

Prescription audit study in a tertiary care hospital using the anatomical therapeutic chemical and defined daily dose classification concept

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ABSTRACT

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Background: In 1996, WHO recognized the need to develop and use the anatomical therapeutic chemical (ATC) and defined daily dose (DDD) as an international standard for utilization studies wherein it appointed experts in clinical pharmacology, drug utilization, drug regulation, drug evaluation, statistics, and medicine to monitor drug utilization on a standard and uniform pattern around the world.

Methods: All the prescriptions presented to main dispensary of the hospital from January-December 2013 in mid-month period of 10 days were taken for the analysis in terms of pharmacovigilance, polypharmacy, drug usage group wise, pharmacoeconomics, and drug utilization in DDD/1000/day along with ATC classification using customized software.

Results: 35,808 prescription forms containing 121,303 drugs were presented to main dispensary of the hospital across which medicines were dispensed. Polypharmacy was calculated to be 3.27 ± 0.44 standard deviation. Approximately, 60% of total prescription forms analyzed were lacking essential and basic such information as age, gender, or diagnosis. Pharmacoeconomics of this study indicates that a major budget was mainly spent on drug groups belonging to - metabolic, antibiotics, hormones, and related drugs. A voluminous consumption of drugs belonged to drug groups - metabolic drugs like statins, hormones, and related drugs such as various oral hypoglycemics, anti-hypertensives, psychotropics, and proton pump inhibitors or H₂ blockers, all indicative toward stressful and sedentary lifestyle of modern era and unhealthy lifestyles, a trend of increasing morbidity and mortality, reducing significantly the productive man-hours.

Conclusions: This study was undertaken in a government tertiary care hospital which gives insight into the day to day functioning status of our health care delivery system. It is an opportunity on self-assessment in further improving the health care delivery by implementing measures which economizes on scarce health care budget as well as minimizing the common prescription errors.

Keywords: Defined daily dose use, Drug utilization, Pharmacovigilance, Prescription audit

INTRODUCTION

Quality assessment of medical care rendered to patients is the need of time and can only be assessed by a prescription audit, because the same is based on documentary evidence which supports elements of prescription writing having there in diagnosis, treatment, and justified details. In principle, it is prudent for physicians to have an objective and systemic way of analysis of treatment and care provided to the patient.¹ Thus, medical audit may be defined as looking at what one is doing with the aim of providing patient care and use of resources.² Prescription audit is a tool designed for the objective assessment by and to the medical care provider of

meeting their own standards in their professional day to day care. Hence, prescription audit is a tool as well as a technique and its application is science as well as an art.³

In order to measure drug use, it is important to have both a classification system and a unit of measurement which preferably should be standardized and acceptable to all the agencies in the world for uniform comparisons of results. This need was formulated in the late sixties, which still was in the infancy period in the field of drug utilization research. Due to an increasing interest in this field, a symposium entitled The Consumption of Drugs took place in Oslo in 1969 organized by the WHO Regional Office for Europe.

It was at this symposium the Drug Utilization Research Group was established tasked with the development of internationally applicable methods for drug utilization research.⁴

Inspired by this interest, a system named anatomical therapeutic chemical (ATC) classification was developed by the Norwegian Medicinal Depot. Norwegian researchers also developed a technical unit of measurement called the defined daily dose (DDD) to be used in drug utilization studies.⁵ The ATC/DDD system has been used in Norway since the early seventies for presenting drug consumption data.

In 1996, when the decision on globalization the ATC/DDD system was taken, the WHO Division of drug Management and Policies in Geneva established the WHO International Working Group for Statistics Methodology. This working group now consists of 12 WHO appointed experts in clinical pharmacology, drug utilization, drug regulation, drug evaluation, statistics, and medicine. The members of the working group represent different users of the ATC/DDD system and different nationalities as they represent the 6 WHO global regions. The following countries are represented in the working group: Australia, Japan, India, Pakistan, Morocco, Ghana, Zimbabwe, USA, Argentina, The Netherlands, and Sweden. The working group meets twice annually. The WHO Collaborating Centre for Drug Statistics Methodology receives expert advice from the working group and acts as secretariat for the group.

In developing countries, the constantly increasing number of medicine list and treatment option serves to increase the irrational medicine use and cost that ultimately leads to poor patient outcome and significant wastage of money and resources. Cost of each such irrational drug use is enormous in terms of both scarce resources and adverse clinical consequences of therapies that may have real risks but no objective benefits. Now a days, the prescribing pattern is changing and has become just an indication for the medicine with some instructions of doses without considering its rationality.⁶ The study of prescribing pattern is that part of medical audit which seeks to monitor, evaluate, and if necessary suggest modifications in prescribing practices of medical professionals so as to make medical care rational and cost effective.^{7,8} Prescription audit also forms part of drug utilization studies which audits prescribing indicators such as average consultation time, average time for dispensing, if all drugs actually are dispensed, drugs if adequately labeled/legibly written, patients' knowledge if has understood the usage of drugs as told by pharmacist and facility indicators like availability of copy of essential drugs list/drug formulary of hospital, VED analysis etc.⁹

Prescription is an order to the pharmacist for scientific medication to the patient at a particular time for the disorder summarized for logical diagnosis.¹⁰ This focuses

at the diagnostic acumen and therapeutic proficiency of the clinician with the necessary instruction for palliation or restoration of the patient's health.¹² Thus, quality-of-life can be improved by enhancing the standards of medical treatment at all levels of health care delivery systems.¹³ Medical audit overseas the observance of these standards. Setting standards and assessing the quality of medical care through performance review should become part of everyday clinical practice.¹⁴

In India, several research groups have initiated efforts to study the drug utilization patterns mostly in the last three decades. Most of these groups are from pharmacology departments of medical institutions, pharmacy colleges, newly established pharmacy practice departments, and other independent agencies.

Various studies in different parts of the World conducted on the prescription auditing¹⁵⁻¹⁸ produced their own database for the future comparative study. Study of drug prescription and its utilization in India have been reported at both hospital¹⁹⁻²¹ and community levels.^{22,23} Most of these studies have measured drug utilization by counting of prescriptions which indicates actually prescribing frequency of drug prescription. To have uniformity of drug utilization studies at various places, DDD is of great value as compared to traditional units of measurements of drugs utilization by weight or cost. DDD is of great value as compared to traditional units of measurement of drugs utilization by weight or cost. Indian studies have used DDD concept and focused on drug utilization in selected groups of patients for a brief period.²⁴⁻²⁶ We undertook this study at one of the tertiary care hospitals for a period of 1 year based on the prescriptions presented to dispensary and classified the drugs by ATC/DDD methodology to determine the prevalence of prescribing pattern of all major groups of drugs. An attempt was also made to associate prescription data with two demographic characteristics, viz, age and sex, to develop patient population database.

Statistics

Student's t-test was applied as a measure of dispersion and other figures are in percentage being a descriptive study.

METHODS

Study place: Tertiary Care Government Hospital, Pune.

Study design: Prospective descriptive study.

Ethical consideration: Approved from Institutional Ethical Committee.

Study duration: 01 January, 2013-31 December, 2013.

Methodology

Prescription forms which were presented to the dispensary of the hospital by the patients for collecting their medicines from January 2013 to December 2013 in the mid-month period of 10 days, i.e., from 11th to 20th of each month were taken into account for analysis. Since the period of study was one full calendar year, the data were consolidated every 03 monthly, i.e., Jan-Mar, Apr-Jun, Jul-Sep, Oct-Dec.

RESULTS

Pharmacovigilance

All prescriptions from 11th to 20th of each month were counted and analyzed for their completeness. The prescriptions which were incomplete on account of lacking elementary parameters such as age, gender, and diagnosis were labeled incomplete. Further, the prescription forms were also categorized into having generic or brand name of drugs in them. This was also further categorized as prescriptions originating from specialists, residents, and general duty medical practitioners (Tables 1-4).

Polypharmacy

All ages of patients were categorized into eight categories as 0-10, 11-20, 21-30, 31-40, 41-50, 51-60, 61-70, and >70 year. The prescriptions forms having different number of drugs in them were tabulated to analyze the most common age group of people having the highest number of drugs prescribed per prescription (Figures 1-4).

Drug usage group wise

All ages of patients were categorized into eight categories as above. The drugs in the prescription forms were subdivided into nine major categories, viz. cardiovascular (CVS), nonsteroidal anti-inflammatory drugs (NSAIDs) including analgesics and antipyretics, anti-microbial agents (AMA), central nervous system, gastrointestinal (GIT), respiratory system and anti-allergics, hormones

and related drugs, nutrition and vitamins, and lastly metabolic group of drugs. This gave an indication of the highest prescribed group of drugs in particular age group of patients (Figures 5-8).

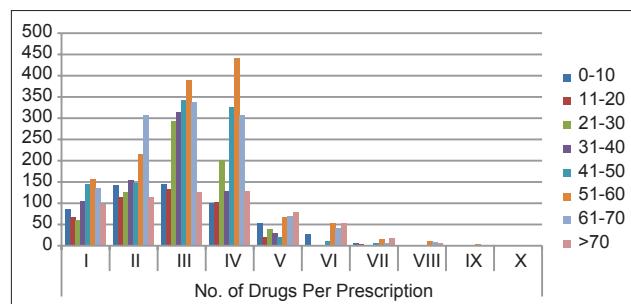


Figure 1: Polypharmacy for period: January 2013-March 2013.

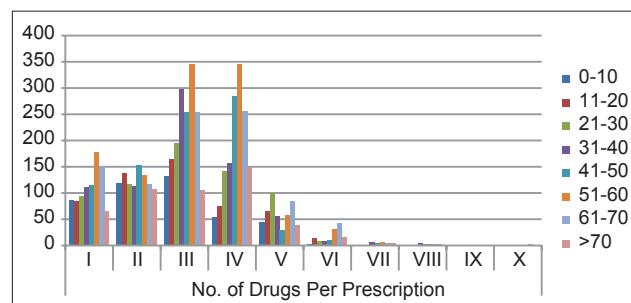


Figure 2: Polypharmacy for period: April 2013-June, 2013.

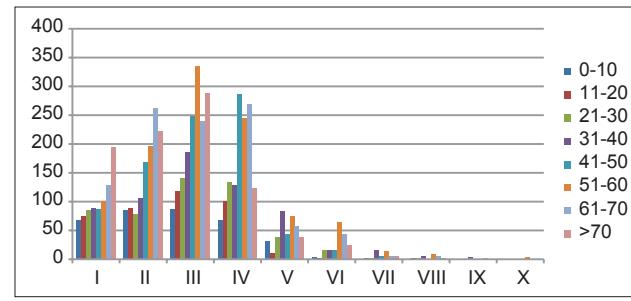


Figure 3: Polypharmacy for period: July 2013-September, 2013

Table 1: Pharmacovigilance for period: January 2013-March 2013.

Total prescription slips	9654	Contains	32464 drugs
Prescription slips complete	3625	Not complete	6029
Specialist (%) n=3803 (63.08)	Residents (%) n=2149 (35.64)	GDMP (%) n=77 (1.28)	
Prescriptions lacking age	1399 (36.79)	883 (41.09)	39 (50.47)
Prescriptions lacking gender	383 (10.07)	241 (11.21)	13 (16.88)
Prescriptions lacking diagnosis	3572 (93.93)	2028 (94.37)	70 (90.91)
Prescriptions in brand name	4160 (36.21)	2007 (34.87)	58 (34.12)
Prescription in generic name	7330 (63.79)	3749 (65.13)	110 (64.71)

GDMP: General duty medical practitioners

Table 2: Pharmacovigilance for period: April 2013-June 2013.

Total prescription slips	8832	Contains	29392 drugs
Prescription slips complete	3104	Not complete	5728
	Specialist (%) n=3574 (61.92)	Residents (%) n=2162 (37.74)	GDMP (%) n=19 (0.33)
Prescriptions lacking age	775 (21.85)	367 (16.98)	4 (21.05)
Prescriptions lacking gender	304 (8.57)	141 (6.52)	3 (15.79)
Prescriptions lacking diagnosis	3219 (90.75)	2063 (95.42)	18 (94.74)
Prescriptions in brand name	3885 (35.81)	1925 (33.13)	12 (33.33)
Prescription in generic name	6963 (64.19)	3886 (66.87)	24 (66.67)

GDMP: General duty medical practitioners

Table 3: Pharmacovigilance for period: July 2013-September 2013.

Total prescription slips	8557	Contains	28916 drugs
Prescription slips complete	3389	Not complete	5168
	Specialist (%) n=3469 (67.12)	Residents (%) n=1659 (32.10)	GDMP (%) n=40 (0.77)
Prescriptions lacking age	883 (24.02)	377 (22.70)	13 (32.10)
Prescriptions lacking gender	128 (3.69)	78 (4.70)	5 (12.50)
Prescriptions lacking diagnosis	3260 (93.98)	1592 (95.96)	37 (92.50)
Prescriptions in brand name	4166 (34.35)	1599 (31.61)	33 (37.08)
Prescription in generic name	7961 (65.65)	3460 (68.39)	55 (61.80)

GDMP: General duty medical practitioners

Table 4: Pharmacovigilance for period: October 2013-December 2013.

Total prescription slips	8765	Contains	30531 drugs
Prescription slips complete	3280	Not complete	5485
	Specialist (%) n=3467 (63.21)	Residents (%) n=2003 (36.52)	GDMP (%) n=15 (0.27)
Prescriptions lacking age	1011 (29.16)	521 (26.01)	7 (46.48)
Prescriptions lacking gender	174 (5.02)	100 (4.99)	2 (13.33)
Prescriptions lacking diagnosis	3271 (94.35)	1915 (95.61)	14 (93.33)
Prescriptions in brand name	3986 (34.67)	1834 (31.87)	13 (39.39)
Prescription in generic name	7510 (65.33)	3920 (68.13)	20 (60.61)

GDMP: General duty medical practitioners

Gender wise usage of drug groups

Furthermore, a comparison of drug groups' usage was done for eight age groups as above to know the morbidity pattern in the population (Tables 5-8).

Pharmacoeconomics

Drugs' price was taken from the procurement cost of our hospital which is, of course, much lower than the retail price as printed on the drug packages. An estimate was made of the cost factor involved on health care based on drug groups, as above, in the four quadrants of a year for 10 days of each month (Figures 9 and 10).

Drug utilization

All the drugs which are considered in the study are classified using ATC concept and their consumption pattern in all the four quadrants are expressed in conventional pattern of percentage as well as scientifically in accordance of WHO recommendations using DDD concept. The recommended formula by WHO for drug utilization pattern is (Table 9):

$$\text{DDD / 1000 / day} = \frac{\text{Total number of dosage strengths of units prescribed} \times \text{each dosage unit} \times 1000}{\text{DDD} \times \text{Duration of study} \times \text{Total sample size}}$$

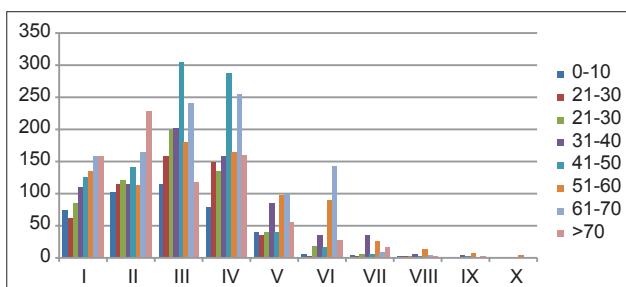


Figure 4: Polypharmacy for period: October 2013-December 2013.

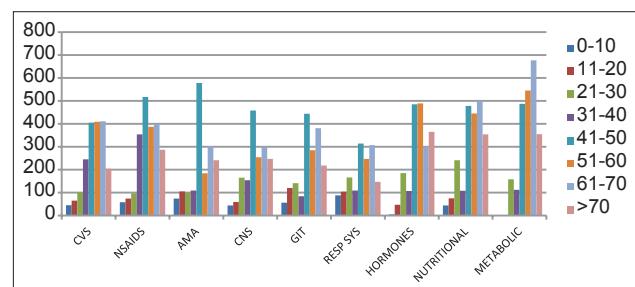


Figure 7: Drug usage groupwise for the period: July 2013-September 2013.

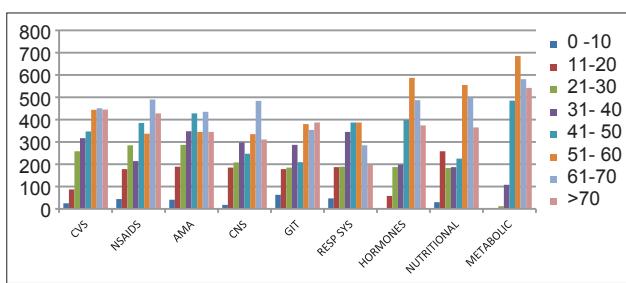


Figure 5: Drug usage groupwise for the period: January 2013-March 2013.

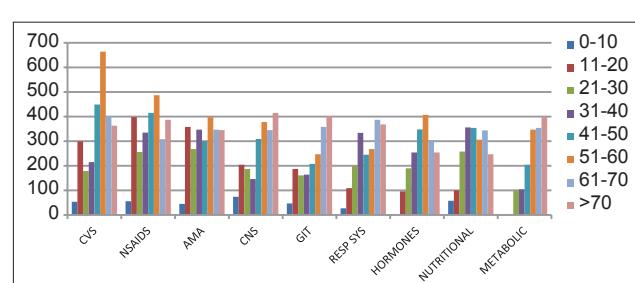


Figure 8: Drug usage group wise for the period: October 2013-December 2013.

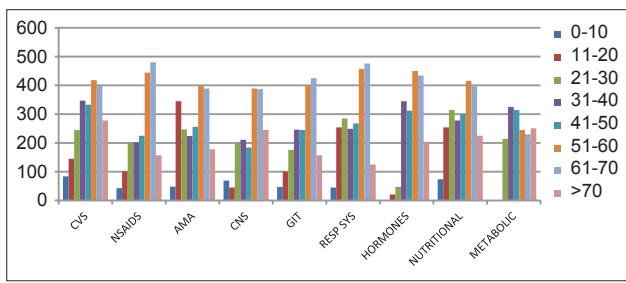


Figure 6: Drug usage groupwise for the period: April 2013-June 2013.

Table 5: Genderwise prescribing frequency for drug groups for a period: January 2013-March 2013.

Drug groups	Number of patients (% of the population)	
	Male n=5480 (60.77)	Female n=3537 (39.23)
CVS	2100 (38.32)	1000 (28.87)
NSAIDs	1987 (36.26)	1117 (31.58)
AMA	1748 (31.90)	1018 (28.78)
CNS	1348 (24.60)	797 (22.53)
GIT	1254 (22.88)	666 (18.83)
Respiratory system and anti-allergics	1645 (30.02)	983 (27.79)
Hormones and related drugs	1452 (26.50)	837 (23.66)
Nutritional	2458 (44.85)	1469 (41.53)
Metabolic	1648 (30.07)	765 (21.63)

CVS: Cardiovascular, NSAIDs: Nonsteroidal anti-inflammatory drugs, AMA: Anti-microbial agents, CNS: Central nervous system, GIT: Gastrointestinal

Table 6: Genderwise prescribing frequency for drug groups for a period: April 2013-June 2013.

Drug groups	Number of patients (% of the population)	
	Male n=5480 (60.77)	Female n=3537 (39.23)
CVS	1645 (36.74)	1048 (27.75)
NSAIDs	1437 (32.10)	1164 (29.79)
AMA	1478 (33.01)	1106 (28.31)
CNS	1141 (25.49)	854 (21.86)
GIT	1314 (29.35)	1101 (28.18)
Respiratory system and anti-allergics	1393 (31.12)	1070 (27.39)
Hormones and related drugs	1154 (25.78)	759 (19.43)
Nutritional	1648 (36.81)	1435 (36.73)
Metabolic	1125 (25.13)	836 (21.40)

CVS: Cardiovascular, NSAIDs: Nonsteroidal anti-inflammatory drugs, AMA: Anti-microbial agents, CNS: Central nervous system, GIT: Gastrointestinal

DISCUSSION

A prescription is a prescriber's order to dispense a specific treatment - Usually, medication for a specific patient. Thus, it needs no further emphasis that for error-free dispensing, all prescription order forms should be legible, unambiguous, dated and signed with a rubber stamp of prescriber bearing his/her registration number for an optimal communication between prescriber, pharmacist, and nurse/health care providers. Many common errors include lack of patients

Table 7: Genderwise prescribing frequency for drug groups for a period: July 2013-September 2013.

Drug groups	Number of patients (% of the population)	
	Male n=4578 (59.51)	Female n=3115 (40.49)
CVS	2054 (44.87)	1112 (35.70)
NSAIDs	1987 (43.40)	1190 (38.20)
AMA	1648 (36.00)	1184 (38.01)
CNS	1456 (31.80)	865 (27.77)
GIT	1289 (28.16)	873 (28.03)
Respiratory system and anti-allergics	1485 (32.44)	1034 (33.19)
Hormones and related drugs	1647 (35.98)	949 (30.47)
Nutritional	1875 (40.96)	1263 (40.55)
Metabolic	1568 (34.25)	910 (29.21)

CVS: Cardiovascular, NSAIDs: Non-steroidal anti-inflammatory drugs, AMA: Anti-microbial agents, GIT: Gastrointestinal, CNS: Central nervous system

Table 8: Genderwise prescribing frequency for drug groups for period: October 2013-December 2013.

Drug groups	Number of patients (% of the population)	
	Male n=4017 (55.59)	Female n=3209 (44.41)
CVS	2154 (47.05)	871 (27.96)
NSAIDs	2147 (46.90)	1268 (40.71)
AMA	1987 (43.40)	912 (29.28)
CNS	1789 (39.08)	906 (29.09)
GIT	1487 (32.48)	662 (21.25)
Respiratory system and anti-allergics	1547 (33.79)	1003 (32.20)
Hormones and related drugs	1385 (30.25)	694 (22.28)
Nutritional	1854 (40.50)	1132 (36.34)
Metabolic	1466 (32.02)	748 (24.01)

CVS: Cardiovascular, NSAIDs: Non-steroidal anti-inflammatory drugs, AMA: Anti-microbial agents, GIT: Gastrointestinal, CNS: Central nervous system

particular such as self or relative, age, gender in the superscription whereas diagnosis, dosage and frequency of drug administration are other few. Prescribing in a generic name offers the pharmacist flexibility in selecting drug or substitutes of the same in case of possibility of near similar drug if available. However, it should not be assumed that every generic drug product is as satisfactory as the trade-named product, though most generics are satisfactory. Bioavailability for the products varies between manufacturers and sometimes between different lots of drug produced by the same manufacturer. It thus produces

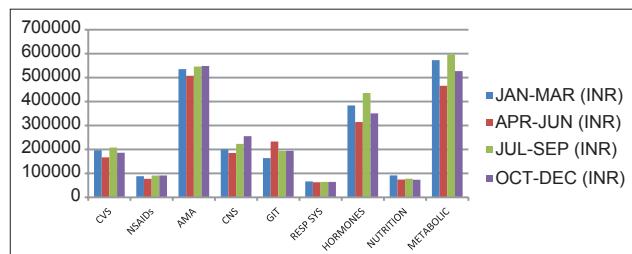


Figure 9: Pharmacoconomics - drug group wise: January 2013-December 2013.

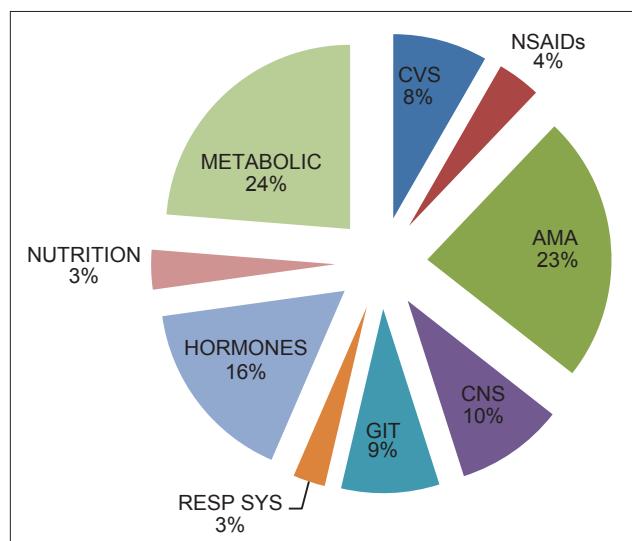


Figure 10: Pharmacoconomics - expense on various drug groups (%): January 2013-December 2013.

same drug but having different biochemical equivalence which is specifically relevant for drugs having a narrow therapeutic margins and some chronic drug therapies like various psychiatric agents and anti-epileptics, etc. Thus, in case of very small number of drugs which usually have a low therapeutic index, poor solubility or a high ratio of inert ingredients to active drug content, a specific manufacturer's product may give a more consistent result. In this study, it was found that approximately two-third of prescriptions was in generic name and rest in brand names optimizing the balance between the two.

Average number of drugs per prescription form is an important index of prescription audit as it forms the basis for review and educational intervention in day to day prescribing practices of doctors. It is preferable to keep the mean number of drugs per prescription as low as possible since higher figures may lead to increased risk of drug interactions, as well as unnecessary economic burden. Most importantly for antimicrobials, such overuses increase the chance of bacterial resistance. In this study, we have found average three drugs per prescription form which is within acceptable limits. Other studies based on community practices reveal a mean of two drugs per prescription form.²⁷ Some hospital-based studies in India reported a figure of 3-5 drugs per prescription form.¹⁹⁻²¹ By and large, it is difficult to keep the number of drug per prescription

Table 9: Drug utilization in percentage and DDD/1000/day.

Drug	ATC	Jan-Mar		Apr-Jun		Jul-Sep		Oct-Dec	
		%	DDD	%	DDD	%	DDD	%	DDD
Category: CVS									
Acenocoumarol	B01AA07	0.15	0.46	0.17	0.51	0.18	0.57	0.15	0.45
Acetazolamide	S01EC01	0.06	0.32	0.05	0.23	0.06	0.30	0.06	0.32
Amiodarone	C01BD01	0.04	0.29	0.04	0.28	0.04	0.32	0.05	0.39
Amlodipine	C08CA01	1.50	49.77	1.49	48.80	1.15	38.31	1.53	50.37
Aspirin	B01AC06	0.69	11.39	0.67	10.92	0.68	11.25	0.65	10.66
Atenolol	C07AB03	1.33	14.73	1.32	14.36	1.30	14.36	1.51	16.51
Carvedilol	C07AG02	0.27	1.11	0.30	1.22	0.28	1.16	0.27	1.11
Clonidine	C02AC01	0.04	0.14	0.04	0.14	0.05	0.16	0.05	0.16
Clopidogrel	B01AC04	0.27	8.86	0.25	8.13	0.27	9.14	0.27	8.86
Colspirin	B01AC06	0.26	8.59	0.27	8.73	0.27	8.86	0.27	8.86
Digoxin	C01AA05	0.14	2.23	0.10	1.48	0.12	1.85	0.14	2.13
Diltiazem	C08DB01	0.14	0.03	0.13	0.02	0.16	0.03	0.13	0.03
Dipyridamole	B01AC07	0.01	0.05	0.02	0.11	0.02	0.10	0.02	0.13
Enalapril	C09AA02	1.05	17.32	1.00	16.34	1.05	17.38	1.03	16.95
Ethamsylate	B02BX01	0.15	7.36	0.11	5.65	0.18	9.00	0.13	6.64
Felodipine	C08CA02	0.27	5.91	0.21	4.52	0.30	6.54	0.26	5.72
Frusemide	C03CA01	1.21	40.09	1.17	38.25	1.29	42.95	1.20	39.44
Hydrochlorthiazide	C03AA03	0.06	0.95	0.07	1.13	0.09	1.50	0.05	0.90
Indapamide	C03BA11	0.14	2.16	0.12	1.83	0.19	2.99	0.13	2.00
Isopropamide	A03AB09	0.02	0.41	0.03	0.45	0.04	0.61	0.03	0.42
ISMO	C01DA08	0.35	2.90	0.34	2.79	0.43	3.58	0.33	2.70
Lisinopril	C09AA03	0.05	0.75	0.08	1.36	0.09	1.50	0.04	0.62
Losartan	C09CA01	0.33	5.52	0.30	4.89	0.35	5.79	0.32	5.19
Metoprolol	C07AB02	0.63	3.45	0.62	3.36	0.68	3.75	0.59	3.23
Mexilitine	C01BB02	0.16	0.49	0.14	0.42	0.16	0.51	0.14	0.43
Nifedipine	C08CA05	0.16	10.36	0.17	11.45	0.16	10.64	0.15	9.96
Pentoxiphylline	C04AD03	0.16	0.31	0.18	0.35	0.16	0.33	0.15	0.29
Prazosin	C02CA01	0.22	1.47	0.23	1.48	0.25	1.64	0.21	1.41
Propranolol	C06AA05	0.93	3.58	0.92	3.51	0.92	3.58	0.93	3.55
Ramipril	C09AA05	1.61	26.59	1.60	26.20	1.68	27.95	1.60	26.29
Ticlopidine	B01AC05	0.33	5.52	0.29	4.67	0.33	5.52	0.29	4.84
Verapamil	C08DA01	0.06	0.32	0.17	0.90	0.07	0.39	0.06	0.32
Category: Musculoskeletal									
ASA	N02BA01	1.93	22.39	1.93	22.14	1.95	22.67	2.00	22.96
Celecoxib	M01AH01	0.25	1.66	0.29	1.93	0.27	1.77	0.28	1.83
Diclofenac sodium	M01AB05	2.42	40.02	0.29	37.50	2.41	40.02	2.50	41.10
HCQS	P01BA02	0.09	1.34	0.11	1.69	0.09	1.34	0.11	1.61
Ibuprofen	M01AE01	2.12	35.05	2.16	35.32	2.35	39.13	2.27	37.37
Indomethacin	M01AB01	0.16	1.17	0.16	1.19	0.15	1.14	0.19	1.40
Ketorolac	M01AB15	0.06	0.21	0.07	0.25	0.06	0.21	0.06	0.23
Mefenamic acid	M01AG01	0.14	0.84	0.13	0.76	0.13	0.74	0.16	0.90
Meloxicam	M01AC06	0.68	3.73	0.66	3.59	0.65	3.59	0.72	3.97
Naproxen	M01AE02	0.51	3.95	0.52	4.01	0.53	4.10	0.55	4.20
Paracetamol	N02BE01	2.40	19.91	2.43	19.92	2.41	20.01	2.48	20.34
Piroxicam	M01AC01	0.51	3.95	0.46	3.55	0.54	4.20	0.55	4.20

Contd...

Table 9: Contd...

Drug	ATC	Jan-Mar		Apr-Jun		Jul-Sep		Oct-Dec	
		%	DDD	%	DDD	%	DDD	%	DDD
Rofecoxib	M01AH02	1.52	11.74	0.74	5.62	1.50	11.68	1.52	11.66
Category: AMA									
Acyclovir	J05AB01	0.06	0.57	0.06	0.63	0.07	0.74	0.06	0.58
Albendazole	P02CA03	0.04	0.44	0.38	0.41	0.41	0.45	0.40	0.44
Amantadine	N04BB01	0.17	2.80	0.16	2.64	0.16	2.73	0.17	2.77
Amoxicillin	J01CA04	1.05	12.99	1.08	13.27	1.05	13.04	1.05	12.97
Azithromycin	L01FA10	0.07	0.55	0.12	0.98	0.08	0.65	0.06	0.52
Cephalexin	J01DA01	0.06	0.36	0.09	0.56	0.06	0.36	0.07	0.42
Chloramphenicol	J01BA01	0.04	0.10	0.04	0.09	0.02	0.05	0.04	0.11
Chloroquine	P01BA01	0.28	4.70	0.32	5.27	0.31	5.11	0.28	4.57
Ciprofloxacin	J01MA02	0.87	10.12	0.92	10.54	0.89	10.40	0.88	10.17
Clofazimine	J04BA01	0.26	12.68	0.28	13.55	0.28	13.91	0.26	12.87
Cloxacillin	J01CF02	0.31	1.30	0.34	1.39	0.32	1.33	0.32	1.30
Dapsone	J04BA02	0.27	8.86	0.26	8.43	0.28	9.27	0.27	9.00
DEC	P02CB02	0.16	1.36	0.16	1.38	0.16	1.36	0.16	1.38
Doxycyline	J01AA02	0.77	17.00	0.77	16.87	0.77	17.00	0.86	18.91
Erythromycin	J01FA01	0.40	1.56	0.44	1.67	0.40	1.54	0.46	1.78
Ethambutol	J04AK02	0.27	4.43	0.30	4.97	0.27	4.50	0.32	5.19
Ethionamide	J04AD03	0.06	1.91	0.09	2.86	0.05	1.50	0.08	2.77
Fluconazole	J02AC01	0.16	0.22	0.16	0.21	0.06	0.08	0.16	0.22
Furazolidone	G01AX06	0.09	1.43	0.16	2.64	0.09	1.43	0.11	1.80
Griseofulvin	D01BA01	0.47	5.44	0.53	6.11	0.46	5.34	0.51	5.91
Isoniazid	J04AC01	0.13	4.23	0.21	6.78	0.12	3.95	0.15	4.84
Ketoconazole	J02AB02	0.08	1.30	0.10	1.66	0.09	1.43	0.08	1.38
Levofloxacin	J01MA12	0.20	4.68	0.20	4.64	0.21	4.77	0.22	5.04
Metronidazole	P01AB01	0.63	2.94	0.62	2.83	0.61	2.83	0.69	3.20
Minocycline	A01AB23	0.26	4.07	0.26	3.94	0.29	4.45	0.29	4.46
Nalidixic acid	J01MB02	0.40	2.31	0.39	2.21	0.43	2.51	0.46	2.64
Nitrofurantoin	J01XE01	0.28	3.25	0.29	3.37	0.32	3.67	0.29	3.34
Norfloxacin	J01MA06	0.87	13.43	0.87	13.28	0.92	14.32	0.95	14.53
Oflloxacin	J01MA01	0.19	1.02	0.18	0.98	0.21	1.16	0.21	1.13
Pefloxacin	J01MA03	0.18	1.40	0.6	1.23	0.17	1.34	0.20	1.55
Primaquine	P01BA03	0.16	0.21	0.16	0.20	0.16	0.20	0.16	0.21
Pyrazinamide	J04AK01	0.15	2.52	0.15	2.48	0.16	2.73	0.17	2.77
Quinine	P01BC01	0.12	1.07	0.17	1.60	0.16	1.45	0.14	1.32
Rifampicin	J04AB02	0.35	8.69	0.34	8.47	0.37	9.20	0.37	9.13
Roxitromycin	J01FA06	0.27	2.07	0.25	1.90	0.28	2.16	0.27	2.10
Secnidazole	P02AB07	0.12	0.38	0.13	0.42	0.12	0.40	0.13	0.42
Sulfamethoxazole	J01EC01	0.26	1.22	0.22	1.01	0.30	1.37	0.29	1.34
Tinidazole	P01AB02	0.55	2.13	0.52	2.00	0.55	2.15	0.59	2.26
Category: CNS									
Alprazolam	N05BA12	0.19	0.72	0.17	0.63	0.21	0.80	0.25	0.95
Amitryptilline	N06AA09	0.06	0.41	0.05	0.33	0.06	0.41	0.10	0.66
Baclofen	M03BX01	0.32	2.13	0.30	1.96	0.31	2.05	0.40	2.63
Betahistine	N07CA01	0.30	5.05	0.30	4.89	0.35	5.79	0.39	6.37
Bromocriptine	N04BC01	0.16	0.91	0.17	1.00	0.20	1.17	0.29	1.65

Table 9: Contd...

Drug	ATC	Jan-Mar		Apr-Jun		Jul-Sep		Oct-Dec	
		%	DDD	%	DDD	%	DDD	%	DDD
Buspirone	N05BE01	0.03	0.18	0.04	0.20	0.06	0.34	0.06	0.35
Carbamazepine	N03AF01	0.73	7.28	0.71	6.96	0.82	8.22	0.18	1.74
Chlordiazepoxide	N05BA02	0.19	2.05	0.17	1.81	0.23	2.50	0.27	3.00
Buspirone	N05BE01	0.03	0.18	0.04	0.20	0.06	0.34	0.06	0.35
Carbamazepine	N03AF01	0.73	7.28	0.71	6.96	0.82	8.22	0.18	1.74
Chlordiazepoxide	N05BA02	0.19	2.05	0.17	1.81	0.23	2.50	0.27	3.00
Cinnarizine	N07CA02	0.06	0.49	0.08	0.70	0.11	0.99	0.08	0.68
Citalopram	N06AB04	0.08	1.94	0.09	2.15	0.10	2.45	0.11	2.70
Clobazam	N05BA09	0.63	5.25	0.63	5.20	0.71	5.86	0.80	6.54
Clomipramine	N06AA04	0.27	2.66	0.27	2.62	0.32	3.19	0.40	3.90
Clonazepam	N03AE01	0.10	0.41	0.09	0.38	0.14	0.58	0.17	0.69
Clozapine	N05AH02	0.07	0.31	0.07	0.28	0.10	0.43	0.13	0.55
Diazepam	N05BA01	0.28	4.64	0.30	4.97	0.28	4.70	0.36	5.88
Doxipine	N06AA12	0.77	0.22	0.05	0.17	0.10	0.33	0.13	0.43
Ergotamine	N02CA02	0.07	0.51	0.07	0.53	0.10	0.76	0.13	1.00
Flunarizine	N07CA03	0.47	7.77	0.44	7.23	0.51	8.45	0.61	10.03
Fluoxetine	N06AB03	0.33	16.16	0.34	16.49	0.34	17.18	0.48	23.46
Fluvoxamine	N06AB08	0.21	3.55	0.20	3.31	0.22	3.68	0.27	4.50
Haloperidol	N05AD01	0.14	0.89	0.20	1.24	0.17	1.07	0.23	1.43
Chlorpromazine	N05AA01	0.22	2.41	0.23	2.46	0.24	2.64	0.33	3.60
Levodopa	N04BA02	0.36	5.93	0.36	5.87	0.38	6.34	0.46	7.61
Lithium carbonate	N05AN01	0.26	8.73	0.30	9.94	0.27	9.00	0.35	11.62
Lorazepam	N05BA06	0.27	3.55	0.30	1.99	0.26	1.75	0.036	9.52
Nimodipine	C08CA06	0.22	0.74	0.27	0.89	0.24	0.79	0.30	0.50
Nitrazepam	N05CD02	0.21	10.64	0.25	12.42	0.23	11.45	0.29	14.12
Nortriptyline	N06AA10	0.27	2.95	0.27	2.91	0.22	2.24	0.35	3.78
Olanzapine	N05AH03	0.09	0.72	0.11	0.87	0.10	0.85	0.15	1.21
Phenobarbitone	N03AA02	0.27	5.32	0.28	5.42	0.29	5.73	0.34	6.64
Phenytoin sodium	N03AB02	0.20	3.27	0.25	4.07	0.21	3.48	0.25	4.08
Risperidone	N05AX08	0.27	2.66	0.27	2.62	0.25	2.50	0.34	3.32
Selegiline	N04BD01	0.20	9.82	0.23	11.07	0.21	10.23	0.34	16.81
Sertraline	N05AE03	0.24	12.57	0.27	13.88	0.19	10.01	0.27	13.84
Sodium valproate	N03AG01	0.52	3.21	2.50	3.04	0.46	2.85	0.67	4.13
Pyridostigmine	N07AA02	0.26	2.82	0.31	3.41	0.30	3.36	0.38	4.20
Zolpidem	N05CF02	0.23	7.64	0.25	8.28	0.23	7.50	0.34	11.07
Category: GIT									
Cisapride	A03FA02	0.06	0.76	0.21	2.71	0.06	0.76	0.08	1.11
Colloidal bismuth subcitrate	A02BX05	0.12	1.48	0.34	4.24	0.14	1.74	0.18	2.18
Digene	A02AD01	1.23	25.08	1.54	31.41	1.23	29.37	1.40	27.22
Diloxanide furoate	P01AC01	0.32	5.32	0.62	10.09	0.33	5.45	0.38	6.23
Domperidone	A03FA03	0.11	0.86	0.64	2.64	0.16	1.21	0.18	1.36
Esmoprazole	A02BC05	0.63	9.80	0.91	13.85	0.73	11.33	0.81	12.40
Famotidine	A02BA03	0.61	10.09	0.80	1318	0.72	11.93	0.74	12.11
Flavonoids	C05CA01	0.14	-	0.34	-	0.20	-	0.20	-
Lansoprazole	A02BC03	0.15	8.45	0.76	12.42	0.57	9.41	0.55	9.00
Loperamide	A07DA03	0.02	0.15	0.05	0.28	0.02	0.15	0.03	0.21

Contd...

Table 9: Contd...

Drug	ATC	Jan-Mar		Apr-Jun		Jul-Sep		Oct-Dec	
		%	DDD	%	DDD	%	DDD	%	DDD
Metoclopramide	AO3FA01	0.32	5.32	0.52	8.43	0.41	6.82	0.37	6.02
Mosapride	-	0.55	6.09	0.72	7.83	0.68	7.50	0.58	6.37
Omeprazole	A02BC01	1.59	18.47	1.78	20.40	1.68	19.57	1.64	18.84
Ondansetron	A04AA01	0.07	0.92	0.12	1.47	0.09	1.12	0.11	1.30
Prochlorperazine	NO5AB04	0.27	1.33	0.35	1.72	0.32	1.62	0.32	1.56
Ranitidine	A02BA02	1.15	57.28	1.44	70.93	1.30	65.04	1.26	62.28
Salazopyrine	A07EC01	0.19	2.30	0.30	3.67	0.24	2.97	0.23	2.85
Category: Respiratory system and anti-allergics									
Bromhexine	R05CB02	0.63	6.53	0.76	7.73	0.63	6.49	0.65	6.63
Etophylline and theophylline	R03DA05	1.60	12.35	1.66	12.65	1.53	11.49	1.63	12.46
Salbutamol	R03CC02	1.02	7.89	1.03	7.84	0.98	7.57	1.00	7.69
Astemizole	R06AX11	0.68	5.25	0.61	4.67	0.62	4.84	0.67	5.17
Cetirizine	R06AE0	0.93	7.16	0.98	7.52	0.83	6.43	0.90	6.91
Ketotifen	R06AX17	1.09	8.43	1.11	8.47	1.01	7.86	1.07	8.20
Terfenadine	R06AX12	1.13	8.75	1.22	9.31	1.15	8.91	1.13	8.65
Pheniramine maleate	R06AB06	1.37	10.60	1.38	10.54	1.29	9.99	1.35	10.37
Cyproheptadine	R06AX02	1.15	8.88	1.18	9.03	1.03	7.99	1.14	8.72
Promethazine	R06AD02	1.22	11.30	1.40	12.86	1.26	11.76	1.16	10.66
Category: Hormones and related drugs									
Acarbose	A10BF01	0.28	2.28	0.23	1.88	0.33	2.73	0.27	2.21
Clomiphene citrate	G03GB02	0.19	2.99	0.16	2.46	0.27	4.14	0.19	2.19
Conjugated estrogen	G03CA57	0.63	14.70	0.57	13.18	0.73	16.89	0.65	15.02
Danazole	G03XA01	0.63	5.25	0.62	5.08	0.72	5.97	0.62	5.12
Dexamethasone	H02AB02	0.69	5.35	0.63	4.78	0.76	5.89	0.62	4.78
Eltroxine	H03AA01	0.49	10.91	0.41	9.04	0.58	12.91	0.43	9.50
Glibenclamide	A10BB01	1.29	14.99	1.27	14.60	1.41	16.46	1.26	14.53
Gliclazide	A10BB09	0.63	7.30	0.57	6.59	0.72	8.35	0.61	7.02
Glimepride	A10BB12	0.51	12.68	0.51	12.42	0.59	14.62	0.47	11.52
Glipizide	A10BB07	1.05	12.99	0.99	12.14	1.13	14.06	0.90	11.11
Metformin	A01BA02	1.06	26.29	0.92	22.70	1.17	29.25	0.95	23.35
Neomercazole	H03BB01	0.56	9.20	0.57	9.34	0.65	10.84	0.53	8.65
Norethisterone	G03AC01	0.09	1.34	0.09	1.41	0.11	1.78	0.08	1.23
Pioglitazone	A10BG03	0.16	2.42	0.14	2.18	0.20	3.05	0.14	2.13
Prednisolone	H02AB06	0.36	3.95	0.32	3.51	0.41	4.55	0.31	3.41
Repaglinide	A10BX02	0.22	0.86	0.21	0.79	0.27	1.03	0.19	0.73
Rosiglitazone	A10BG02	0.58	6.45	0.57	6.22	0.61	6.73	0.53	5.77
Category: Vitamins and nutrition									
Calcium carbonate	A12AA03	2.69	5.95	2.33	5.09	2.69	5.97	2.67	5.86
Ferrous sulphate	B03AA07	2.87	47.59	2.31	37.80	2.05	34.02	2.30	37.78
Alfacalcidiol	M05A12A	0.58	2.26	0.57	2.18	0.39	1.51	0.41	1.58
B complex	A11EB03	3.90	-	3.62	-	3.26	-	3.24	-
Folic acid	B03BB01	2.26	34.87	1.79	27.41	1.55	24.12	1.40	21.51
Vitamin C	A11GA01	1.95	10.75	1.72	9.41	1.37	7.57	1.21	6.64
Vitamin E	A11HA03	1.91	44.39	1.83	41.96	1.56	36.27	1.25	28.87

Contd...

Table 9: Contd...

Drug	ATC	Jan-Mar		Apr-Jun		Jul-Sep		Oct-Dec	
		%	DDD	%	DDD	%	DDD	%	DDD
Category: Metabolism									
Alendronate	M05BA04	1.23	13.64	1.14	12.40	1.23	13.64	1.13	12.41
Atorvastatin	C10AA05	1.58	52.37	1.40	45.78	1.58	52.50	1.51	49.54
Gemfibrozil	C10AB04	1.63	18.90	1.43	16.45	1.62	18.85	1.53	17.63
Lovastatin	C10AA02	1.59	17.59	1.54	16.82	1.58	17.50	1.47	16.05
Simvastatin	C10AA01	2.46	54.37	2.15	46.99	2.53	55.99	2.32	50.74
UDCA	A5AA02	1.43	33.22	1.36	31.10	1.63	37.89	1.37	31.48

CVS: Cardiovascular, GIT: Gastrointestinal, CNS: Central nervous system, UDCA: Ursodeoxycholic acid

below two but higher figure is always fraught with increased risk of drug interaction and errors of prescribing.

The increase in a number of patients with advancing age especially above 50-year suggests a need of time to address the requirement of the geriatric population separately. A voluminous increase of prescription for drug groups like - CVS, nutrition and vitamins, metabolic drugs, hormones and related drugs, NSAIDs, psychotropics, and AMAs is pointer to prevailing morbidity pattern of society. There is a sharp decrease in no. of patients >70 year age group probably due to age-related in-capabilities to collect medicines at the hospital dispensary.

Between the genders, common group of drugs highly prescribed to both are from nutrition and vitamins. However, the second highest drug prescribed for males has been CVS drugs and likewise for females been NSAIDs. Furthermore, these findings suggest of high prevalence of CVS related morbidity such as MI, CAD, and hypertension prevalent in the male subpopulation and females of pain related ailments.

Pharmaco-economically, the budget spent in all four quarters of the period of study, suggests more or less a uniform pattern without significant variation. Among the various drug groups, the order of the expense has been on metabolic drugs, AMAs, and hormones and related drugs. Majority of such drug groups are recently been available on the shelf and are of specific needs of the elderly population. This indicates a rising trend of metabolic disorders as well as endocrinial disorders, all pointers toward diseases of sedentary lifestyles with reduced fitness levels. AMAs per say are not the only one with a high cost per tablet but also been the most misused and over prescribed group of drugs.

The basic definition of the DDD is: The assumed average maintenance dose per day for a drug used for its main indication in adults.²⁹ The DDD methodology does not indicate the exact number of patients who have been treated with a drug. This concept assumes that every person prescribed a particular drug is taking the specified dose of the said drug every day, ignoring alterations of dose of the drug by disease or if any patient-related factors be there or

any other variable. With this concept of drug utilization, a drawback may be that drugs prescribed for a brief period can have their prescribing prevalence underestimated too. Variation in the pattern of dosage, duration of the study, the scatter of population over different ages, gender and races, all can contribute to differences in audit reports using DDD methodology to determine the prevalence of prescribing.

An important basis for the WHO recommendation to use the ATC/DDD methodology in drug utilization studies is to improve comparison of drug utilization studies in making international comparisons and to study long-term trends in drug use. Due to continuous change and development of the drug assortment and drug uses, a regular revision of the ATC system is also necessary. Some ATC and DDD alterations are made annually, and an updated version of the list that includes all officially assigned ATC codes and DDDs is issued every year.²⁹ In order to compare data from different countries and different periods, it is essential to know which ATC codes and DDDs are used. Consumption figures may look very different when different ATC/DDD versions are applied.

To have uniformity in drug utilization studies world over, DDD is of great value as compared to any traditional method of use determination like in percentage, or weight or cost wise. Study of drug utilization in India has been reported both in hospital¹⁹⁻²¹ and community levels.^{22,23,27} Most of these studies have measured drug utilization by counting of prescriptions which actually indicates the prescribing frequency. The study of drug utilization is made more meaningful if the frequency of prescribing is studied with a total quantity of drugs per prescription. Indian studies have used DDD concept and focused on drug utilization in selected groups of patients for a brief period,²⁴⁻²⁶ whereas this study was undertaken the study on a vast scale for the first time the country.

Drugs were classified using ATC classification system which mandates the drugs to divided into different groups according to the organ or system on which they act and their chemical, pharmacological and therapeutic property.²⁸ Drugs are classified in groups at five different levels. There are fourteen main groups (first level), with two therapeutic/

pharmacological subgroups (second and third levels), the fourth level is a therapeutic/pharmacological chemical subgroup, and the fifth level is the chemical substance.

The complete classification of metformin illustrates the structure of the code:

A: Alimentary tract and metabolism (first level, anatomical main group).

A10: Drugs used in diabetes (second level, pharmacological/therapeutic main group).

A10B: Oral blood glucose lowering drugs (third level, chemical/pharmacological/therapeutic subgroup).

A1013A: Biguanides (fourth level, chemical/pharmacoiological/therapeutic subgroup).

A10BA02: Metformin (fifth level, subgroup for the chemical substance).

Thus, in the ATC system, all plain metformin preparations are given the code A10BA02. Prescribing prevalence studies are also useful to determine the prevailing morbidity patterns. To quote an example from our study, the DDD/1000/day of amlodipine (49.77), frusemide (40.09), and ramipril (26.59) in the first quarter, and likewise high value in all other quarters of year points toward a pandemic state of CVS disorders like hypertension and myocardial ischemia, etc. Similarly, the high prescription pattern for antipsychotics like fluoxetine (16.16), sertraline (12.57), nitrazepam (10.64) is an index of high psychiatric morbidity in the society. Also in a drug group of GIT, it is realized a high consumption pattern of ranitidine (57.28) and likewise in the other quarter of years too, is an indicator toward urbanization and its related disorders of anxiety and poor hygiene sanitation. A high usage of OHAs like metformin (26.29), glimepride (12.68), glipizide (12.99), and metabolic drugs like atorvastatin (52.37), simvastatin (54.37) are indicators toward growing prevalence of metabolic disorders in the society.

Studies elsewhere in India on prescription audit employing the DDD concept²⁴⁻²⁶ have focused on a select group of patients which could not be meaningfully compared to our data because the present study covers a general outpatient department (OPD) population. Present study is done on a sample population of an urban demographic profile dependent for their health care needs through OPDs on a government tertiary care hospital. This may be giving insight into the day to day functioning status of our health care delivery system and opportunity on self-assessment in further improving the delivery of health care by implementing measures which economizes on scarce health care budget and facility and optimize the needful help to patients by minimizing errors in prescription writing by physicians. This study may serve as one of the works upon which further studies in prescription

audit can be taken up to improve the scope for improvement in prescribing practices among the physicians.

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