 2008).

87 Despite these gains, poorly controlled hypertension remains a health
88 problem of major proportions, particularly among minority populations, the poor,
89 those with lower levels of education, and those with limited access to medical care.
90 Many people with high blood pressure in low- and middle-income countries are not
91 aware of their hypertensive status, and lack access to treatment, and management
92 services. Vietnam has been undergoing health transition where populations are
93 suffering from a “double burden” of communicable diseases and non-
94 communicable ones, including raised blood pressure (Khan and Khoi, 2008). There
95 were a few surveys aiming to assess the level of raised blood pressure in Vietnam
96 but they were only covering a small number of provinces, with different sampling
97 methods and using different age ranges (Hoang et al., 2019). Vietnam, a lower
98 middle-income country, has undergone an epidemiological transition from
99 communicable diseases to non-communicable diseases. The prevalence of
100 hypertension in adults aged 25–64 years increased from 15% in 2002 to 18.9% in
101 2015, and is predicted to rise in the coming decades. A 2008 national survey
102 conducted in eight Vietnamese provinces and cities reported that only 48% of
103 hypertensive individuals knew their disease status, 30% were on treatment
104 programs, and 11% had their hypertension under control (Quoc Cuong et al., 2019;
105 Nguyen et al., 2020). Vietnam established its national HTN program in 2008. Since
106 then, studies on the prevalence of HTN, including few national samples, have been
107 published and varied in their estimates. In 2012, a time trends analysis for blood
108 pressure was performed using a dataset compiled by multiple cross-sectional
109 studies; the analysis showed a significant increase of HTN prevalence of 0.9% per
110 year between 2001 and 2009 (Meiqari, 2019).

111 Unlike most medical conditions, community surveillance has been the most
112 common approach to evaluating the success of efforts to treat and control high BP.
113 During the past decade, many countries have conducted large-scale, national health
114 surveys to determine the prevalence and treatment of hypertension. This
115 information might provide insights into ways to improve public health strategies to
116 prevent target organ damage. The aim of this study was to provide up-to-date
117 assessment of the evidence on the magnitude of HTN in Vietnam. Consolidating
118 available data in the literature into a national estimate of the prevalence of HTN is
119 crucial to provide information for future policy regarding population need,
120 management, and control of HTN in Vietnam.

121 **MATERIAL AND METHODS**

122 **Study Design**

123
124
125
126 The study was conducted at Military Hospital 121. Consisting of 166
127 patients who came to the Clinic of Cardiovascular – Endocrinology (A1), they
128 were randomly selected and divided into two groups including: a) primary

129 hypertension group (136 patients); b) control group without hypertension (30
130 patients). Research period: February 2017 to January 2018.

131 Selection criteria:

132 • Control group. Being healthy, without hypertension, have similarities in
133 age, gender distribution, and anthropometric indicators with the hypertension group.
134 Fasting blood glucose was tested at the biochemistry department <7.0mmol
135 / L. Agree to join the research.

136 • Hypertension group. Patients with primary hypertension who were
137 screened at random were eligible for the study. Diagnosed with primary
138 hypertension according to the standard of JNC VII with systolic blood pressure
139 >140 mmHg and/or diastolic blood pressure >90 mmHg as measured by mercury
140 sphygmomanometer, or patients under control blood pressure is equal to one of
141 the common antihypertensive drugs, with a fasting blood glucose index tested at
142 the biochemistry department <7.0 mmol / L. Agree to join the research.
143

144 146 Exclusion criteria for both disease and control groups:

145 2047. • Diabetes has been diagnosed and being treated according to ADA
148

149 • Endocrine diseases that affect blood glucose (Cushing, Basedow
150 maximal limb etc.).

151 • Fever, severe water and electrolyte disorders.

152 • Acute bacterial infections.

153 • Acute liver disease or severe kidney failure.

154 • Patients who are taking drugs that affect glucose metabolism: steroids,
155 catecholamines, beta blockers, insulin, birth control pill etc.

156 • Severe comorbidities: stroke, coma, liver failure, heart failure, severe
157 kidney failure.

158 • Secondary hypertension group (renal artery stenosis, renal
159 parenchyma disease, adrenal medullary tumor etc.).

160 • Pregnant.

161 • Patients who did not cooperate in the study.
162

163 Sample size

164 166 The sample size was calculated following the formula:

167

168

169

170

171

172

$$n = \frac{Z_{(1-\alpha/2)}^2 * p(1-p)}{d^2} \quad (1)$$

173

174

175

177 where p is insulin resistance rate in hypertensive patients estimated by Huynh Van
 178 Minh's research in 1996 with; $p = 41.7\%$. So $n = 130$ patients. The minimum
 179 number of patients required in the study was 130, in fact we studied over 136
 180 patients.

181

182 RESEARCH DESIGN

183

184 The prospective study, cross-sectional description had a comparative control
 185 between the research group and the control group.

186

187 RESULTS

188

189 Among participants, there were 66.2% women with HPN and almost the
 190 same amount in control group (CG) (Table 1). The mean age was from 57 to 60
 191 years. All of participants belonged to the group with risk habits, such as smoking,
 192 drinking alcohol, eating salty foods and sweets without the hard physical exercises.
 193 As HPN as well as CG almost do not have any comorbidities, such as gestational
 194 diabetes or bad history of HPN. The average SBP and DBP for HPN group was
 195 150/82 mmHg, where in CG – 110/64 mmHg. Average BMI, waist and WHR
 196 parameters and central obesity perimeter were slightly higher among subjects of
 197 the HPN group.

198

199 Baseline biochemical characteristics of participants are presented in Table 2.
 200 The mean age \pm SD of hypertensive patients and normotensives were 47.6 ± 4.2 and
 201 38.4 ± 3.7 years, respectively. Levels of fasting blood sugar, HbA_{1C} and fasting
 202 plasma insulin were 5.74 ± 4.3 mmol/L, 5.71 ± 0.52 % and 86.01 ± 60.08 pmol/L,
 203 respectively, in hypertensive subjects while in normotensive subjects, they were
 204 5.21 ± 0.61 mmol/L, 5.55 ± 0.55 % and 51.19 ± 23.63 pmol/L respectively, which
 were significantly higher in hypertensive patients.

205

Table 1. General characteristics of the study respondents

	Hypertension group (n=136)	Control group (n = 30)	p
Sex: male, n (%)	46 (33.8)	13 (43.3)	0.325
female, n (%)	90 (66.2)	17 (56.7)	
Age: max	84	82	0.28
min	24	25	
mean	60.33 ± 12.33	57.56 ± 14.61	
Daily habits:			
smoking, n (%)	35 (25.7)	10 (33.3)	0.397
drinking alcohol, n (%)	30 (22.1)	6 (20)	0.804
eating salty food, n (%)	47 (34.6)	4 (13.3)	0.039
eating sweets, n (%)	48 (35.3)	4 (13.3)	0.019
Physical exercise:			
light, n (%)	19 (14)	4 (13.3)	< 0.001
moderate, n (%)	27 (19.9)	15 (50)	
hard, n (%)	33 (24.3)	9 (30)	
none, n (%)	57 (41.9)	2 (6.7)	
Family health history of diabetes:			

yes, n (%)	24 (17.6)	5 (16.7)	0.898
no, n (%)	112 (82.4)	25 (83.3)	
Family health history of hypertension:			
yes, n (%)	50 (36.8)	11 (36.7)	0.992
no, n (%)	86 (63.2)	19 (63.3)	
Gestational diabetes:			
yes, n (%)	0	0	n.d.
no, n (%)	136 (100)	30 (100)	
Birth weight > 4 kg:			
yes, n (%)	2 (1.5)	0	0.504
no, n (%)	134 (98.5)	30 (100)	
Time detection of hypertension:			
< 5 years, n (%)	59 (43.4)		
5-10 years, n (%)	37 (27.2)		
>10 years, n (%)	40 (29.4)		
Systolic blood pressure SBP (mmHg)	150.03 ± 27.15	110.33 ± 11.29	<0.001
Diastolic blood pressure DBP (mmHg)	82.82 ± 18.27	64.40 ± 12.70	<0.001
Average (body mass index) BMI	23.49 ± 3.86	22.52 ± 3.19	0.201
Classification BMI:			
under weight, n (%)	10 (7.4)	2 (6.7)	
normal range, n (%)	56 (41.2)	17 (56.7)	0.535
over weight, n (%)	28 (20.6)	6 (20)	
obesity, class I, (n%)	35 (25.7)	4 (13.3)	0.107
obesity, class II, (n%)	7 (5.1)	1 (3.3)	0.266
Waist circumference (cm)	85.82 ± 11.60	82.66 ± 9,04	0.158
Buttocks circumference	95.27 ± 9.51	93.60 ± 6.00	
waist-hip ratio (WHR)	0.90 ± 0.86	0.88 ± 0.061	
Central obesity:			
yes, n (%)	94 (69.1)	18 (60)	0.335
no, n (%)	42 (30.9)	12 (40)	

206

207

208 The cholesterol, triglyceride and serum HDL was significantly higher in
 209 hypertensive patients (5.55 ± 1.27 , 2.94 ± 2.17 and 2.43 ± 0.87 mmol/L
 210 respectively) than in normotensive subjects (4.56 ± 0.71 , 1.62 ± 0.93 and
 211 1.36 ± 0.33 mmol/L).

211

212

213

Table 2. Baseline biochemical characteristics of patients investigated in the current study

	Hypertension group (n=136)	Control group (n = 30)	p
Characteristics of blood sugar level, fasting plasma insulin, HbA1C, and HOMA indicators			
Fasting blood sugar level (mmol/L)	5.74 ± 4.3	5.21 ± 0.61	0.505
HbA1C (%)	5.71 ± 0.52	5.55 ± 0.55	0.138
Fasting plasma insulin (pmol/L)	86.01 ± 60.08	51.19 ± 23.63	<0.001
	1.61 ± 1.09	0.98 ± 0.41	<0.001
	111.56 ± 51.45	86.68 ± 41.63	0.014

HOMA2-IR	86.30 ± 46.62	119.42 ± 54.74	0.001
HOMA2-%B			
HOMA2-%S			
Blood lipid characteristics			
Cholesterol (mmol/L)	5.55 ± 1.27	4.56 ± 0.71	< 0.001
High cholesterol, n (%)	84(61.8)	8 (26.7)	< 0.001
Triglyceride (mmol/L)	2.94 ± 2.17	1.62 ± 0.93	< 0.001
High triglyceride, n (%)	88(64.7)	9 (30)	< 0.001
LDL-C (mmol/L)	3.83 ± 1.42	2.64 ± 0.59	< 0.001
High LDL-C, n (%)	86(63.2)	3 (10)	< 0.001
HDL-C (mmol/L)	2.43 ± 0.87	1.36 ± 0.33	0.592
Low HDL-C, n (%)	28(20.6)	5 (16.7)	0.626
Metabolic lipid disorders, n (%)	125 (91.9)	15 (50)	< 0.001
The rate of insulin resistance and beta cell function			
Insulin resistance:			
No, n (%)	64(47.1)	22 (73.3)	<0.01
Yes, n (%)	72 (52.9)	8 (26.7)	
Beta cell function:			
diminished, n (%)	5(3.7)	4 (13.3)	0.033
normal, n (%)	97(71.3)	23 (76.7)	
enhance, n (%)	34 (25)	3 (10)	
Characteristics of beta cell function in hypertension group			
	Insulin resistance (n=72)	Not insulin resistance (n=64)	
Reduced, n (%)	0 (0)	5 (7.8)	< 0.001
Normal, n (%)	41 (56.9)	56 (87.5)	< 0.001
Enhance, n (%)	31 (43.1)	3 (4.7)	< 0.001

214

215

216

217

218

219

220

221

222

223

224

225

Table 3 presents the prevalence of hypertension by some sociodemographic factors and physical parameters. Data showed that the prevalence of hypertension increased significantly with overweight and obesity (59.7 % vs 40.3 % and 76.4 % vs 23.6 %, respectively). Female population had higher prevalence of hypertension compared to male population (75 % vs. 25 %). The prevalence of hypertension also varied as a function of three NCD risk factors, smoking, high BMI, and comorbidities. Specifically, among population with BMI at normal range, the prevalence of hypertension was 40.3 % while among population with BMI with over weight, this figure was 59.7 %. About 25 % people currently smoking, 22 % people currently drinking, 34-35 % those who eating salty foods and sweets were reported to have hypertension.

226

227

Table 3. Prevalence of hypertension by sociodemographic factors and NCD risk factors

	Insulin resistance			Reduced beta cell function		
	Yes (n=72)	No (n=64)	p	Yes (n=5)	No (n= 131)	p
Age	59.90 ± 10.5	60.82 ± 13.6	0.66	71.4 ± 10.2	59.91 ± 12.2	0.04

		1	4	1	4	1
physical activity:	42 (58.3)	37 (57.8)	0.95	3 (60)	76 (58)	0.93
yes, n (%)	30 (41.7)	27 (42.2)	1	2 (40)	55 (42)	
no, n (%)						
Sex:						
male, n(%)	18 (25)	28 (43.8)	0.02	3 (60)	43 (32.82)	0.20
female, n(%)	54 (75)	36 (56.3)	1	2 (40)	88 (67.72)	7
Family health history of diabetes:			0.09	0	24 (18.32)	0.29
yes, n (%)	9 (12.5)	15 (23.4)	5	5 (100)	107 (81.68)	
no, n (%)	63 (87.5)	49 (76.6)				
Family health history of hypertension:			0.85	1 (20)	49 (37.40)	0.42
yes, n (%)	27 (37.5)	23 (35.9)	0	4 (80)	82 (62.60)	
no, n (%)	45 (62.5)	41 (64.1)				
Gestational diabetes:	0	0	-	0	0	-
yes, n (%)	72 (100)	64 (100)		5 (100)	131 (100)	
no, n (%)						
Birth weight >4 kg:	0	0	-	0	2 (1.52)	0.78
yes, n (%)	72 (100)	64 (100)		5 (100)	129 (98.48)	
no, n (%)						
Smoking:			0.07			0.07
yes, n (%)	21(32.8)	14 (19.4)	5	3 (60)	32 (24.43)	
no, n (%)	43 (67.2)	58 (80.6)		2 (40)	99 (75.57)	
SBP, (mmHg)	148.88 ± 28.90	151.25 ± 25.20	0.60	144 ± 23.02	150.26 ± 27.34	0.61
DBP, (mmHg)	82.55 ± 20.58	83.12 ± 15.41	0.85	80 ± 12.24	82.93 ± 18.49	0.72
Waist circumference (cm)	88.29 ± 11.50	83.05 ± 11.16	0.00	76.80 ± 13.41	85.81 ± 11.64	0.07
BMI	24.52 ± 4.17	22.34 ± 3.15	0.29	22.09 ± 4.94	23.55 ± 3.84	0.41
WHR	0.91 ± 0.08	0.89 ± 0.08	0.12	0.87 ± 0.74	0.90 ± 0.09	0.37
BMI classification:	29 (40.3)	37 (58.8)		3 (60)	64(48.9)	
	43 (59.7)	27 (42.2)		2 (40)	67(51.1)	

normal range, n(%) over weight, n (%)			0.04 1			0.62 5
Central obesity: yes, n (%) no, n (%)	55 (76.4) 17 (23.6)	34 (53.1) 30 (46.8)	0.04 4	3 (60) 2 (40)	91 (69.6) 40 (30.4)	0.64 4

228

229

230

231

232

233

234

235

236

237

238

239

All studied patients in HPN group have high levels of investigated biochemical characteristics at the condition of insulin resistance and high beta-cell function (Table 4). For instance, the level of fasting blood sugar increasing on 17% at insulin resistance, compared to the absence of it. Simultaneously, such characteristics as triglyceride, high cholesterol, high triglyceride and metabolic lipid disorders also higher on 13-18 %. All of these characteristics also significantly raising at reduced beta-cell function condition. For example, a high triglyceride level increased to 20%; were low HDL-C parameter – 60%.

Table 4. Relationship between insulin resistance and beta cell function and biochemical characteristics in hypertension group

	Insulin resistance			Reduced beta cell function		
	Yes (n=72)	No (n=64)	p	Yes (n=5)	No (n= 131)	p
Fasting blood sugar level (mmol/L)	6.24 ± 5.85	5.17 ± 0.61	0.15	6.1 ± 0.69	5.73 ± 4.38	0.850
HbA ₁ C (%)	5.83 ± 0.57	5.58 ± 0.43	0.005	5.46 ± 0.53	5.73 ± 0.52	0.267
Cholesterol (mmol/L)	5.56 ± 1.35	5.53 ± 1.17	0.889	5.61 ± 1.00	5.55 ± 1.28	0.821
Triglyceride (mmol/L)	3.04 ± 1.29	2.82 ± 1.03	0.573	4.06 ± 2.69	2.89 ± 1.13	0.242
LDL-C (mmol/L)	3.97 ± 1.56	3.67 ± 1.24	0.221	3.96 ± 1.42	3.83 ± 1.43	0.840
HDL-C (mmol/L)	1.46 ± 0.51	3.52 ± 1.83	0.277	1.27 ± 0.45	2.45 ± 0.07	0.808
High cholesterol, n (%)	46 (63.9)	38 (59.4)	0.589	3 (60)	81 (61.8)	0.934
High triglyceride, n (%)	47 (65.3)	41 (64.1)	0.882	4 (80)	84 (64.1)	0.466
High LDL-C, n (%)	47 (65.3)	39 (60.9)	0.6	3 (60)	83 (63.3)	0.878
Low HDL-C, n (%)	17 (23.6)	11 (17.2)	0.355	4 (80)	11 (20.6)	0.971

Metabolic lipiddisorders, n (%)	68 (99.4)	57 (89.1)	0.251	68 (99.4)	121 (92.4)	0.32
---------------------------------	-----------	-----------	-------	-----------	------------	------

240

241 Patients (n = 18) with increased LVMI have higher HOMA2-%B score
 242 compared to those (n = 118) LVMI is not increased (127.75 ± 53.26 vs $108.76 \pm$
 243 51.38) (Table 5). However, all patients in the group without increase LVMI have
 244 lower values of HOMA2-IR>1.22, HOMA2-IR<1.22, HOMA2-%B<45.05 and
 245 HOMA2-%B>45.05. However, retinopathy and hypertension complications almost
 246 do not correlate with biochemical characteristics and stay practically the same in
 247 all cases.

248

249 DISCUSSION

250

251 Among a lot of risk factors affected the HPN such as tobacco use, unhealthy
 252 diet, harmful use of alcohol, low physical activity, overweight, and persistent stress
 253 – are the prevalent. These findings are consistent with, and extend, those of other
 254 reports, relating healthy lifestyle and behaviors to better BP control. Participation
 255 in moderate-intensity physical activity, consumption of a diet rich in fruits and
 256 vegetables, and maintenance of a healthy body weight have all been described as
 257 lowering BP. In addition, counselling interventions promoting healthy behaviors
 258 are effective in reducing BP in individuals who do not have hypertension or known
 259 cardiovascular risk factors. However, determining the independent effects of the
 260 individual components of a healthy lifestyle on BP is challenging, as these risk
 261 factors are highly interrelated. Many positive effects may be due to described
 262 physiological mechanisms. Additional benefits may be realized through a greater
 263 tendency to adhere to medical advice and treatment among individuals who engage
 264 in healthylifestyles.

265 Table 5. Correlation of left ventricular mass (LVMI), retinopathy and
 266 hypertension complications with laboratory and parameters in hypertension group.

Relationship between insulin resistance and beta cell function and LVMI (echocardiography)			
	increase LVMI (n = 18)	not increase LVMI (n = 118)	p
HOMA2-IR	1.61 ± 0.73	1.61 ± 1.14	0.996
HOMA2-%B	127.75 ± 53.26	108.76 ± 51.38	0.148
LVMI (g/m^2)	235.51 ± 60.16	122.47 ± 28.2	< 0.001
HOMA2-IR>1.22	12 (66.67%)	60 (50.85%)	0.21
HOMA2-IR<1.22	6 (33.33%)	58 (29.15%)	
HOMA2-%B<45.05	1 (5.56%)	4 (3.39%)	0.514
HOMA2-%B>45.05	17 (94.44%)	114 (96.61%)	
Relationship between insulin resistance and beta cell function and hypertension retinopathy			
	retinopathy (n = 37)	not retinopathy(n = 99)	p
HOMA2-IR	1.42 ± 0.73	1.68 ± 1.2	0.219
HOMA2-%B	110.94 ± 53.57	111.4 ± 51.44	0.963

HOMA2-IR>1.22	19 (51.35%)	53 (53.54%)	0.82
HOMA2-IR<1.22	18 (48.64%)	46 (46.46%)	
HOMA2-%B<45.05	2 (5.41%)	3 (3.03%)	0.613
HOMA2-%B>45.05	35 (94.59%)	96 (96.96%)	
Relationship between insulin resistance and beta cell function and hypertension complications			
	GRF < 60 (ml/min/1,73 m ²) (n = 25)	GRF > 60 (ml/min/1,73 m ²) (n = 111)	p
HOMA2-IR	1.84 ± 1.49	1.55 ± 0.99	0.371
HOMA2-%B	111.78 ± 71.17	111.16 ± 46.82	0.967
GRF (ml/min/1.73m ²)	50.64 ± 8.05	86.26 ± 29.05	< 0.001
HOMA2-IR>1.22	13 (48.15%)	59 (53.15%)	0.91
HOMA2-IR<1.22	12 (51.85%)	52 (46.85%)	
HOMA2-%B<45.05	2 (8%)	3 (2.71%)	0.204
HOMA2-%B>45.05	23 (92%)	108 (97.29%)	

267

268

CONCLUSION

269

270

271

272

273

274

275

276

This study found that physical inactivity, a diet low in fruits and vegetables, being overweight or obese, the presence of diabetes were strong risk factors for high blood pressure, and the risk of hypertension increased linearly with each additional exposure. Many of these risk factors are modifiable. Therefore, these findings may be important for health policy and clinical practice. Further study is needed to determine whether hypertension can be delayed or even prevented with early interventions targeting these risk factors.

277

278

279

280

281

282

283

284

285

The results of this study also demonstrate that patients with hypertension are more likely than normotensive patients to exhibit dyslipidemia, including elevated TC, LDL, TG, and reduced HDL cholesterol levels. Our results suggest that elevated BP may predict certain disturbances in lipoprotein metabolism. This association will help to develop future strategies for preventing both hypertension and dyslipidemia through proper lifestyle changes or medical management or by the combination of both. Hypertensive patients need measurement of BP and lipid profile at regular intervals throughout their primary healthcare.

286

ACKNOWLEDGEMENTS

287

288

289

The authors are grateful to Military Hospital 121 for financial support.

290

CONFLICTS OF INTEREST

291

292

The authors declare no conflict of interest.

293

294

REFERENCES

295

296

297

298

Burnier M. and Egan B.M. (2019). Adherence in hypertension: A review of prevalence, risk factors, impact, and management. *Circ. Res.*, 124:1124- 1140.

Chu D.-T., Nguyet N.T.M., Dinh T.C., Lien N.V.T., et al. (2018). An update on physical health

- 299 and economic consequences of overweight and obesity. *Diabetes Metabol. Syndr. Clin.*
300 *Res. Rev.*, 12:1095-1100.
- 301 Chu D.-T., Nguyet N.T.M., Nga V.T., Lien N.V.T., et al. (2019). An update on obesity: Mental
302 consequences and psychological interventions. *Diabetes Metabol. Syndr. Clin. Res. Rev.*,
303 13:155-160.
- 304 Cuffee Y., Ogedegbe C., Williams N.J., Ogedegbe G., et al. (2014). Psychosocial risk factors for
305 hypertension: An update of the literature. *Curr. Hypert. Rep.*, 16:483.
- 306 Ettehad D., Emdin C.A., Kiran A., Anderson S.G., et al. (2016). Blood pressure lowering for
307 prevention of cardiovascular disease and death: A systematic review and meta-analysis.
308 *The Lancet*, 387:957-967.
- 309 Gordon M.O., Beiser J.A., Brandt J.D., Heuer D.K., et al. (2002). The ocular hypertension
310 treatment study: Baseline factors that predict the onset of primary open-angle glaucoma.
311 *Arch. Ophthalmol.*, 120:714-720.
- 312 Hoang V.M., Tran Q.B., Vu T.H.L., Nguyen T.K.N., et al. (2019). Patterns of raised blood
313 pressure in Vietnam: Findings from the who steps survey 2015. *Int. J. Hypertension.*,
314 2019: 1219783.
- 315 Johnson C.A., Keltner J.L., Cello K.E., Edwards M., et al. (2002). Baseline visual field
316 characteristics in the ocular hypertension treatment study. *Ophthalmology*, 109:432-437.
- 317 Khan N.C. and Khoi H.H. (2008). Double burden of malnutrition: The Vietnamese
318 perspective. *Asia Pac. J. Clin. Nutr.*, 17 (Suppl 1):116-118.
- 319 Lelong H., Blacher J., Baudry J., Adriouch S., et al. (2019). Combination of healthy lifestyle
320 factors on the risk of hypertension in a large cohort of French adults. *Nutrients*, 11:1687.
- 321 Leung A.A., Bushnik T., Hennessy D., McAlister F.A., et al. (2019). Risk factors for
322 hypertension in Canada. *Health Rep. Stat. Canada*, 30:3-13.
- 323 Lim S.S., Vos T., Flaxman A.D., Danaei G., et al. (2012). A comparative risk assessment of
324 burden of disease and injury attributable to 67 risk factors and risk factor clusters in 21
325 regions, 1990–2010: A systematic analysis for the global burden of disease study 2010.
326 *The Lancet*, 380:2224-2260.
- 327 Margolis K.L., Buchner D.M., LaMonte M.J., Zhang Y., et al. (2019). Hypertension treatment
328 and control and risk of falls in older women. *J. Am. Geriatr. Soc.*, 67:726-733.
- 329 Meiqari L., Essink D., Wright P., and Scheele F. (2019). Prevalence of hypertension in Vietnam:
330 A systematic review and meta-analysis. *Asia Pac. J. Publ. Health*, 31:101-112.
- 331 Naghavi M., Abajobir A.A., Abbafati C., Abbas K.M., et al. (2017). Global, regional, and
332 national age-sex specific mortality for 264 causes of death, 1980–2016: A systematic
333 analysis for the global burden of disease study 2016. *The Lancet*, 390:1151-1210.
- 334 Nguyen S.M., Tran H.T.T., Tran B.Q., Van Hoang M., et al. (2020). Compliance to dietary
335 guidelines on fruit and vegetable intake and prevalence of hypertension among
336 Vietnamese adults, 2015. *Eur. J. Prevent. Cardiol.*, 27:39-46.
- 337 Olsen M.H., Angell S.Y., Asma S., Boutouyrie P., et al. (2016). A call to action and a lifecourse
338 strategy to address the global burden of raised blood pressure on current and future
339 generations: The lancet commission on hypertension. *The Lancet*, 388:2665-2712.
- 340 Quoc Cuong T., Van Bao L., Anh Tuan N., Van Thang V., et al. (2019). Associated factors of
341 hypertension in women and men in Vietnam: A cross-sectional study. *Int. J. Env. Res.*
342 *Publ. Health*, 16:4714.
- 343 Ramli A., Ahmad N.S., and Paraidathathu T. (2012). Medication adherence among hypertensive
344 patients of primary health clinics in Malaysia. *Patient Pref. Adher.*, 6:613.
- 345 Rubio-Ruiz M.E., Perez-Torres I., Soto M.E., Pastelin G., et al. (2014). Aging in blood vessels.
346 Medicinal agents for systemic arterial hypertension in the elderly. *Ageing Res. Rev.*,
347 18:132-147.
- 348 Stafylas P.C. and Sarafidis P.A. (2008). Carvedilol in hypertension treatment. *Vasc. Health*
349 *Risk Manag.*, 4:23.
- 350 Sun Z. (2015). Aging, arterial stiffness, and hypertension. *Hypertension*, 65:252-
351 256.
- 352 Trudel X., Brisson C., Gilbert-Ouimet M., Duchaine C.S., et al. (2019). Masked hypertension
353 incidence and risk factors in a prospective cohort study. *Eur. J. Prevent. Cardiol.*,
354 26:231-237.

- 355 Van N.B., Hoang L.V., Van T.B., Anh H.N.S., et al. (2019). Prevalence and risk factors of
356 hypertension in the Vietnamese elderly. *High Blood Press. Cardiovasc. Prevent.*,
357 26:239-246.
- 358 Wolf-Maier K, Cooper R.S., Kramer H., Banegas J.R., et al. (2004). Hypertension treatment and
359 control in five European countries, Canada, and the United States. *Hypertension*, 43:10-
360 17.
- 361 World Health Organization. (2013). *A global brief on hypertension: silent killer, global public*
362 *health crisis: World Health Day 2013* (No. WHO/DCO/WHD/2013.2). World Health
363 Organization.
- 364 World Health Organization. (2014). *Global status report on noncommunicable diseases*
365 *2014* (No. WHO/NMH/NVI/15.1). World Health Organization.
- 366 Zhou B., Danaei G., Stevens G.A., Bixby H., et al. (2019). Long-term and recent trends in
367 hypertension awareness, treatment, and control in 12 high-income countries: An analysis
368 of 123 nationally representative surveys. *The Lancet*, 394:639-651.