

Drug Utilization Evaluation and Cost of Illness Analysis in Diabetes and Cardiovascular Disease in Tertiary Care Hospital

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ABSTRACT:

Background: Cardiovascular disease (CVD) is a major cause of death and disability among people with diabetes. Diabetes mellitus (DM) has become a rising epidemic in the last century, more pressing in the last few decades with the exponential rise of obesity, and has become one of the leading causes of death worldwide.

Aim: The aim of this study is to Evaluate Drug Utilization Pattern and Cost of Illness Analysis of Diabetes and Cardiovascular Disease.

Methodology: A Prospective observational study With 102 Inpatients from Department of Cardiology and Endocrinology.

Results and Observation: Over all study exhibits 230 drug-drug interaction, most of the drug-drug interaction present in Antiplatelets + Anticoagulants, Antiplatelets + Diuretics and ACE inhibitors + Diuretics etc. In over all COI study expressed, Myocardial infarction patients were more expensive than other CVD and DM patients. 80 percent of medicines were prescribed in National List of Essential Medicine. Drug Utilization Evaluation of Cardiovascular disease and diabetes mellitus as analyzed through the WHO core prescribing indicators like Number of medicines per prescription in this study population as 713(100%) medicines, Number of generic name per prescription in this total study population as 109(15%) medicines, Number of antibiotics per prescriptions in this total study population as 15(2%) , Number of injections per prescriptions in this study population as 114(16%) and Number of essential drugs in this study population as 524(%).

Conclusion: Over all patients were 102 and their prescription had 230 drug-drug interactions. Serious Drug-Drug interactions were 11%, Moderate Drug-Drug interactions were 84%, Minor Drug-Drug interactions were 5%. Most drug-drug interaction were founded in Pharmacodynamic mechanism compared with pharmacokinetic mechanism and the Drug-Drug interaction were

founded in Category-C.WHO Core prescribing indicators were used to assess these prescription and these patients prescription were prescribed by the WHO Core prescribing indicators. In CVD and DM patients were prescribed most of the drugs as Antiplatelets, Statin, Anticoagulants like Clopidogrel, Aspirin, Atorvastatin, Rosuvastatin, Heparin sodium etc.

KEYWORDS: Cardiovascular Disease, WHO core Prescribing Indicators, Diabetes Mellitus (DM), Drug Utilization Evaluation (DUE), Potential Drug-Drug Interactions, Cost of Illness Analysis.

I. INTRODUCTION:

"CARDIOVASCULAR DISEASES"

(CVDs) refers to a variety of heart and blood vessel conditions. There are other additional underlying factors that contribute to CVDs. These are a reflection of the three main causes — urbanisation, population ageing, and globalisation that are causing social, economic, and cultural transformation. Poverty, stress, and inherited factors are other CVD risk factors. The cardiovascular system can experience a wide range of issues, including endocarditis, rheumatic heart disease, irregularities in the conduction system, and cardiovascular disease (CVD) or heart disease, among others.⁽¹⁾

Additionally, pharmacological therapy for diabetes, high blood lipids, and hypertension is necessary to lessen cardiovascular risk and minimize heart attacks and strokes in individuals who suffer from these diseases.

The economic cost of CVD include both the costs of general healthcare for the economy as well as the loss of productivity brought on by the condition. Given the high prevalence of CVD, this economic impact could be significant. Especially in nations with higher rates of CHD and stroke morbidity. The WHO projects that between 2010 and 2030, the cost to the US healthcare budget for people between the ages of 50 and 65 would

increase from 15% to 25% in 2030. ⁽³⁾ Although CVD may directly result from a variety of aetiologies, including rheumatic fever that causes valvular heart disease and emboli in patients with atrial fibrillation that result in ischemic stroke, addressing risk factors associated with the development of atherosclerosis is crucial because it is a common factor in the pathophysiology of CVD. ⁽²⁾ Hypertension (HTN), Coronary Artery Disease (CAD), Cerebral Vascular Accident (CVA), Myocardial Infarction (MI), Congestive Heart Failure (CHF), Angina Pectoris, Atherosclerosis, Aortic Stenosis, Rheumatic Heart Disease (RHD), Ischemic Heart Disease (IHD), Cardiac Arrhythmia, Mitral Valve Regurgitation and Pericarditis are all included in the category of cardiovascular disease. ⁽²⁾

DIABETES MELLITUS:

According to the WHO, Diabetes Mellitus (DM) is classified as a heterogeneous metabolic condition with disturbances in the metabolism of carbohydrates, fats, and proteins as well as a common trait of persistent hyperglycemia. Diabetes mellitus (DM) is a state of hyperglycemia that has a variety of underlying causes. It is categorized generally into Type 1 "insulin-dependent" (IDDM) and Type 2 "non-insulin-dependent" (NIDDM). ⁽⁴⁾

It results from either an insulin deficit, an insulin resistance, or both. Beta cells in the pancreas release insulin to regulate blood sugar levels. Some of the symptoms that diabetic people frequently experience include blurred vision, excessive thirst, fatigue, frequent urination, hunger, and weight loss.

Diabetes also increases the likelihood of developing other illnesses, such as non-alcoholic fatty liver disease, cataracts, erectile dysfunction, peripheral artery and cerebrovascular disease, heart disease, and obesity. They also have a higher chance of contracting some infectious diseases, such as tuberculosis. Blood glucose (or blood sugar) levels that are elevated in people with diabetes are chronic metabolic conditions that over time cause substantial harm to the heart, blood vessels, eyes, kidneys, and nerves. ⁽⁵⁾

TYPE 1: Insulin Dependent Diabetes Mellitus (IDDM)

T1DM, also called as the insulin-dependent diabetes mellitus (IDDM), manifests due to the autoimmune damage of the B-cells which then

leads to the suppression or cessation of insulin production. T1DM is also called the "Juvenile Onset Diabetes Mellitus".

There is beta cell destruction in pancreatic islets; majority of cases are autoimmune (type 1A) antibodies that destroy beta cells are detectable in blood, but some are idiopathic (type 1B) no beta cell antibody is found. In all type 1 cases circulating insulin are more prone to ketosis. This type is less common and has a low degree of genetic predisposition. ⁽⁶⁾

Type 2: Non-Insulin dependent diabetes mellitus (NIDDM)

There is no loss or only moderate reduction in beta cell mass; insulin in circulation is low, normal or even high, no anti beta cell antibody is demonstrable, has a high degree of genetic predisposition; generally has a late onset (past middle age). Over 90% cases of diabetes are type 2 diabetes mellitus. ⁽⁶⁾

There are two issues because the glucose is not adequate entering the body's cells.

- A build-up of glucose in the blood.
- Glucose, which the cells require for energy and growth, is not being delivered to them.

Gestational Diabetes Mellitus:

Gestational diabetes is a condition in which a pregnant woman has elevated glucose levels and other symptoms of diabetes. During pregnancy, you make hormones that cause glucose to build up in blood. Usually, pancreas can send out enough insulin to handle it. But body can't make enough insulin or stops using insulin as it should rise blood sugar levels, and get gestational diabetes. Gestational diabetes are a high risk of developing type 2 diabetes later in life. ⁽⁶⁾

DRUG INTERACTIONS:

"Drug interaction is defined as the pharmacological activity of one drug is altered by the concomitant use of another drug or by the presence of some other substance. The drug whose activity is affected by such an interaction is called as a "Object drug". The agent which precipitates such an interaction is referred to as the "Precipitant". ⁽⁷⁾

Types of Drug Interactions:

- Drug-Drug interactions
- Drug-food interactions
- Chemical-drug interactions
- Drug-laboratory test interaction
- Drug-disease interaction

COST OF ILLNESS ANALYSIS:

Cost of illness (COI), known as burden of disease (BOD), is a definition that encompasses various aspects of the disease impact on the health outcomes in a country, specific regions, communities, and even individuals. A determination of the economic impact of an illness or condition. [Changik Jo et al.,]⁽⁸⁾

Cost of Illness includes:

- Medical care for prevention, treatment & Social services for rehabilitation.
- Productivity loss

The objective of Cost-of-illness (COI) analysis is to evaluate the economic burden of illness on society as a whole in terms of the consumption of health care resources and production losses. The economic cost of illness represents the economic benefit of a health care intervention that eradicated such illness.

Types of Costs:

Direct, indirect, and intangible costs are the three categories into which COI studies often divide costs. Due to the fact that intangible expenses, as measurement issues and related disputes make it difficult to quantify in COI investigations, we here primarily focus on the first two price ranges.

DIRECT COST:

The direct costs, which are borne by the health care system, society, families, and individual patients, include medical and nonmedical expenses. The former is described as the costs of medical care for a diagnosis, treatment, and rehabilitation, among other things, whereas the last one is connected to the use of non-health care resources such as transportation, living expenses, moving, and property losses and unrecognized cares of any kind.

IN-DIRECT COST:

In COI studies, the term "indirect" occasionally refers to productivity losses caused by morbidity and mortality, which are borne by the individual, family, society, or the employer. In contrast to accounting and most business disciplines, where "indirect" costs indicate the supporting and overhead activities that need to be shared among the users. It has been

advised to replace the word with "productivity losses" or "productivity costs" to eliminate any confusion or uncertainty to readers who may experience.⁽⁸⁾

DRUG UTILIZATION EVALUATION:

According to WHO, Drug Utilization evaluation is defined as the marketing, distribution, prescription and use of drugs in society, with special emphasis on the resulting medical, social and economic consequences.⁽²⁾

Drug Utilization Reviews (DUR), also referred to as Drug Utilization Evaluations (DUE) or Medication Utilization Evaluations (MUE), are defined as an authorized, structured, ongoing review of health care provider prescribing, pharmacist dispensing, and patient use of medication. DURs involve a comprehensive review

of patients' prescription and medication data before, during, and after dispensing to ensure appropriate medication use in decision making and positive patient outcomes.⁽¹⁰⁾

Drug Utilization Evaluation (DUE) is an ongoing authorized and systematic quality improvement process, designed to optimize drug use by developing criteria and standards.

- To educate clinicians and other Health Care Professionals (HCP), to increase appropriate drug use.
- To provide feedback of results obtained during study to clinicians and other Health Care Professionals.
- To review drug use.
- To analyze prescription pattern.
- To develop and standards which describe optimal drug use.
- To promote appropriate drug use through education and other interventions.⁽¹¹⁾

OBJECTIVE:

- To understand the Prescribing Pattern of Drugs used in Cardiovascular Disease and T2DM, Promote patient awareness about the diseases.
- To Evaluate The Potential Drug – Drug Interactions (PDDIs).
- To Identify The Rationality of Drug Prescribed For CVD And DM Patients.
- To Estimate Cost of Illness Analysis of Cardiovascular Disease and Diabetes Mellitus Individuals.
- To Determine Whether Providing Essential Medicine.
- To Understand The Pattern of Drug Utilization Using WHO Core Prescribing Indicators.

II. METHODOLOGY:

- **Study Design:**

A prospective Observational Study

- **Study Site:**

The study was carried out at Cardiology and Endocrinology department in Sudha Institute of Medical Science (SIMS).

- **Sample Size:**

A total number of 102 people were included according to inclusion and exclusion criteria.

- **Study Duration:**

The study was conducted for duration of 6 months (March 2022 – August 2022).

- **Study Subjects:**

Patients diagnosed with T2DM accompanying Cardiovascular Disease

- **Study Criteria:**

- **Inclusion Criteria:**

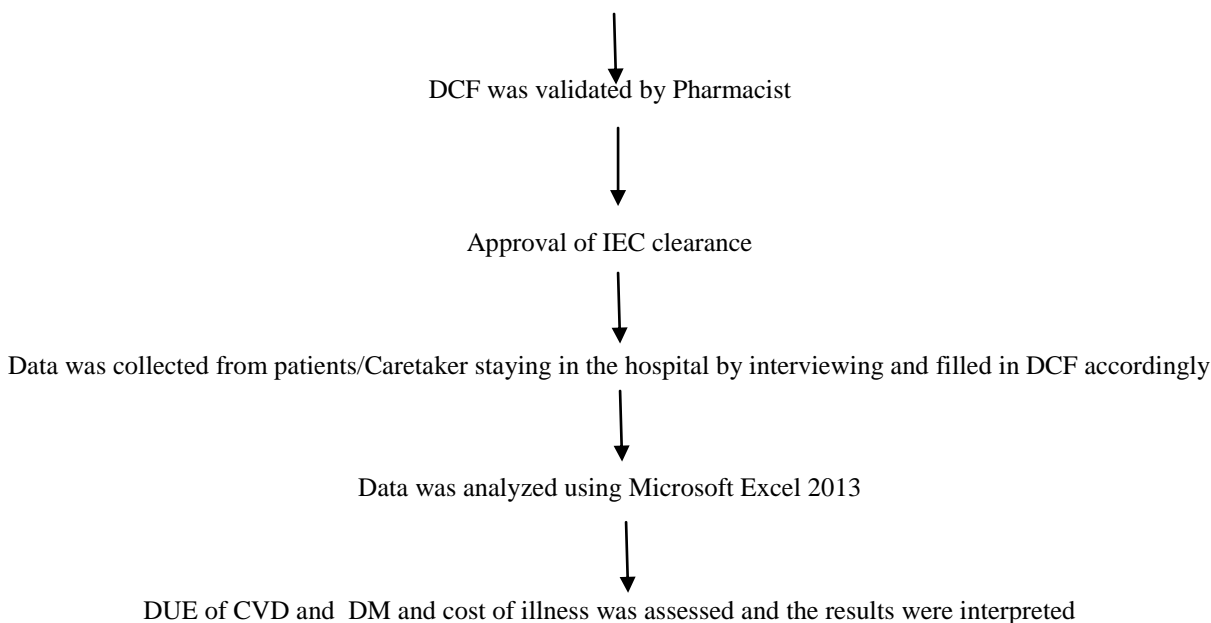
- 25 - 90 years old both male and female patients.
- Patient diagnosed with Type-2 DM and T2DM along with its CVD complications (HTN, Atherosclerosis, CHD, HF, Angina, MI and stroke) during the study period.
- Patients who are willing to participate in the study.

- **Exclusion Criteria:**

- Patient below 25 years.
- Pregnant women and lactating women.
- Patient diagnosed with CVD and Type-2 DM Combined with other co- morbidities.
- Vulnerable populations without geriatrics.
- Patients who did not agree to participate in the study.
- Observation with illegible information were excluded

PLAN OF THE STUDY:

Preparation of protocol and Data collection form



METHODS OF EVALUATION DIRECT AND INDIRECT HEALTHCARE COSTS:

- Doctor consultation fees
- Laboratory investigation cost
- Medicine cost
- Room services cost

- Food charges
- Surgery cost&Travelling charges of patient's Attender

STATISTICAL ANALYSIS:

Statistical Analysis was done by entering the data into Microsoft Excel spreadsheet (version 2013) and analysed using Microsoft Excel and represented as number and percentage, and the Mean, Standard Deviation was also performed.

III. RESULTS & DISCUSSION:

There are many variations in prescribing patterns of Diabetes mellitus with hypertension

which requires lifelong treatment as enormously increased the burden of chronic diseases and needs much care while choosing drugs. In a tertiary care centre, prescribing pattern are powerful tools to ascertain the role of drugs in society. Hence, there is a need for appropriate, safe, effective and economical study to find out the patterns of drug therapy among diabetic hypertensive patients with other complications.

Table 1: Age wise distribution of cardiovascular disease and Diabetes Mellitus:

Age in years	Demographic Data		
	Number of CV and DM Patients (n=102)		
	Male	Female	Total
31 to 40	2(3%)	3(11%)	5(5%)
41 to 50	20(27%)	6(22%)	26(25%)
51 to 60	25(33%)	7(26%)	32(31%)
61 to 70	18(24%)	8(30%)	26(25%)
71 to 80	8(11%)	3(11%)	11(11%)
81 to 90	2(3%)	0%	2(2%)
Total	75	27	102

In this study total no of population were 102 and the male patient were 75 and female patient were 27. The Age wise distribution represents in Cardiovascular disease and Diabetes mellitus, the male patient were affected in 33% under 51 to 60

age group and female patient were affected in 30% under 61 to 70 age group. Most of the CVD and DM patients were affected at the age group of 51 to 60. In that males (33%) patient more than the female patient (26%).

Table 2: Gender wise Distribution:

Gender	Total Number of Study Population (n=102)	Percentage (%)
Male	75	74%
Female	27	26%

Table 2 represents the total no of study population were 102 and male patients were 75 (74%) and female patients were 27 (26%) in Cardiovascular disease and Diabetes mellitus.

Table 3: Co-morbidities in the study population

Co-Morbidities	Number. of patients (n=102)	Percentage (%)
DM	16	16%
HTN	15	15%
DM, HTN	9	9%

HTN,CAD	1	1%
CAD	3	3%
CAD,DM	1	1%
DM,HTN,CAD	4	4%
NoCo-Morbidities	53	52%

In this study represents the Co-Morbidities as DM (16%), Hypertension (15%), Coronary artery disease (3%), DM + Hypertension (9%), Hypertension + Coronary artery disease (1%), Coronary artery disease

+ Diabetes Mellitus (1%), DM + Hypertension + Coronary artery disease (4%) and No Co-Morbidities as 52% of patients. Most of the patients affected the DM Co-Morbidities (16%)

Table:4 Classification of drugs in the cardiovascular drugs used

Classifications Of Drugs	Number of Cardiovascular drugs (n=470)		Total (%)
	Males	Females	
Diuretics	37(10%)	19(18%)	56(12%)
Beta-Blockers	37(10%)	11(10%)	48(10%)
Ace Inhibitors	17(5%)	9(8%)	26(6%)
Calcium channel blockers	9(2%)	2(2%)	11(2%)
Alpha-Blockers	1(0%)	0(0%)	1(0%)
Angiotensin-2 Inhibitors	11(3%)	5(5%)	16(3%)
Statins	69(19%)	13(12%)	82(17%)
Vasodilators	6(2%)	4(4%)	10(2%)
Antiplatelet Drugs	127(35%)	22(21%)	149(32%)
Anticoagulant	48(13%)	16(15%)	64(14%)
Digitalis Glycosides	0(0%)	2(2%)	2(0%)
Others	2(1%)	3(3%)	5(2%)
Total	364	106	470

Table-4 represents the diuretics drugs prescribed in male (10%) and female (18%), Beta-blockers prescribed in male (10%) and female (10%), ACE inhibitors prescribed in male (5%) and female (8%), CCB prescribed in male (2%) and female (2%), Alpha-blockers prescribed in male (0%) and female (0%), Angiotensin-2 Inhibitors prescribed in male (3%) and female (5%), St

atins prescribed in male (19%) and female (12%), Vasodilators prescribed in male (2%) and female (4%), Antiplatelet prescribed in male (35%) and female (21%), Anticoagulants prescribed in male (13%) and female (15%), Digitalis glycosides didn't prescribed in male and prescribed in female (3%), Other cardiovascular drugs are prescribed in male (1%) and female (3%)

Table:5 Classification of Diabetes mellitus drugs

Classifications Of Drugs	Number of Antidiabetic Drugs (n=35)	
	Males	Females

Sulphonylureas	4(14%)	1(17%)
Biguanides	21(72%)	3(50%)
Insulin	4(14%)	2(33%)

Table-5 represents most of the DM patients treated to the Biguanides drugs prescribed in Male (72%) and Female (50%), Sulphonylurease prescribed in

Male (14%) and Female (17%) ,Insulin prescribed in Male(14%) and Female(33%).

Table:6 Classification of Non-Cardiovascular Drugs

Class of Non-Cardiovascular drugs	Number of Non –Cardiovascular Drugs(n=189)
NSAID's	21(11%)
Antibiotics	15(8%)
Antihistamines	54(29%)
PPI	22(12%)
Sedative drugs	16(8%)
Antiemetic	7(4%)
Others	54(29%)

Table 6 represents the non-cardiovascular drugs prescribed in Cardiovascular disease and Diabetes mellitus patients like NSAIDs were 11%(21 drugs), Antibiotics were 8% (15 drugs) , Antihistamine were 29%(54 drugs), PPI were

12%(22 drugs), Sedative drugs were 8%(16 drugs) , Antiemetic were 4%(7 drugs) and others were 29%(54 drugs) and total no of non-cardiovascular drugs used 189.

Table: 7 Cost Of Illness Analysis both Direct and Indirect Cost of Myocardial Infarction in Cardiovascular Disease:

Disease		Average per patient (in rupees)	Myocardial Infarction (in rupees)
Direct Cost	Consultation Fee	3896	171400
	Diagnostic Cost	16627	731590
	Room Cost	7381	324750
	Medicine Cost	1816	79,919
	Food Cost	2607	114700
	Surgery	839386	36933000
	Others	15886	698993
	Travel Cost	890	39,160
Indirect Cost		4034	177500
Total Cost		892523	39271012

In this study represents cost-of-illness analysis of Cardiovascular disease patients and their total direct costs in rupees (39,09,3512) like consultation fees (1,71,400), Diagnostic cost (7,31,590), Room cost (3,24,750), Medicine

cost (79,919), Food cost (1,14,700), Surgery cost (3,69,33,000), Travel cost (39,160) and others like duration of drug usage wages (6,98,993) and indirect cost of DM patients were 1,775,00 rupees.

Table: 8 Type of Drug-drug interaction

Drug Interaction	Outcomes	Mechanisms	Number of drug interaction (%)
Antiplatelet drugs + Anticoagulant	Increased risk of bleeding	Pharmacodynamic mechanism	39(29%)
Antiplatelet drugs + Diuretics	Clopidogrel increase level of Torsamide by slowing the drug metabolism	Pharmacokinetic mechanism	29(19%)
ACE-inhibitors + Antiplatelet Therapy	Combining this drugs increase the risk of high potassium blood levels	Pharmacodynamic mechanism	17(11%)
ACE Inhibitors + Diuretics	Combining this both drugs may cause low blood pressure and reduce kidney function	Pharmacodynamic mechanism	15(10%)
Betablockers + Diuretics	Atenolol increase and Torsamide decrease the potassium levels in the blood	Pharmacodynamic mechanism	15(10%)
PPI + Antiplatelets	Clopidogrel taken in this combination may inhibit platelet aggregation (Increased risk of bleeding)	Pharmacokinetic drug metabolism	8(5%)

Antiplatelet +ARB	Telmisartan and aspirin both increase the potassium level in the blood, may reduce the kidney function and particularly in elderly or volume depleted individuals	Pharmacodynamic mechanism	7(5%)
Anticoagulant ACE Inhibitors	Heparin and Enalapril drugs increase the risk of high potassium levels in the blood	Pharmacodynamic mechanism	7(5%)
ARB+Statins	Telmisartan and Atorvastatin may	Pharmacodynamic mechanism	7(5%)

In this study represents most of the CVD and DM patients were affected. Antiplatelet+Anticoagulant drug-drug interaction (39), in addition Antiplatelet+Diuretic

s drug-drug interaction (29) and least drug-drug interaction class were ACE inhibitors+Biguanides (6).

Table:9 Scheme Utilization

Scheme utilization	Male	Female	Total	Percentage
Cash	54	18	72	71%
Scheme	21	9	30	29%

Table 9 represents the total study population is cash usage is 71% and scheme utilization is 29%.

Table:10 various drugs prescribed for diabetes with Cardiovascular Disease

Class of Drugs	Drugs	Number of Drugs (N=429)	Percentage (%)
Beta blockers	Atenolol	2	0.5%
	Metoprolol	29	7%
	Carvedilol	10	2%
	Bisoprolol	5	1%
	Labetalol	1	0.23%
ARB	Losartan	2	0.5%
	Telmisartan	14	3%
Vasodilators	Isosorbide mononitrite	1	0.23%
	Nitroglycerine	1	0.23%
Antiplatelet	Clopidogrel	40	9%
	Aspirin	75	17%
	Prasugrel	13	3%
	Ticagrelor	27	6%

Statin	Rosuvastatin	8	2%
	Atorvastatin	7	2%
AntiHyperlipedimic Agents	Ezetimibe	1	0.23%
Diuretics	Torseמידe	27	6%
	Furosemide	8	2%
Cardiacglycosides	Digoxin	2	0.5%
ACEinhibitors	Enalapril	24	6%
	Ramipril	1	0.23%

Table 10 represents the Diabetes mellitus with cardiovascular disease patientsprescribed drugs are Beta blockers, ARB ,Vitamins, Antiplatelet ,Statin, Antibiotics,NSAIDs,PPI,Anti-

histamines,Anti-psychiatric,Anticoagulant,Antiemetic,Vasodilators ,Diuretics also.

Table:11CategorywisedistributionofDrugInteractions

CategoryOfDrugInteraction	Numberof DrugInteractions(n=230)(%)
Serious	25(11%)
Moderate	194(84%)
Minor	11(5%)

Over all this study represents most of the patients affected in moderate Druginteractionswere84%(194),inaddition seriousdruginteractionswere11%(25)andfinallyMinor druginteractions were5%(11).

Table:12TotalcostofdirectandIndirectinCardiovascularDisease

Disease	Direct cost(inrupees)	Indirect cost(inrupees)	Total(inrupees)
Hypertension	35,139		35,139
CompleteHeartBlock	64,004		64,004
Myocardialinfarction	5,893,132	198,000	6,091,132
MyocardialInfarction+CompleteHeart Block	355,744		355,744
Atherosclerosis	37,059		37,059
AcuteCoronarySyndrome	536,554	13,800	550,354
Angina	255,090	10,700	265,790
CoronaryArteryDisease	3,569,365	328,000	3,897,365
DilatedCardioMyopathy	130,330		130,330
SupraventricularTachycardia	23,886	3,500	27,386
CongestiveCardiacFailure	127,438	7,000	134,438
Total			11,390,741
Average			1,898,457

Median		134,438
StandardDeviation		1966359.545

Table 12 represents cost of illness analysis direct and indirect cost (in rupees) of Cardiovascular disease study population, mean \pm SD of this COA of cardiovascular disease study

population is **1,898,457 \pm 1,966,359**. COA of this cardiovascular disease study population most cost utilized is the Myocardial infarction patients compared with other cardiovascular disease patients.

Table:13 Total Costs of Direct and Indirect in Diabetes Mellitus

Disease		Diabetes Mellitus (n=4) (In Rupees)
Direct Cost	Consultation Fee	12,600
	Diagnostic Cost	15,120
	Room Cost	22,750
	Medicine Cost	49,525
	Food Cost	8,000
	Surgery	18,000
	Others	19,953
	Travel Cost	1,400
Indirect Cost		9,000
Total Cost		1,56,348

In this study represents cost-of-illness analysis of Diabetes mellitus patients and their total direct costs (1,47,348 rupees) like consultation fees (12,600 rupees), Diagnostic cost (15,120 rupees), Room cost (22,750 rupees), Medicine

cost (49,525 rupees), Food cost (8,000 rupees), Surgery cost (18,000 rupees), Travel cost (1,400 rupees) and others like duration of drug usage wages (1,400 rupees) and indirect cost of DM patients were 9,000 rupees.

Table:14 Total costs of Direct and Indirect of cardiovascular with Diabetes Mellitus

Disease	Direct cost (In Rupees)	Indirect cost (In Rupees)	Total (In Rupees)
Diabetes Mellitus+ Hypertension	3,000	3,000	6,000
Diabetes Mellitus+ Mitral Valve Replacement	158,992	3,000	161,992
Diabetes Mellitus+ Angina	55,117	4,800	59,917

Table 14 represents as DM+HTN cost is 6000 rupees, DM+Mitral valve replacement cost is 161,992 rupees and DM+Angina cost is 59,917 rupees.

Table:15 Drug Utilization Evaluation of Cardiovascular disease

WHO core prescribing indicators	Cardiovascular Diseases
No. of Medicines/Prescriptions	671
No. of Generic name/ prescriptions	106
No. of Antibiotics/prescriptions	9

No.ofInjections/prescriptions	95
No.ofEssentialdrugs	491

In this study represents Drug Utilization Evaluation of Cardiovascular disease as analyzed through the WHO core prescribing indicators like No of medicines perprescription in CVD study population as 671 medicines, No of generic name

perprescriptioninCVDstudypopulationas106medicines, NoofantibioticsperprescriptionsinCVDstudypopulationas9, NoofinjectionsperprescriptionsinCVDstudypopulation as 95and No ofessential drugsin CVDstudypopulation as 491 .

Table:16DrugutilizationevaluationsofDiabetesmellitusinstudyPopulation

WHOcoreprescribingindicators	Diabetesmellitus
No.ofMedicines/Prescriptions	24
No.ofGenericname/ prescriptions	0
No.ofAntibiotics/prescriptions	4
No.ofInjections/prescriptions	10
No.ofEssentialdrugs	17

Table16representsDrugUtilizationEvaluationofdiabetesmellitusasanalyzedthroughtheWHOcoreprescribingindicatorslikeNumberofmedicinesperprescriptioninDMstudypopulationas24drugs, Number

ofantibioticsperprescriptionsinDMstudypopulationas4drugs, Numberofinjectionsperprescriptionsin DM study population as 10 drugs and Number of essential drugs in DM studypopulationas 17drugs.

Table:17DrugUtilizationEvaluationofCardiovasculardiseaseandDiabetes Mellitusinthis

WHOcoreprescribingindicators	Numberofdrugs	Percentage
No.ofMedicines/Prescriptions	713	100%
No.ofGenericname/prescriptions	109	15%
No.ofAntibiotics/prescriptions	15	2%
No.ofInjections/prescriptions	114	16%
No.ofEssentialdrugs	524	73%

Table 17 represents Drug Utilization Evaluation of Cardiovascular disease anddiabetesmellitusasanalyzedthroughtheWHOcoreprescribingindicatorslikeNumberofmedicinesperprescriptioninthisstudypopulationas713(100%)medicines, Numberof generic name per prescription in this total study population as 109(15%) medicines, Number of antibiotics per prescriptions in this total study population as 15(2%) ,Number of injections per prescriptions in this study population as 114(16%) andNumberofessential drugs in this studypopulationas 524(%) .

study, the prevalence of coronary artery disease was high.

- The most commonly prescribed categories of cardiovascular drugs were Anti hypertensives and Antiplatelets indicating a high prevalence of hypertension and coronary artery disease in patients. Extensive polypharmacy was noticed in the prescriptions.
- Diabetes is a chronic and potentially disabling disease that represents an important public health and clinical concern. This study reflected the actual utilization of different anti-diabetic drug classes in diabetes patients. It was observed that among the anti-diabetic drugs, most frequently prescribed drug was Glimpiride followed by Metformin, Gliclazide, and least prescribed drug was Human Insulin in monotherapy. Surprisingly, none of the prescriptions were in accordance to

IV. CONCLUSION:

- In the present study it was observed that cardiovascular disease was more common in male compared to female and the risk for CVD increased with increasing age. In the present

the WHO/ American Diabetic Association (ADA) guidelines for the utilization of anti-diabetics as a monotherapy. It is needless to mention that such guidelines are aimed to achieve a cost effective therapy and for choosing treatment alternatives. Physicians should follow such guidelines so as to attain therapeutic goals.

RECOMENDAT IONS & FUTURE DIRECTIONS

- In future, similar studies can be planned in Government hospital settings. Studies for evaluating defined daily doses (DDD) for anti-hyperglycemic & cardiovascular drugs agents can also be envisaged.
- Further studies from time to time are required in drug prescription pattern and standard treatment guidelines should be circulated amongst practicing physicians to encourage rational prescription.
- This analysis and previous cost effectiveness studies have focused on the influence of traditional risk factors in the development of cardiovascular events, but—because of the increasing prevalence of diabetes—an integrated diabetes-cardiovascular disease approach should be pursued in future.

LIMITATIONS:

- The sample was not random and could have been a potential source of bias.
- This is only Prospective Observational Study if a cross-sectional study in Multiple center studies can demonstrated a Clear idea
- Glycosylated HbA1c was not recorded but recommended in the study

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