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2 Prescription pattern study of oral glucose lowering agents in  
3 patients with type 2 diabetes in outpatient department of a  
4 tertiary care teaching hospital.

5  
6 Abstract

7 **Background:** Type II diabetes mellitus (T2DM) is a chronic metabolic disorder  
8 characterised by persistent hyperglycaemia. Globally, an estimated 537 million adults (20–  
9 79 years) were living with diabetes in 2021 – about 10.5 % of the world’s population in this  
10 age group, and the number is projected to reach 643 million by 2030 and 783 million  
11 by 2045.[1] In India alone, approximately 77 million people had diabetes in 2019 and this is  
12 expected to increase to 134 million by 2045.[2] Appropriate selection of glucose-lowering  
13 agents and rational prescribing are therefore essential to control blood glucose and  
14 prevent complications.

15 **Objectives:** To evaluate the demographic profile, antidiabetic prescription patterns,  
16 common drug combinations, and adherence to WHO prescribing indicators among T2DM  
17 outpatients at a tertiary care teaching hospital.

18 **Methods:** A descriptive, observational, cross-sectional study was conducted on 600  
19 prescriptions of patients with T2DM attending the general medicine OPD of MGM Hospital,  
20 Chhatrapati Sambhaji Nagar. Data was extracted from prescriptions into a spreadsheet. For  
21 each encounter we recorded the patient’s age and gender, the number of oral  
22 glucose-lowering agents, the total number of drugs, the number of medicines prescribed by  
23 generic name, the number of drugs from the Indian essential medicines list, and whether an  
24 injectable drug or antibiotic was prescribed. WHO core prescribing indicators were  
25 computed according to standard definitions.[3]

26 **Statistical analysis:** - Data were entered in Microsoft Excel and analysed using descriptive  
27 statistics. Continuous variables are presented as mean  $\pm$  standard deviation (SD) and range,  
28 and categorical variables as numbers and percentages.

29 **Results:** The mean age of the patients was  $53.0 \pm 11.6$  years (range 22–80), and 51.1%  
30 were male and 48.9% female. Most patients belonged to the 41–60 year age group (55.5%),  
31 followed by 61–80 years (29.2%) and less than 40 years (15.3%). Polytherapy ( $\geq 4$  classes  
32 of glucose-lowering agents) was observed in 39.3% of prescriptions and triple therapy in  
33 27.5%, whereas dual therapy and monotherapy accounted for 17.8% and 15.3%,  
34 respectively. DPP-4 inhibitors were the most frequently prescribed class (61.1%), followed  
35 by sulfonylureas (58.1%) and biguanides (56.8%); insulin was used in 38.7% of  
36 encounters. The average number of drugs per prescription was 3.59. Drugs were  
37 prescribed by generic name in 86.4% of cases, injections were used in 23.3% and

38 antibiotics in 6.0% of encounters, and 55.5% of the drugs were from the essential  
39 medicines list. The most frequent oral combination was biguanide + DPP-4 inhibitor +  
40 sulfonylurea (8.64%), followed by biguanide + DPP-4 inhibitor + SGLT2 inhibitor +  
41 sulfonylurea (6.15%)

42 **Conclusions:** The present study highlights extensive use of combination therapy in T2DM,  
43 with a preference for newer agents such as DPP-4 and SGLT2 inhibitors. The mean number  
44 of medicines per encounter exceeded the WHO reference value of less than 2, and the  
45 injection use rate was slightly above the recommended 20 %.[3] Although the percentage of  
46 drugs prescribed by generic name was high, adherence to the essential medicines list needs  
47 improvement. Regular prescription audits and adherence to national and international  
48 guidelines are necessary to promote rational use of antidiabetic agents.

49 **Keywords:** Type II diabetes mellitus, prescription pattern, oral antidiabetic drugs, WHO  
50 prescribing indicators, rational drug use

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52

## 53 Introduction

54 Type II diabetes mellitus (T2DM) is a chronic metabolic disorder characterised by  
55 persistent hyperglycaemia. Globally, an estimated 537 million adults (20–79 years) were  
56 living with diabetes in 2021 – about 10.5 % of the world’s population in this age group, and  
57 the number is projected to reach 643 million by 2030 and 783 million by 2045. [1] It is  
58 among the leading global health emergencies, accounting for significant morbidity,  
59 mortality and health-care expenditure. [2] India bears a substantial share of this burden: an  
60 estimated 77 million people were living with diabetes in 2019, with projections of  
61 134 million by 2045. Alarmingly, over half of these cases remain undiagnosed. Type II  
62 diabetes mellitus (T2DM) accounts for about 90 % of all cases and is strongly associated  
63 with lifestyle factors such as sedentary behaviour, unhealthy diet and obesity. [2] Recent  
64 systematic reviews indicate that polypharmacy—defined as the use of five or more  
65 medications—is present in nearly 59% of older adults with diabetes and is associated with  
66 increased healthcare utilisation and adverse drug events [3].

67 T2DM is a progressive disorder; maintenance of glycaemic goals often requires  
68 combination therapy. The American Diabetes Association (ADA) recommends stepwise  
69 addition of medications to metformin and acknowledges that early combination therapy  
70 may be needed to achieve glycaemic targets. Newer agents like dipeptidyl-peptidase 4  
71 (DPP-4) inhibitors, sodium–glucose cotransporter 2 (SGLT2) inhibitors and glucagon-like  
72 peptide-1 receptor agonists offer cardiovascular and renal benefits and are being  
73 increasingly used. [4] At the same time, inappropriate prescribing practices such as  
74 polypharmacy, overuse of injections and antibiotics, and deviation from essential  
75 medicines lists can jeopardize patient safety and increase costs. To address these issues the  
76 WHO, in collaboration with the International Network for Rational Use of Drugs, developed  
77 a set of core drug use indicators to evaluate prescribing patterns. The five prescribing

78 indicators assess the average number of medicines per encounter, the percentage of drugs  
79 prescribed by generic name, the percentage of encounters with an antibiotic prescribed,  
80 the percentage of encounters with an injection prescribed and the percentage of medicines  
81 prescribed from the essential medicines list. The WHO proposes reference values for these  
82 indicators (<2 medicines per encounter, 100 % generic prescribing, <30 % antibiotic  
83 encounters, <20 % injections and 100 % essential medicines). [5] Guidelines from  
84 international bodies now emphasise early use of SGLT2 inhibitors and GLP-1 receptor  
85 agonists in patients with cardiovascular or renal comorbidities due to their cardiorenal  
86 benefits [6]. Prescription patterns in developed countries show that metformin  
87 monotherapy has stabilised while SGLT2 inhibitors and GLP-1 receptor agonists are being  
88 prescribed more frequently, reflecting these guideline changes [7]. However, observational  
89 studies from Ethiopia and geriatric clinics have documented widespread polypharmacy,  
90 low rates of generic prescribing and limited adherence to essential medicines lists,  
91 highlighting the need for local audits [8][9].

92 The present study was undertaken to describe the current trends in prescribing oral  
93 glucose-lowering agents and insulin for T2DM in a tertiary care teaching hospital, to  
94 analyse adherence to WHO prescribing indicators and to identify common drug  
95 combinations. Such information is vital for promoting rational drug use and aligning  
96 practice with guidelines. Hospital-based audits in central Maharashtra and southern India  
97 have reported an average of 1.7–3.0 antidiabetic drugs per prescription, with combination  
98 therapy employed in approximately 70% of encounters and nearly universal generic  
99 prescribing [10][11]. These findings provided a benchmark for the present evaluation.

## 100 Methods

### 101 Study design and setting

102 This was a descriptive, observational, non-interventional, cross-sectional study carried out  
103 in the general medicine OPD of MGM Medical College & Hospital,  
104 Chhatrapati Sambhaji Nagar (Aurangabad), a tertiary care teaching hospital. The study  
105 commenced after obtaining approval from the Institutional Ethics Committee.  
106 Prescriptions were collected until the target sample size of 600 encounters was reached; a  
107 total of 600 prescriptions were analysed. Only prescriptions of adult patients (>18 years)  
108 diagnosed with T2DM and receiving at least one glucose-lowering agent were included.  
109 Prescriptions for pregnant women, type I diabetes or gestational diabetes were excluded.

### 110 Data collection

111 Data from each eligible prescription was extracted into a structured Excel sheet. Variables  
112 included the patient's age and gender, the number of drugs belonging to each class of  
113 glucose-lowering agents (insulin, biguanides, SGLT2 inhibitors, thiazolidinediones,  
114 sulfonylureas and DPP-4 inhibitors), the total number of drugs in the prescription  
115 (including drugs for comorbidities), the number of medicines prescribed by generic name,  
116 the number of drugs from the essential medicines list, and whether any injection or  
117 antibiotic was prescribed. The presence of a class was defined as at least one drug from  
118 that class in the prescription. Monotherapy was defined as the use of a single class of

119 glucose-lowering agent, dual therapy as two classes, triple therapy as three classes and  
120 polytherapy as four or more classes. Age groups were categorized as <40, 41–60, 61–80  
121 and >80 years. Therapy categories (monotherapy, dual therapy, triple therapy and  
122 polytherapy) were derived from the number of classes prescribed. WHO core prescribing  
123 indicators were computed as follows: [5]

- 124 (i) average number of drugs per encounter = total number of drugs prescribed ÷  
125 total number of encounters
- 126 (ii) percentage of drugs prescribed by generic name = (total number of drugs  
127 prescribed by generic name ÷ total number of drugs prescribed) × 100
- 128 (iii) percentage of encounters with an injection or antibiotic = (number of  
129 encounters with ≥1 injection/antibiotic ÷ total encounters) × 100
- 130 (iv) percentage of drugs prescribed from the essential medicines list = (total number  
131 of drugs from the essential medicines list ÷ total number of drugs  
132 prescribed) × 100.
- 133 (v) Drug combinations were summarized by counting prescriptions containing ≥2  
134 classes and listing the most common combinations.

135

## 136 Statistical analysis

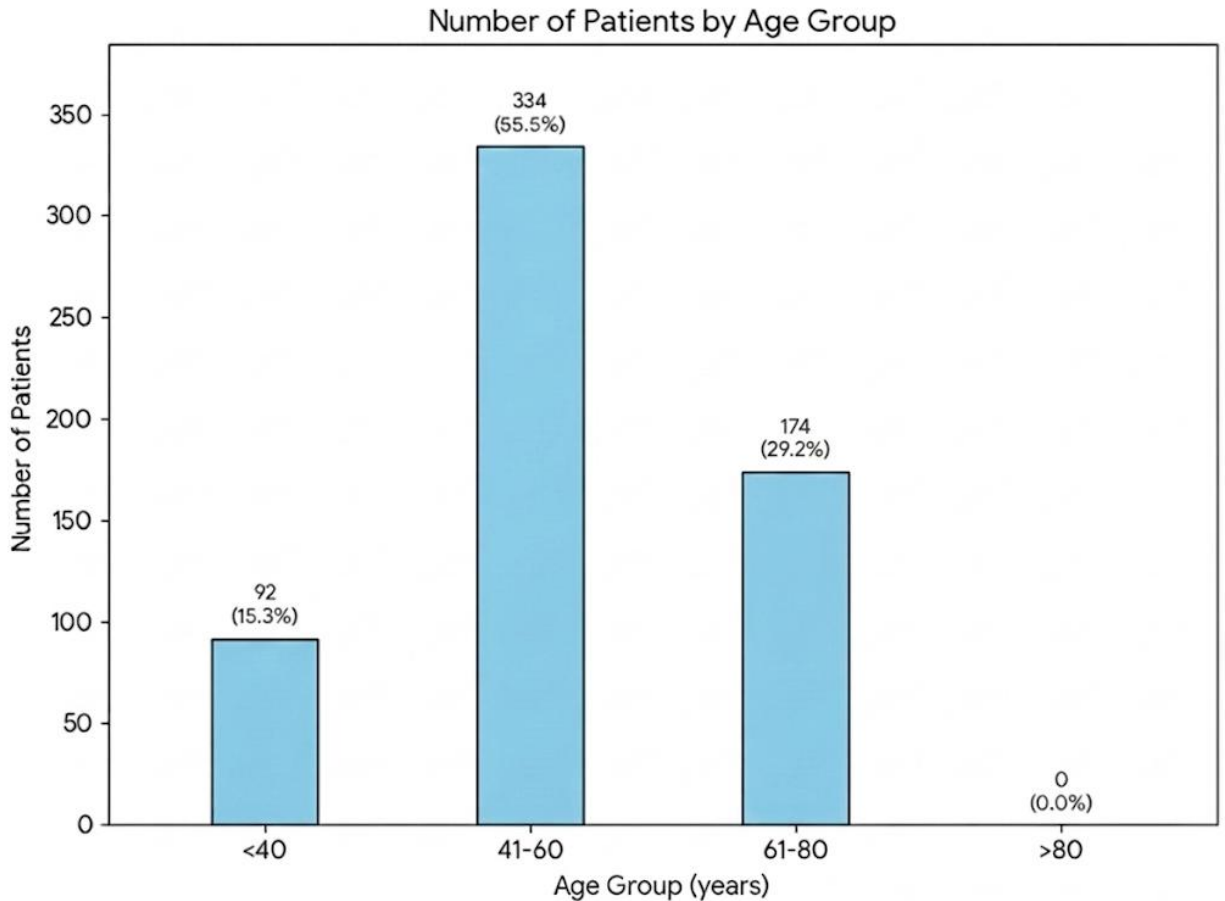
137 Data were entered in Microsoft Excel and analysed using descriptive statistics. Continuous  
138 variables are presented as mean ± standard deviation (SD) and range, and categorical  
139 variables as numbers and percentages.

## 140 Results

### 141 Demographic characteristics

142 The study included 600 prescriptions. The mean age of patients was 53.0 ± 11.6 years  
143 (median 53 years; range 22–80 years). Age distribution is shown in **Figure 1**; most patients  
144 (55.5 %) were between 41–60 years. Gender distribution is summarized in **Table 1** –  
145 51.1 % were male and 48.9 % were female.

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**Figure 1: Age distribution**

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**Table 1 - Gender distribution of patients (n = 600)**

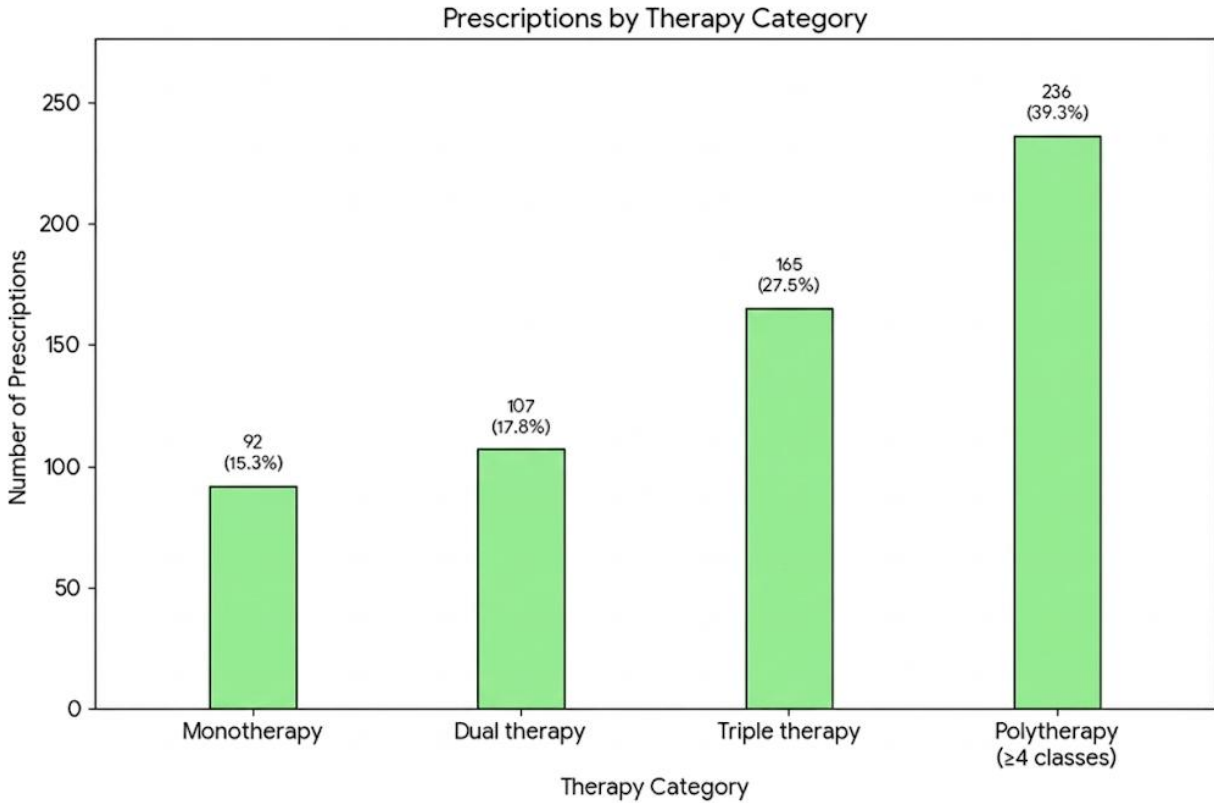
Gender	Number of patients	Percentage
Male	307	51.1 %
Female	293	48.9 %

151

## 152 Drug therapy distribution

153 For each prescription the number of different classes of glucose-lowering agents was  
 154 counted. Monotherapy was observed in 92 prescriptions (15.3 %), dual therapy in 107  
 155 prescriptions (17.8 %), triple therapy in 165 prescriptions (27.5 %) and polytherapy ( $\geq 4$   
 156 classes) in 236 prescriptions (39.3 %). The distribution of therapy categories is shown in  
 157 **Figure 2.**

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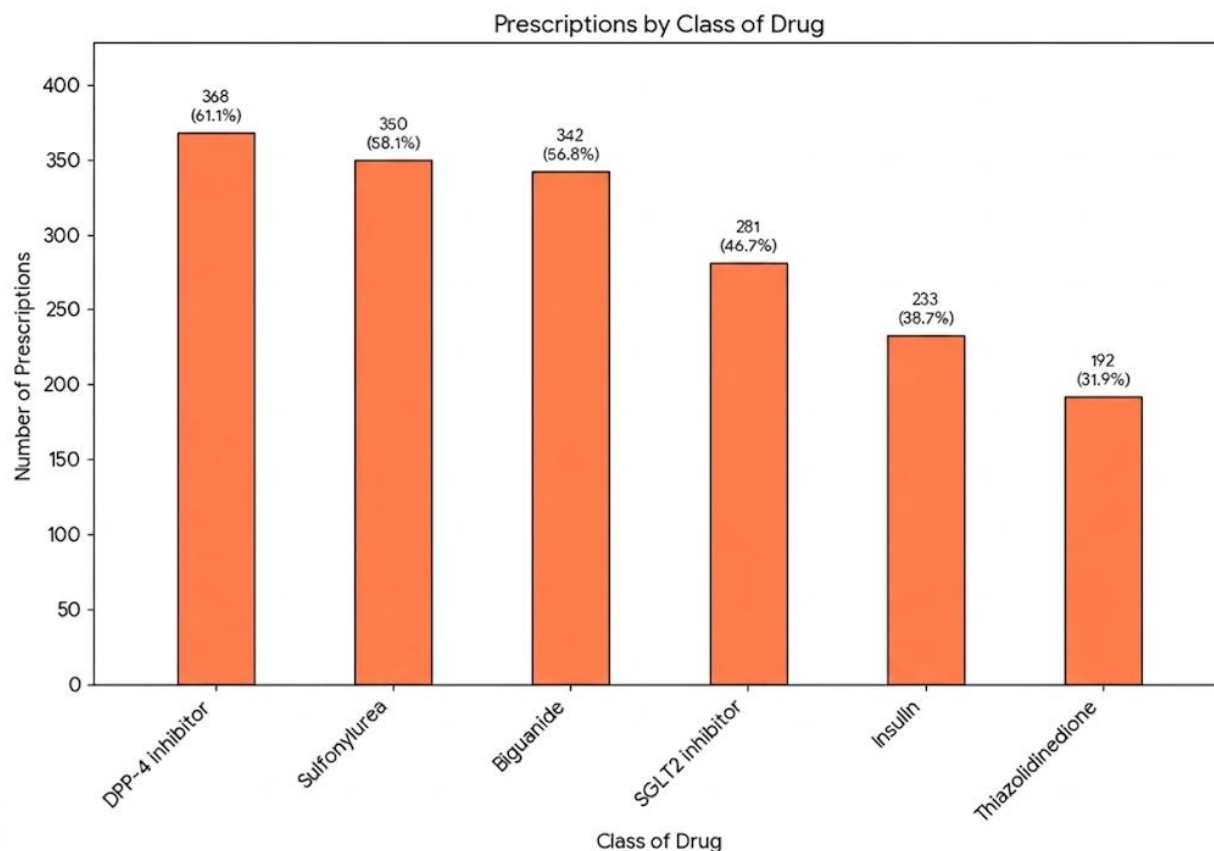
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**Figure 2: Distribution of drug therapy**

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162 Utilization pattern of glucose-lowering agents

163 The utilization of individual classes of glucose-lowering agents is summarized in **Figure 3**.  
 164 DPP-4 inhibitors were the most frequently prescribed class (61.1 % of encounters),  
 165 followed by sulfonylureas (58.1 %) and biguanides (metformin) (56.8 %). SGLT2 inhibitors  
 166 were prescribed in 46.7 % of encounters, while insulin and thiazolidinediones were used in  
 167 38.7 % and 31.9 % of encounters, respectively.



**Figure 3: Utilization by drug class**

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171 WHO prescribing indicators

172 The average number of drugs per prescription was 3.59 (SD 1.61; median 3). Drugs were  
 173 prescribed by generic name in 86.4 % of cases. Injections were prescribed in 23.3 % of  
 174 encounters and antibiotics in 6.0 %. Overall, 55.5 % of all drugs belonged to the Indian  
 175 essential medicines list. These findings are compared with the WHO reference values in  
 176 **Table 2**.

177 **Table 2 - WHO core prescribing indicators**

Indicator	Study value	WHO reference value
Average number of medicines per encounter	3.59	< 2 per encounter
Percentage of drugs prescribed by generic name	86.4 %	100 %
Percentage of encounters with an antibiotic prescribed	6.0 %	< 30 %
Percentage of encounters with an injection prescribed	23.3 %	< 20 %
Percentage of drugs prescribed from essential medicines list	55.5 %	100 %

178

## 179 Common drug combinations

180 Among the 600 prescriptions, 508 (84.7 %) contained  $\geq 2$  classes of glucose-lowering  
181 agents. The five most common combinations are presented in **Table 3**. The most frequent  
182 combination was biguanide + DPP-4 inhibitor + sulfonylurea (8.64 % of all prescriptions),  
183 followed by biguanide + DPP-4 inhibitor + SGLT2 inhibitor + sulfonylurea (6.15 %).  
184 Dual-class combinations such as biguanide + DPP-4 inhibitor and DPP-4  
185 inhibitor + sulfonylurea each accounted for 3.65 % of prescriptions.

186 **Table 3 – Top combinations of glucose-lowering agents (n = 600)**

Combination of classes	Number of prescriptions	Percentage of prescriptions
Biguanide + DPP-4 inhibitor + sulfonylurea	52	8.64 %
Biguanide + DPP-4 inhibitor + SGLT2 inhibitor + sulfonylurea	37	6.15 %
DPP-4 inhibitor + SGLT2 inhibitor + insulin + sulfonylurea	26	4.32 %
Biguanide + DPP-4 inhibitor	22	3.65 %
DPP-4 inhibitor + sulfonylurea	22	3.65 %

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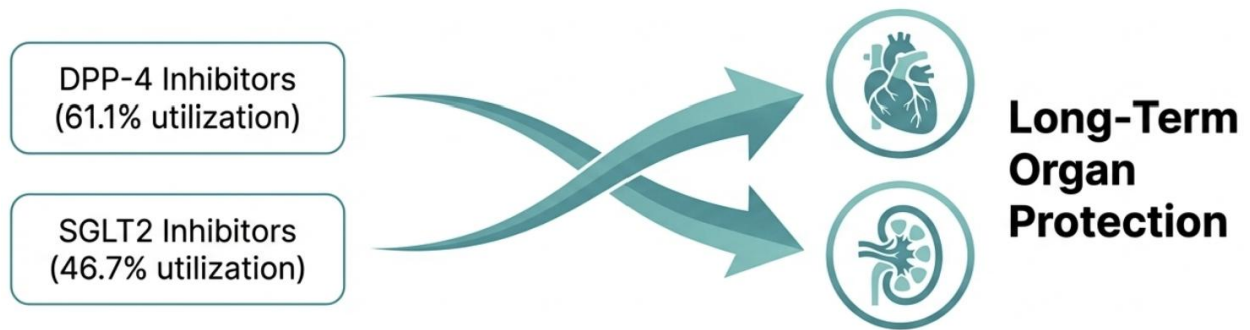
## 188 Discussion

189 This cross-sectional study analyzed 600 prescriptions of patients with T2DM attending the  
190 general medicine OPD of a tertiary care teaching hospital in India. The mean age of patients  
191 (53 years) and predominance of the 41–60 year age group reflect the middle-aged nature  
192 of T2DM, consistent with global and Indian reports. IDF estimates suggest that diabetes  
193 prevalence increases markedly after 40 years and peaks in older adults. [1] We observed a  
194 nearly equal gender distribution (male 51.1 %, female 48.9 %), whereas other studies have  
195 reported a slight male preponderance. The high prevalence of polytherapy (multiple  
196 glucose-lowering classes) observed in our cohort aligns with the broader global challenge  
197 of polypharmacy (total medication burden): a meta-analysis of 21 studies involving more  
198 than 520,000 participants found that approximately 59% of older adults with diabetes  
199 experience polypharmacy, noting significant associations with longer disease duration and  
200 multiple comorbidities [3][8]. Comparable reports from geriatric clinics document an  
201 average of 5.22 medications per patient and limited use of generic prescribing [9].

202 Polytherapy and triple therapy together accounted for 66.8 % of prescriptions,  
203 underscoring the progressive nature of T2DM and the need for multiple agents. The ADA  
204 emphasizes that maintenance of glycaemic targets often requires combination therapy. [4]  
205 In our study, DPP-4 inhibitors were the most commonly prescribed agents, followed by

206 sulfonylureas and metformin (biguanide). Newer agents such as SGLT2 inhibitors were  
207 used in nearly half of the encounters. The high utilization of DPP-4 inhibitors and SGLT2  
208 inhibitors suggests a shift towards medications with favourable cardiovascular and renal  
209 profiles, as advocated by recent guidelines. **(Figure 4)** [4]

## High utilization of newer agents reflects a positive shift toward cardiorenal protection

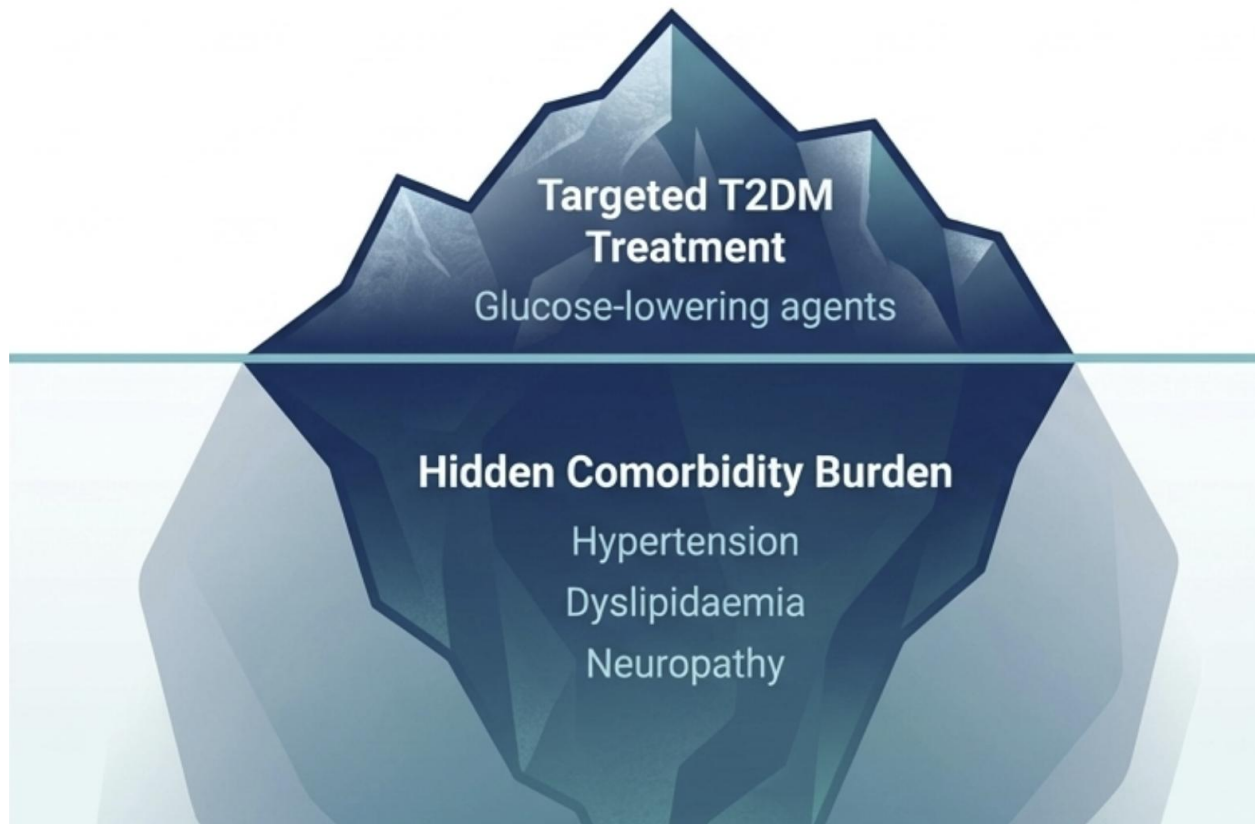


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211 **Figure 4: - Prescription shift toward cardiorenal protection**

212 Hospital-based audits in central Maharashtra and southern India have reported lower  
213 mean numbers of antidiabetic drugs per prescription (1.7–3.0), high reliance on metformin  
214 and sulfonylureas, and nearly universal generic prescribing [10][11]. Conversely,  
215 population-based studies in Spain indicate that prescriptions for SGLT2 inhibitors and  
216 GLP-1 receptor agonists have increased significantly over the past decade due to evidence  
217 of cardiovascular benefit [7]. Despite this trend, real-world data from the United States  
218 reveal that only 7.4% of older adults with T2DM receive SGLT2 inhibitors or GLP-1  
219 receptor agonists, even though 83.4% of high-risk patients might benefit [12]. These  
220 discrepancies highlight variability in adoption of newer agents across settings.

221 The average number of drugs per prescription (3.59) exceeded the WHO reference value of  
222 <2. While this average falls just below the strict definition of polypharmacy (five or more  
223 medications), it indicates a high overall pill burden and a strong tendency toward  
224 polypharmacy in this population. However, patients with T2DM often have comorbidities  
225 such as hypertension and dyslipidaemia, necessitating additional medications. **(Figure 5)**



226

227 **Figure 5: -Progressive disease and co-morbidities drive average prescription beyond**  
 228 **normal limits**

229 The rate of generic prescribing (86.4 %) was high but still below the optimal 100 %.  
 230 Generic prescription promotes affordability and should be encouraged. Injections were  
 231 used in 23.3 % of encounters, slightly exceeding the WHO's suggested upper limit of  
 232 20 %.[5] Many of these injections were insulin or vitamin B12 preparations; nevertheless,  
 233 prescribers should be cautious about over-use of injectables. Antibiotics were prescribed in  
 234 only 6.0 % of encounters, well within the recommended limit of 30 %, indicating judicious  
 235 use. Only 55.5 % of the drugs were from the essential medicines list, suggesting room for  
 236 improvement in aligning prescriptions with national formularies. Polytherapy with three  
 237 or more classes accounted for most prescriptions, with biguanide + DPP-4  
 238 inhibitor + sulfonylurea being the most common combination. Our average number of  
 239 drugs per encounter exceeded that reported in a family medicine primary care study (2.89)  
 240 and cross-sectional audits from central India [13][10], reflecting the burden of  
 241 comorbidities in our tertiary care cohort. Similarly, our essential medicines list adherence  
 242 (55.5%) was lower than the 98.3% and 89.80% reported in central Maharashtra and  
 243 Srikakulam studies respectively [10][11]. Emerging evidence supports combination  
 244 therapy with SGLT2 inhibitors and GLP-1 receptor agonists: a recent meta-analysis  
 245 involving over 42,000 participants found that such combinations reduced hospitalisations  
 246 for heart failure and major adverse cardiovascular events compared with monotherapy and  
 247 produced greater HbA1c and weight reductions, albeit with an increase in gastrointestinal  
 248 side-effects [14]. Cost considerations may limit uptake of these agents; a primary care

249 study reported that financial constraints were a major barrier to prescribing SGLT2  
250 inhibitors [13].

## 251 Strengths and limitations

252 The strengths of this study include a relatively large sample size and a comprehensive  
253 evaluation of WHO prescribing indicators. Data were collected from actual prescriptions,  
254 minimizing recall bias. The study is limited by its single-centre, cross-sectional design;  
255 findings may not be generalizable to other settings. Nevertheless, the study provides  
256 valuable insights into current prescribing patterns and highlights areas for quality  
257 improvement.

## 258 Conclusions

259 This study highlights the complex reality of managing T2DM in a tertiary care setting,  
260 where the progressive nature of the disease frequently necessitates combination therapy.  
261 Encouragingly, prescribing patterns reflect a modernizing shift; clinicians are increasingly  
262 utilizing newer agents like DPP-4 and SGLT2 inhibitors to align with recent guidelines  
263 advocating for cardiovascular and renal protection. Furthermore, the high rate of generic  
264 prescribing and judicious use of antibiotics indicate positive prescribing behaviours.  
265 However, these clinical benefits are offset by significant challenges regarding rational drug  
266 use. The average number of medications per encounter notably exceeded WHO reference  
267 values, pointing to a high burden of polypharmacy. Additionally, with only about half of the  
268 prescribed drugs belonging to the essential medicines list, there is a clear missed  
269 opportunity for cost-effective care. Moving forward, targeted clinician education and  
270 regular prescription audits are essential. Healthcare providers must strive to balance  
271 aggressive, guideline-directed combination therapy with the principles of rational  
272 prescribing to minimize polypharmacy and enhance the affordability of diabetes care

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