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**“A ONE YEAR CROSS SECTIONAL STUDY ON  
ATTAINMENT OF TARGET BLOOD PRESSURE  
IN HYPERTENSIVES AT KLE DR. PRABHAKAR  
KORE HOSPITAL, BELAGAVI”**

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**BY**

**REG.NO: BG0121015**

# **Dissertation**

*Submitted to*

*KAHER, Belagavi, Karnataka,*

*In partial fulfilment of the requirements for the degree of*

**M.D.**

**IN**

**GENERAL MEDICINE**

**DEPARTMENT OF GENERAL MEDICINE  
JAWAHARLAL NEHRU MEDICAL COLLEGE,  
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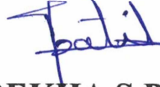
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**A ONE YEAR CROSS SECTIONAL STUDY ON ATTAINMENT OF TARGET BLOOD  
PRESSURE IN HYPERTENSIVES AT KLE DR. PRABHAKAR KORE HOSPITAL,  
BELAGAVI**

This is to certify that the dissertation entitled  
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## ABBREVIATION

NFHS	National Family Health Survey
BP	Blood pressure
SBP	Systolic blood pressure
DBP	Diastolic blood pressure
CKD	Chronic kidney disease
CVD	Cardiovascular disease
ASCVD	Atherosclerotic cardiovascular disease
DM	Diabetes mellitus
IR	Insulin Resistance
LDL	Low density lipoprotein
HDL	High density lipoprotein
CRP	C-reactive protein
ACC	American College of Cardiology
AHA	American Heart Association
ESC	European Society of Cardiology
ESH	European Society of Hypertension
TOD	Target-organ damage
HTN	Hypertension
CVS	Cardiovascular system
ICP	Intracranial pressure

JNC	Joint National Committee
CCB	Calcium channel blockers
MI	Myocardial Infarction
ACE	Angiotensin-converting enzyme
ARB	Angiotensin receptor blocker
GRF	Glomerular filtration rate
MSPSS	Multidimensional scale of perceived social support
IQR	Interquartile range
IPAQ	International physical activity questionnaire

## **ABSTRACT**

### **A ONE YEAR CROSS SECTIONAL STUDY ON ATTAINMENT OF TARGET BLOOD PRESSURE IN HYPERTENSIVES AT KLE DR. PRABHAKAR KORE HOSPITAL, BELAGAVI**

#### **Introduction**

Currently 1.3 billion people have hypertension globally. Almost half of the population with hypertension is not aware of their condition. In India, the prevalence of hypertension among men and women were reported as 24% and 21% respectively from the National Family Health Survey - 5 (NFHS-5) which was conducted in the year 2019-2020.

The study aimed to estimate the proportion of patients with hypertension attaining target blood pressure. This study also aimed to compare the profile of two groups of patients who have and have not attained target blood pressure in aspects of patient factor, socio-economic status and education in order to prevent hypertensive retinopathy and stroke.

#### **Methods**

An observational cross-sectional study was conducted among patients admitted to a tertiary care centre in Karnataka. The study period was between 1<sup>st</sup> January 2023 to 31<sup>st</sup> December 2023. The individuals who satisfy the inclusion and exclusion criteria were included in the study.

#### **Results**

A total of 400 study participants were included in this study. The mean age of the study participants was  $58.97 \pm 12.83$  years. Majority of the study participants were in the age group of 31 to 60 years (n=212, 53%). Most of the study participants were

male (n=259, 64.8%). Among the study participants, 44.5% (n=178) had smoking habit and 33.25% (n=133) had alcohol habits. Adequate amount of physical activity was followed by only 26.5% (n=106) of the study participants only. Single, double and triple drugs for treatment of hypertension was taken by 55%, 30.75% and 14.25% respectively. Among the study participants, knowledge and adherence questions were asked. Among 400 individuals, 77% (n=308) were aware of the adverse effects of increased blood pressure. Among the study participants, 70.25% (n=281) were taking their anti-hypertensive medications regularly. Among the study participants, 55.75% (n=223) of the study participants were aware of the benefits of taking anti-hypertensives and 62.5% (n=250) were aware of the adverse effects of them. Majority of their physicians (88.5%, n=354) explained the benefits of taking anti-hypertensives.

### **Conclusion**

In our study, only 21.5% achieved target blood pressure. The mean age of the study participants was  $58.97 \pm 12.83$  years. Study variables significantly associated with control of hypertension were education, socio-economic status, heart rate, treatment of hypertension, newly diagnosed hypertension, total cholesterol, urea levels and urine protein in our study.

## INDEX

<b>Sr. No</b>	<b>Content</b>	<b>Page. no</b>
<b>1</b>	<b>INTRODUCTION</b>	<b>1-2</b>
<b>2</b>	<b>AIM AND OBJECTIVES</b>	<b>3</b>
<b>3</b>	<b>REVIEW OF LITERATURE</b>	<b>4-41</b>
<b>4</b>	<b>MATERIAL AND METHODS</b>	<b>42-44</b>
<b>5</b>	<b>RESULTS</b>	<b>45-63</b>
<b>6</b>	<b>DISCUSSION</b>	<b>64-70</b>
<b>7</b>	<b>CONCLUSION</b>	<b>71-72</b>
<b>8</b>	<b>REFERENCES</b>	<b>73-76</b>
<b>9</b>	<b>ANNEXURES</b>	<b>77-93</b>
	<b>1. CONSENT FORM</b>	<b>94-104</b>
	<b>2. PROFORMA</b>	
	<b>3. MASTER CHART</b>	

## **LIST OF TABLES**

<b>Sr. No.</b>	<b>Table</b>	<b>Page no.</b>
<b>1</b>	Modifiable and non-modifiable risk factors of hypertension	<b>6</b>
<b>2</b>	Cardiovascular risk factors and hypertension	<b>7</b>
<b>3</b>	JNC8 Hypertension classification	<b>8</b>
<b>4</b>	ACC/ AHA hypertension classification	<b>9</b>
<b>5</b>	Classification of blood pressure based on ESC/ ESH guidelines	<b>10</b>
<b>6</b>	Antihypertensive drugs	<b>16</b>
<b>7</b>	Distribution of age among study participants	<b>25</b>
<b>8</b>	Distribution of gender among study participants	<b>26</b>
<b>9</b>	Distribution of BMI among study participants	<b>27</b>
<b>10</b>	Smoking and Alcohol habits of the study participants	<b>29</b>
<b>11</b>	Physical activity among study participants	<b>30</b>
<b>12</b>	Number of drugs for control of hypertension	<b>33</b>
<b>13</b>	Mean value of lipid profile	<b>34</b>
<b>14</b>	Comparison of socio-demographic variables with control of hypertension	<b>37</b>
<b>15</b>	Comparison of Education and locality	<b>38</b>
<b>16</b>	Comparison of personal habits, diet and physical activity of study participants	<b>39</b>
<b>17</b>	Comparison of BMI and heart rate of study participants	<b>40</b>

<b>18</b>	Comparison of hypertension profile	<b>41</b>
<b>19</b>	Comparison of biochemical parameters and urine protein among study participants	<b>42</b>
<b>20</b>	Knowledge and adherence of study participants and the control of hypertension	<b>43</b>

## **LIST OF FIGURES**

<b>Sr. No.</b>	<b>Figure</b>	<b>Page. no</b>
<b>1</b>	Pathophysiology of primary hypertension	<b>5</b>
<b>2</b>	Hypertensive retinopathy	<b>12</b>
<b>3</b>	Hypertension and stroke	<b>13</b>
<b>4</b>	Age distribution of study participants	<b>25</b>
<b>5</b>	Gender distribution of study participants	<b>26</b>
<b>6</b>	BMI distribution among study participants	<b>27</b>
<b>7</b>	Salt intake of study participants	<b>28</b>
<b>8</b>	Personal habits of study participants	<b>29</b>
<b>9</b>	Physical activity among study participants	<b>30</b>
<b>10</b>	Diet followed by the study participants	<b>31</b>
<b>11</b>	Number of drugs taken for control of hypertension	<b>33</b>
<b>12</b>	Urine protein among study participants	<b>35</b>
<b>13</b>	Knowledge and adherence of study participants regarding hypertension and its medication	<b>36</b>

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## INTRODUCTION

Hypertension is characterized as increased blood pressure in the systemic arteries in a persistent way. Hypertension is also known as systemic arterial hypertension. Hypertension is calculated as systolic blood pressure divided by the DBP (diastolic blood pressure). Systolic blood pressure (SBP) is the pressure that exerts on the arterial walls when heart contracts and the diastolic blood pressure is the pressure when the heart relaxes (1). Majority of the people with hypertension has primary hypertension or essential hypertension. The etiology of this primary or essential hypertension is usually multifactorial gene-environment cause. The most prevalent preventable risk factor for chronic kidney disease (CKD), cognitive impairment, myocardial infarction, heart failure, stroke, and cardiovascular disease (CVD) is hypertension. It also serves as the primary cause of all-cause death and disability globally. Gradient and continuous, the association between increased BP and the risk of CVD begins as low as 115/75 mmHg, well within the range generally accepted as normotensive (2).

Reducing the burden of disease and increasing life expectancy among people worldwide depend on the effective prevention and treatment of hypertension. Predicted atherosclerotic CVD (ASCVD) risk should be taken into account while treating hypertension, as individuals with high CVD risk benefit most from blood pressure reducing therapy (3).

Currently 1.3 billion people have hypertension globally. Almost half of the population with hypertension is not aware of their condition. In the year 1990, the people with disease were 650 million. Whereas, in 2019, the disease affected 1.3

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billion people which is double the population (4). Prevalence of hypertension among men and women were reported as 24% and 21% respectively from the National Family Health Survey - 5 (NFHS-5) which was conducted in the year 2019-2020 in India (5).

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## **AIM AND OBJECTIVES**

This study aimed to estimate the proportion of patients with hypertension attaining target blood pressure.

This study also aimed to compare the profile of two groups of patients who have and have not attained target blood pressure in aspects of patient factor, socio-economic status and education in order to prevent hypertensive retinopathy and stroke.

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## REVIEW OF LITERATURE

Hypertension is the most important risk factor for morbidity and mortality in India (6). There are different stages of hypertension according to the progression of disease. This can be used to assess the cardiovascular risk in the individuals. Cardiovascular risk assessment helps in predicting the future likelihood of occurrence of cardiovascular events like myocardial infarction and stroke.

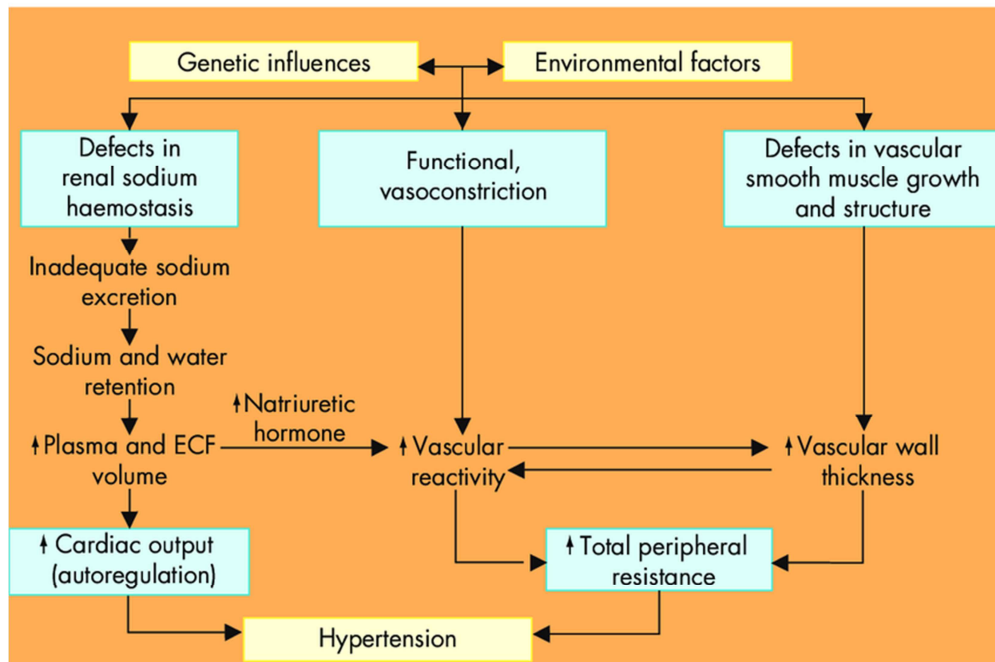
The progression of hypertension in an order of early to advanced was represented as Stage 1, Stage 2 and Stage 3 hypertension. It is characterized by the cumulative presence or absence of markers of hypertensive cardiovascular disease and evidence of target organ damage regardless of the blood pressure levels. Blood pressure is considered as a biomarker for hypertension (7).

### **Pathophysiology of hypertension**

Normal blood pressure is linked to the lowest risk of cardiovascular disease, but people in the "high-normal" range are at danger of hypertension unless they change their diet to include more calcium, magnesium, and potassium as well as less fat. A shift in the pressure–natriuresis relationship and a rise in vascular tone are brought on by the dysfunction of these hormones as well as changes in the sympathetic nervous system and the renin–angiotensin–aldosterone system. Retention of sodium causes water to be retained as well, increasing blood volume and raising blood pressure. Tissue ischaemia and renal vasoconstriction accompany the subtle renal damage. In addition to contributing to renal inflammation, tissue ischaemia also affects the kidneys' internal structure—specifically, the glomeruli and tubules—which in turn encourages the kidneys to retain more sodium. Increases in resting blood pressure eventually result from this vicious cycle, which also causes hypertension.

One aspect of the pathophysiology of hypertension is inflammation. Vasoactive inflammatory cytokines are released as a result of endothelial damage and tissue ischaemia. While several of these cytokines—like histamine—have vasodilatory effects during acute inflammatory damage, chronic inflammation plays a role in smooth muscle contraction and vascular remodelling. Reduced release of vasodilators, like nitric oxide, and increased release of vasoconstrictors, like endothelin, are further characteristics of endothelial damage and dysfunction in primary hypertension. Finally, those without clinical DM, IR is frequently associated with hypertension. Thirteen Reduced endothelial release of nitric oxide and other vasodilators is linked to insulin resistance. It also makes the kidneys retain water and salt, which impairs renal function. The renin-angiotensin-aldosterone system and the sympathetic nervous system hyperactivity are linked to insulin resistance (8).

**Figure 1: Pathophysiology of primary hypertension**



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## Risk factors for development of hypertension

Risk factors of hypertension is classified as Modifiable and non-modifiable risk factors of hypertension. Unhealthy diet including excessive salt consumption, diet high in saturated and trans-fat, and low intake of fruits and vegetables, physical inactivity, consumption of tobacco and alcohol and overweight or obese are considered as modifiable risk factors.

Non-modifiable risk factors are age (above 65 years), family history of hypertension, co-existing diseases like diabetes or kidney disease (Table 1) (4).

**Table 1: Modifiable and non-modifiable risk factors of hypertension**

Modifiable risk factors	Non-modifiable risk factors
<ol style="list-style-type: none"><li>1. Diet Excessive salt intake High in saturated fat and trans fat Low intake of fruits and vegetables</li></ol>	<ol style="list-style-type: none"><li>1. Age above 65 years</li><li>2. Family history of hypertension</li><li>3. Co-morbidities like diabetes or kidney disease</li></ol>
<ol style="list-style-type: none"><li>2. Physical activity</li><li>3. Consumption of tobacco and alcohol</li><li>4. Overweight or obesity</li></ol>	

### Cardiovascular risk factors and classification of hypertension

The below table describes the risk factors associated in each stage of hypertension (Table 2).

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**Table 2: Cardiovascular risk factors and hypertension**

Stages	Cardiovascular risk factors
Normal	None or few
Stage 1	Several risk factors present
Stage 2	Many risk factors present
Stage 3	Many risk factors present

Cardiovascular risk factors are:

1. Increased age
2. Elevated blood pressure
3. High heart rate
4. Overweight or obese
5. Central obesity (increased abdominal circumference, increased abdominal adiposity)
6. Dyslipidemia (elevated LDL or non-HDL cholesterol, low HDL cholesterol, and elevated triglycerides)
7. Elevated blood glucose, insulin resistance and/ or diabetes
8. Chronic kidney disease
9. Smoking
10. Family history of premature CVD (Men less than 50 years, women less than 60 years)
11. Sedentary lifestyle
12. Psychosocial stressors
13. Elevated high sensitivity-CRP (C-reactive protein)

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These markers are either absent or few of them are present in normal individuals. In stage 1 hypertension, several risk factors may be present. In stage 2 and stage 3 several risk factors mentioned above will be prevalent (7).

#### Blood pressure and hypertension

There are few guidelines worldwide for diagnosis and staging of hypertension. The first, Joint National Committee (Eighth) classification as below Table 3.

**Table 3: JNC8 Hypertension classification**

Blood pressure	Systolic blood pressure	Diastolic blood pressure
Normal	<120	And <80
Prehypertension	120-139	Or 80-90
Stage 1 hypertension	140-159	Or 90-99
Stage 2 hypertension	≥160	Or ≥100

The normal blood pressure according to JNC8 is less than 120 and 80 of systolic and diastolic blood pressure. Prehypertension is determined as either systolic blood pressure of 120-139 or diastolic blood pressure of 80-90. Stage 1 hypertension is considered as 140-159 of systolic blood pressure or 90-99 of diastolic blood pressure. Stage 2 hypertension is considered as either more than 160 of systolic blood pressure or more than 100 of diastolic blood pressure (9).

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The ACC/ AHA guidelines for classification of hypertension is mentioned below in Table 4.

**Table 4: ACC/ AHA hypertension classification**

Blood pressure	Systolic blood pressure	Diastolic blood pressure
Normal	<120	And <80
Elevated	120-129	Or <80
Stage 1 hypertension	130-139	Or 80-89
Stage 2 hypertension	≥140	Or ≥90

ACC (American College of Cardiology)/ AHA (American Heart Association) recommendations for normal blood pressure is less than 120 systolic and less than 80 diastolic blood pressure. A person has elevated blood pressure when the systolic is between 120 and 129 or the diastolic above 80. Stage 1 hypertension is considered as 130 to 139 systolic blood pressure or 80 to 89 DBP. Stage 2 hypertension is considered as SBP is 140 or above and DBP 90 or above (10).

The ESC (European Society of Cardiology)/ ESH (European Society of Hypertension) guidelines for classification of hypertension is mentioned below in Table 5.

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**Table 5: Classification of blood pressure based on ESC/ ESH guidelines**

Blood pressure	Systolic blood pressure	Diastolic blood pressure
Optimal	<120	And <80
Normal	120-129	And 80-84
Elevated	130-139	Or 85-89
Grade 1 hypertension	140-159	Or 90-99
Grade 2 hypertension	160-179	Or 100-109
Grade 3 hypertension	≥180	Or ≥110

In ESC/ ESH guidelines, optimal blood pressure is considered as SBP of less than 120 and diastolic blood pressure of less than 80. Normal blood pressure is 120-129 SBP and 80-84 DBP. Elevated or prehypertensive is considered as 130-139 SBP or 85-89 DBP. Grade 1 hypertension is 140 to 159 SBP or 90-99 DBP. Grade 2 hypertension is 160 to 179 SBP or 100 to 109 of DBP. Grade 3 hypertension is considered as SBP of 180 or above Or DBP of 110 or above (11).

#### Complications of hypertension

Complications of hypertension includes ventricular hypertrophy, heart failure, accelerated atherosclerosis, cerebrovascular disease, stroke, renal failure, retinopathy, dissecting aneurysm, retinal hemorrhages, hypertensive encephalopathy (12, 13).

#### Hypertensive retinopathy

Retinal, cardiovascular, renal, and cerebrovascular systems are among the systems that are impacted by poorly managed hypertension. Target-organ damage (TOD) is the term for the harm done to these systems. Three forms of visual injury are brought on by hypertension: optic neuropathy, retinopathy, and choroidopathy (14). Since retinal vessels are the only blood vessels visible during a regular examination,

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screening for hypertensive retinopathy is primarily done for this reason. Hypertensive retinopathy and choroidopathy are two obvious symptoms of persistently increased HTN, and they are a reflection of vascular alterations taking place in other systems. Together, general practitioners and ophthalmologists can guarantee that patients with hypertension are effectively screened and treated promptly to lower the risk of both ocular and systemic morbidity and death (15).

The severity of the condition determines how hypertensive retinopathy is managed:

**Mild:**

Treatment for mild hypertensive retinopathy involves regular blood pressure monitoring.

**Moderate:**

A doctor's referral is necessary for moderate hypertensive retinopathy in order to rule out other possible causes, like DM, and check in the CVS. Normal care, such as blood pressure monitoring and control, is essential.

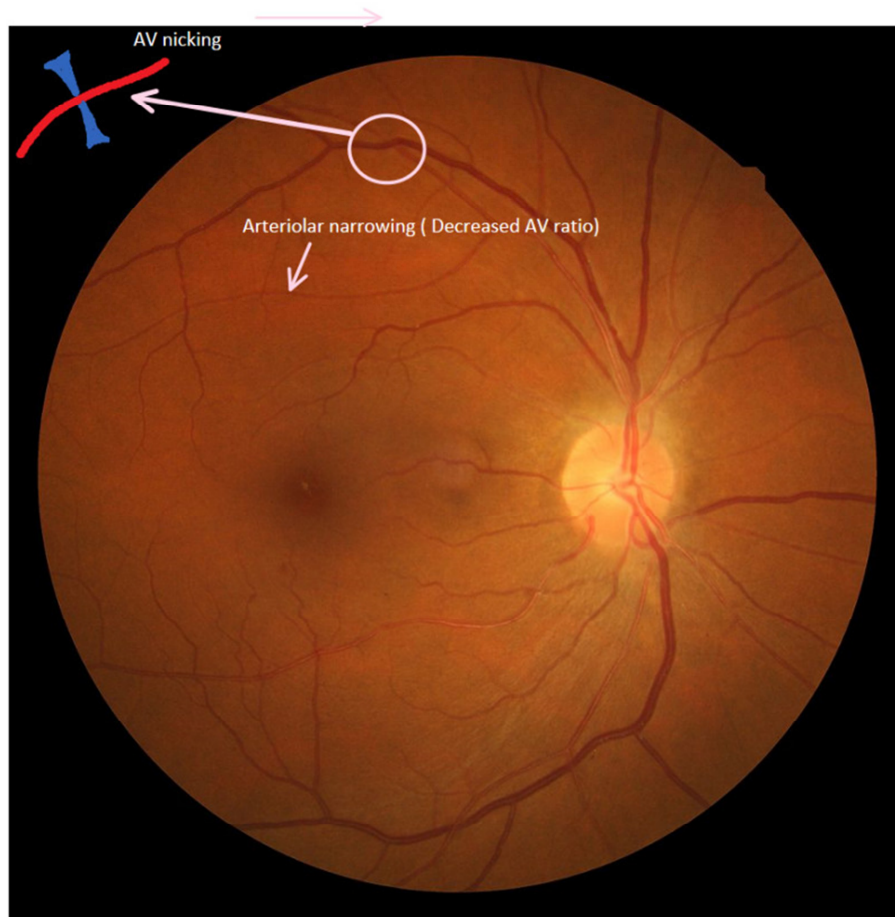
**Severe:**

Because severe hypertensive retinopathy has the highest fatality rate, it needs to be treated and sent to a specialist immediately. The kidney, heart, and brain are among the other systems that need to be watched for indications of TOD (14).

Only rarely may persistent hypertensive retinopathy result in noticeable vision loss. Treatment of hypertension can stop the alterations in the retina. Arteriolar constriction and AV alterations nevertheless continue. The death rate for untreated

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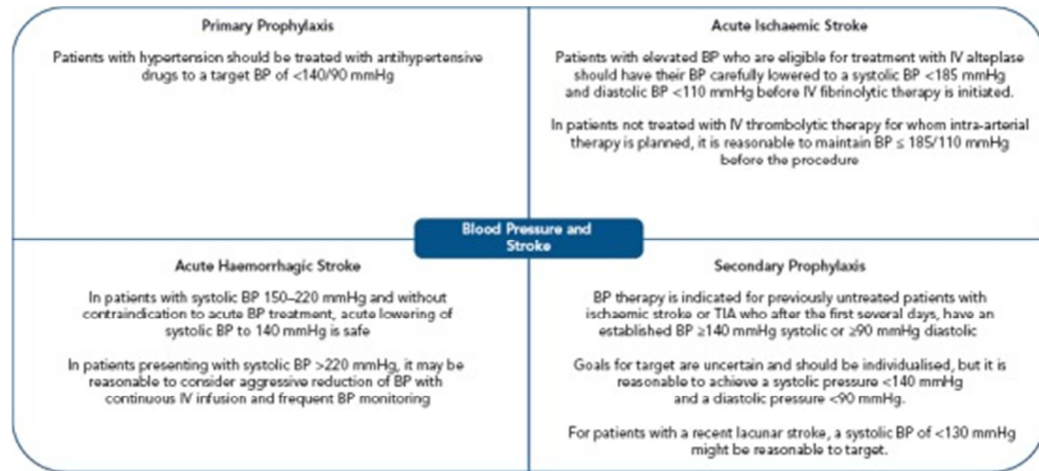
malignant hypertension is as high as 50% after two months of diagnosis and over 90% after a year. Either secondary optic atrophy following protracted papilloedema or retinal pigmentary alterations following exudative retinal detachment induce vision loss in hypertensive retinopathy.



**Figure 2: Hypertensive retinopathy**

### Hypertensive stroke

Depending on the stroke subtype and the stage at which the illness manifests, there are differences in the cause of stroke and its hemodynamic effects. As a result, controlling blood pressure (BP) in stroke patients is difficult and necessitates a clear diagnosis as well as the establishment of treatment objectives (16) (Figure 3).



**Figure 3: Hypertension and stroke**

When an intracranial or cervical artery is blocked, a portion of the brain is deprived of blood and oxygen, which leads to acute ischemic strokes. A central ischemic lesion forms in the brain a few minutes after an artery occlusion; however, if recanalization therapies are used, a greater area at risk of hypoperfusion may be salvageable. Acute blood pressure drops can jeopardise perfusion in vital locations, which is why the salvageable area, also known as the ischemic penumbra, is mostly dependent on collateral blood flow (17).

Early antihypertensive medication commencement or restart is recommended only in patients receiving recombinant tissue-type plasminogen activator treatment or in cases of severe hypertension during the immediate phase of an ischemic stroke. Antihypertensive medication is advised for patients who qualify for IV thrombolysis so long as their systolic and diastolic blood pressures are  $\leq$ 185 mmHg and  $\leq$ 110 mmHg, respectively, before to therapy and for the first 24 hours following treatment (18). Although there were no long-term gains in independence or mortality from a recently published clinical trial assessing stricter blood pressure objectives following

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IV thrombolysis for acute ischemic stroke, lower blood pressure levels were linked to a lower risk of hemorrhagic transformation (19).

Reduction of acute blood pressure is unclear among individuals with acute ischemic stroke in which the thrombolysis was not yet developed. Treatment for these people should only be started if the patient has another obvious indication or if the SBP or DBP is greater than 220 mmHg or 120 mmHg. Even lowering blood pressure quickly to less than hypertensive levels can have negative effects. Therefore, within the first 24 hours following the beginning of stroke, BP should be carefully decreased by approximately 15% if indicated. When BP is lower than 180/105 mmHg within the three days following an acute ischemic stroke, patients do not appear to benefit from starting or continuing to use blood pressure-lowering medication. After an acute ischemic stroke, stable patients who maintain hypertension ( $\geq 140/90$  mmHg) for longer than three days may benefit from starting or restarting blood pressure-lowering medication. For stable hypertension individuals, it is reasonable to resume blood pressure control after the first twenty-four hours (18).

After ischemic stroke, spontaneous, non-traumatic intracerebral haemorrhage is next reason for development of stroke. Frequent reasons include drug abuse, vascular abnormalities, amyloid angiopathy, hypertension, and bleeding diatheses (20). Another variety of hemorrhagic stroke is subarachnoid haemorrhage. The two main causes of subarachnoid haemorrhage are bleeding from vascular abnormalities located close to the pial surface and rupture of arterial aneurysms located at the base of brain (21).

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In a major randomised clinical trial, intensive BP lowering (<140 mmHg) was safe and related with a little better functional recovery in stroke survivors; nevertheless, it did not clearly improve clinical prognosis in these patients. Additionally, there was a positive trend towards a decline in the traditional clinical end goal of death and significant impairment. In a different clinical trial utilising intravenous nicardipine, however, more severe BP lowering (<120 mmHg) was linked to more renal side events in addition to not demonstrating any therapeutic advantages (22). According to American Heart Association guidelines, it is safe and beneficial to drop a patient's SBP to 140 mmHg immediately after experiencing an intracerebral haemorrhage if they do not have a contraindication to receiving acute blood pressure medication. This can improve the patient's functional prognosis. Another crucial factor to take into account while treating patients who have experienced intracerebral bleeding is intracranial pressure. CPNs should be kept between 61 and 80 mmHg if the systolic blood pressure is greater than 180 mmHg and there is evidence or suspicion of excessive ICP. A slight drop in blood pressure (160/90 mmHg) is advised if there is no indication or reason to suspect elevated intracerebral pressure. It is probably safe to abruptly drop blood pressure to 140 mmHg if it is 150–200 mmHg (20).

#### Treatment for hypertension

List of antihypertensive drugs are listed below in Table 6.

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**Table 6: Antihypertensive drugs**

Class of antihypertensive drugs	
Diuretics	Thiazides (Hydrochlorothiazide, Chlorthalidone, Indapamide).  High ceiling (Furosemide)  Potassium sparing (Spironolactone, Amiloride)
ACE inhibitors	Captopril, Enalapril, Lisinopril, Ramipril, Perindopril
ARBs	Losartan, Telmisartan, Valsartan, Candesartan, Irbesartan
Calcium channel blockers	Amlodipine, Cilnidipine, Diltiazem, Verapamil, Nifedipine
Beta adrenergic blockers	Propranolol, Metoprolol, Atenolol
Alpha adrenergic blockers	Prazosin, Phenoxybenzamine, Phentolamine
Alpha + Beta blockers	Labetalol, Carvedilol
Central sympatholytic	Clonidine, Methyl dopa
Vasodilators	Arteriolar (Hydralazine, Minoxidil, Diazoxide)  Arteriolar + Venous (Sodium nitroprusside)
Direct renin inhibitor	Aliskerin

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Thiazide:

Eighth Joint National Committee recommended thiazide to be given as preferred medicine for hypertension. It can be given as monotherapy or as one of the drugs in multitherapy for everyone regardless of age, ethnicity. Only contraindication was to avoid in CKD patients, in which case an angiotensin-converting enzyme inhibitor or angiotensin II receptor blocker is indicated. Lipid-Lowering and Antihypertensive Treatment Trial to Prevent Heart Attack Unless there are contraindications, the ALLHAT research advised starting treatment for hypertension with thiazide diuretics.

### **Calcium channel blockers (CCB)**

Calcium channel blockers were advised by Eighth Joint National Committee guidelines to be given as preferred medicine for hypertension. It can be given as monotherapy or as one of the drugs in multitherapy for everyone regardless of age, ethnicity, patients with chronic kidney disease, on the other hand, should use ACE inhibitors or ARBs as recommended as the first-line treatment. Like thiazide diuretics, CCBs have been demonstrated to reduce all cardiovascular outcomes except heart failure. When patients are unable to tolerate thiazides, they might be utilised as the most effective substitute. Dihydropyridines and non-dihydropyridines are the two categories of CCBs.

Dihydropyridines are more frequently used to treat hypertension since they are stronger vasodilators. Their impact on cardiac conduction and contractility is minimal. They are therefore more frequently utilised in the treatment of HTN. In this group, amlodipine and nifedipine are the most commonly utilised drugs. Non-

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dihydropyridines have a greater impact on cardiac conduction and contractility but are less effective as vasodilators. They are used less to treat HTN and more as antiarrhythmic drugs.

### **ACE Inhibitors and ARBs**

The preferred antihypertensives for people with heart failure and chronic renal disease are ACE inhibitors and ARBs. For those with chronic renal disease who exhibit proteinuria, they are recommended as the initial line of treatment. With first and second drugs, these two kinds of antihypertensive drugs are listed in JNC8 guidelines as the first-line treatment for HTN in non-Black individuals. They have been shown to have a cardioprotective impact in people at high risk of cardiovascular disease, independent of their antihypertensive action. Both classes are advised as first-line treatments for individuals with left ventricular failure and either ST-elevation MI or non-ST elevation MI combined with diabetes, systolic dysfunction, or anterior infarct. They also have comparable efficacies and treatment indications. When it comes to lowering blood pressure, preventing stroke, and preventing heart failure, thiazide is superior to ACE inhibitors; CCBs are superior to ACE inhibitors in these areas.

### **Beta-blockers**

It is not recommended to use beta-blockers as the sole treatment for hypertension unless there is a clear sign of myocardial infarction and heart failure. When administered in younger individuals, beta-blockers are linked to lower cardiovascular morbidity and mortality; however, in older patients, they are less protective and have been linked to an increased risk of strokes (23).

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A study conducted in Latvia, Europe, was evaluated the patients who were treated for hypertension on whether they achieved adequacy of blood pressure control. This study included 455 individuals between age 18 to 80 years with history of hypertension of at least 1 year duration. The target blood pressure for this study participants were less than 140/ 90 mm Hg with low or moderate cardiovascular risk and less than 135/85 to 125/75 mm Hg for high-risk individuals. Among the study participants, almost half of them (46.2%) had attained the targeted blood pressure values. Individuals in the low and moderate risk group were higher in proportion in achieving the target blood pressure when compared to high-risk group (61.7% vs 34.4%,  $p < 0.0001$ ). In this study, majority of the individuals were given combination of two (26.2%) or three (31.6%) anti-hypertensive drugs. Overall conclusion was in the sample of treated hypertension patients, the rate of effective blood pressure management was less than 50%, and it was considerably lower (34.4%) in patients with high or very high increased cardiovascular risk (24).

A retrospective cohort study conducted in Turkey included 437 individuals with hypertension diagnosed during the study. The study aimed to estimate the proportion of study participants diagnosed as hypertensive on achievement of target blood pressure. The study also aimed to determine the factors responsible for not achieving the target blood pressure among the study participants. The follow-up data of 63.1% of individuals were available and collected. In the baseline visit, 18.1% and in the follow-up visit 48.6% of the individuals achieved target blood pressure (according to JNC8 guidelines). Diabetes mellitus and baseline systolic blood pressure were positively impacted in achieving systolic blood pressure. Age, addition of more anti-hypertensive medications impacted negatively on the systolic blood pressure target. Diabetes mellitus, baseline diastolic blood pressure, duration of

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hypertension was positively associated with diastolic blood pressure improvement whereas GFR values and additional anti-hypertensive drugs negatively impacted the diastolic blood pressure target. Study concluded only 50% of patients had attained the hypertension objectives, despite significant improvement in meeting BP targets. Patients with hypertension should get both comprehensive management techniques and appropriate pharmaceutical treatments (25).

A cross-sectional study conducted in Ethiopia during 2021 aimed to explore the determinants of blood pressure control status of individuals with history of hypertension. The study was aimed to conduct during the COVID-19 pandemic period. The study included individuals diagnosed with hypertension at least 3 months of treatment and included individuals above 18 years of age. A total of 360 individuals were included in the study. Multidimensional scale of perceived social support (MSPSS), a 12-item check-list was also administered to the patients to understand the perceived social support. Prevalence of uncontrolled hypertension among the study participants was 55.8%. Poor medication adherence, male gender, secondary education, and low social support were the determinants of uncontrolled blood pressure among the study participants (26).

Another cross-sectional study conducted in Japan aimed to determine the rate of controlled blood pressure according to Japanese hypertension management guidelines and the factors associated in achieving the same. The study was conducted between 2012 and 2015. The residents, the study participants were grouped into 6 groups. First group includes young, middle-aged, and early-phase elderly patients (G1). Second group includes patients with cerebrovascular disease (G2). Third group includes patients with cardiovascular diseases (G3). Fourth group includes individuals

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with chronic kidney disease with proteinuria (G4). Fifth group includes individuals with diabetes (G5). Sixth group includes individuals with chronic kidney disease without proteinuria (G6). Target blood pressure was achieved in 52.6%, 84.3%, 50.6%, 45.6%, 48.7% and 75% in G1, G2, G3, G4, G5 and G6 respectively. Body mass index and intake of anti-lipidemic medication were associated factors with uncontrolled blood pressure (27).

A study conducted in India to examine the hypertension care continuum among 1.7 million individuals between 18 to 98 years. The nationally representative data from the National Family Health Survey (NFHS) was conducted in 2 phases; one in 2019-2020 and another in 2020-2021. From the study, 28.1% had hypertension, of which 36.9% had already diagnosed with the disease. among the individuals with known hypertension, 44.7% were taking medication for it. Among the individuals who were under treatment, 52.5% had their hypertension under control (28).

A cross-sectional study from South India aimed to evaluate the prescription pattern for hypertension and factors affecting controlled blood pressure. The study included 650 individuals with known hypertension and are on treatment. Among 650 individuals, 39.54% achieved target blood pressure levels. Age, occupational status, monthly family income, locality, physical activity, diet was significantly associated in achieving target blood pressure. Among individuals who achieved target blood pressure, 37.35% were on monotherapy and 48.25% were on multiple drug therapy (29).

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## METHODOLOGY

An observational cross-sectional study was conducted among patients visiting the Department of General Medicine (outpatient department) and admitted to the medical ward of KLES Dr Prabhakar Kore Hospital and Medical Research Centre, Belagavi. The study period was between 1<sup>st</sup> January 2023 to 31<sup>st</sup> December 2023. The individuals who satisfy the inclusion and exclusion criteria were included in the study.

### **Inclusion criteria**

- Patients age 18 or above
- History of known hypertension
- Duration of hypertension minimum 3 months
- Provide consent to participate in the study

### **Exclusion criteria**

- Critically ill and mentally unstable individuals
- Patients with chronic kidney disease

Patients who were attending the Outpatient Department and Inpatient ward of Department of General Medicine of KLES Dr Prabhakar Kore Hospital and Medical Research Centre, Belagavi between the period of January 2023 to December 2023 were checked for the inclusion and exclusion criteria of our study. If the patient satisfied the inclusion and exclusion criteria, the individual was considered as eligible to participate in the study. Eligible participants who provided informed consent were included in the study. Blood pressure of each individual was recorded. Steps followed while measuring the blood pressure were as below:

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1. The study participant was asked to sit, relax and stay quiet for 5 minutes.
  2. Had the patient bladder empty, and checked for intake of caffeine, exercise and smoking in the past 30 minutes.
  3. Removed the clothing from the arm.
  4. Used a properly validated and calibrated blood pressure measurement device.
  5. Supported patient's arm and position cuff on bare arm at the level of right atrium.
  6. Used the correct cuff size.
  7. For first record, recorded BP in both the arms, and used the one with higher reading.
  8. Separate similar readings after 1 to 2 minutes.
  9. Recorded systolic and diastolic blood pressure.
  10. Used an average of 2 readings obtained on 2 occasions to estimate the individual's BP.
  11. Recorded the systolic and diastolic blood pressure.

The blood pressure, personal details, and other study variables were recorded from the patient and the previous records as required in the pre-designed study proforma.

The individuals were categorized based on the JNC8 guidelines. The patients who attained the target blood pressure and those who didn't achieve the target blood pressure were compared for socio-economic and education.

#### Sample size calculation

The sample size formula used to calculate the sample size of our current study was:

$$\text{Sample size formula} = \frac{Z^2(pq)}{d^2}$$

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Where ,  $Z^2$  - standard normal variant at a 95%

degree of confidence = 1.96

p - Expected proportion from population prevalence

q = 100-p

d - margin of error = 5%.

Sample size n = 384 should be taken.

Hence, the sample size is 384.

Statistical analysis

Data was collected and entered in the Excel sheet. IBM SPSS Version 25 was used to analyse the data. The continuous variables were reported as mean and standard deviation. The categorical variables were reported as number and percentages. Multiple logistic regression was carried out. Chi-square was done to see the association. ANOVA was carried out to see the significant difference.

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## RESULTS

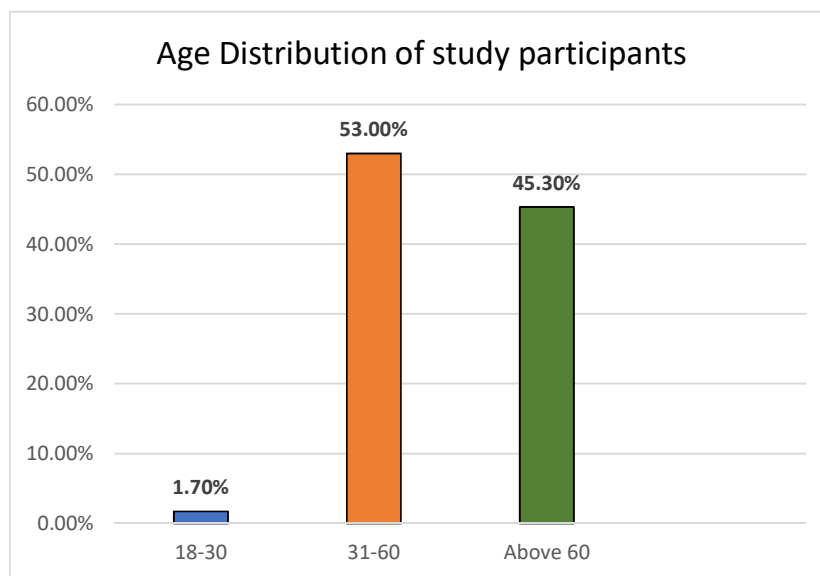
A total of 400 study participants were included in this study.

### *Age and Gender distribution*

The mean age of the study participants was  $58.97 \pm 12.83$  years. The age distribution is tabulated as Table 7 (and Figure 4). Majority of the study participants were in the age group of 31 to 60 years (n=212, 53%).

**Table 7: Distribution of age among study participants**

Age category	Number (n=400)	Percentage
18-30 Years	7	1.7%
31-60 Years	212	53%
Above 60 Years	181	45.3%



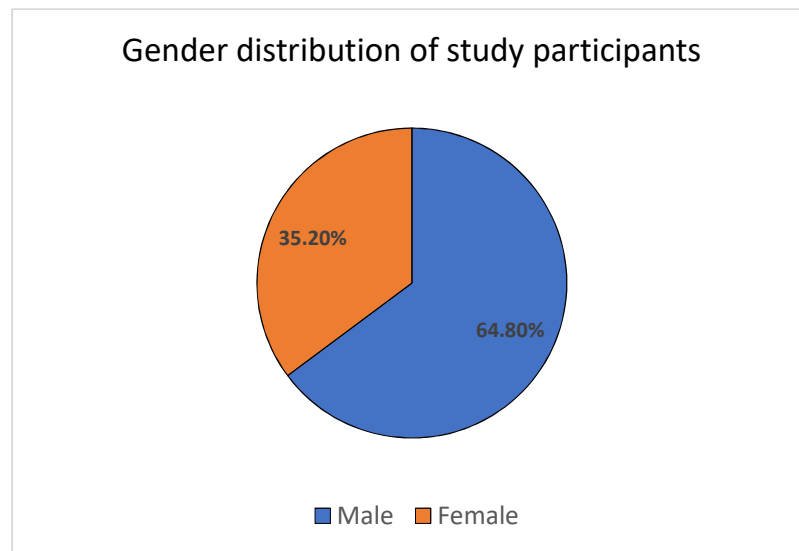
**Figure 4: Age distribution of study participants**

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Majority of the study participants were male (n=259, 64.8%) (Table 8) (Figure 5).

**Table 8: Distribution of gender among study participants**

Gender	Number (n=400)	Percentage
Male	259	64.8%
Female	141	35.2%



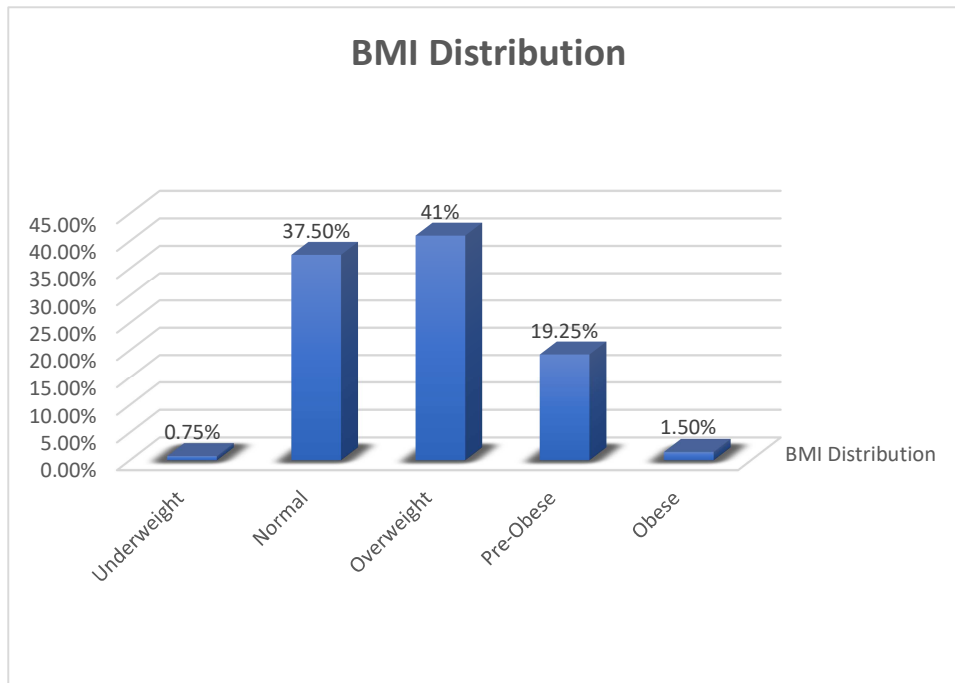
**Figure 5: Gender distribution of study participants**

*Body Mass Index*

The mean BMI of the study participants was  $23.52 \pm 2.38$ . The majority of the study participants were categorized as overweight (BMI of 23 to 24.9) based on their BMI (n=164, 41%) (Table 9 and Figure 6).

**Table 9: Distribution of BMI among study participants**

Body Mass Index		Number (n=400)	Percentage
Less than 18.5	Underweight	3	0.75%
18.5 to 22.9	Normal	150	37.5%
23 to 24.9	Overweight	164	41%
25 to 29.9	Pre-Obese	77	19.25%
30 to 40	Obese (Type 1)	6	1.5%

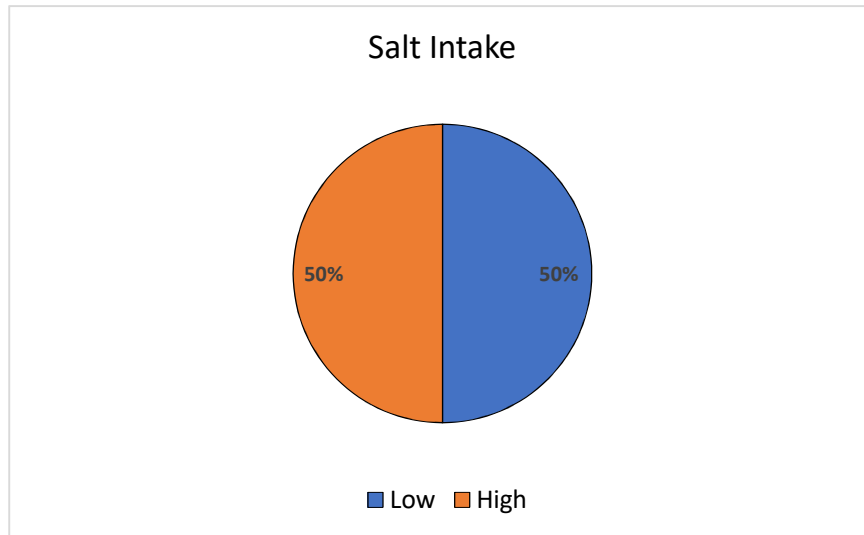


**Figure 6: BMI distribution among study participants**

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*Salt Intake of study participants*

Salt intake of the study participants was similar (that is, low intake: n=200, 50% and high intake n=200, 50%) (Figure 7).



**Figure 7: Salt intake of study participants**

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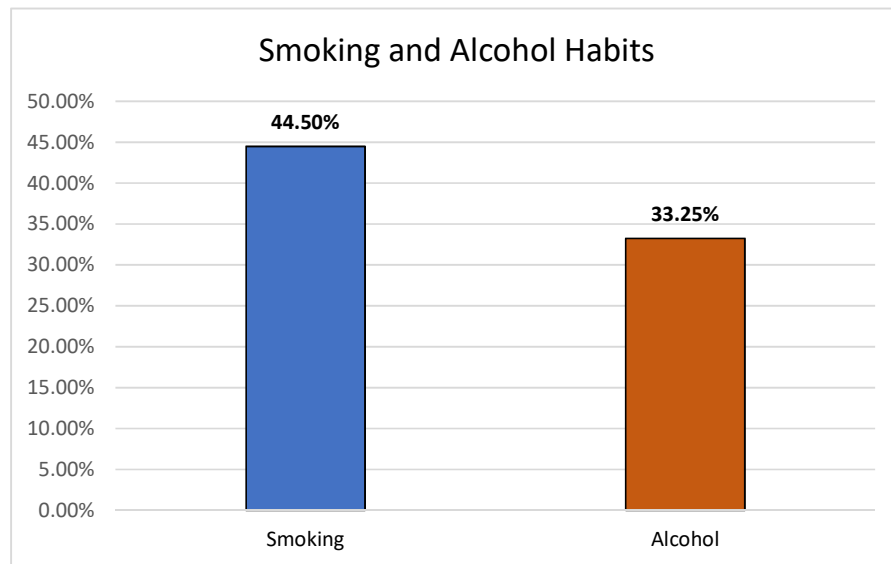
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***Personal Habits – Smoking and Alcohol***

Among the study participants, 44.5% (n=178) had smoking habit and 33.25% (n=133) had alcohol habits (Table 10 and Figure 8).

**Table 10: Smoking and Alcohol habits of the study participants**

<b>Habit</b>	<b>Number (n=400)</b>	<b>Percentage</b>
Smoking	178	44.5%
Alcohol	133	33.25%



**Figure 8: Personal habits of study participants**

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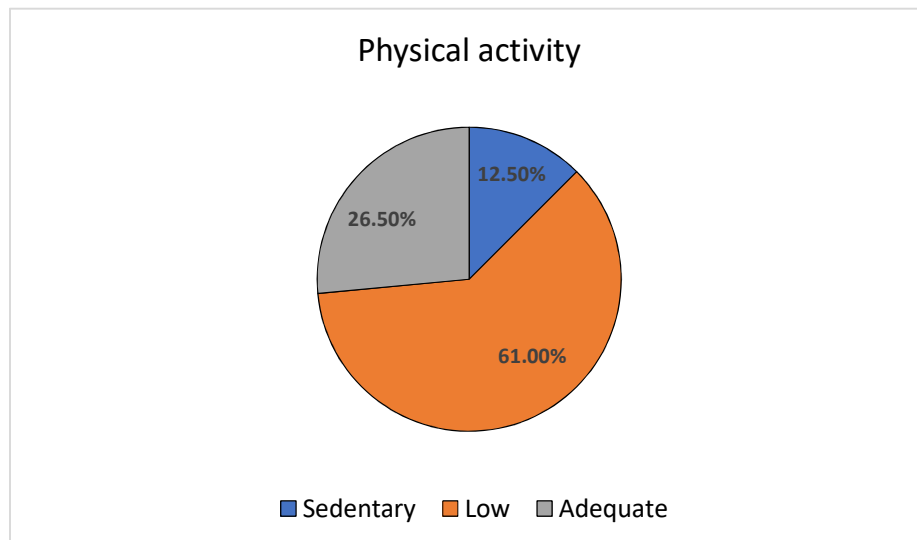
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***Physical activity***

Adequate amount of physical activity was followed by only 26.5% (n=106) of the study participants. Low amount of physical activity was reported in 61% (n=244) of the study participants. No physical activity or sedentary lifestyle was recorded in 12.5% (n=50) of the study participants (Table 11 and Figure 9).

**Table 11: Physical activity among study participants**

<b>Physical activity</b>	<b>Number (n=400)</b>	<b>Percentage</b>
Sedentary	50	12.5%
Low	244	61%
Adequate	106	26.5%



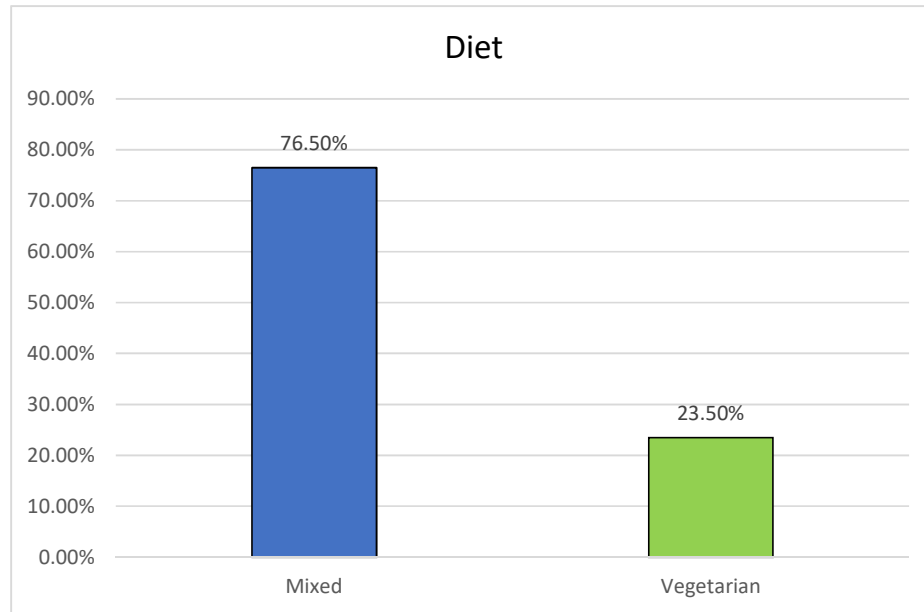
**Figure 9: Physical activity among study participants**

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*Diet of study participants*

Majority of the study participants had mixed diet (n=306, 76.5%). Only 23.5% (n=94) of the study participants followed vegetarian diet (Figure 10).



**Figure 10: Diet followed by the study participants**

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### ***Blood pressure and Heart rate***

Mean systolic blood pressure of the study participants was  $167.4 \pm 15.9$ . Mean diastolic blood pressure of the study participants was  $97.5 \pm 6.5$ . The mean heart rate of the study participants was  $89.8 \pm 7.4$ .

### ***Hypertension profile***

Among the study participants, 184 (46%) were newly diagnosed hypertension cases and the rest 216 (54%) were known cases of hypertension.

Among the study participants, only 49.5% (n=198) of them were under treatment for hypertension. Majority of the study participants had uncontrolled hypertension (n=314, 78.5%).

### ***Number of drugs taken for hypertension***

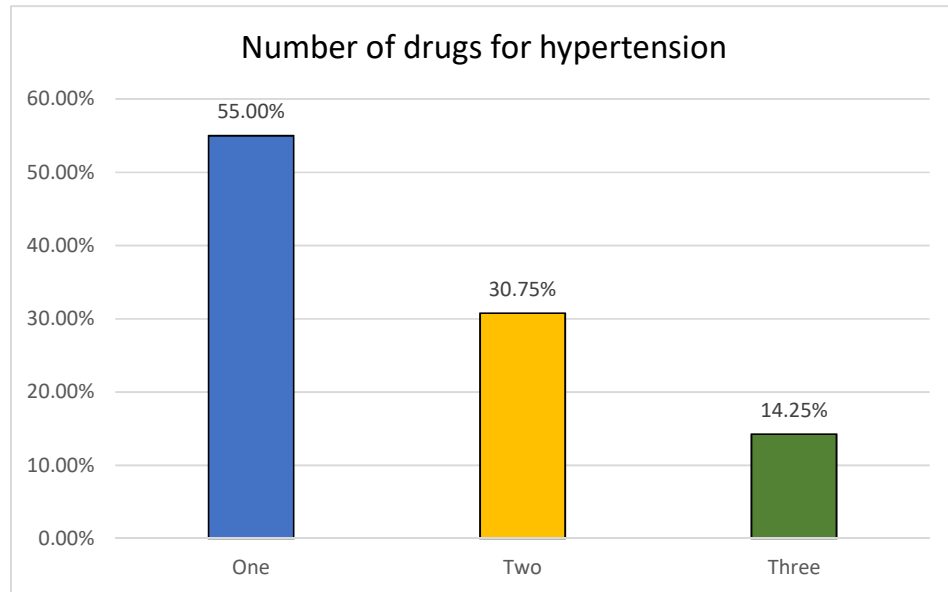
Majority of the study participants take single drug for hypertension (n=220, 55%). Two drugs for control of hypertension were taken by 30.75% (n=123) study participants. Three drugs for control of hypertension were taken by 14.25% (n=57) of the study participants (Table 12 and Figure 11).

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**Table 12: Number of drugs for control of hypertension**

Number of drugs	Number (n=400)	Percentage
1	220	55%
2	123	30.75%
3	57	14.25%



**Figure 11: Number of drugs taken for control of hypertension**

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***Lipid profile***

Mean value of total cholesterol, triglycerides, low-density lipoprotein and high-density lipoprotein of the study participants were  $244.4 \pm 44.1$ ,  $261.6 \pm 90.4$ ,  $150.7 \pm 44.5$ , and  $34.1 \pm 6.1$  respectively (Table 13).

**Table 13: Mean value of lipid profile**

<b>Lipid profile</b>	<b>Mean</b>	<b>Standard deviation</b>
Total cholesterol	244.4	44.1
Triglycerides	261.6	90.4
Low density lipoprotein	150.7	44.5
High density lipoprotein	34.1	6.1

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### *Serum Creatinine and Urea*

Mean value of serum creatinine and urea among the study participants were  $0.96 \pm 0.37$  and  $28.39 \pm 8.1$  respectively.

### *Urine protein*

Urine protein was not found in 40.25% (n=161) study participants. Trace protein was found in (1) 34.5% (n=138) study participants. Urine protein of 1+ was found in 25.25% (n=101) of the study participants (Figure 12).

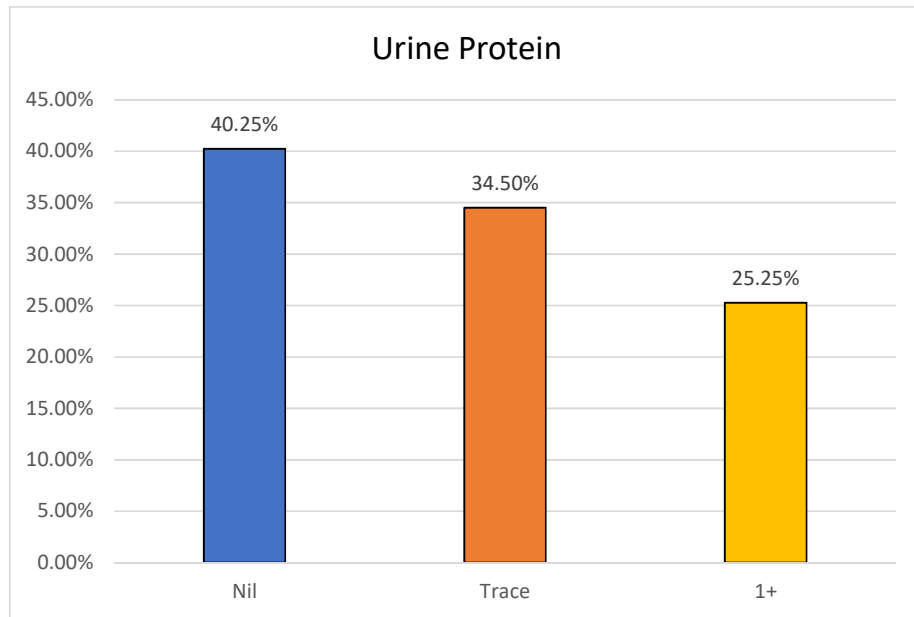


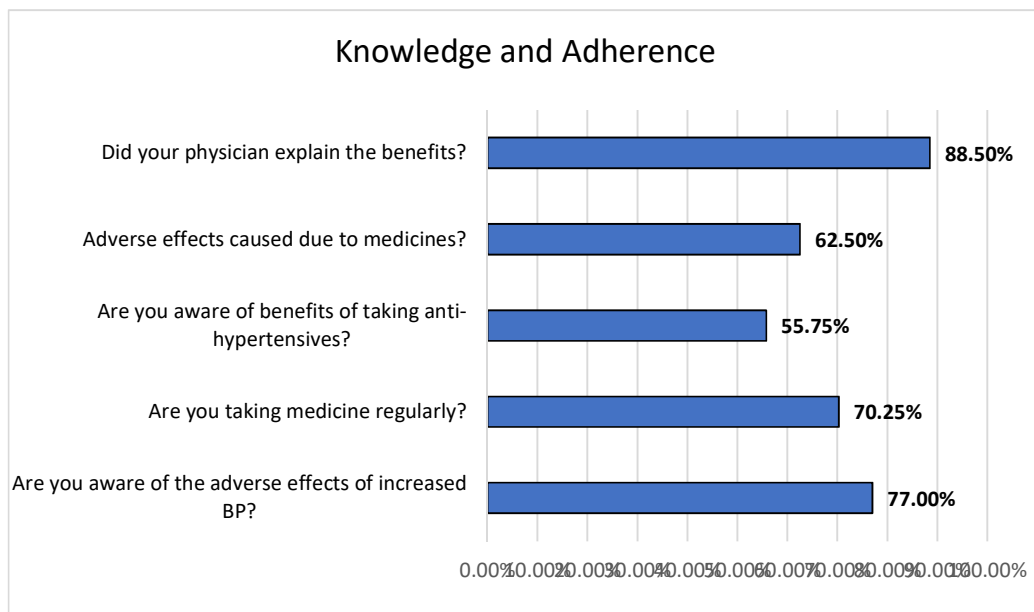
Figure 12: Urine protein among study participants

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***Knowledge and adherence of the study participants***

Among the study participants, knowledge and adherence questions were asked. Among 400 individuals, 77% (n=308) were aware of the adverse effects of increased blood pressure in village setup. Among the study participants, 70.25% (n=281) were taking their anti-hypertensive medications regularly. Among the study participants, 55.75% (n=223) of the study participants were aware of the benefits of taking anti-hypertensives and 62.5% (n=250) were aware of the adverse effects of them. Majority of their physicians (88.5%, n=354) explained the benefits of taking anti-hypertensives (Figure 13).



**Figure 13: Knowledge and adherence of study participants regarding hypertension and its medication**

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*Comparison of socio-demographic profile and control of hypertension*

On comparison of socio-demographic profile with controlled and uncontrolled hypertension groups, education and socio-economic status were significantly associated (p= 0.003 and 0.003 respectively) (Table 14).

**Table 14: Comparison of socio-demographic variables with control of hypertension**

<b>Variables</b>	<b>Controlled BP (n=86)</b>	<b>Uncontrolled BP (n=314)</b>	<b>P value</b>
<b>Age</b>			
18-30 Years	1 (1.2%)	6 (1.9%)	0.051
31-60 Years	47 (54.7%)	165 (52.5%)	
Above 60 years	38 (43.9%)	143 (45.5%)	
<b>Gender</b>			
Male	55 (64%)	204 (65%)	0.478
Female	31 (36%)	110 (35%)	
<b>Education</b>			
School	18 (20.9%)	27 (8.6%)	<b>0.003</b>
High school	34 (39.5%)	100 (31.8%)	
Diploma	19 (22.1%)	113 (36%)	
Undergrad	12 (14%)	63 (20.1%)	
Postgrad	3 (3.5%)	11 (3.5%)	
<b>Socioeconomic status</b>			
Upper	6 (7%)	9 (2.9%)	<b>0.003</b>
Middle	49 (57%)	236 (75.1%)	
Lower	31 (36%)	69 (22%)	

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*Education and locality*

Educational status was significantly associated with locality of the study participants (rural or urban) ( $p=0.0003$ ). Study participants who went to some school in rural and urban locality were 11.6% ( $n=27$ ) and 10.7% ( $n=18$ ) respectively. Study participants who went to high school and residing in rural and urban settings were 40.1% ( $n=93$ ) and 24.4% ( $n=41$ ) respectively. Study participants who went to high school and residing in rural and urban settings were 32.8% ( $n=76$ ) and 33.4% ( $n=56$ ) respectively. Individuals with under-graduation among the rural and urban settings were 14.2% ( $n=33$ ) and 25% ( $n=42$ ) respectively. Study participants who had completed post-graduation and residing in rural and urban settings were 1.3% ( $n=3$ ) and 6.5% ( $n=11$ ) respectively (Table 15).

**Table 15: Comparison of Education and locality**

<b>Education</b>	<b>Rural (n=232)</b>	<b>Urban (n=168)</b>	<b>P value</b>
School	27 (11.6%)	18 (10.7%)	0.0003
High school	93 (40.1%)	41 (24.4%)	
Diploma	76 (32.8%)	56 (33.4%)	
Undergrad	33 (14.2%)	42 (25%)	
Postgrad	3 (1.3%)	11 (6.5%)	

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*Comparison of personal habits, diet & physical activity and control of hypertension*

Personal habits like smoking and alcohol habits, salt intake, diet and physical activity were compared with controlled and uncontrolled blood pressure groups. There were no significant association with control of blood pressure (Table 16).

**Table 16: Comparison of personal habits, diet and physical activity of study participants**

<b>Variables</b>	<b>Controlled BP (n=86)</b>	<b>Uncontrolled BP (n=314)</b>	<b>P value</b>
<b>Salt intake</b>			
Low	48 (55.8%)	152 (48.4%)	0.137
High	38 (44.2%)	162 (51.6%)	
Smoking	38 (44.2%)	140 (44.6%)	0.523
Alcohol	23 (26.7%)	110 (35%)	0.093
<b>Physical activity</b>			
Sedentary	7 (8.1%)	43 (13.7%)	0.368
Low	54 (62.8%)	190 (60.5%)	
Adequate	25 (29.1%)	81 (25.8%)	
<b>Diet</b>			
Vegetarian	21 (24.4%)	73 (23.2%)	0.461
Mixed	65 (75.6%)	241 (76.8%)	

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*Comparison of BMI & Heart rate and control of hypertension*

BMI and heart rate was compared with controlled and uncontrolled blood pressure groups. Heart rate was significantly associated with control of blood pressure (p=0.000) (Table17).

**Table 17: Comparison of BMI and heart rate of study participants**

<b>Variables</b>	<b>Controlled BP (n=86)</b>	<b>Uncontrolled BP (n=314)</b>	<b>P value</b>
<b>BMI</b>			
Less than 18.5	1 (1.2%)	2 (0.6%)	0.434
18.5 to 22.9	32 (37.2%)	118 (37.6%)	
23 to 24.9	32 (37.2%)	132 (42%)	
25 to 29.9	21 (24.4%)	56 (17.8%)	
More than 30	0 (0%)	6 (1.9%)	
Heart Rate	87.74 ± 6.6	93.07 ± 7.3	<b>0.000</b>

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*Comparison of hypertension profile and control of hypertension*

Known or newly diagnosed hypertension, treatment of hypertension and number of drugs taken to control hypertension were compared with controlled and uncontrolled blood pressure groups. Known/ newly diagnosed hypertension, and treatment taken for hypertension were significantly associated with control of blood pressure in the study participants ( $p=0.003$ , and  $0.000$  respectively) (Table 18).

**Table 18: Comparison of hypertension profile**

<b>Variables</b>	<b>Controlled BP (n=86)</b>	<b>Uncontrolled BP (n=314)</b>	<b>P value</b>
<b>Hypertension</b>			
Newly diagnosed	28 (32.6%)	156 (49.7%)	<b>0.003</b>
Known HTN	58 (67.4%)	158 (50.3%)	
Treatment for HTN	57 (66.3%)	141 (44.9%)	<b>0.000</b>
<b>No. of drugs taken</b>			
1	49 (57%)	171 (54.5%)	0.321
2	29 (33.7%)	94 (29.9%)	
3	8 (9.3%)	49 (15.6%)	

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*Comparison of biochemical parameters and control of hypertension*

Total cholesterol, triglycerides, low-density lipoprotein, high-density lipoprotein, creatinine, urea and urine protein were compared with controlled and uncontrolled hypertension. Total cholesterol (p=0.021), urea (p=0.017) and urine protein (p=0.009) were significantly associated with control of blood pressure (Table 19).

**Table 19: Comparison of biochemical parameters and urine protein among study participants**

<b>Variables</b>	<b>Controlled BP (n=86)</b>	<b>Uncontrolled BP (n=314)</b>	<b>P value</b>
TC	241.79 ± 41.96	246.12 ± 45.33	<b>0.021</b>
TG	262.15 ± 88.91	261.27 ± 91.56	0.097
LDL	151.62 ± 43.87	150.22 ± 45.04	0.280
HDL	33.80 ± 5.83	34.31 ± 6.32	0.079
Creatinine	1.01 ± 0.37	0.96 ± 0.37	0.152
Urea	29.07 ± 8.23	27.95 ± 8.01	<b>0.017</b>
<b>Urine Protein</b>			
Nil	33 (38.4%)	128 (40.8%)	<b>0.009</b>
Trace	21 (24.4%)	117 (37.3%)	
1+	32 (37.2%)	69 (22%)	

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***Knowledge and Adherence compared with status of hypertension***

Awareness of adverse effects of blood pressure, regular intake of medication, awareness of benefit of intake of anti-hypertensive, awareness of adverse effects due to intake of medicines and explanation of adverse effects due to medications were not significantly associated with control of blood pressure (Table 20).

**Table 20: Knowledge and adherence of study participants and the control of hypertension**

<b>Variables</b>	<b>Controlled BP  (n=86)</b>	<b>Uncontrolled BP  (n=314)</b>	<b>P value</b>
Are you aware of the adverse effects of increased BP?	71 (82.6%)	237 (75.5%)	0.106
Are you taking medicine regularly?	59 (68.6%)	222 (70.7%)	0.400
Are you aware of benefits of taking anti-hypertensives?	45 (52.3%)	178 (56.7%)	0.274
Are you aware of adverse effects caused due to medicines?	50 (58.1%)	200 (63.7%)	0.206
Did your physician explain the benefits?	76 (88.4%)	278 (88.5%)	0.548

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## DISCUSSION

This cross-sectional study includes 400 study participants.

### *Prevalence of controlled hypertension*

In our study, only 21.5% of the study participants achieved target blood pressure. Few studies showed higher rates of achieving target blood pressure. In a study conducted in South Africa, 61.2% achieved target blood pressure (30). In a nationwide cohort study from United Kingdom, 52.8% of the study participants had achieved target blood pressure (31). In a study conducted in Singapore reported 49.7% achieved target blood pressure in their study (32). Target blood pressure rate of 39.54% was reported in a cross-sectional study conducted in Tamil Nadu, India (29). In an observational study conducted in India, the target blood pressure was attained in 25.3% of the study participants (32).

According to Eight Joint National Committee (JNC8), the normal blood pressure is less than 120 and 80 of systolic and diastolic blood pressure. Prehypertension is determined as either systolic blood pressure of 120-139 or diastolic blood pressure of 80-90. Stage 1 hypertension is considered as 140-159 of systolic blood pressure or 90-99 of diastolic blood pressure. Stage 2 hypertension is considered as either more than 160 of systolic blood pressure or more than 100 of diastolic blood pressure (9).

### *Age and gender*

Mean age of the study participants was 60.7 years Rayner B et al (30). Median age of the study participants was 71 (IQR 68 to 77) years in Todd et al (31). The mean

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age of study participants was  $57.12 \pm 12.23$  years in Anusha C et al (33). In our study, mean age of the study participants was  $58.97 \pm 12.83$  years.

Comparison of age between the study participants who had achieved blood pressure and who didn't achieve blood pressure was not significant in our study. Contradictory results were seen in other studies (29, 34).

In our study, 64.8% of the study participants were male. Majority of the study participants were male in (55.85%) Shanmugapriya et al study (29) and in Anusha C et al (Male: Female ratio of 2.06:1) (33). In Koh KH study, majority of the study participants were females (59.9%) (32). Similar results were found in Rayner B et al with 56.3% of female study participants (30).

Gender was not associated with target achievement of blood pressure in our study. Similar results were seen in other studies (29, 32).

### ***Education and socio-economic status***

Education and socio-economic status were significantly associated with control of blood pressure in our study. Among the study participants, 5.5%, 74.8% and 8.3% were illiterate, primary education and diploma/ undergrad or postgrad respectively. Educational status was significantly associated with control of blood pressure (34). Educational status was not statistically significant in control of hypertension in Shanmugapriya et al (29). Monthly family income was statistically significant in control of hypertension in Shanmugapriya et al (29).

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### ***Body mass index***

The mean BMI of the study participants in our study was  $23.52 \pm 2.38$ . Percentage of underweight, normal, overweight, pre-obese and obese in our study participants were 0.75%, 37.5%, 41%, 19.25% and 1.5% respectively. Normal BMI was reported in 25.1% and overweight was reported in 74.8% of the study participants (34). BMI of less than 23 was found in 14.1% and more than 23 was found in 85.9% of the study participants (32). Prevalence of underweight, normal, overweight and obese were found in 3.37%, 50.19%, 34.08% and 12.36% respectively (29). Mean BMI was  $28.2 \pm 5.15$  in Todd et al (31). BMI was not associated with control of hypertension in our study. Similar results were found in Shanmugapriya et al (29). Contradicting results were found in a few studies (32, 33, 34).

### ***Salt consumption and dietary habits***

Salt consumption in our study was similar in low and high intake category (n=200, 50% in each category). Salt intake was not significantly associated with control of hypertension in our study. Contradictory result was recorded in Shanmugapriya et al study (29).

Majority of the study participants had mixed diet (n=306, 76.5%). Only 23.5% (n=94) of the study participants followed vegetarian diet in our study. Diet was significantly associated with control of hypertension in our study. Diet was not significant in Shanmugapriya et al study (29).

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### *Personal habits*

Among the study participants, 44.5% (n=178) had smoking habit and 33.25% (n=133) had alcohol habits. Prevalence of smoking and alcohol were 5.6% and 1.8% (32). Prevalence of smoking was 15.7% in Rayner B et al (30) and 8.8% in Todd et al (31). Smoking and alcohol were prevalent in 22% and 24.7% of the study participants (33).

Smoking and alcohol were significantly not associated with control of hypertension in our study. Similar results, smoking or alcohol was not significantly associated in a cross-sectional study (32). Contradictory results were found in Shanmugapriya et al (29). None of the study participants who had a habit of smoking achieved target blood pressure in Anusha C et al (33).

Adequate amount of physical activity was followed by only 26.5% (n=106) of the study participants. Low amount of physical activity was reported in 61% (n=244) of the study participants. No physical activity or sedentary lifestyle was recorded in 12.5% (n=50) of the study participants. High IPAQ score was found in 65.85% of the study population and moderate IPAQ scores was reported in 32.77% of the study participants (29).

Physical activity was not significant in achieving target blood pressure in our study. In Shanmugapriya et al, physical activity was significantly associated with control of blood pressure (29).

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### *Antihypertensive drugs*

Majority of the study participants take single drug for hypertension (n=220, 55%). Two drugs for control of hypertension were taken by 30.75% (n=123) study participants. Three drugs for control of hypertension were taken by 14.25% (n=57) of the study participants in our study. In Kandasamy G et al, 60.9% of the study participants had taken single drug and 39.1% had taken combination of two or more drugs for hypertension (34). Among study participants, intake of one, two, three, and four drug combination were taken by 43.38%, 34.92%, 16.61% and 4.15% respectively (29). Among study participants, 30.7%, 42.8% and 26.5% were taking single, double and more than three drugs for control of hypertension (30). Average drugs taken by the study participants in controlled and uncontrolled blood pressure group were  $1.78 \pm 0.73$  and  $1.92 \pm 0.91$  respectively (33).

Intake of number of drugs was not significantly associated with control of blood pressure in our study. Similar result was also found in literature (33). Contradictory results were found in Shanmugapriya et al study (29).

Long-term research on antihypertensive medications show that monotherapy usually lowers the risk of CVD in a sizable number of patients while keeping a good safety record. Choosing the right monotherapy for different kinds of patients could be the main obstacle. The most sensible course of action is frequently to choose a drug that has the greatest potential to lower blood pressure while having the fewest negative effects on a certain patient (34).

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### **Biochemical parameters**

Mean value of total cholesterol, triglycerides, low-density lipoprotein and high-density lipoprotein of the study participants were  $244.4 \pm 44.1$ ,  $261.6 \pm 90.4$ ,  $150.7 \pm 44.5$ , and  $34.1 \pm 6.1$  respectively.

Total cholesterol was significantly associated with control of blood pressure in our study. Dyslipidemia was significantly associated in Koh KH et al (32) and not significantly associated with control of blood pressure in Anusha C et al (33).

Mean value of serum creatinine and urea among the study participants were  $0.96 \pm 0.37$  and  $28.39 \pm 8.1$  respectively.

Urea was significantly associated with control of blood pressure in our study. Urine protein was not found in 40.25% (n=161) study participants. Trace protein was found in (1) 34.5% (n=138) study participants. Urine protein of 1+ was found in 25.25% (n=101) of the study participants. There were not relevant studies to compare these results with to our knowledge.

### ***Knowledge and adherence***

Among the study participants, knowledge and adherence questions were asked.

Among 400 individuals, 77% (n=308) were aware of the adverse effects of increased blood pressure. Knowledge of adverse reactions was found in 8.6% of study participants (35).

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Among the study participants, 55.75% (n=223) of the study participants were aware of the benefits of taking anti-hypertensives and 62.5% (n=250) were aware of the adverse effects of them. Among the study participants, 70.25% (n=281) were taking their anti-hypertensive medications regularly. Only 28.4% of the individuals had taken medications regularly in the past two weeks (35). Among good knowledge individuals, majority were taking medications regularly (36).

There are few limitations in the current study. Prevalence of comorbid diseases, locality (urban or poor) and duration of hypertension were few important factors which were not included in the data collection.

Main strength of our study was, inclusion of education and socio-economic status.

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## CONCLUSION

In our study, only 21.5% of the study participants achieved target blood pressure. The mean age of the study participants was  $58.97 \pm 12.83$  years. Study variables significantly associated with control of hypertension were education, socio-economic status, heart rate, treatment of hypertension, newly diagnosed hypertension, total cholesterol, urea levels and urine protein in our study.

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**ANNEXURES-I**

**KAHERs JNMC BELAGAVI**

**INFORMED CONSENT FORM**

**Title Of Research Study: “A ONE YEAR CROSS SECTIONAL STUDY ON ATTAINMENT OF TARGET BLOOD PRESSURE IN HYPERTENSIVES AT KLE HOSPITAL BELAGAVI.**

**Principal Investigator:-**

**Introduction and Purpose:-** study of attainment of target blood pressure in both attained and non attained groups and further reduce morbidity and mortality associated

**Procedure:**

If you agree to be part of the research study, you will be asked the relevant history and will be subjected to relevant clinical examination and investigations. You will also have to give blood and urine samples for the necessary investigations.

**-Risk and Benefits:**

The only risk and possible discomfort you might get is while taking blood from your arm for the investigations. It may cause swelling, pain, redness (rarely happens) at the site from where the blood is drawn. You may not be benefitted by these investigations but you will be part of this study which is going to be useful to others in the future.

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**Alternatives:**

Taking part in this study is voluntary. You may choose not to take part in this study.

If you decide to take part you can later change your mind and withdraw from the study. Your decision will not change the present or future health care or other services that you receive. The study doctor or sponsor may stop your participation in this study at any time. If you choose not to take part in the study, you will receive the standard treatment for patients with your condition.

**Privacy and Confidentiality:**

All information collected about you during the course of this study will be kept confidential to the extent permitted by law. The code numbers will identify you in this research record. Information from this study may be published but your identity will be confidential in any publication.

**Institution / Sponsor's policy:**

Does not apply to this research

**Financial incentives for participation:**

You will not be paid / offered any gifts /incentives for participating in the study.

**Authorization to publish the results:**

The results of the study would be forwarded to the KLE University, Belgaum as part of requirement towards the completion of MD degree, review and publishing.

**In case of the queries during study or in future you may contact following persons,** Dr. HARSHA HEGDE Chairman, College Ethical Dissertation Research Committee J. N. Medical. College Nehru Nagar, Belagavi 590010.

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**ANNEXURES-II**

**A ONE YEAR CROSS SECTIONAL STUDY ON ATTAINMENT OF TARGET  
BLOOD PRESSURE IN HYPERTENSIVES AT KLE DR. PRABHAKAR  
KORE HOSPITAL, NEHRUNAGAR, BELAGAVI**

**Case Proforma**

<b>Name</b>	<b>Date</b>
<b>Age</b>	<b>Phone Number</b>
<b>Sex</b>	<b>IPD/OPD Number</b>

Area of residence: urban/rural/semi urban

Marital status:

Level of education: uneducated/Primary/Secondary/Graduation/Masters

Occupation:

Nature of employment:

Monthly family income:

Height:

Weight:

BMI:

**Comorbidities**

Diabetes Mellitus: Yes/No

Left ventricular heart failure: Yes/No

Right ventricular heart failure: Yes/No

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Asthma: Yes/No

Chronic kidney disease: Yes/No

Hyperlipidemia: Yes/No

Metabolic Syndrome: Yes/No

**Habit/diet history**

History of smoking

History of alcohol intake:

Diet: Vegetarian/Non-vegetarian

Salt intake:

Physical activity: no activity/ low /adequate

AHA guidelines	
Adequate	At least 150 minutes of moderate intense aerobic activity per week (approx. 30 min brisk walk for 5 days)
Low physical activity	Less than 150 of intense activity. Eg: Leisure activity, light swimming, housework etc.
No physical activity	(Sedentary lifestyle)

**Details related to hypertension**

Hypertension: previously diagnosed/ newly diagnosed

Treatment for hypertension: yes/no

Hypertension control at baseline: yes/no

Antihypertensive medication at baseline: (drug name and dose)

Number of drugs

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Medication adherence

**Blood Pressure**

Systolic BP

Diastolic BP

Hear rate

Blood pressure categories based on JNC 8.

Normal	<120/<80mmHg
Prehypertension	1220-139/80-89 mmHg
Stage 1 Hypertension	140-159/90-99 mmHg
Stage 2 Hypertension	≥160/≥100 mmHg

Please note: the above table is just for reference. While collecting data please add the systolic and diastolic BP values separately in the excel sheet so that it will be easier for categorization and analysis.

**Laboratory Investigations**

LDL

HDL

TG

VLDL

Renal function tests findings:

Routine urine and Microscopy findings

**Knowledge of Hypertension and related effects**

1. Are you aware of adverse effects of increased blood pressure related to heart, kidney and stroke?
2. Are you taking your medications regularly?
3. Are you aware of benefits of taking antihypertensive medication?
4. Are you aware of adverse effects caused due to medication non-compliance?

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5. Did your physician explain the benefits of antihypertensive medication and effects of medication non-compliance?

### ANNEXURE –III MASTER CHART

Age	Age_Cat	Gender	Height	WEIGHT	BMI	SALT INTAKE	Smoking	Alcohol	PA	Diet	SBP	DBP	HR	HR	HTN	Treatment of HTN	CONTROLLED HTN	BORDERLINE HTN	Medication	No. of Drugs	Medication adherence	LDL	HDL	TRIGLYCERIDE	TOTAL CHOLESTROL	S-CREATININE	UREA	Urine Protein	Aware of adverse effects of BP	Medication regularly	Aware of benefits of taking antihypertensive	Adverse effect caused due to medicines	Physician explain about benefits of medication
50	2	1	154	57	24	1	2	2	2	2	150	96		88	2	1	2	YES	AMLODIPINE -2.5 mg	1	YES	60	27	120	102	0.5	13	0	YES	YES	YES	NO	YES
67	3	1	173	74	24.7	1	1	1	2	2	150	100		80	1	2	2	YES	AMLODIPINE -5 mg	1	NO	176	39	241	274	0.7	34	1	YES	YES	YES	NO	YES
71	3	1	153	48	20.5	1	1	2	2	2	176	104		98	2	1	2	NO	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	YES	180	48	351	246	1.4	38	1	NO	NO	NO	NO	YES
62	3	2	170	78	27	1	2	2	2	2	180	100		84	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	150	38	204	287	0.9	12	0	NO	YES	NO	YES	YES
45	2	1	166	57	20.7	1	1	2	2	2	154	98		84	2	1	1	YES	NIFEDIPINE -15 mg	1	YES	166	48	199	215	1.4	29	1+	YES	YES	YES	NO	YES
47	2	1	170	75	26	1	1	1	2	2	180	106		100	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	NO	240	49	500	300	1.4	27	1	YES	YES	YES	YES	YES
60	2	2	178	70	22.1	1	2	2	3	2	166	94	0	84	2	2	2	NO	TELMISARTAN -40mg	1	YES	145	40	200	217	1.4	29	1+	YES	NO	YES	YES	YES
71	3	1	168	65	23	1	1	1	3	2	150	90		84	1	1	2	YES	AMLODIPINE -2.5 mg	1	NO	175	39	195	287	0.5	39	0	YES	NO	YES	YES	YES
56	2	2	158	59	23.6	2	2	2	2	2	160	98		82	1	2	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	YES	150	41	462	250	1.2	29	1+	NO	NO	NO	YES	NO

83	3	2	171	75	25.6	1	1	1	1	2	178	104			100	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	NO	149	32	210	108	1.5	27	0	YES	YES	YES	YES	YES
27	1	2	150	48	21.3	1	2	2	1	2	140	90			80	1	2	1	YES	AMLODIPINE -2.5 mg	1	NO	90	30	241	254	1.1	10	0	YES	YES	NO	NO	YES
57	2	1	166	74	26.8	2	1	1	3	1	146	90			86	2	1	1	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	YES	116	45	211	265	1.4	19	1	NO	NO	NO	NO	NO
52	2	1	170	59	20.4	1	2	1	3	2	156	110			84	2	1	2	YES	TELMISARTAN+HYDR OCHLOROTHIAZIDE (	2	NO	120	36	330	230	0.7	17	0	YES	YES	YES	YES	YES
78	3	1	159	57	22.5	1	1	2	2	2	160	104			84	1	2	2	NO	TELMISARTAN+HYDR OCHLOROTHIAZIDE (	2	NO	202	47	196	256	1	11	1	NO	NO	NO	NO	YES
82	3	1	174	70	23.1	2	1	1	2	2	170	100			90	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	NO	170	39	250	254	0.9	15	0	NO	NO	NO	NO	NO
68	3	1	157	68	27.5	1	1	2	2	1	156	98			100	2	1	1	YES	TELMISARTAN+HYDR OCHLOROTHIAZIDE (	2	YES	160	28	196	203	1.2	19	1+	YES	YES	YES	YES	YES
84	3	1	160	60	23.4	2	1	1	2	2	162	94			92	2	2	2	YES	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	NO	150	38	323	214	0.9	12	0	NO	YES	NO	YES	YES
78	3	1	172	69	23.3	1	1	2	2	2	180	100			82	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	NO	175	34	248	234	0.9	25	0	YES	NO	YES	NO	YES
45	2	2	154	56	23.6	1	2	2	3	1	156	96			90	2	1	1	YES	TELMISARTAN - 20mg	1	YES	174	29	210	298	1.4	29	1	YES	YES	YES	YES	YES
24	1	2	159	57	22.5	1	2	2	3	2	150	90			80	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	150	40	352	300	1.4	35	0	YES	YES	YES	YES	YES
47	2	2	163	68	25.6	2	2	2	3	2	190	110			100	2	1	2	NO	AMLODIPINE+ATENOL OL (5mg/50mg)	2	NO	180	39	245	245	1.5	38	1	NO	YES	NO	NO	YES
59	2	1	180	80	24.6	2	2	2	1	1	154	96			80	1	2	2	YES	TELMISARTAN - 20mg	1	YES	105	28	354	256	0.3	35	0	NO	YES	NO	NO	YES
65	3	1	150	50	22.2	1	1	1	2	2	186	110			98	2	1	2	NO	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	NO	98	29	129	200	0.9	13	0	YES	YES	YES	NO	YES
46	2	1	178	79	24.9	2	1	2	2	2	150	90			88	1	2	2	YES	TELMISARTAN + AMLODIPINE ( 40	2	NO	220	48	410	300	0.7	25	1	YES	YES	YES	YES	YES

mg/5 mg )																																		
57	2	1	154	54	22.7	1	2	2	2	2	152	98			80	2	2	2	YES	TELMISARTAN - 20mg	1	YES	97	28	198	210	1.4	19	0	YES	YES	NO	NO	YES
51	2	2	155	57	23.7	2	2	2	2	2	154	96			86	2	1	1	YES	NIFEDIPINE -10 mg	1	NO	110	34	215	247	0.9	35	1	YES	NO	NO	NO	YES
55	2	1	174	69	22.7	1	1	1	2	2	188	100			94	2	1	2	NO	NIFEDIPINE -20 mg	1	YES	174	27	245	297	1.2	29	0	YES	YES	YES	YES	YES
72	3	2	152	71	30.7	2	2	2	2	1	166	96			82	2	1	2	NO	CLONIDINE -0.1mg twice a day to thrice a day	1	NO	150	29	259	245	0.8	35	1+	NO	YES	NO	NO	YES
51	2	2	151	49	21.4	2	2	2	3	1	154	94			90	1	2	2	YES	TELMISARTAN - 20mg	1	NO	93	28	240	280	0.6	15	1	YES	NO	NO	NO	YES
71	3	1	169	76	26.6	2	1	2	1	2	170	100			80	2	1	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	90	35	195	275	0.9	11	1+	NO	YES	NO	NO	YES
65	3	1	168	75	26.5	1	1	1	2	2	144	90			86	2	1	1	YES	TELMISARTAN+HYDR OCHLOROTHIAZIDE ( 40mg/12.5 mg)	2	YES	175	39	350	300	0.5	19	0	YES	YES	YES	YES	YES
63	3	2	158	59	23.6	2	2	2	3	1	160	98			84	1	2	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	NO	150	50	400	289	1.1	29	1+	NO	NO	NO	NO	NO
46	2	1	164	65	24.1	2	1	2	3	1	162	98			100	1	2	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	YES	180	40	243	278	0.8	27	0	NO	NO	NO	NO	YES
40	2	1	151	90	39.4	1	2	2	3	2	154	88			84	1	2	2	YES	TELMISARTAN -40mg	1	YES	174	29	250	264	0.6	15	0	YES	YES	YES	YES	YES
32	2	1	149	46	20.7	1	2	2	1	2	154	98			80	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	152	34	299	245	0.9	37	1	NO	NO	NO	NO	NO
74	3	2	158	63	25.2	2	2	2	2	1	166	102			82	1	2	2	NO	METOPROLOL +TELMISARTAN (25mg/40mg)	1	NO	167	36	222	256	0.9	15	0	NO	NO	NO	NO	NO
58	2	2	167	69	24.7	1	2	2	2	2	170	90			84	2	1	2	NO	TELMISARTAN - 20mg	1	NO	154	29	241	239	1.4	12	1+	YES	YES	YES	YES	NO
19	1	2	151	47	20.6	1	2	2	3	2	166	90			86	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	150	39	254	245	0.3	35	1	YES	YES	NO	YES	YES
38	2	1	163	49	18.4	2	1	1	1	1	150	90			88	1	2	2	YES	TELMISARTAN - 20mg	1	NO	250	36	451	287	0.6	38	0	YES	YES	NO	NO	YES
75	3	1	159	57	22.5	2	2	1	2	2	184	94			88	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	163	40	195	248	0.7	12	1+	YES	YES	YES	YES	YES
75	3	2	152	51	22	1	1	1	2	2	180	98			92	2	1	1	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	148	35	222	236	0.7	31	1	NO	NO	NO	NO	YES
45	2	2	178	79	24.9	2	2	1	2	1	174	92			82	1	2	2	YES	METOPROLOL +TELMISARTAN (25mg/40mg)	1	YES	147	37	245	289	0.3	25	0	YES	YES	YES	YES	YES
68	3	1	157	55	22.3	1	1	1	2	2	150	90			88	1	2	2	YES	METOPROLOL	1	NO	151	31	233	256	1.4	19	0	NO	NO	NO	NO	NO

65	3	2	172	57	19.2	2	2	2	2	2	178	100			92	1	2	2	YES	+TELMISARTAN (25mg/40mg) TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIAZIDE (40mg/5mg/12.5)	3	NO	147	29	210	263	0.3	33	0	YES	NO	NO	NO	YES
70	3	1	166	65	23.5	1	1	2	2	2	174	98			96	2	1	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIAZIDE (40mg/5mg/12.5)	1	YES	170	39	241	126	1.4	39	1	YES	NO	YES	YES	YES
60	2	1	165	71	26	1	2	2	1	1	178	101			82	2	1	2	NO	TELMISARTAN -40mg	1	NO	180	36	320	290	1.4	12	1+	NO	YES	NO	YES	YES
52	2	2	150	50	22.2	1	2	2	2	1	152	90			80	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	80	41	350	200	0.9	37	1	YES	YES	YES	NO	YES
74	3	1	171	70	23.9	2	1	1	2	2	176	94			82	2	1	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	NO	65	28	159	198	0.3	35	1	NO	YES	YES	NO	YES
53	2	1	167	65	23.3	1	1	2	2	2	156	90			80	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	89	29	170	300	0.7	12	1+	YES	YES	YES	YES	YES
87	3	1	178	80	25.2	2	1	1	2	2	180	110			94	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIAZIDE (40mg/5mg/12.5)	3	NO	165	39	365	287	1.5	38	1+	NO	NO	YES	YES	YES
56	2	1	155	55	22.8	1	1	1	2	2	160	100			90	2	2	2	YES	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	NO	150	35	279	300	0.7	12	1+	YES	YES	NO	NO	YES
67	3	2	150	50	22.2	1	2	2	2	2	174	96			88	2	1	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIAZIDE (40mg/5mg/12.5)	1	YES	166	28	299	271	1.4	12	1+	NO	NO	NO	NO	NO
56	2	1	168	57	20.2	2	1	2	2	2	166	92			82	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	130	27	385	278	0.3	35	1	YES	YES	YES	NO	YES
47	2	1	170	72	24.9	2	1	1	2	2	154	92			86	1	2	2	YES	TELMISARTAN - 40mg	1	NO	245	32	187	224	0.5	32	1+	YES	YES	NO	YES	YES
50	2	2	150	48	21.3	1	2	2	2	1	160	98			94	1	2	2	YES	TELMISARTAN + AMLODIPINE ( 20 mg/5 mg )	2	YES	90	36	201	180	0.9	12	1	YES	YES	NO	YES	YES
46	2	1	175	70	22.8	2	1	1	2	2	174	100			92	2	1	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	NO	99	27	176	245	1.4	39	0	YES	YES	NO	NO	YES
64	3	1	152	51	22	1	2	2	2	1	158	88			90	1	2	2	YES	TELMISARTAN -20mg	1	YES	174	36	245	290	0.5	28	1	NO	YES	NO	YES	YES
61	3	1	178	76	23.9	2	2	2	2	2	170	100			92	2	1	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	NO	99	39	241	183	0.9	11	1	YES	YES	NO	YES	YES
50	2	1	168	78	27.6	2	1	1	3	2	158	90			88	1	2	2	YES	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	100	26	299	300	0.5	19	0	YES	NO	NO	YES	YES
44	2	2	161	60	23.1	1	2	2	3	2	176	100			98	2	1	2	YES	TELMISARTAN -40mg	1	YES	174	29	250	297	0.9	35	1	YES	NO	NO	YES	YES

71	3	1	170	75	25.9	1	2	2	2	2	166	100			86	2	1	1	YES	AMLODIPINE -2.5 mg	1	NO	152	32	199	245	1.2	29	0	YES	YES	YES	YES	NO
86	3	1	182	81	24.4	2	1	1	2	2	140	90			80	1	2	2	YES	METOPROLOL +TELMISARTAN (25mg/40mg)	1	YES	167	36	222	210	0.8	35	1+	NO	YES	NO	NO	YES
75	3	1	169	59	20.6	1	1	2	2	2	170	102			98	1	2	2	NO	TELMISARTAN - 20mg	1	NO	154	29	241	102	0.7	34	1	YES	NO	YES	YES	NO
40	2	2	153	50	21.3	1	2	2	2	1	146	90			84	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	101	28	230	150	0.3	35	1	YES	YES	YES	YES	NO
87	3	1	152	51	22	1	1	1	2	2	178	98			94	2	1	1	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	NO	148	35	222	199	0.7	31	1	NO	NO	NO	NO	YES
70	3	1	161	61	23.5	1	1	1	2	2	174	94			94	2	1	1	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	69	27	179	245	1.4	19	0	NO	NO	NO	NO	YES
39	2	1	155	55	22.8	2	2	2	2	2	190	100			98	2	1	2	NO	TELMISARTAN - 40mg	1	NO	106	27	190	275	0.7	39	0	NO	YES	YES	YES	YES
80	3	2	150	48	21.3	1	2	2	2	2	184	98			82	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	158	39	350	250	1.4	29	0	NO	YES	NO	YES	YES
60	2	1	175	74	24.1	2	1	1	3	2	156	92			84	1	2	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	YES	181	35	200	256	0.7	25	1	YES	YES	YES	YES	YES
63	3	1	165	65	23.8	2	2	2	3	1	154	88			90	1	2	2	YES	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	241	49	485	300	1.4	19	0	NO	NO	NO	NO	NO
63	3	1	153	50	21.3	1	1	1	2	1	180	106			94	2	1	2	NO	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	YES	195	34	204	254	0.9	35	1	YES	YES	YES	YES	YES
60	2	2	169	65	22.7	1	2	2	2	2	160	100			86	1	2	2	YES	NIFEDIPINE – 15 mg	1	NO	166	44	199	215	0.7	25	1	YES	YES	NO	NO	YES
59	2	1	173	57	19	1	1	2	1	2	178	104			88	1	2	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	NO	120	28	204	289	1.5	38	1	YES	YES	YES	YES	YES
68	3	1	170	68	23.5	2	2	2	2	2	166	94			86	1	2	2	YES	TELMISARTAN - 20mg	1	YES	119	28	187	215	0.3	25	1	YES	NO	NO	YES	YES
54	2	2	166	65	23.5	2	2	2	3	1	152	100			96	2	1	1	YES	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	NO	95	32	150	200	0.3	19	1	YES	YES	NO	NO	NO
63	3	1	178	79	24.9	2	2	1	2	1	174	92			82	1	2	2	NO	METOPROLOL +TELMISARTAN (25mg/40mg)	1	YES	147	37	245	289	0.3	25	0	YES	YES	YES	YES	YES
88	3	1	157	55	22.3	1	1	1	2	2	150	90			88	1	2	2	YES	METOPROLOL +TELMISARTAN (25mg/40mg)	1	NO	151	31	233	182	1.4	19	0	NO	NO	NO	NO	NO

53	2	1	172	57	19.2	2	2	2	2	2	180	100			92	1	2	2	NO	TELMISARTAN + AMLODIPINE ( 20 mg/5 mg )	2	YES	147	29	210	178	0.3	33	1	YES	YES	YES	NO	YES
56	2	1	166	65	23.5	1	2	2	2	2	174	94			86	1	2	2	NO	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	NO	95	50	200	249	1.4	19	1+	YES	YES	YES	YES	NO
56	2	1	173	74	24.7	2	1	1	2	2	156	90			84	2	1	1	YES	TELMISARTAN -40mg	1	YES	99	27	247	240	0.4	29	1+	YES	YES	YES	NO	NO
76	3	1	151	49	21.4	1	2	2	2	2	180	110			98	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	179	37	250	264	1.2	29	0	YES	YES	YES	YES	YES
47	2	2	161	60	23.1	1	2	2	1	2	174	100			100	2	1	2	NO	TELMISARTAN -40mg	1	YES	174	29	250	279	0.9	35	1	YES	NO	NO	YES	YES
70	3	1	170	75	25.9	1	2	2	2	2	168	98			86	2	1	1	YES	AMLODIPINE -2.5 mg	1	YES	240	27	199	245	1.2	29	0	YES	YES	YES	YES	NO
39	2	1	151	60	26.3	1	2	2	3	2	154	90			84	1	2	2	YES	TELMISARTAN -40mg	1	YES	174	29	250	187	0.6	15	0	YES	YES	YES	YES	YES
83	3	1	182	80	24.1	2	1	2	1	2	140	98			80	1	2	2	YES	METOPROLOL +TELMISARTAN (25mg/40mg)	1	YES	174	36	250	160	0.8	35	1+	NO	YES	YES	NO	YES
42	2	1	153	54	23	1	1	2	2	1	200	110			98	2	2	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	NO	101	28	230	265	0.3	25	1	YES	NO	YES	YES	NO
67	3	2	166	70	25.4	1	2	2	3	1	166	96	0		88	1	2	2	YES	AMLODIPINE+ATEN OLOL (5mg/50mg)	1	NO	150	29	200	245	1.4	29	1+	YES	NO	YES	YES	YES
53	2	1	164	67	24.9	1	1	2	2	2	152	100			88	2	1	1	YES	AMLODIPINE -5 mg	1	YES	98	39	178	260	0.5	19	0	YES	YES	YES	NO	NO
56	2	1	151	52	22.8	2	2	2	3	2	156	90			90	1	2	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	NO	103	50	210	274	0.6	38	0	YES	NO	NO	NO	YES
53	2	1	160	60	23.4	2	1	2	2	1	170	90			98	1	2	2	NO	TELMISARTAN -40mg	1	YES	90	28	200	275	0.9	31	1+	NO	YES	NO	YES	YES
60	2	1	178	74	23.3	2	1	1	3	2	170	96			98	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	117	27	254	223	1.4	29	1+	YES	YES	NO	NO	YES
49	2	1	180	85	26.2	2	1	1	3	2	140	90			80	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	119	32	145	228	0.9	19	1	NO	NO	NO	NO	NO
66	3	2	161	79	30.4	2	2	2	2	1	170	100			98	1	2	2	NO	AMLODIPINE - 5 mg	1	NO	110	31	201	290	0.8	11	1	YES	YES	YES	YES	YES
65	3	1	178	85	26.8	2	1	1	3	2	174	100			98	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	110	28	150	245	1.1	11	1	NO	NO	NO	YES	YES
30	1	1	174	70	23.1	2	1	1	3	2	150	92			88	1	2	2	YES	TELMISARTAN - 40mg	1	NO	100	30	158	224	0.5	35	1+	YES	YES	YES	YES	YES
47	2	2	162	59	22.4	1	2	2	2	1	162	98			98	2	1	1	YES	TELMISARTAN + AMLODIPINE ( 20 mg/5 mg )	2	NO	240	29	185	246	0.3	12	1	YES	YES	NO	NO	YES
70	3	1	175	70	22.8	2	1	1	2	2	174	100			96	2	1	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	189	27	176	245	1.4	39	0	YES	YES	NO	NO	YES
60	2	2	155	57	23.7	1	2	2	2	1	190	102			88	2	1	2	NO	TELMISARTAN - 40mg	1	NO	106	24	190	245	0.6	39	1	YES	YES	YES	YES	YES
40	2	2	160	61	23.8	1	2	2	2	2	172	100			86	2	1	2	NO	TELMISARTAN +	3	YES	150	28	205	233	1.4	29	1	YES	YES	YES	NO	YES



48	2	1	153	54	23	1	1	1	1	2	162	92			98	1	2	2	YES	TELMISARTAN - 20mg	1	NO	87	36	210	245	0.3	25	0	YES	YES	YES	YES	YES
81	3	1	166	70	25.4	1	1	2	2	2	190	108	0		86	2	1	2	NO	AMLODIPINE+ATEN LOL (5mg/50mg)	2	YES	150	29	244	178	0.7	29	0	YES	YES	YES	YES	YES
59	2	1	165	60	22	1	1	1	2	2	170	102			96	1	2	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	107	34	245	312	0.8	37	1	NO	NO	NO	NO	YES
65	3	2	162	64	24.3	1	2	2	3	2	152	94	0		84	1	2	1	YES	TELMISARTAN -40mg	1	YES	165	35	211	288	1.4	39	1	NO	YES	NO	YES	YES
48	2	1	178	69	21.7	1	1	2	3	2	150	90			80	1	2	1	YES	TELMISARTAN -20mg	1	YES	142	34	224	173	1.4	30	1+	YES	YES	YES	YES	YES
72	3	2	162	61	23.2	1	2	2	1	2	180	110			96	2	1	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	120	33	204	289	1.5	38	0	NO	NO	NO	YES	YES
71	3	2	154	57	24	2	2	2	2	1	160	98			86	1	2	2	YES	TELMISARTAN - 20mg	1	NO	99	28	178	300	0.3	25	0	YES	YES	NO	YES	YES
55	2	1	165	68	24.9	1	1	1	3	2	178	108			92	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	NO	137	35	500	147	0.7	35	1	YES	YES	YES	NO	YES
62	3	2	152	53	22.9	2	2	2	3	2	162	96			88	2	2	1	YES	METOPROLOL +TELMISARTAN (25mg/40mg)	1	YES	136	28	201	270	0.3	32	1	NO	YES	YES	YES	YES
51	2	2	160	57	22.2	1	2	2	3	2	154	88			86	1	2	1	YES	TELMISARTAN - 20mg	1	NO	78	50	201	247	1.4	29	0	YES	YES	NO	NO	YES
58	2	1	173	57	19	2	1	1	3	2	178	104			90	1	2	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	120	28	120	289	1.5	38	1	YES	YES	YES	YES	YES
57	2	1	170	74	25.6	1	2	2	1	2	162	94			82	2	1	1	YES	TELMISARTAN - 40mg	1	YES	119	29	187	300	0.3	35	1	YES	NO	YES	YES	YES
55	2	1	152	51	22	2	2	1	2	2	150	88			94	1	2	2	YES	TELMISARTAN - 20mg	1	YES	111	32	222	299	0.7	28	1	YES	YES	YES	NO	YES
73	3	1	178	79	24.9	1	2	2	2	1	174	92			80	2	1	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	NO	147	34	241	172	0.7	37	0	YES	YES	YES	YES	YES
46	2	1	180	80	24.6	2	1	1	2	2	166	96			92	2	1	1	YES	TELMISARTAN + AMLODIPINE ( 20 mg/5 mg )	2	YES	152	26	211	269	0.3	33	0	YES	NO	NO	NO	YES
54	2	1	166	65	23.5	1	2	2	2	2	174	90			88	1	2	2	NO	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	NO	95	39	470	281	1.4	19	1+	YES	YES	YES	YES	NO
30	1	1	173	71	23.7	2	1	1	2	2	168	98			92	2	1	2	NO	TELMISARTAN -40mg	1	YES	170	33	247	249	0.4	39	0	YES	YES	YES	NO	YES
70	3	1	178	76	23.9	2	1	2	2	2	180	108	0		100	1	2	2	NO	AMLODIPINE+ATEN LOL (5mg/50mg)	2	YES	241	48	250	233	1.2	29	0	YES	YES	YES	YES	YES
48	2	2	154	57	24	1	2	2	2	2	166	98			84	1	2	2	YES	TELMISARTAN -40mg	1	NO	175	27	241	248	0.9	35	0	YES	NO	YES	YES	YES
56	2	2	150	47	20.8	1	2	2	2	1	150	90			82	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	140	28	174	245	1.2	32	1+	YES	YES	NO	YES	YES
65	3	2	164	60	22.3	1	2	2	2	1	154	90			80	1	2	2	YES	TELMISARTAN -20mg	1	NO	144	24	262	236	0.9	15	0	YES	YES	NO	YES	NO
64	3	1	180	82	25.3	2	1	1	3	2	150	90			82	1	1	1	YES	AMLODIPINE -2.5 mg	1	YES	68	24	204	152	0.8	35	0	YES	YES	YES	YES	YES
53	2	1	168	67	23.7	1	1	1	3	2	200	110			100	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE	3	NO	180	40	250	263	1.4	39	1+	NO	YES	NO	YES	YES

56	2	2	163	67	25.2	1	2	2	1	2	152	96	0	88	1	2	1	YES	(40mg/5mg/12.5) AMLODIPINE+ATEN OLOL (5mg/50mg)	2	NO	150	30	241	123	0.9	29	0	YES	NO	YES	YES	YES
45	2	1	164	60	22.3	2	1	1	3	2	174	106		98	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	102	35	178	261	0.5	28	1	YES	NO	NO	YES	YES
55	2	1	174	75	24.7	2	2	2	2	2	182	104		92	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	200	32	419	277	0.6	39	1	YES	YES	YES	YES	YES
60	2	2	160	60	23.4	2	2	2	3	2	170	106		100	2	1	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	178	39	369	262	0.9	40	0	NO	NO	YES	NO	YES
51	2	1	176	74	23.8	2	1	1	2	2	184	108		100	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	225	39	195	287	0.5	39	0	YES	NO	YES	YES	YES
59	2	1	158	59	23.6	2	2	2	3	1	160	94		82	1	2	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	NO	150	32	452	249	1.1	29	0	YES	YES	YES	YES	YES
54	2	1	171	80	27.3	1	2	2	3	1	180	106		102	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	180	40	250	301	1.5	40	1+	YES	YES	YES	YES	YES
79	3	2	155	52	21.6	2	2	2	3	1	170	100		88	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	NO	102	32	180	299	1.1	21	1	YES	YES	YES	YES	YES
58	2	1	153	50	21.3	1	1	1	3	2	180	96		86	2	1	2	NO	AMLODIPINE -2.5 mg	1	YES	156	32	187	240	0.9	13	0	YES	YES	YES	YES	YES
45	2	1	163	65	24.4	2	1	1	1	2	196	108		98	2	1	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	179	37	250	108	1.2	29	0	YES	YES	YES	YES	YES
75	3	2	153	45	19.2	1	2	2	2	1	150	90		80	2	1	1	YES	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	NO	97	27	248	190	0.8	37	1+	YES	YES	NO	NO	YES
68	3	2	178	70	22	1	2	2	2	2	166	96	0	88	2	2	2	YES	TELMISARTAN -40mg	1	NO	150	40	200	193	1.4	29	1+	YES	NO	YES	YES	YES
43	2	1	178	74	23.3	2	1	1	2	2	170	98		98	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	117	27	254	248	1.4	29	1+	YES	YES	NO	NO	YES
37	2	2	157	53	21.5	2	2	2	2	2	182	104		100	2	1	2	NO	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	NO	180	40	243	279	1.4	19	0	YES	NO	YES	NO	YES
64	3	1	158	57	22.8	2	1	2	1	2	170	104		98	1	2	2	NO	PRAZOSIN (2.5 mg) twice a day to thrice a	1	YES	71	39	209	246	1	29	1+	YES	YES	YES	YES	YES







72	3	2	151	87	38.1	1	2	2	2	2	178	104			96	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	178	35	470	222	1.5	35	0	YES	NO	YES	YES	YES	YES
48	2	2	149	47	21.1	1	2	2	2	2	156	90			80	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	147	35	197	275	0.9	37	1	NO	NO	YES	YES	YES	
72	3	2	162	61	23.2	2	2	2	2	2	172	106			96	2	1	2	NO	METOPROLOL +TELMISARTAN (25mg/40mg)	1	NO	167	36	445	256	0.9	15	0	NO	NO	NO	NO	NO	
60	2	2	167	69	24.7	1	2	2	2	2	170	90			84	2	2	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	NO	154	36	241	230	1.4	12	0	YES	YES	YES	YES	YES	
56	2	1	159	59	23.3	2	1	1	3	2	166	98			86	1	2	2	YES	AMLODIPINE -5 mg	1	YES	147	39	299	288	0.3	35	1+	YES	YES	NO	YES	YES	
60	2	1	174	78	25.7	2	1	2	3	2	182	108			96	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	177	32	241	182	0.9	27	0	YES	NO	YES	NO	YES	
70	3	2	152	47	20.3	2	2	2	1	2	174	98			84	2	1	2	NO	METOPROLOL +TELMISARTAN (25mg/40mg)	2	YES	145	38	163	223	1.4	29	1	YES	YES	YES	YES	YES	
36	2	1	174	74	24.4	1	1	1	2	2	154	92			82	1	2	2	YES	TELMISARTAN -40mg	1	YES	144	29	488	262	0.6	34	0	YES	YES	NO	NO	YES	
76	3	1	182	85	25.6	2	1	1	3	2	200	110			100	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	180	36	250	200	0.8	37	1+	YES	YES	NO	YES	YES	
56	2	1	169	67	23.4	1	1	1	1	2	190	108			100	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	90	28	201	210	0.9	31	1	YES	YES	NO	YES	YES	
53	2	2	152	47	20.3	2	2	2	2	1	170	98	0		102	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	1	NO	150	29	418	345	1.4	29	1+	YES	NO	YES	YES	YES	
76	3	1	164	66	24.5	2	1	1	3	2	178	100			98	2	1	2	NO	NIFEDIPINE – 20 mg	1	YES	174	39	241	199	0.5	29	0	NO	NO	NO	NO	NO	
72	3	1	157	54	21.9	1	2	2	2	1	150	90			82	1	2	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	YES	98	34	210	287	0.6	38	1+	YES	YES	YES	YES	YES	
78	3	2	169	70	24.5	1	2	2	2	2	174	98			82	2	2	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	173	39	240	187	1.4	39	1	YES	NO	YES	NO	YES	
41	2	2	149	45	20.2	1	2	2	2	2	162	90			84	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	120	39	195	287	0.5	39	1	YES	YES	YES	YES	YES	
48	2	1	167	59	21.1	2	1	1	3	2	150	94			82	1	2	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	YES	163	35	250	122	1.2	35	1+	NO	YES	YES	YES	YES	
47	2	2	158	60	24	2	2	2	2	2	202	102			96	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	179	35	210	280	1.5	37	0	YES	YES	YES	YES	YES	

50	2	1	178	85	26.8	1	2	2	1	2	182	108			98	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	102	41	145	245	1.1	31	1	YES	YES	YES	YES	YES
68	3	1	174	70	23.1	2	1	1	3	2	150	92			88	1	2	1	YES	AMLODIPINE -5 mg	1	NO	103	32	180	264	0.6	35	1+	YES	YES	YES	YES	YES
56	2	1	162	65	24.7	1	1	2	3	2	178	104			104	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	90	29	198	287	0.7	32	0	YES	NO	NO	YES	YES
40	2	2	159	47	18.5	1	2	2	2	2	160	98			100	2	1	1	YES	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	103	35	482	245	1.4	40	1	YES	YES	YES	NO	YES
65	3	1	155	61	25.3	2	1	2	2	2	180	106			102	2	1	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	110	38	411	220	0.6	39	0	YES	YES	YES	YES	YES
67	3	1	160	65	25.3	2	1	1	2	2	170	104			98	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	154	32	210	239	1.1	33	1+	NO	NO	NO	NO	YES
42	2	1	176	78	25.1	2	2	2	2	1	200	110			98	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (80mg/5mg/12.5)	3	YES	179	41	210	274	1.2	39	0	YES	YES	NO	YES	YES
55	2	1	155	56	23.3	2	1	1	2	2	170	108			82	1	2	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	NO	107	26	175	365	0.8	30	0	YES	NO	NO	YES	YES
51	2	1	170	75	25.9	1	1	1	3	2	170	98	0		84	2	2	2	NO	TELMISARTAN -40mg	1	YES	150	40	473	274	1.4	29	1+	YES	NO	YES	YES	YES
72	3	2	159	53	20.9	1	2	2	3	1	152	98			98	1	2	2	YES	TELMISARTAN - 20mg	1	YES	174	39	254	269	1.4	30	0	YES	YES	YES	YES	YES
65	3	2	158	59	23.6	1	2	2	1	2	160	94			82	1	2	2	YES	TELMISARTAN -20mg	1	YES	142	34	211	175	1.4	34	1+	YES	YES	YES	YES	YES
64	3	1	162	66	25.1	1	1	1	2	2	174	104			96	2	1	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	NO	140	33	241	249	1.5	38	1	YES	YES	YES	YES	YES
71	3	1	154	57	24	2	2	2	2	2	166	92			82	1	2	2	YES	TELMISARTAN - 40mg	1	YES	130	33	199	301	0.3	26	1+	YES	NO	YES	NO	YES
62	3	1	178	80	25.2	1	2	2	3	1	174	108			96	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (80mg/5mg/12.5)	3	YES	65	26	121	120	0.3	25	0	YES	YES	YES	YES	YES
71	3	2	157	55	22.3	1	1	1	2	2	152	90			84	1	2	2	YES	METOPROLOL +TELMISARTAN (25mg/40mg)	1	NO	151	33	422	284	1.7	19	0	YES	YES	YES	YES	NO
77	3	1	172	71	24	2	1	1	3	2	180	102			106	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	NO	147	33	380	298	0.3	33	0	YES	NO	YES	NO	YES
54	2	2	166	65	23.5	1	2	2	1	2	174	100			96	2	1	2	NO	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA	1	YES	140	39	229	160	1.2	22	0	NO	NO	NO	NO	YES

57	2	2	148	44	20	1	2	2	2	1	152	90			80	1	2	1	YES	ZIDE (40mg/5mg/12.5) TELMISARTAN -20mg	1	NO	90	30	166	274	0.4	25	0	YES	YES	NO	NO	YES
75	3	1	168	74	26.2	1	1	1	2	2	180	110			104	2	2	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	179	37	250	226	1.2	29	0	YES	YES	YES	YES	YES
44	2	1	161	60	23.1	1	2	2	2	2	174	108			102	2	1	2	NO	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	YES	240	47	490	290	0.9	34	0	YES	YES	NO	YES	YES
44	2	1	159	57	22.5	1	2	2	2	2	160	92			84	1	2	2	YES	TELMISARTAN+HYDR OCHLOROTHIAZIDE ( 40mg/12.5 mg )	2	NO	109	32	196	223	1	21	0	YES	YES	YES	YES	YES
41	2	1	174	78	25.7	1	2	1	3	2	170	102			98	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	120	28	214	210	0.8	16	1	YES	YES	YES	YES	YES
77	3	1	166	68	24.6	2	1	1	2	2	156	98			84	1	2	2	YES	TELMISARTAN+HYDR OCHLOROTHIAZIDE ( 40mg/12.5 mg )	2	YES	120	30	199	298	1	29	0	NO	NO	NO	NO	YES
63	3	1	160	66	25.7	1	1	2	3	2	154	92			98	2	1	1	YES	AMLODIPINE+ATENOL OL (5mg/50mg)	2	YES	162	33	207	174	0.9	22	1	YES	YES	NO	NO	YES
55	2	1	172	70	23.6	2	2	2	1	2	180	106			98	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	NO	175	38	248	228	0.9	35	1+	YES	YES	YES	YES	YES
78	3	1	177	78	24.9	1	1	2	2	2	180	110			86	2	1	2	NO	TELMISARTAN - 40mg	1	YES	180	33	210	270	1	29	1	YES	YES	YES	YES	NO
55	2	1	159	59	23.3	1	2	2	3	2	150	90			80	1	2	2	YES	AMLODIPINE -2.5 mg	1	YES	150	29	411	301	1.4	22	1	YES	NO	NO	NO	YES
51	2	1	163	66	24.8	2	1	1	2	2	190	110			104	2	1	2	NO	AMLODIPINE+ATENOL OL (5mg/50mg)	2	NO	180	26	133	299	1.5	38	1	YES	YES	NO	YES	YES
50	2	2	152	50	21.6	2	2	2	1	1	154	96			88	1	2	2	YES	TELMISARTAN - 20mg	1	YES	111	28	174	220	0.8	35	1+	YES	YES	YES	YES	YES
66	3	1	150	47	20.8	1	1	1	2	2	148	88			86	1	2	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	NO	147	28	150	200	0.9	15	0	YES	YES	YES	YES	YES
50	2	1	178	81	25.5	2	1	1	2	2	150	90			80	1	2	2	YES	TELMISARTAN + AMLODIPINE ( 40 mg/5 mg )	2	NO	106	26	165	254	0.7	35	1	YES	YES	YES	YES	YES
68	3	1	165	67	24.6	2	1	1	3	2	170	104			98	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	108	29	229	274	1.4	28	0	YES	YES	YES	YES	YES
52	2	2	155	55	22.8	2	2	2	2	2	154	96			86	1	2	2	YES	NIFEDIPINE - 10 mg	1	YES	111	33	222	133	1.1	35	0	YES	YES	YES	NO	YES







72	3	1	174	69	22.7	2	1	1	2	2	188	92			88	2	1	1	NO	ZIDE (40mg/5mg/12.5) TELMISARTAN + AMLODIPINE (40 mg/5 mg)	2	NO	198	35	259	236	1.2	39	1+	YES	NO	NO	YES	YES
71	3	1	167	62	22.2	1	1	2	2	2	204	106			98	2	1	1	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	NO	149	27	188	298	0.6	24	1+	NO	NO	NO	NO	YES
60	2	1	163	67	25.2	1	1	1	3	2	170	104			96	2	1	1	NO	METOPROLOL +TELMISARTAN (25mg/40mg)	2	YES	110	27	401	187	1.4	31	0	YES	NO	NO	NO	YES
63	3	1	172	73	24.6	2	1	1	1	2	156	90			90	1	2	2	YES	TELMISARTAN - 40mg	1	NO	180	32	246	258	1.4	27	1	YES	YES	YES	YES	YES
54	2	1	174	77	25.4	2	2	1	2	2	182	100			90	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	170	37	250	241	0.9	25	0	NO	NO	NO	NO	NO
78	3	1	162	66	25.1	1	1	1	2	1	150	92			88	1	2	2	YES	AMLODIPINE -5 mg	1	YES	120	32	154	245	1.2	28	0	YES	YES	YES	YES	YES
71	3	1	160	67	26.1	1	1	2	2	2	204	110			102	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	172	38	245	260	1.4	38	1+	YES	YES	YES	YES	YES
53	2	1	167	62	22.2	1	2	2	2	2	180	104			100	2	1	1	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	NO	103	32	180	240	0.9	25	0	YES	YES	YES	NO	NO
60	2	1	178	80	25.2	2	1	1	2	2	150	88			88	1	2	2	YES	TELMISARTAN - 20mg	1	NO	220	30	258	245	1.4	29	0	NO	NO	NO	NO	YES
46	2	1	159	61	24.1	1	2	2	1	2	150	92			82	2	1	1	YES	AMLODIPINE -2.5 mg	1	YES	142	28	200	267	1.4	40	0	YES	YES	YES	YES	YES
56	2	1	174	70	23.1	2	1	1	2	2	170	100			90	2	1	2	YES	CLONIDINE -(0.1mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	YES	110	26	180	200	0.9	25	0	YES	YES	YES	YES	YES
58	2	2	153	52	22.2	1	2	2	2	1	156	88			82	1	2	1	YES	AMLODIPINE -2.5 mg	1	YES	120	29	187	256	0.5	28	1+	YES	NO	YES	YES	YES
72	3	1	174	78	25.7	2	1	1	3	2	174	108			98	2	1	1	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	NO	130	32	215	220	0.9	25	0	YES	YES	YES	YES	YES
65	3	1	168	66	23.3	2	1	1	2	1	158	92			82	1	2	2	YES	TELMISARTAN - 40mg	1	YES	126	35	220	274	1.3	30	1+	NO	NO	NO	NO	NO
72	3	1	160	60	23.4	2	1	1	2	2	204	106			100	2	1	2	NO	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	180	38	245	259	1.4	39	0	YES	YES	YES	YES	YES
37	2	1	173	72	24	1	2	2	3	2	140	90			82	1	2	1	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN +	1	YES	91	27	339	205	0.3	10	0	YES	YES	YES	YES	YES



70	3	2	158	57	22.8	2	2	2	2	2	178	98			100	1	2	2	NO	TELMISARTAN + AMLODIPINE (40 mg/5 mg)	2	YES	190	50	289	187	1.4	34	1	YES	YES	YES	YES	YES	YES
67	3	2	160	62	24.2	1	2	2	3	2	160	94			90	2	1	1	YES	AMLODIPINE -5 mg	1	YES	199	48	369	244	1.1	30	0	YES	YES	NO	YES	YES	
50	2	2	147	41	18.9	2	2	2	2	1	152	92			82	1	2	2	YES	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	NO	229	47	293	241	0.5	29	1	NO	NO	NO	NO	YES	
52	2	2	155	52	21.6	2	2	2	2	1	150	90			80	1	2	2	YES	NIFEDIPINE - 10 mg	1	NO	113	30	170	210	1.1	32	1	NO	NO	NO	NO	YES	
48	2	1	174	70	23.1	1	1	1	2	2	188	106			100	2	1	2	NO	NIFEDIPINE - 30 mg	1	YES	166	36	245	150	1.2	29	0	YES	NO	NO	NO	YES	
80	3	1	161	62	23.9	2	1	1	2	2	168	94			90	2	1	2	YES	TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	3	YES	172	32	241	266	0.8	35	0	YES	YES	YES	YES	YES	
38	2	1	180	80	24.6	2	1	1	1	2	178	104			106	2	1	1	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	125	39	320	271	1.1	29	1+	YES	YES	NO	YES	YES	
65	3	2	152	51	22	1	2	2	2	2	150	94			82	1	2	1	YES	TELMISARTAN -40mg	1	YES	110	28	271	289	0.4	21	0	YES	YES	YES	YES	YES	
77	3	1	140	48	24.4	2	1	1	2	2	196	105			100	2	1	1	NO	TELMISARTAN + AMLODIPINE (40 mg/5 mg)	2	NO	210	38	293	247	1.2	29	0	YES	YES	YES	YES	YES	
60	2	1	168	64	22.6	1	2	2	2	2	192	100			82	2	1	1	NO	TELMISARTAN + AMLODIPINE (40 mg/5 mg)	2	YES	149	36	241	229	0.8	37	1+	YES	YES	NO	NO	YES	
88	3	1	180	70	21.6	1	2	2	2	1	140	90	0		80	1	2	2	YES	TELMISARTAN -40mg	1	NO	148	28	229	247	0.8	17	1+	YES	YES	NO	YES	YES	
49	2	1	140	44	22.4	2	2	2	3	2	188	98			86	2	1	2	NO	TELMISARTAN + AMLODIPINE (40 mg/5 mg)	2	NO	145	27	247	300	1.2	38	1+	YES	NO	NO	YES	YES	
56	2	1	171	62	21.2	1	1	2	2	1	164	92			80	2	1	1	YES	AMLODIPINE -5 mg	1	YES	249	48	419	204	0.8	37	1+	YES	YES	NO	YES	YES	
67	3	2	149	40	18	1	2	2	2	1	148	88	0		88	1	2	2	YES	TELMISARTAN -20mg	1	YES	112	31	203	223	0.7	29	1+	YES	YES	NO	YES	YES	
68	3	2	155	50	20.8	2	2	2	2	2	152	92			84	2	1	1	YES	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	NO	210	40	360	223	1.4	29	1+	YES	YES	NO	NO	YES	
53	2	1	180	78	24	1	2	2	1	2	168	92			80	1	2	2	YES	TELMISARTAN + AMLODIPINE (20 mg/5 mg)	2	YES	120	49	422	276	1.5	38	1	YES	YES	YES	YES	YES	
39	2	1	170	77	26.6	2	1	1	2	2	178	96			86	2	2	2	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	199	38	443	284	1.1	34	1	YES	YES	YES	YES	YES	
67	3	2	162	60	22.8	1	2	2	2	2	188	98			94	2	1	1	NO	AMLODIPINE+ATEN OLOL (5mg/50mg)	2	YES	114	28	247	298	1.1	30	1+	YES	NO	NO	NO	YES	
50	2	1	172	71	24	2	1	2	3	2	174	98			82	1	2	2	NO	PRAZOSIN (2.5 mg) twice a day to thrice a day + TELMISARTAN + AMLODIPINE +HYDROCHLOROTHIA ZIDE (40mg/5mg/12.5)	1	YES	176	33	233	248	1.4	38	1	YES	YES	NO	YES	YES	
61	3	2	152	57	24.6	1	2	2	2	2	152	90			88	1	2	1	YES	TELMISARTAN - 40mg	1	NO	250	35	407	293	0.7	17	0	YES	YES	YES	YES	YES	
65	3	1	169	61	21.3	2	2	2	2	2	170	98			100	2	1	2	NO	TELMISARTAN+HYDR	2	YES	215	29	240	285	1	33	1	YES	YES	YES	NO	YES	



