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Analysis of Prescribing Pattern among Cardiovascular Patients at National Institute of Cardiovascular Disease, Dhaka, Bangladesh

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Authors' contributions

This work was carried out in collaboration between all authors. Author MSI designed and supervised the study. Authors IAHS and LR performed the laboratory tests and prepared the manuscript. Author AAH revised the final manuscript. All authors read and approved the final manuscript.

Article Information

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Original Research Article

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ABSTRACT

Objective: The aim of this study was to comprehend the prescribing pattern among cardiovascular patients at the national institute of Cardiovascular Disease (NICVD), Dhaka, Bangladesh. **Methodology:** The current study was carried out in a national institute of Cardiovascular Disease (NICVD), Dhaka, Bangladesh from 13th February, 2017 to 2nd December, 2017. Cross-sectional types of descriptive research were performed in the cardiac outpatient department through the period of study. A total of 1,000 patients meeting the inclusion and exclusion criteria, were interviewed with the structured questionnaire. The questionnaire includes the demographic details of the patients, prescribing trend and the written permission was also taken from administrative offices as well as patients in this hospital.

Results: The findings of this study suggest that out of 1,000 patients, only 13.1% of patients were

used monotherapy, whereas 86.9% of patients were taken combination therapy for the treatment of CVD. Among the combination therapy, penta therapy was the most popular. **Conclusions:** Drug prescription pattern in our study observed that combination therapy was more commonly prescribed than monotherapy.

Keywords: Prescribing pattern; cardiovascular patients; CVD; national institute of cardiovascular disease.

1. INTRODUCTION

CVD continues to be the most common cause of mortality all over the world, in the 2013 Global Burden of Disease (GBD) study claimed that 17.3 million deaths were caused by CVD. It is responsible for 31.5% of all deaths and 45% of all non-infectious disease mortality worldwide, and two times higher than that caused by cancer, furthermore greater than totally communicable, maternal, neonatal and nutritional disorders combined. The 2013 GBD also observed that CVD caused a higher figure of deaths and was accountable for a greater number of all deaths than in 1990 when 12.3 million deaths were attributed to CVD, corresponding to 25.9% of to overall deaths [1]. The economic influence of various types of CVD is huge. Conventionally, Bangladesh is experienced with infectious diseases. Similar to many other low-income countries in the global, she has been suffering from epidemiological transition; the predominating disease pattern is converting from infectious diseases to non-infectious diseases. [2-3].

CVD is accounted for 75% of the deaths happening in developing countries like Bangladesh [4]. Studying the present situation, it has been expected that at least 25 million individuals will face death because of CVDs by 2030 [5-6]. Whereas the prevalence and death because of CHD is decreasing in the developed countries but it is not happening in lower-income countries.

There has been a horrific rise in the past twenty years in the prevalence of CHD and cardiovascular mortality in India and other south Asian nations [7]. The incidence of cardiovascular diseases has been rising in the new decade [8].

In the majority of nations, the absolute number of mortalities from CVD is rising due to rising life expectancy and related population longevity and deaths of individuals over than 70 years of age [9]. If the account is taken of population longevity, cardiovascular mortality rates have decreased worldwide by 16% between 2000 and 2012 [10]. A trend is driven by decreasing in smoking cigarette, population-level blood pressure improvements and best treatment for CVD. Declines are more in higher –income countries than in low- and middle – income countries [11-12].

Recently, important medicines to treat CVD are accessible, e.g. pharmacological treatment of high low-density lipoprotein (LDL) cholesterol levels increased blood pressure and inhibiting platelet function with statins, anti-hypertensive agents (thiazides, beta blockers, calcium channel blockers, angiotensin-converting enzyme (ACE) inhibitors, or angiotensin II receptor blockers (ARBs), etc. and antiplatelet agents e.g. Low dose aspirin, respectively [13-14].

The objective of this study was to understand the prevalence of cardiovascular disease and the different prescribing patterns of the cardiovascular drug in tertiary care hospital in Bangladesh.

2. METHODS

2.1 Participants and Study Site

The current study was carried out in a National Institute of Cardiovascular Disease (NICVD), Dhaka, Bangladesh from 13th February, 2017 to 2nd December, 2017. Types of the descriptive study were performed in the cardiac outpatient department through the period of study a total of 1,000 patients who meets the criteria (inclusion and exclusion criteria) were interviewed with the structured questionnaire. The main target of the study was to find out the prevalence of CVD and several patterns among 1,000 patients with CVD diseases. The questionnaire includes the demographic details of the patients. The written consent was approved by administrative offices in the hospital.

2.2 Statistical Analysis

The statistical calculation was carried out on collected and compiled data with the help of Microsoft Excel 2010 (Roselle, IL, USA).

3. RESULTS

Prescribing status of cardiovascular drugs among 1,000 patients with CVD is given in Table 1. From the 1,000 patients, 26.3% of patients were taken penta therapy whereas 13.1% of patients were used monotherapy for the treatment of CVD. The outcomes of the cardiovascular drug prescribed with monotherapy are shown in Table 2. The proportion of patients who took bisoprolol as a monotherapy was the highest while the lowest was azilsartan medoxomil, ramipril, prazosin and carvedilol used as a monotherapy. The figures were 29.77% and 0.77% respectively. 28.20% of patients were administered bisoprolol + atorvastatin as a dual therapy and 11.11% of patients were taken clopidogrel + atorvastatin as a dual therapy for the treatment of CVD. The consequences of the cardiovascular drug prescribed with dual therapy are presented in Table 3.

The percentage of patients who used clopidogrel + atorvastatin + glyceryl trinitrate (nitroglycerine) as a triple therapy was the maximum but the Clopidogrel+Atorvastatine+ minimum was Azilsartan Medoxomil used as a triple therapy. percentages were 17.77 and The 0.4 successively. The results of the cardiovascular drug prescribed with triple therapy are displayed in Table 4. The percentage of patients who were administered atorvastatin + bisoprolol + ramipril + glyceryl trinitrate (nitroglycerine) as a quadruple therapy and used glyceryl trinitrate (nitroglycerine) + propranolol hydrochloride + frusemide + spironolactone was the same. The percentages were 1.93. The outcomes of the cardiovascular drug prescribed with quadruple therapy are demonstrated in Table 5. 52.47% of patients used clopidogrel + bisoprolol + atorvastatin + glyceryl trinitrate(nitroglycerine) + trimetazidine dihydrochloride as a penta therapy whereas 2.28% of patients administered clopidogrel + rosuvastatin + bisoprolol + trinitrate (nitroglycerine) alvcervl azisartanmedoxomil as a penta therapy for the management of CVD. The results of the cardiovascular drug prescribed with penta therapy are demonstrated in Table 6.

Status	Frequency	Percent	
Monotherapy	131	13.1%	
Dual Therapy	117	11.7%	
Triple Therapy	135	13.5%	
Quadruple Therapy	155	15.5%	
Penta Therapy	263	26.3%	
Hexa Therapy	117	11.7%	
Hepta Therapy	67	6.7%	
Octa Therapy	15	1.5%	

Table 2. 0	Cardiovascular	drug prescribed	with monotherapy
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Prescribed Medicines	Frequency	Percent
Bisoprolol	39	29.77%
Losartan Potassium	3	2.29%
Propranolol Hydrochloride	36	27.49%
Metoprolol Tartrate	5	3.82%
Atorvastatine	26	19.84%
Azilsartan Medoxomil	1	0.77%
Rosuvastatin	7	5.34%
Trimetazidine Dihydrochloride	2	1.52%
Ramopril	1	0.77%
Prazosin	1	0.77%
Atenolol	2	1.52%
Carvedilol	1	0.77%
Glyceryl Trinitrate (nitroglycerine)	4	3.05%
OlmesartanMedoxomil	3	2.29%

Prescribed Medicines	Frequency	Percent
Bisoprolol+Atorvastatine	33	28.20%
Amlodipine+Olmesartan Medoxomil	3	2.57%
Clopidogrel+Atorvastatine	13	11.11%
Atorvastatine+Losartan Potassium	6	5.12%
TrimetazidineDihydrochloride+Losartan Potassium	1	0.85%
Clopidogrel+Bisoprolol	5	4.28%
Frusemide+Spironolactone	6	5.12%
Bisoprolol+Losartan Potassium	7	5.98%
Clopidogrel+Rosuvastatin	9	7.69%
Ramopril+Carvedilol	1	0.85%
Propranolol Hydrochloride +Atorvastatine	1	0.85%
Bisoprolol+OlmesartanMedoxomil	5	4.28%
Rosuvastatin+Bisoprolol	9	7.69%
Bisoprolol+AzilsartanMedoxomil	1	0.85%
Atorvastatine+GlycerylTrinitrate (nitroglycerine)	1	0.85%
Rosuvastatin+Losartan Potassium	2	1.70%
Bisoprolol +Hydrochlorothiazide	1	0.85%
Hydrochlorothiazide+OlmesartanMedoxomil	2	1.70%
Rosuvastatin+Metoprolol	1	0.85%
OlmesartanMedoxomil+Diltiazem Hydrochloride	1	0.85%
Rosuvastatin+AzilsartanMedoxomil	1	0.85%
Glyceryl Trinitrate (nitroglycerine+Propranolol	1	0.85%
Hydrochloride		
Amlodipine+Metoprolol	1	0.85%
Bisoprolol+Ramopril	1	0.85%
Atorvastatine +OlmesartanMedoxomil	1	0.85%
TrimetazidineDihydrochloride+Ramopril	2	1.70%
Glyceryl Trinitrate (nitroglycerine+Clopidogrel	2	1.70%

Table 3. Cardiovascular drug prescribed with dual therapy

Table 4. Cardiovascular drug prescribed with triple therapy

Prescribed Medicines	Frequency	Percent
Clopidogrel+ Atorvastatine+ Diltiazem Hydrochloride	4	2.96%
Atorvastatine+Bisoprolol+GlycerylTrinitrate (nitroglycerine)	9	6.67%
Atorvastatine+TrimetazidineDihydrochloride+Ramipril	1	0.74%
Clopidogrel+Bisoprolol+GlyceryITrinitrate (nitroglycerine)	3	2.22%
Clopidogrel+Trinitrate (nitroglycerine)+TrimetazidineDihydrochloride	1	0.74%
Clopidogrel+Atorvastatine+GlycerylTrinitrate (nitroglycerine)	24	17.77%
Clopidogrel+Atorvastatine+Cilnidipine	1	0.74%
Bisoprolol+ Atorvastatine+OlmesartanMedoxomil	5	3.70%
Atorvastatine+Amlodipine+Atenolol	1	0.74%
Atorvastatine+Bisoprolol+Losartan Potassium	5	3.70%
TrimetazidineDihydrochloride+Ramipril+Carvedilol	1	0.74%
Atorvastatine+Clopidogrel+Bisoprolol	19	14.08%
Clopidogrel+Bisoprolol+Rosuvastatin	9	6.67%
Clopidogrel+Rosuvastatin+Glyceryl Trinitrate (nitroglycerine)	3	2.22%
Bisoprolol+LosartanPotassium+GlycerylTrinitrate (nitroglycerine)	1	0.74%
Bisoprolol+Atorvastatine+TrimetazidineDihydrochloride	5	3.70%
Rosuvastatin+Bisoprolol+AzilsartanMedoxomil	1	0.74%
Rosuvastatin+LosartanPotassium+Carvedilol	2	1.48%
Clopidogrel+Atorvastatine+OlmesartanMedoxomil	1	0.74%
Clopidogrel+Atorvastatine+TrimetazidineDihydrochloride	2	1.48%
TrimetazidineDihydrochloride+Frusemide+Spironolactone	1	0.74%
Bisoprolol+OlmesartanMedoxomil+Rosuvastatin	2	1.48%

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Prescribed Medicines	Frequency	Percent
Clopidogrel+Atorvastatine+AzilsartanMedoxomil	4	0.4%
Clopidogrel+Atorvastatine+Verapamil Hydrochloride	1	0.74%
Bisoprolol+Atorvastatine+AzilsartanMedoxomil	3	2.22%
Ramopril+GlycerylTrinitrate (nitroglycerine)+Propranolol Hydrochloride	4	2.97%
Atorvastatine+PropranololHydrochloride+Losartan Potassium	1	0.74%
Carvedilol+Frusemide+Spironolactone	1	0.74%
Clopidogrel+GlycerylTrinitrate (nitroglycerine)+OlmesartanMedoxomil	1	0.74%
Clopidogrel+Bisoprolol+OlmesartanMedoxomil	1	0.74%
Bisoprolol+Frusemide+Spironolactone	2	1.48%
Clopidogrel+Atorvastatine+Metoprolol Tartrate	1	0.74%
Glyceryl Trinitrate (nitroglycerine)+Rosuvastatine+Metoprolol Tartrate	3	2.22%
Bisoprolol+GlycerylTrinitrate (nitroglycerine)+OlmesartanMedoxomil	1	0.74%
Bisoprolol+Amlodipine + OlmesartanMedoxomil	1	0.74%
Bisoprolol+Atorvastatine+Losartan Potassium	2	1.48%
Clopidogrel+Rosuvastatin+Losartan Potassium	1	0.74%
Propranolol Hydrochloride+Frusemide+Spironolactone	1	0.74%
TrimetazidineDihydrochloride+GlycerylTrinitrate	1	0.74%
(nitroglycerine)+Atorvastatine		
Diltiazem Hydrochloride+Amlodipine+ OlmesartanMedoxomil	1	0.74%
Clopidogrel+Atorvastatine+Propranolol Hydrochloride	2	1.48%
Glyceryl Trinitrate (nitroglycerine)+Atorvastatine+Losartan Potassium	1	0.74%
Atorvastatine+Frusemide+Spironolactone	1	0.74%

26.49% of patients were used clopidogrel + atorvastatin + glyceryl trinitrate (nitroglycerine) + trimetazidinedihydrochloride bisoprolol+losartan potassium as a hexa therapy while 7.69% of patients were administered clopidogrel + rosuvastatin + glyceryl trinitrate (nitroglycerine) + trimetazidinedihydrochloride + frusemide + spironolactone for the management of CVD. The results of the cardiovascular drug prescribed with hexa therapy are presented in Table 7. 34.32% of patients were taken clopidogrel + atorvastatin + bisoprolol + glyceryl trinitrate (nitroglycerine) trimetazidinedihydrochloride + frusemide+ spironolactone as a hepta therapy whereas 7.46% of patients were used Clopidogrel + Atorvastatin + Glyceryl Trinitrate (nitroglycerine) + Trimetazidine Dihydrochloride + Frusemide + Spironolactone + Ramipril as a hepta therapy for the mitigation of CVD. The outcomes of the cardiovascular drug prescribed with hepta therapy are shown in Table 8. 40% of patients were used Clopidogrel + atorvastatin + bisoprolol alvcervl trinitrate (nitroalvcerine) + trimetazidinedihvdrochloride frusemide + spironolactone + ramipril as an octa therapy for the treatment of CVD. The findings of the cardiovascular drug prescribed with octa therapy are shown in Table 9.

4. DISCUSSION

By 2020, cardiovascular disease burden is predicted to be 85% of the worldwide in the

developing countries, and the increse in CAD mortality in developing nations between 1990 and 2020 is projected to be 120% in female and 137% in male [15-16]. Full study has been done on the epidemiology of CVDs and the related risk factors have been well published which several factors like include age, high blood pressure (hypertension), smoking, high blood cholesterol, diabetes, overweight or obesity, lack of exercise and family history of heart disease [17-18]. In our study, we found that the presence of hypertension was higher in males compared to females, which has the similarity to the results of Gupta R et al. [19] and Guang Hui Dong et al. [20]. The percentage of patients on combination therapy was 86.9% in this study. This observation is higher than finding reported in Ibadan, Nigeria (73%) [21] and also higher than the finding (56%) reported by Ezuo and Njoku [22]. The great number of prescriptions of combination therapy might point out the great prevalence of patients suffering from severe and moderate hypertension, and the presence of comorbid diseases.

In our study among patients who were on monotherapy, the majority of them (58.78) were treated with the beta blocker. However, this result corresponding with the study done in India that found frequently medication for the hypertensive a patient on monotherapy was beta blocker [23]. Beta-blockers have become the first recommended treatment for hypertension, affirmed currently by the guidelines of JNC VI [24]. Bisoprolol is probably the most beta1 selective agent mostly available [25]. There is some proof that it may be more efficient than atenolol in the control and management of hypertension [25-28]. Whereas diuretics were found to be the fundamentally prescribed class of antihypertensive drugs in the United Kingdom and Sub-Saharan countries [29-30].

This is in correspondence with the published international guidelines for antihypertensive treatment. JNC-VII guidelines provide diuretics should be prescribed as initial therapy for the majority of patients with either alone or in combination with drugs from other classes. As the diuretic is familiar to boost the antihypertensive efficacy of multidrug regimen, they can be helpful in attaining BP control and are less costly than other antihypertensive agents [31]. In spite of these recommendations, diuretics were reported to be prescribed less often to patients as monotherapy treatment in our study. Over the last two decades, there has been an abundant increase of new antihypertensive like ACE inhibitors, ARBs and, CCBs [32].

Where statins were the highest drugs prescribed for hypolipidaemic. The prescription of a single hypolipidaemic drug was popular. The preference of the statin as monotherapy depends on the disease process and atorvastatin was prescribed as a major hypolipidaemic drug. Also, it was found that atorvastatin was choice as a single hypolipidaemic agent. Many findings have shown first line lipid-lowering drugs used for primary preventions of coronary artery diseases (CAD) [33].

According to our study, the prescribing status was based on different therapies i.e 13.1% were on monotherapy and 86.9% were on combinations therapies i.e. dual. triple. quadruple, penta, hexa, hepta and octa therapies. And among the combination therapy penta therapy was the most common combination therapy prescribed by doctor with 26.3% followed by quadruple therapy with 15.5% and then triple therapy was the third most common combination therapy with 13.5% and dual therapy with 11.7% while hexa therapy was slightly lower than dual therapy and percentage was 11.7%, finally hepta therapy and octa therapy were 6.7% 1.4% respectively.

But Iqbal Arain et a., reported that dual therapy dominated the highest percentage of combination therapy with 42.64%, while triple therapy was the second highest with 14.40% and lowest number was quadruple therapy with 6.84%. Whereas the number of patients who prescribed monotherapy was 36.11% [34].While Krunal [35] reported that 49.50% of patients were on dual therapy, 33.16% of the patients were on monotherapy and 15.5% of patients were on triple therapy.

In our study we observed that in most prescribing monotherapy was Bisoprolol with 29.77% followed by Propranolol Hydrochloride with 27.49% and dual therapy was Bisoprolol+ Atorvastatin with 28.20% followed by Clopidogrel+Atorvastatine with 11.11%. In triple therapy was Clopidogrel+Atrovastatin+Glycery Trinitrate (nitroglycerine) with 17.77% followed by Atorvastatine+Clopidogrel+Bisoprolol with 14.08% and quadruple therapy was Clopidogrel+Atorvastatine+Bisoprolol+Glyceryl Trinitrate (nitroglycerine) with 23.22% followed by Clopidogrel+Atorvastatine+Glyceryl Trinitrate (nitroglycerine)+TrimetazidineDihydrochloride 9.03% with and penta therapy was Clopidogrel+Bisoprolol+Atorvastatine+Glyceryl Trinitrate+Trimetazidine Dihydrochloride with 52.47% and followed bv Clopidogrel+Bisoprolol+Trimetazidine Dihydrochloride+Rosuvastatin+Glyceryl Trinitrate with 14.82% and Hexa therapy was Clopidogrel+Atorvastatine+Glyceryl Trinitrate (nitroglycerine)+TrimetazidineDihydrochloride+Bi soprolol+Losartan Potassium with 26.49% Clopidogrel+GlycerylTrinitrate followed by (nitroglycerine)+TrimetazidineDihydrochloride+At orvastatine+Bisoprolol+Olmesartan Medoxomil 10.25% and 7 therapy was Clopidogrel+Atorvastatine+Bisoprolol+Glyceryl Trinitrate (nitroglycerine)+Trimetazidine Dihydrochloride+Frusemide+Spironolactone with 34.32% followed by Clopidogrel+Rosuvastatin+ Bisoprolol+Glyceryl Trinitrate (nitroglycerine)+ Trimetazidine Dihydrochloride+ Frusemide+ Spironolactone with 13.43% and 8 therapy was Clopidogrel+Atorvastatine+ Bisoprolol+Glyceryl Trinitrate (nitroglycerine)+ Trimetazidine Dihydrochloride+Frusemide+ Spironolactone+ Ramipril with 40% followed by Clopidogrel+ Rosuvastatin+Bisoprolol+Trimetazidine Dihvdrochloride+Amlodipine+Olmesartan Medoxomil+ Frusemide+Spironolactone With 13.33%.

Prescribed Medicines	Frequency	Percent
Atorvastatine+ Bisoprolol+Ramipril+GlycerylTrinitrate (nitroglycerine)	3	1.93%
Clopidogrel+Atorvastatine+ Bisoprolol +TrimetazidineDihydrochloride	12	7.74%
Clopidogrel+Bisoprolol+GlycerylTrinitrate (nitroglycerine)+TrimetazidineDihydrochloride	8	5.16%
Clopidogrel+Atorvastatine+GlycerylTrinitrate (nitroglycerine)+TrimetazidineDihydrochloride	14	9.03%
Clopidogrel+Atorvastatine+Bisoprolol+GlyceryITrinitrate (nitroglycerine)+	36	23.22%
Clopidogrel+Bisoprolol+LosartanPotassium+Atorvastatine	11	0.09%
Losartan Potassium+Atorvastatine+TrimetazidineDihydrochloride+	1	0.64%
Glyceryl Trinitrate (nitroglycerine)		
Bisoprolol+LosartanPotassium+TrimetazidineDihydrochloride+	1	0.64%
Glyceryl Trinitrate (nitroglycerine)		
Clopidogrel+Bisoprolol+Rosuvastatin+GlycerylTrinitrate (nitroglycerine)	9	5.80%
Atorvastatine+Rosuvastatin+Bisoprolol+AzisartanMedoxomil	2	1.29%
Clopidogrel+Bisoprolol+Rosuvastatin+TrimetazidineDihydrochloride	1	0.64%
Clopidogrel+Bisoprolol+Atorvastatine+TrimetazidineDihydrochloride	1	0.64%
Bisoprolol+Atorvastatine+TrimetazidineDihydrochloride+AzilsartanMedoxomil	4	2.58%
Clopidogrel+Atorvastatine+GlyceryITrinitrate (nitroglycerine)+Carvedilol	8	5.16%
Clopidogrel+Atorvastatine+Amlodipine+Atenolol	1	0.64%
Clopidogrel+Atorvastatine+GlyceryITrinitrate (nitroglycerine)+OlmesartanMedoxomil	2	1.29%
Clopidogrel+Atorvastatine+Amlodipine+OlmesartanMedoxomil	7	4.51%
Rosuvastatin+Bisoprolol+TrimetazidineDihydrochloride+GlycerylTrinitrate (nitroglycerine)	2	1.29%
Clopidogrel+Atorvastatine+Frusemide+Spironolactone	1	0.64%
Glyceryl Trinitrate (nitroglycerine)+Losartan Potassium+Atorvastatine+	1	0.64%
Verapamil Hydrochloride		
Clopidogrel+Rosuvastin+TrimetazidineDihydrochloride+GlycerylTrinitrate (nitroglycerine)	2	1.29%
Clopidogrel+Atorvastatine+TrimetazidineDihydrochloride+Losartan Potassium	1	0.64%
Atorvastatine+Bisoprolol+GlycerylTrinitrate (nitroglycerine)+Losartan Potassium	3	1.93%
Atorvastatine+Bisoprolol+GlycerylTrinitrate (nitroglycerine)+Losartan Potassium	1	0.64%
Atorvastatine+Bisoprolol+GlycerylTrinitrate (nitroglycerine)+Losartan Potassium	2	1.29%
Atorvastatine+Bisoprolol+GlycerylTrinitrate (nitroglycerine)+Losartan Potassium	1	0.64%
Atorvastatine+Bisoprolol+TrimetazidineDihydrochloride+GlycerylTrinitrate (nitroglycerine)	1	0.64%
Clopidogrel+Bisoprolol+Atorvastatine+Ramopril	5	3.22%
Clopidogrel+Rosuvastatin+Bisoprolol+Losartan Potassium	4	2.58%

Table 5. Cardiovascular drug prescribed with quadruple therapy

Prescribed Medicines	Frequency	Percent
Clopidogrel+TrimetazidineDihydrochloride+Atorvastatine+Carvedilol	1	0.64%
Clopidogrel+GlycerylTrinitrate (nitroglycerine)+Atorvastatine+Metoprolol Tartrate	1	0.64%
Clopidogrel+GlycerylTrinitrate (nitroglycerine)+Atorvastatine+Propranolol Hydrochloride	4	2.58%
Atorvastatine+Amlodipine+Atenolol+TrimetazidineDihydrochloride	1	0.64%
Glyceryl Trinitrate (nitroglycerine)+Propranolol Hydrochloride+Frusemide+Spironolactone	3	1.93%

Table 6. Cardiovascular drug prescribed with penta therapy

Penta	Frequency	Percent
Clopidogrel+Bisoprolol+Atorvastatine+GlycerylTrinitrate(nitroglycerine)+Trimetazidine Dihydrochloride	138	52.47%
Clopidogrel+Bisoprolol+TrimetazidineDihydrochloride+Rosuvastatin+GlycerylTrinitrate(nitroglycerine)	39	14.82%
Clopidogrel+Atorvastatine+GlycerylTrinitrate(nitroglycerine)+Trimetazidine Dihydrochloride+Carvedilol	9	3.42%
Clopidogrel+Atorvastatine+Bisoprolol+GlycerylTrinitrate(nitroglycerine)+Losartan Potassium	1	0.38%
Clopidogrel+Rosuvastatin+Bisoprolol+GlycerylTrinitrate (nitroglycerine)+AzisartanMedoxomil	6	2.28%
Clopidogrel+Atorvastatine+Carvedilol+Frusemide+Spironolactone	5	1.90%
Clopidogrel+GlycerylTrinitrate(nitroglycerine)+Atorvastatine+Carvedilol+Ramopril	3	1.14%
Clopidogrel+Atorvastatine+Bisoprolol+Ticagrelor+Ramopril	2	0.76%
Bisoprolo+GlycerylTrinitrate(nitroglycerine)+Losartan Potassium+Trimetazidine Dihydrochloride+Rosuvastatin	3	1.14%
Losartan Potassium+Clopidogrel+Bisoprolol+Atorvastatine+TrimetazidineDihydrochloride	2	0.76%
Clopidogrel+GlycerylTrinitrate(nitroglycerine)+Carvedilol+Atorvastatine+Losartan Potassium	1	0.38%
Clopidogrel+Atorvastatine+GlycerylTrinitrate(nitroglycerine)+TrimetazidineDihydrochloride+Metoprolol Tartrate	2	0.76%
Atorvastatine+GlycerylTrinitrate(nitroglycerine)+Metoprolol Tartrate+Trimetazidine Dihydrochloride+Ramipril	1	0.38%
Rosuvastatin+Bisoprolol+LosartanPotassium+Ticagrelor+OlmesartanMedoxomil	1	0.38%
Clopidogrel+Atorvastatine+TrimetazidineDihydrochloride+Carvedilol+Ramipril	2	0.76%
Clopidogrel+Bisoprolol+GlycerylTrinitrate(nitroglycerine)+Atorvastatine+Ramopril	11	4.18%
Clopidogrel+Atorvastatine+GlyceryITrinitrate(nitroglycerine)+CarvediloI+Ramopril	1	0.38%
Clopidogrel+Atorvastatine+GlyceryITrinitrate +TrimetazidineDihydrochloride+Carvedilol	16	6.08%
Clopidogrel+Rosuvastatin+Atorvastatine+GlycerylTrinitrate(nitroglycerine) +Trimetazidine Dihydrochloride	2	0.76%
Clopidogrel+Rosuvastatin+Bisoprolol+TrimetazidineDihydrochloride+OlmesartanMedoxomil	3	1.14%
Clopidogrel+Bisoprolol+GlycerylTrinitrate(nitroglycerine)+TrimetazidineDihydrochloride+Metoprolol Tartrate	1	0.38%
Clopidogrel+Atorvastatine+Ranolazine+Bisoprolol+GlycerylTrinitrate(nitroglycerine)	4	1.52%
Clopidogrel+Atorvastatine+Amlodipine+GlycerylTrinitrate(nitroglycerine)+Losartan Potassium	1	0.38%
Clopidogrel+Atorvastatine+TrimetazidineDihydrochloride+GlycerylTrinitrate(nitroglycerine) +Ramopril	3	1.14%
Clopidogrel+Bisoprolol+Atorvastatine+LosartanPotassium+GlycerylTrinitrate(nitroglycerine)	2	0.76%

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Penta	Frequency	Percent
Clopidogrel+Bisoprolol+Rosuvastatin+LosartanPotassium+GlycerylTrinitrate(nitroglycerine)	1	0.38%
Trimetazidine+Dihydrochloride+Carvedilol+Ramopril+Frusemide+Spironolactone	1	0.38%
Atorvastatine+TrimetazidineDihydrochloride+Frusemide+Spironolactone+Carvedilol	2	0.76%
Bisoprolol+TrimetazidineDihydrochlori+Ramopril+Frusemide+Spironolactone	1	0.38%
Clopidogrel+Bisoprolol+Atorvastatine+Ramopril+Cilnidipine	1	0.38%
Clopidogrel+Bisoprolol+Rosuvastatine+Frusemide+Spironolactone	1	0.38%
Atorvastatine+Carvedilol+Frusemide+Spironolactone+Losartan Potassium	1	0.38%
Clopidogrel+Atorvastatine+Bisoprolol+GlycerylTrinitrate (nitroglycerine)+Trimetazidine Dihydrochloride	1	0.38%
Clopidogrel+Atorvastatine+Bisoprolol+Amlodipine+GlycerylTrinitrate(nitroglycerine)	1	0.38%
Clopidogrel+GlycerylTrinitrate (nitroglycerine)+Atorvastatine+Carvedilol+Losartan Potassium	5	1.90%

Table 7. Cardiovascular drug prescribed with hexa therapy

Prescribed medicines	Frequency	Percent
Clopidogrel+Atorvastatine+GlycerylTrinitrate (nitroglycerine)+TrimetazidineDihydrochloride+Bisoprolol+Losartan Potassium	31	26.49%
Clopidogrel+Glyceryl Trinitrate (nitroglycerine)+Trimetazidine Dihydrochloride+Atorvastatine+Bisoprolol+OlmesartanMedoxomil	12	10.25%
Clopidogrel+Atorvastatine+GlyceryITrinitrate (nitroglycerine)+TrimetazidineDihydrochloride+Carvedilol+Ramipril	3	2.56%
Clopidogrel+Bisoprolol+Atorvastatine+Trimetazidine Dihydrochloride+Amlodipine +OlmesartanMedoxomil	5	4.27%
Clopidogrel+Atorvastatine+TrimetazidineDihydrochloride+Bisoprolol+Glyceryl Trinitrate (nitroglycerine)+Amlodipine	2	1.70%
Atorvastatine+Losartan Potassium+Glyceryl Trinitrate (nitroglycerine)+	1	0.85%
Trimetazidine Dihydrochloride+Amlodipine +OlmesartanMedoxomil		
AzisartanMedoxomil+Clopidogrel+Bisoprolol+RosuvastatinGlyceryl Trinitrate (nitroglycerine)+TrimetazidineDihydrochloride	3	2.56%
Clopidogrel+Atorvastatine+Frusemide+Spironolactone+Ramipril+Carvedilol	2	1.70%
Clopidogrel+Atorvastatine+GlycerylTrinitrate (nitroglycerine)+TrimetazidineDihydrochloride+Carvedilol+Losartan Potassium	1	0.85%
Atorvastatin+TrimetazidineDihydrochloride+Carvedilol+LosartanPotassium+Frusemide+Spironolactone	1	0.85%
Clopidogrel+Atorvastatine+Rosuvastatin+TrimetazidineDihydrochloride+Losartan Potassium+Carvedilol	2	1.70%
Clopidogrel+Bisoprolol+Atorvastatine+TrimetazidineDihydrochloride+Glyceryl Trinitrate (nitroglycerine)+Valsartan	2	1.70%
Clopidogrel+Atorvastatine+Bisoprolol+TrimetazidineDihydrochloride+Frusemide+Spironolactone	2	1.70%
Clopidogrel+Atorvastatine+Bisoprolol+TrimetazidineDihydrochloride+Glyceryl Trinitrate (nitroglycerine)+Ramipril	4	3.41%
Clopidogrel+Rosuvastatin+GlycerylTrinitrate (nitroglycerine)+TrimetazidineDihydrochloride+Frusemide+Spironolactone	9	7.69%
Clopidogrel+Rosuvastatin+Bisoprolol+Amlodipine+OlmesartanMedoxomil+Losartan Potassium	1	0.85%
Clopidogrel+Rosuvastatin+Bisoprolol+Atorvastatine+TrimetazidineDihydrochloride+GlycerylTrinitrate (nitroglycerine)	1	0.85%
Frusemide+Spironolactone+Bisoprolol+Clopidogrel+TrimetazidineDihydrochloride+Ramopril	1	0.85%
Clopidogrel+Atorvastatine+Rosuvastatin+TrimetazidineDihydrochlorideGlyceryl Trinitrate (nitroglycerine)+Indapamide	4	3.41%

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Prescribed medicines	Frequency	Percent
Clopidogrel+Rosuvastatin+Bisoprolol+TrimetazidineDihydrochloride+Amlodipine+Telmisartan	5	4.27%
Clopidogrel+Atorvastatine+Bisoprolol+TrimetazidineDihydrochloride+Glyceryl Trinitrate (nitroglycerine)+Carvedilol	1	0.85%
Clopidogrel+Atorvastatine+TrimetazidineDihydrochloride+GlycerylTrinitrate (nitroglycerine)+Carvedilol+Ramopril	9	7.69%
Clopidogrel+GlycerylTrinitrate (nitroglycerine)+Atorvastatine+Carvedilol+Frusemide+Spironolactone	5	4.27%
Clopidogrel+GlycerylTrinitrate (nitroglycerine)+Atorvastatine+PropranololHydrochloride+Hydrochlorothiazide+Losartan Potassium	1	0.85%
Clopidogrel+TrimetazidineDihydrochloride+GlycerylTrinitrate (nitroglycerine)+ Ramopril+Propranolol Hydrochloride	2	1.70%
Clopidogrel+Atorvastatine+Hydrochlorothiazide+Losartan Potassium+Frusemide+Spironolactone	1	0.85%
Clopidogrel+Atorvastatine+PropranololHydrochloride+Frusemide+Spironolactone+TrimetazidineDihydrochloride	1	0.85%
Clopidogrel+Bisoprolol+TrimetazidineDihydrochloride+GlycerylTrinitrate (nitroglycerine)+Atorvastatine+Perindopril Erbumibe	1	0.85%

Prescribed medicines	Frequency	Percent
Clopidogrel+Atorvastatine+Glyceryl Trinitrate	5	7.46%
(nitroglycerine)+TrimetazidineDihydrochloride+Frusemide+Spironolactone+Ra		
mipril		
Clopidogrel+Rosuvastatin+Bisoprolol+GlycerylTrinitrate (nitroglycerine)+	9	13.43%
TrimetazidineDihydrochloride+Frusemide+Spironolactone		
Clopidogrel+Atorvastatine+GlyceryITrinitrate (nitroglycerine)+	3	4.48%
TrimetazidineDihydrochloride+CarvediloI+Amlodipine+OlmesartanMedoxomil		
Amlodipine+Atenolol+OlmesartanMedoxomil+GlyceryITrinitrate	1	0.49%
(nitroglycerine)+TrimetazidineDihydrochloride+Frusemide+Spironolactone		
Clopidogrel+Atorvastatine+GlyceryITrinitrate (nitroglycerine)	2	2.95%
TrimetazidineDihydrochloride+Carvedilol+Frusemide+Spironolactone		
Aspirin+Clopidogrel+Atorvastatine+GlyceryITrinitrate (nitroglycerine)	4	5.97%
TrimetazidineDihydrochloride+Carvedilol+Amlodipine + Valsartan		
Clopidogrel+Atorvastatine+Bisoprolol+GlycerylTrinitrate (nitroglycerine)+	23	34.32%
TrimetazidineDihydrochloride+Frusemide+Spironolactone		
Clopidogrel+Atorvastatine+TrimetazidineDihydrochloride+GlycerylTrinitrate	1	0.49%
(nitroglycerine)+Losartan Potassium+Frusemide+Spironolactone		
Clopidogrel+Bisoprolol+TrimetazidineDihydrochloride	2	2.95%
+Frusemide+Spironolactone+Rosuvastatin+Ramipril		
Clopidogrel+Bisoprolol+TrimetazidineDihydrochloride+	1	0.49%
Glyceryl Trinitrate (nitroglycerine)+AzilsartanMedoxomil		
+Frusemide+Spironolactone		
Clopidogrel+Atorvastatine+Bisoprolol+TrimetazidineDihydrochloride+	6	8.95%
Glyceryl Trinitrate (nitroglycerine)+Amlodipine+OlmesartanMedoxomil		
Aspirin+Clopidogrel+Atorvastatine+Bisoprolol+TrimetazidineDihydrochloride+	4	5.97%
GlycerylTrinitrate (nitroglycerine)+Hydrochlorothiazide+Losartan Potassium		
Clopidogrel+Atorvastatine+GlyceryITrinitrate (nitroglycerine)+	1	0.49%
+Losartan Potassium+Carvedilol+Frusemide+Spironolactone		
Ticagrelor+Rosuvastatin+Bisoprolol+TrimetazidineDihydrochloride+	1	0.49%
Glyceryl Trinitrate (nitroglycerine)+Carvedilol+Ramopril		
Clopidogrel+GlycerylTrinitrate (nitroglycerine)+Atorvastatine+	2	2.95%
TrimetazidineDihydrochloride+Frusemide+Spironolactone+Metoprolol Tartrate		
Clopidogrel+GlyceryITrinitrate	2	2.95%
(nitroglycerine)+Atorvastatine+TrimetazidineDihydrochloride+Hydrochlorothiazi	İ	
de+OlmesartanMedoxomil+Bisoprolol		

Table 8. Cardiovascular drug prescribed with hepta therapy

Table 9. Cardiovascular drug prescribed with octa therapy

Prescribed medicines	Frequency	Percent
Clopidogrel+Rosuvastatin+Bisoprolol+TrimetazidineDihydrochloride+	2	13.33%
Amlodipine+Olmesartan Medoxomil+Frusemide+Spironolactone		
Clopidogrel+Atorvastatine+Bisoprolol+GlycerylTrinitrate (nitroglycerine)+	6	40%
TrimetazidineDihydrochloride+ Frusemide+Spironolactone+Ramipril		
Clopidogrel+Atorvastatine+GlyceryITrinitrate	1	6.67%
(nitroglycerine)+TrimetazidineDihydrochloride+Frusemide+Spironolactone	e	
+Ramipril+Carvedilol		
Clopidogrel+Bisoprolol+Atorvastatine+TrimetazidineDihydrochloride+	2	13.33%
Losartan Potassium+Frusemide+Spironolactone+Cilnidipine		
Clopidogrel+Rosuvastatin+Carvedilol+TrimetazidineDihydrochloride+	1	6.67%
Glyceryl Trinitrate (nitroglycerine)+Frusemide+Spironolactone+		
Losartan Potassium		
Clopidogrel+Carvedilol+Ramopril+Trimetazidine Dihydrochloride+Glycery	11	6.67%
Trinitrate (nitroglycerine)+AzilsartanMedoxomilFrusemide+Spironolactone	;	

Clopidogrel+Atorvastatine+Trimetazidine Dihydrochloride+Glyceryl	1	6.67%
Trinitrate		
(nitroglycerine)+Carvedilol+Ramopril+Frusemide+Spironolactone		
Clopidogrel+Rosuvastatin+Bisoprolol+TrimetazidineDihydrochloride	1	6.67%
+Frusemide+Spironolactone+Amlodipine+OlmesartanMedoxomil		

But Iqbal Arain et al. reported the most monotherapy was Telmisartan with 11.11% whereas most common dual therapy was Telmisartan+Hydrochlorothiazide with 14.76% moreover among triple therapy most prescribing combination was Atenolol +Chlorthalidone+ Telmisartan with 15.30% and finally, Atenolol+ Chlorthalidone+Telmisartan+αMD was most frequently drug prescribed for quadruple therapy with 26.44%. While Krunal reported the greatest percentage of combination dual therapy was enalapril+atenolol with 22% followed bv enalapril+amlodipine with 10.83% and greatest percentage of triple therapy were enalapril+ atenolol+amlodipine with 8% followed by enalapril+atenolol+frusemide with 4.16% and enalapril+atenolol, amlodpine +furosemide were described the most frequent therapy prescribed for quadruple therapy with 1.66%.

And another study by V Pavani et al. observed that the majority of patients were prescribed with dual therapy with (48.3%), followed by monotherapy with (30.2%), and triple therapy with (13%) and the lowest percentage of patients were with (8.3%) who prescribed with more than 3 drugs [36]. In our study where the maximum number of patients was found in the age group of 40-50 years. While in India the average group of 40-50 years in men and 60 -64 in women in the city whilst in the village an age increase in both gender in the age group 50-64 [37].

Drug prescription pattern in our study observed that combination therapy was more commonly prescribed than monotherapy this result match with previous studied conducted by Kumar et.al [38] and also higher than those recorded in similar studies conducted in southwestern and northern regions of Nigeria [39,40].

5. CONCLUSION

CVDs are the most common and affected diseases in the present time. CVDs are not only affected alone but also caused due to the complications of other diseases. To treat these complex and well-known diseases doctors prescribed various types of drugs either alone or in combination therapy. Our study results suggested that doctors are prescribing combination therapy most frequently than monotherapy which completely matched with the results published regarding this issues.

CONSENT

The written consent was approved by administrative offices in the hospital.

ETHICAL APPROVAL

It is not applicable.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- Naghavi M, Wang H, Lozano R, Davis A, Liang X, Zhou M, Vollset SE, Ozgoren AA, Abdalla S, Abd-Allah F, Aziz MI. Global, regional and national age-sex specific allcause and cause-specific mortality for 240 causes of death, 1990-2013: A systematic analysis for the Global Burden of Disease Study 2013. Lancet. 2015;385:117–71.
- 2. Karar ZA, Alam N, Streatfield K. Epidemiological transition in rural Bangladesh, 1986–2006. Glob Health Action. 2009;2(Supplements):1-9.
- Omran AR. The epidemiologic transition. A theory of the epidemiology of population change. Milbank Memorial Fund Quarterly. 1971;49(4):509-38.
- 4. Bangladesh bureau of statistics. Available:<u>http://www.bbs.gov.bd</u> (Accessed January 2009)
- 5. American Heart Association. Heart and Stroke Statistical Update. Dallas, Tex: American Heart Association; 2001.
- World Health Organization. Cardiovascular diseases. Available:<u>http://www.who.int/mediacentre/f</u> actsheets/fs317/en/index.html
- 7. Krishnan MN. Coronary heart disease and risk factors in India on the brink of an

epidemic. Indian Heart J. 2012;64(4):364-7.

- WHO Mediam center. Fact sheet: Cardiovascular diseases (CVDs); 2016. Available:<u>http://www.who.int/mediacentre/factsheets/fs317/en</u>
- Moran AE, Forouzanfar MH, Roth GA, Mensah GA, Ezzati M, Murray CJ, Naghavi M. Temporal trends in ischemic heart disease mortality in 21 world regions, 1980 to 2010: The Global Burden of Disease 2010 study. Circulation. 2014;129(14): 1483-92.
- Global health estimates: Deaths, disabilityadjusted life year (DALYs), years of life lost (YLL) and years lost due to disability (YLD) by cause, age and sex, 2000–2012. Geneva: World Health Organization. Available:<u>http://www.who.int/healthinfo/glo bal_burden_disease/estimates/en/</u>
- 11. Di Cesare M, Bennett JE, Best N, Stevens GA, Danaei G, Ezzati M. Thecontributions of risk factor trends to cardiometabolic mortality decline in 26industrialized countries. Int J Epidemiol. 2013;42(3):838-48.
- Ezzati M, Obermeyer Z, Tzoulaki I, Mayosi BM, Elliott P, Leon DA. Contributions of risk factors and medical care to cardiovascular mortalitytrends. Nat Rev Cardiol. 2015;12(9):508-30.
- Law MR, Wald NJ, Rudnicka AR. Quantifying effect of statins on low densitylipoprotein cholesterol, ischaemic heart disease, and stroke: Systematic reviewand meta-analysis. BMJ. 2003; 326(7404):1423.
- Antiplatelet Trialists' Collaboration. Collaborative overview of randomized trials of antiplatelet therapy--I: Prevention of death, myocardial infarction, and stroke by prolonged antiplatelet therapy in various categories of patients. BMJ. 1994; 308(6921):81-10.
- 15. Reddy KS, Yusuf S. Emerging epidemic of cardiovascular disease in developing countries. Circulation. 1998;97(6):596-601.
- 16. Bulato RA, Stephens PW. Global estimates and projections of mortality by cause. Washington DC: Population, Health and Nutrition Department: World Bank. 1992;1007.
- World Heart Federation. Cardiovascular disease risk factors. (Accessed on 2015 May 20)

Available:<u>http://www.world-heart-</u> federation.org/press/factsheets/cardiovascular-disease-risk-factors

- 18. NIH. Who is at risk for heart disease? (Accessed on 2015 May 20) Available:<u>http://www.nhlbi.nih.gov/health/h</u> ealth-topics/topics/hdw/atrisk
- Gupta R, Prakash H, Gupta VP, Gupta KD. Prevalence and determinants of coronary heart disease in a rural population of India. J Clin Epidemiol. 1997;50(2):203-9.
- Dong GH, Sun ZQ, Zhang XZ, Li JJ, Zheng LQ, Li J, et al. Prevalence, awareness, treatment and control of hypertension in a rural Liaoning Province, China. Indian J Med Res. 2008;128:122-27.
- 21. Yusuff KB, Balogun O. Physicians' prescribing of anti hypertensive combinations in a tertiary care setting in southwestern Nigeria. J Pharm Pharm Sci. 2005;8(2):235-42.
- 22. Isezuo AS, Njoku CH. Blood pressure control among hypertensives managed in aspecialised health care setting in Nigeria. Afr J Med Med Sci. 2003;32(1):65-70.
- 23. Khurshid F, Aqil M, Alam MS, Kapur P, Pllai KK. Antihypertensive medication prescribing patterns in auniversity teaching hospital in south Delhi. IJPSR. 2012;3(7): 2057-63.
- 24. The sixth report of the Joint National Committee on prevention, detection, evaluation, and treatment of high blood pressure. Arch Intern Med. 1997;157(21): 2413-46.
- Weir MR, Bolli P, Prichard BNC, Weber MA. Bisoprolol. In: Messerli FH (ed). Cardiovascular drug therapy, 2nd edition. W B Saunders Co: Philadelphia. 1996;557– 568.
- 26. Smith C, Teitler M. Beta-blocker selectivity at cloned human beta 1- and beta 2adrenergic receptors. Cardiovasc Drugs Ther. 1999;13(2):123-6.
- Bühler FR, Berglund G, Anderson OK, Brunner HR, Scherrer U, van Brummelen P, Distler A, Philipp T, Fogari R, Mimran A, et al. Double-blind comparison of the cardioselective beta-blockers bisoprolol and atenolol in hypertension: The Bisoprolol International Multicenter Study (BIMS). J Cardiovasc Pharmacol. 1986; 8(Suppl 11):S122-7.
- Prichard BN, Gillam PM. Use of propranolol (inderal) in treatment of hypertension. Br Med J. 1964;2(5411): 725-7.

- 29. Walley T, Duggan AK, Haycox AR, Niziol CJ. Treatment for newly diagnose hypertension: Patterns of prescribing and antihypertensive effectiveness in the UK. J R Soc Med. 2003;96:525 -31.
- Olanrewaju TO, Aderibigbe A, Busari OA, Sanya EO. Antihypertensive drug utilization and conformity to guidelines in a sub-Saharan African hypertensive population. Int J Clin Pharmacol Ther. 2010;48(1):68-75.
- Dahlöf B, Sever PS, Poulter NR, Wedel H, 31. Beevers DG, Caulfield M, Collins R, Kjeldsen SE, Kristinsson A, McInnes GT, Mehlsen J, Nieminen M, O'Brien E, Osteraren J. ASCOT Investigators. Prevention of cardiovascular events with an antihypertensive regimen of amlodipine adding perindopril as required versus atenolol adding bendroflumethiazide as required, in the Anglo-Scandinavian Cardiac Outcomes Trial-Blood Pressure Arm (ASCOT-BPLA): Lowering Α multicenter randomised controlled trial. Lancet. 2005;366(9489):895-906.
- Jhaj R, Goel NK, Gautam CS, Hota D, Sangeeta B, Sood A, Sachdev A. Prescribing patterns and cost of antihypertensive drugs in an internal medicine clinic. Indian Heart J. 2001;53(3):323-7.
- 33. Ravi GR, Pradeepa R, Mohan V. Hypertriglyceridemia and coronary artery-

disease-an update. Indian Heart J. 2004;56(1):21-6.

- Arain MI, Choto MA, Dayo A, Parveen R. Pharmacovigilance studies of antihyperte teaching hospital of Hyderabad, Sindh. J Young Pharm. 2016;8(3):259-265.
- 35. Krunal CS, RAM, Anil P, Singh. Drug utilization study of anti-hypertensive drugs and their adverse effects in patients of a tertiary care hospital. J Clin Exp Res. 2013;1(3):58-67.
- Pavani V, Cidda M, Ramya Krishna T, Parmar MY, Nalini M. Study of prescribing patterns of antihypertensive drugs. IJPBS. 2012;2(2):317-327.
- Dalal PM. Hypertension: A report on community survey on casual hypertension in old Bombay, Sir. H N Hospital Research Society; 1980.
- Kaur S, Gupta S, Kumar D, Lal M, Gilani Z. Prescribing pattern of antihypertensive drugs in a tertiary care hospital in Jammu-A Descriptive study. JK-Practitioner. 2012; 17(4):38-41.
- Yusuff KB, Balogun O. Physician's prescribing of antihypertensive combinations in a tertiary care setting in southwestern Nigeria. J Pharm Pharm Sci. 2005;8(2):235-242.
- Isezuo AS, Njoku CH. Blood pressure control among hypertensives managed in aspecialised health care setting in Nigeria. Afr J Med Med Sci. 2003;32(1):65-70.

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