

REAL-world evidence of risk factors and comorbidities in YOUNG Indian adults with type 2 diabetes mellitus: A REAL YOUNG (diabetes) study

Banshi Saboo¹, Sanjay Agarwal², Sunil Gupta³, Brij Makkar⁴, A Panneerselvam⁵, Abhay Kumar Sahoo⁶, G. D. Ramchandani⁷, Sambit Das⁸, Suhas Erande⁹, Yogesh Kadam¹⁰, Mahesh V. Abhyankar¹¹, Santosh Revankar¹¹

 ¹Department of Diabetology, Diabetes Care and Hormone Clinic, Ambawadi, Ahmedabad, Gujarat, ²Department of Internal Medicine, Dr. Sanjay Agarwal's Aegle Clinic, City, Pune, Maharashtra, ³Department of Diabetology, Sunil's Diabetes Care Research Centre, Nagpur, Maharashtra, ⁴Department of Diabetology, Dr. Makkar's Diabetes and Obesity Centre, Delhi,
⁵Department of Diabetology, Aruna Diabetes Centre, Chennai, Tamil Nadu, ⁶Department of Endocrinology, IMS and SUM Hospital, Bhubaneshwar, Odisha, ⁷Ramachandani Diabetes Care and Research Centre, Kota, Rajasthan, ⁶Department of Endocrinology, Endeavour Clinics, Bhubaneswar, Odisha, ⁶Department of Diabetology, Akshay Hospital and Diabetic Speciality Centre and Insulin Pump Centre, Pune, Maharashtra, ¹⁰Department of Diabetology, Poona Diabetes Centre, Pune, Maharashtra, ¹¹Department of Scientific Services, Scientific Services, USV Private Limited, Mumbai, Maharashtra, India

Abstract

Objective: To assess the clinical characteristics, risk factors, and comorbidities associated with type 2 diabetes mellitus (T2DM) in young adult patients. **Methods:** This is a retrospective, multicentric real-world study that included young adults (18–45 years) with T2DM. Primary information including demographics, medical and family history, biochemical measures (pre-and post-prandial blood glucose levels, glycosylated hemoglobin [HbA1c] and blood pressure, and lipid parameters) smoking and drinking habits were collected retrospectively from the medical records of the respective hospitals/clinics. Data were analyzed using descriptive and appropriate comparative statistics. **Results:** A total of 22,921 patients from 623 sites were included. The median age was 37.0 years and the majority were men (61.6%). The proportion of patients from the age group >35–≤45 years was 62.7%. Among all patients, 46.9% had only T2DM; however, 53.1% of patients had T2DM with other comorbidities (T2DM with hypertension, dyslipidemia, and both). The majority of patients had elevated body mass index (BMI) (overweight, 46.6%; and obese, 22.9%). Family history of T2DM (68.1%) were the most common in overall population. Sedentary lifestyle (63.1%), alcohol consumption (38.9%), and regular smoking (23.1%) were the most common associations in patients with T2DM with dyslipidemia and hypertension. Uncontrolled HbA1c level (≥7%) were observed in 79.2% of patients. The level of HbA1c was significantly increased with the duration of T2DM and sedentary lifestyle (p < 0.001). **Conclusion:** Higher BMI, family history of T2DM, sedentary lifestyle, alcohol consumption, and smoking were the most common risk facors, while hypertension and dyslipidemia were the most prevalent comorbidities associated with T2DM in young Indian adults.

Keywords: Diabetes, hypertension, risk factors, sedentary lifestyle, young

Address for correspondence: Dr. Mahesh V. Abhyankar, Vice President, Scientific Services, USV Pvt Ltd, Mumbai, Maharashtra, India. E-mail: dr.mabhyankar@gmail.com d: 28-09-2020 Accepted: 01-03-2021

Received: 28-09-2020 Published: 30-09-2021

Access this article online				
Quick Response Code:	Website: www.jfmpc.com			
	DOI: 10.4103/jfmpc.jfmpc_2010_20			

This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

For reprints contact: WKHLRPMedknow_reprints@wolterskluwer.com

How to cite this article: Saboo B, Agarwal S, Gupta S, Makkar B, Panneerselvam A, Sahoo AK, *et al.* REAL-world evidence of risk factors and comorbidities in YOUNG Indian adults with type 2 diabetes mellitus: A REAL YOUNG (diabetes) study. J Family Med Prim Care 2021;10:3444-52.

Introduction

Diabetes mellitus (DM) is one of the most prevalent noncommunicable diseases that leads to significant morbidity and mortality. The pandemic of DM had affected 9.3% of global population in 2019 and is projected to increase to more than 10% by 2030. In India, the estimated number of people living with diabetes was 77 million in 2019 and it will reach 101 and 134 million by 2030 and 2045, respectively.^[1]

The Indian Council of Medical Research-India Diabetes (ICMR-INDIAB) study reported that around 51% of the adult population (with mean age ranging from 35.8 to 43.9 years) from 15 states of India have diabetes or prediabetes.^[2] This study also highlighted that the take-off point for diabetes was in the age group of 25-34 years in both urban and rural areas of India indicating the early onset of diabetes from the young age.^[2] Indian patients present a high rate of progression from prediabetes to diabetes mainly due to the aggressive nature of the disease.^[3,4] Evidence suggests an early onset of type 2 diabetes mellitus (T2DM) in Indians as compared to Western populations.^[3-5] Several studies have revealed a high prevalence of T2DM in young population.^[6,7] Therefore, it is of paramount importance to screen the young Indian population for early diagnosis of prediabetes to prevent progression to T2DM and the risk of associated comorbidities.

Major risk factors associated with T2DM are obesity, family history, sedentary lifestyle, and comorbidities.^[8] Consumption of energy-dense foods with reduced physical activity is the key contributor to obesity among young people. Family history is another factor associated with the development of T2DM in offsprings. Consequently, the offspring of a parent with T2DM has high chances of having T2DM in adulthood and the risk is even higher when both the parents have T2DM.^[9] Insulin resistance is a common key mechanism in the pathogenesis of T2DM and other comorbid conditions include hypertension, dyslipidemia, polycystic ovary syndrome, and rheumatoid arthritis. These risk factors predominantly affect the young population which could lead to longer disease exposure and increased chronic complications.^[10-12] Hence, the early identification of risk factors is necessary to avoid further complications and comorbidities.

With the view of limited real-world evidence about the prevalence of various risk factors and associated comorbidities among the young adult populations from India, the present real-world study was aimed to determine the patterns of T2DM, risk factors, and comorbidities in young Indian adults.

Materials and Methods

This is a retrospective, multicentric, and real-world study conducted across 623 sites in India. The study protocol was approved by an Independent Ethics Committee. The study was conducted in accordance with the ethical principles that are outlined in the Declaration of Helsinki.

Study population

The present study included young adults of either sex, age within the range of 18 to 45 years who were diagnosed with T2DM. Demographic details (age, sex, weight, height, and sedentary lifestyle), medical and family history, biochemical measures (pre-and post-prandial blood glucose levels, HbA1c, blood pressure, and lipid parameters), smoking and alcohol habits were extracted from hospital/clinical records and a single dataset was created.

Definitions

According to the Standards of Medical Care in Diabetes by the American Diabetes Association (ADA) 2020, optimal glycemic control in nonpregnant adults was defined as HbA1c \geq 7%, while uncontrolled diabetes was defined as HbA1c \geq 7%; fasting blood glucose (FBG) was defined as \geq 130 mg/dL, and postprandial blood glucose (PPG) was defined as \geq 200 mg/dL.^[13]

Obesity: Generalized obesity was defined as BMI $\geq 25 \text{ kg/m}^2$; overweight as BMI 23–25 kg/m², and abdominal obesity was defined as waist $\geq 90 \text{ cm}$ (males) and $\geq 80 \text{ cm}$ (females).^[14]

Statistical analysis

Data were analyzed using the Statistical Package for Social Sciences (SPSS) software, version 23.0. Data were presented as number and percentages or median (interquartile range). Normal distribution of quantitative data was assessed by Shapiro–Wilk test. The comparison of quantitative and qualitative variables between the groups was done using the Mann–Whitney U test and Chi-square test, respectively. A *P* value < 0.05 was considered statistically significant.

Results

A total of 22,921 patients with a median (IQR) age of 37.0 (33.0–39.0) years were included in this study. The proportion of men (61.6%) was higher than women (38.4%). A total of 46.6% of the patients were categorized as overweight, 22.9% of patients were obese, 28.9% were normal weight, and 1.5% were underweight. Among the patients, 46.9% had only T2DM, while the incidence of patients diagnosed with T2DM and hypertension was 27.8% followed by T2DM with dyslipidemia and hypertension (13.5%), and T2DM with dyslipidemia (11.7%). Family history of T2DM was present in 15,368 (68.1%) patients. The median levels of HbA1c, fasting plasma glucose (FPG), and 215.0 mg/dL across all the population, respectively [Table 1]. The correlation coefficient of FPG level was found to be significantly positive with the PPG level ($r^2 = 0.5463$) [Figure 1a]

Age group-wise observations

The majority of patients belonged to the age group of >35 to \leq 45 years (n = 14376). The proportion of patients diagnosed

Parameters	Number of patients (n=22921)
Age (years)	37.0 (33.0-39.0)
Age group (years), n (%)	
≥18-≤25	881 (3.8)
>25-≤35	7664 (33.4)
>35-≤45	14376 (62.7)
Sex, n (%)	
Men	14129 (61.6)
Women	8792 (38.4)
Anthropometric parameters	
Height (cm) [n=22327]	164.0 (158.0-170.0)
Weight (kg) [n=22748]	72.0 (65.0-80.0)
BMI, (kg/m^2) [n=22303]	26.8 (24.3-29.6)
Underweight	336 (1.5)
Normal weight	6453 (28.9)
Overweight	10396 (46.6)
Obese	5118 (22.9)
Waist circumference (cm) [n=16272]	90.0 (74.0-198.0)
Diagnosed with n (%)	
T2DM	10760 (46.9)
T2DM with dyslipidemia	2691 (11.7)
T2DM with HTN	6369 (27.8)
T2DM with HTN and dyslipidemia	3101 (13.5)
Family history, n (%) [$n=22581$]	
T2DM	15368 (68.1)
T2DM with HTN	7578 (33.1)
T2DM with dyslipidemia	4294 (19.2)
T2DM with HTN and dyslipidemia	3451 (15.1)
Smoking habits, n (%) [n=21831]	
Former	1123 (5.1)
Occasional	3470 (15.9)
Regular	3800 (17.4)
No	13438 (61.6)
Alcohol consumption, n (%) [n=22139]	6380 (28.8)
Sedentary lifestyle, n (%) [n=21817]	11348 (52.0)
Duration (days) [n=21275]	600.0 (180-1080)
Biochemical parameters	
FPG (mg/dL) [<i>n</i> =21272]	140.0 (120.0-170.0)
PPG (mg/dL) $[n=21517]$	215.0 (180.0-265.0)
HbA1c (%) $[n=17404]$	7.8 (7.0-8.6)
SBP (mm Hg) [n=9065]	142.0 (130.0-158.0)
DBP (mm Hg) $[p=9034]$	90.0 (83.0-98.0)

Data shown as median (IQR), unless otherwise specified. BMI, body mass index; DBP, diastolic blood pressure; T2DM, type 2 diabetes mellitus; FPG, fasting plasma glucose; HbA1c, hemoglobin A1c; HTN, hypertension; IQR, interquartile range; PPG, postprandial plasma glucose; SBP, systolic blood pressure

with T2DM only was significantly higher in the youngest age group (\geq 18 to \leq 25 years, 62.1%), while the proportion of patients having T2DM with hypertension was significantly higher in the age group of >35 to \leq 45 years (67.8%) (p < 0.001). The incidence of risk factors showed an increasing trend from the \geq 18 to \leq 25 years age group to the >35 to \leq 45 years age group. The median waist circumference was significantly higher in the patients of age group >35 to \leq 45 years (90 cm) as compared to the other two age groups (\geq 18– \leq 25 years: 87cm; and >25– \leq 35 years: 89 cm) (p < 0.001) [Table 2].

A family history of T2DM was reported in more than 60% of the patients across all the age groups (≥ 18 to ≤ 25 years, 61.1%; ≥ 25



Figure 1: Scatter plots showing the correlation between a) FPG and PPG, b) HbA1c and age, and c) HbA1c and BMI. BMI, body mass index; FPG, fasting plasma glucose; HbA1c, hemoglobin A1c; HTN, hypertension; PPG, postprandial plasma glucose

to \leq 35 years, 66.7%; and > 35 to \leq 45 years, 69.2%) [Figure 2a]. Regular smoking and alcohol consumption were more common in patients of the age group >35 to \leq 45 years (p < 0.001), while a sedentary lifestyle was observed in over 45% of the population across all the age groups. The mean BMI showed a significantly increasing trend from the youngest age group to the oldest group (p < 0.001) [Figure 3a]. A significantly higher level of mean FPG was observed in patients of the age group >35 to \leq 45 years (142.0 mg/dL) compared to the other age groups [Figure 3b, *P* < 0.001]. The mean PPG and HbA1c concentrations were significantly higher in the patients of age group >35 to \leq 45 years (216 mg/dL and 7.8%, respectively) as compared to patients of the age group >25– \leq 35 years (p < 0.001) [Figure 3c and 3d].

Table 2: Age group-wise analysis									
Parameters	Group A (≥18-≤25 years) (<i>n</i> =881)*	Group B (>25-≤35 years) (<i>n</i> =7664)**	Group C (>35-≤45 years) (<i>n</i> =14376) [#]	Р					
Age (years)	24.0 (21.0-25.0)	32.0 (30.0-34.0)	39.0 (28.0-40.0)	<0.001 ^{a-c}					
Sex, n (%)				< 0.001					
Men	496 (56.3)	4505 (58.8)	9128 (63.5)						
Women	385 (43.7)	3159 (41.2)	5248 (36.5)						
Waist circumference (cm)	[n=656]	[n=5501]	[n=10115]	$0.005^{a}, < 0.001^{b,c}$					
	87.0 (79.0-95.7)	89.0 (80.0-96.5)	90.0 (80.0-98.0)						
Diagnosed with, n (%)									
T2DM	547 (62.1)	4083 (53.3)	6130 (42.6)	< 0.001					
T2DM with dyslipidemia	81 (9.2)	869 (11.3)	1741 (12.1)						
T2DM with HTN	162 (18.4)	1892 (24.7)	4315 (67.8)						
T2DM with HTN, and dyslipidemia	91 (10.3)	820 (10.7)	2190 (15.2)						
Smoking habits, n (%)	[n=831]	[<i>n</i> =7279]	[n=13721]						
Former	72 (8.7)	404 (5.6)	647 (4.7)	< 0.001					
Occasional	68 (8.2)	1081 (14.9)	2321 (16.9)						
Regular	66 (7.9)	1107 (15.2)	2627 (19.1)						
No	625 (75.2)	4687 (64.4)	8126 (59.2)						
Alcohol consumption, n (%)	[n=841]	[<i>n</i> =7412]	[n=13886]	< 0.001					
	132 (15.7)	2017 (27.2)	4231 (30.5)						
Sedentary lifestyle, n (%)	[n=825]	[n=7334]	[n=13658]	< 0.001					
	388 (47.0)	3639 (49.6)	7321 (53.6)						
Duration (days)	[n=835]	[n=7088]	[n=13352]	0.552^{a} , $< 0.001^{b,c}$					
	420.0 (180.0-1080.0)	480.0 (240.0-1020.0)	720.0 (360.0-1080.0)						

Data shown as median (IQR), unless otherwise specified. *n=881; **n=7664; "n=14376 unless otherwise specified. *group A vs B; ^bgroup B vs C; ^cgroup A vs C. T2DM, type 2 diabetes mellitus; HTN, hypertension; IQR, interquartile range.



Figure 2: Distribution of patients according to family history (a) and age group and (b) diagnosis. DM, diabetes mellitus; HTN, hypertension

Diagnosis-wise observation

The majority of patients belonged to the age group of >35 to \leq 45 years among all the diagnosis groups (T2DM, 57.0%; T2DM with hypertension, 64.7%; T2 DM with dyslipidemia, 67.8%; and T2DM with hypertension and dyslipidemia, 70.6%). Alcohol consumption was highest in the patients diagnosed with T2DM with dyslipidemia and hypertension (p < 0.001). Regular smoking was highest in patients having T2DM with dyslipidemia and hypertension (23.1%) and T2DM with dyslipidemia (22.4%). A sedentary lifestyle was more common in patients having T2DM with hypertension and dyslipidemia (63.1%) compared to other diagnosis (p < 0.001) [Table 3]. The median PPG level was higher in the patients diagnosed with T2DM along with hypertension and dyslipidemia compared to patients with T2DM, T2DM with hypertension, and T2DM with dyslipidemia. The

median HbA1c levels showed poor glycemic control across all the groups (7.6%–8.0%) [Table 3]. A family history of T2DM was reported in majority of the patients (range, 63.7%-74.6%) across all the diagnosis-wise groups. A family history of hypertension was most common in patients diagnosed with T2DM with hypertension (56.1%) and T2DM with dyslipidemia and hypertension (60.2%) (p < 0.001) [Figure 2b].

Glycosylated hemoglobin HbA1c level-wise observation

The majority of patients diagnosed with T2DM (n = 13796) had uncontrolled HbA1c levels (\geq 7%). Of patients with uncontrolled T2DM, a substantial proportion (n = 10793, 78.2%) had levels of HbA1c in the range of \geq 7% to \leq 9% while 3003 (21.8%) patients had HbA1c levels >9%. Only

3608 (20.7%) patients had controlled HbA1c levels (<7%). Regression analysis showed a large variability between age and BMI with HbA1c [Figure 1b and 1c]. The levels of HbA1c were significantly increased with increasing duration of T2DM (<7%, 540 days; \geq 7 to \leq 9%, 600 days; \geq 9%, 720 days; P < 0.001) and sedentary lifestyle (<7%, 51.3%; \geq 7 to \leq 9%, 51.5%; \geq 9%, 55.3%; P = 0.001).



Figure 3: Age group-wise analysis. (a) BMI, (b) FPG, (c) PPG, and (d) HbA1c. *Group A vs B, Group B vs C, Group A vs C. Group A, $\geq 18-\leq 25$ years; Group B, $>25-\leq 35$ years; Group C, $>35-\leq 45$ years. BMI, body mass index; FPG, fasting plasma glucose; HbA1c, hemoglobin A1c; PPG, postprandial plasma glucose

Discussion

The present study evaluated the risk factors associated with T2DM along with comorbidities like hypertension and dyslipidemia in young Indians diagnosed with T2DM in the real-world setting. The key findings were as follows: majority of patients were men in the age group of >35 to \leq 45 years; around 70% of population had elevated BMI, more than half of the patients had additional associated one or two comorbidities (hypertension, dyslipidemia or both); family history of T2DM, sedentary lifestyle, elevated BMI, alcohol consumption, and regular smoking were the common risk factors associated with T2DM; the incidence of uncontrolled HbA1c level was very common (79.2%); and the risk of elevated HbA1c increased with increasing duration of T2DM and sedentary lifestyle.

Early diagnosis of T2DM has become a fundamental step in the management and reducing the complications and mortality risk associated with T2DM. The present study showed that the incidence of T2DM was strongly associated with nonmodifiable risk factors including age and family history. Around 60% of patients in the youngest age group were diagnosed with T2DM alone. Similarly, an evidence-based STEPS study reported that the prevalence of diabetes was 51% in patients of the age group 25–44 years, suggesting that half of the population were young adults.^[15] The burden of DM in India has increased with the

Table 3: Diagnosis-wise analysis								
Parameters	Group A (T2DM only) (<i>n</i> =10760)*	Group B (T2DM with dyslipidemia) (n=2691)**	Group C (T2DM with HTN) (<i>n</i> =6369) [#]	Group D (T2DM with Dyslipidemia and HTN) (n=3101) ^{##}	Р			
Age (years)	36.0 (32.0-39.0)	38.0 (34.0-40.0)	37.0 (34.0-39.0)	38.0 (35.0-40.0)	<0.001 ^{a-f} , 0.009 ^d			
Age group (years), n (%)					< 0.001			
≥18-≤25	547 (5.1)	162 (2.5)	81 (3.0)	91 (2.9)				
>25-≤35	4083 (37.9)	1892 (29.7)	869 (32.3)	820 (26.4)				
>35-≤45	6130 (57.0)	4315 (67.8)	1741 (64.7)	2190 (70.6)				
Sex, n (%)					< 0.001			
Men	6393 (59.4)	4012 (63.0)	1697 (63.1)	2027 (65.4)				
Women	4367 (40.6)	2357 (37.0)	994 (36.9)	1074 (34.6)				
BMI (kg/m ²)	[n=10457]	[n=6182]	[n=2629]	[n=3035]	<0.001 ^{a-f} , 0.041 ^d			
	26.2 (23.8-28.9)	27.1 (24.8-29.8)	27.4 (25.0-30.0)	27.8 (25.3-30.8)				
Waist circumference (cm)	[n=7345]	[n=4491]	[n=2041]	[n=2395]	$0.166^{a}, 0.007^{b}, < 0.001^{c},$			
	89.0 (80.0-96.0)	90.0 (77.0-96.5)	92.0 (83.0-98.0)	91.0 (80.0-100.0)	$0.689^{d}, 0.158^{e}, 0.132^{f}$			
Smoking habits, n (%)	[n=10312]	[n=6127]	[n=2555]	[n=2837]				
Former	487 (4.7)	351 (5.7)	119 (4.7)	166 (5.9)	< 0.001			
Occasional	1443 (14.0)	1128 (18.4)	400 (15.7)	499 (17.6)				
Regular	1360 (13.2)	1374 (22.4)	411 (16.1)	655 (23.1)				
No	7022 (68.1)	3274 (53.4)	1625 (63.6)	1517 (53.5)				
Alcohol consumption, n (%)	[n=10440]	[n=6209]	[n=2608]	[n=2882]	< 0.001			
	2271 (21.8)	2121 (34.2)	868 (33.3)	1120 (38.9)				
Sedentary lifestyle, n (%)	[n=10268]	[n=6096]	[n=2589]	[n=2864]	< 0.001			
	4849 (47.2)	3196 (52.4)	1497 (57.8)	1806 (63.1)				
FPG (mg/dL)	[n=9929]	[n=5856]	[n=2573]	[n=2914]	0.011^{a} , $< 0.001^{b,c}$, 0.500^{d} ,			
	140.0 (120.0-170.0)	142.0 (124.0-170.0)	140.0 (120.0-175.0)	141.0 (121.0-180.0)	0.148°, 0.244 ^f			
PPG (mg/dL)	[n=10055]	[n=5947]	[n=2568]	[n=2947]	0.006^{a} , $< 0.001^{b,c}$, 0.146^{d} ,			
	210.0 (177.0-262.0)	219.0 (188.0-262.0)	217.0 (180.0-270.0)	220.0 (183.0-278.0)	0.009°, 0.091 ^f			
HbA1c (%)	[n=8038]	[n=4624]	[n=2223]	[n=2519]	<0.001 ^{a,c,d,f} , 0.048 ^b , 0.006 ^e			
· ·	7.6 (7.0-8.5)	7.8 (7.1-8.4)	7.9 (7.1-8.9)	8.0 (7.2-9.0)				

Data shown as median (IQR), unless otherwise specified. *n=10760; **n=2691; "n=6369; ""n=3101, unless otherwise specified. 'group A vs B; 'group A vs D; 'group A vs D; 'group B vs D; 'group B vs D; 'group B vs D; 'group C vs D. BMI, body mass index; T2DM, type 2 diabetes mellitus; FPG, fasting plasma glucose; HbA1c, hemoglobin A1c; HTN, hypertension; IQR, interquartile range; PPG, postprandial plasma glucose

increasing incidences of T2DM diagnosis (from 5.5% to 7.7%) in the adult population (20 years and older) in the last few decades.^[16]

A family history of T2DM is the most important nonmodifiable risk factor responsible for the early occurrence of T2DM. This study showed that more than 60% of patients across all age groups had a family history of T2DM. Similar findings were observed in studies done by Patel et al. in Gujarat and Geetha et al. in Tamil Nadu which reported positive family history of DM in around 60% of the population.^[17,18] People with a positive family history are more prone to develop DM at the early stages of life.^[17] This showed that family history of T2DM is highly associated with the increasing risk of T2DM in young population. Early screening in young population can be helpful in identifying the people with a positive history of DM and can be made aware of the early risk of diabetes and modify the lifestyle accordingly. Behavioral modification including proper physical activity and healthy diet will be advised to these patients for delaying the early onset of this disease.

In the present study, among patients diagnosed with T2DM, hypertension (27.8%) was the most common comorbidity followed by dyslipidemia (11.7%) and the triad of T2DM, dyslipidemia, and hypertension (13.5%). Dyslipidemia and hypertension are the major risk factors associated with macrovascular diseases and emphasizing strict glycemic control may help in delaying or preventing macrovascular disease. The comorbid association of T2DM and dyslipidemia was strongly associated with the age group of young adults. Similarly, a STEPS survey reported the prevalence of T2DM-associated comorbidities and among patients with T2DM as 60% of them had hypertension and 36% patients had dyslipidemia.^[15]

The present study is a large-scale study involving a large cohort of the young Indian adults and therefore could provide the evidence with respect to the strong correlation between these risk factors.^[19,20] A recently published cross-sectional study evaluated that lifestyle factors including alcoholism, smoking, obesity, and family history were the risk factors related to diabetes.^[21] In the present study, the prevalence of T2DM was more common in young adult patients of age ranging from 25 to 45 years. Further, family history of T2DM, regular smoking, and sedentary lifestyle were the other risk factors observed in our population. A recent meta-analysis also demonstrated supporting evidence with respect to common risk factors associated with T2DM that include smoking, sedentary lifestyle, and obesity.^[22]

The target level of HbA1c in patients with T2DM is usually less than 7%.^[13] A high level of HbA1c increases the risk of T2DM-related complications. Therefore, an early intensive management of uncontrolled HbA1c is needed to reduce these complications. In the current study, a majority of patients had uncontrolled HbA1c levels (\geq 7%). Also, a rising trend was observed with respect to FPG and PPG levels with the severity of HbA1c. This is in accordance with an observational study on patients diagnosed with T2DM who had high mean levels of HbA1c (8.5%) with increased FPG (165.4 mg/dL) and PPG (258.4 mg/dL).^[23] A recent TIGHT (The Investigation of Glycosylated Hemoglobin on Therapy in Indian Diabetics) study evaluated glycemic control in adult Indians and their association with microvascular complications. The authors revealed that more than half (53.1%) of the patients were young adults (<55 years) and poor glycemic control was very common in Indian population with T2DM (76.6%).^[19] These observations are in concordance with the present study and together indicate a high burden of poor glycemic control in Indian adults.

Several limitations of this study should be considered. This study did not record the socioeconomic and educational statuses of the patients, medications consumed by the patients which could have added valuable data while inferring the observations.

Conclusion

The overall observations indicate that men from the age group of 25–45 years were most commonly affected by T2DM. Elevated BMI, family history of T2DM, sedentary lifestyle, alcohol consumption, and smoking are the risk factors associated with DM in young Indian adults. Hypertension and dyslipidemia are the prevalent comorbidities associated with T2DM. Hence, early diagnosis of diabetes and the associated comorbidities and treating hyperglycemia and its comorbidities to target levels early in the course of disease is necessary for alleviating the long-term risk of T2DM-related complications.

Acknowledgements

We acknowledge Ms. Farida Hussain, Mr. A Thamburaj and Ms. Shashikala Borhade of USV Pvt. Ltd. for their assistance in carrying out the project. The medical writing support was provided by Dr. Pradip Mate from the Scientific Services Team of USV Pvt. Ltd. and Ms. Snehal Khanolkar from Sqarona Medical Communications LLP (Mumbai). We acknowledge BioQuest Solutions Pvt. Ltd for their services in the conduction of the real-world study.

Contributors: Dr. A Balachandran, Dr. A K Gupta, Dr. A Kamlesh, Dr. A M Rao, Dr. A Muthukumaran, Dr. A Shahul Hameed, Dr. A Shanmugam, Dr. A Siddharth Prasad, Dr. A Syed Sultan Ibrahim, Dr. A Vidya Sagar, Dr. A Viswanathan, Dr. Abani Kumar Patro, Dr. Achal Tiwari, Dr. Ajay H Kantharia, Dr. Ajay Jain, Dr. Ajay Kumar Sinha, Dr. Ajay Patwari, Dr. Ajay Sharma, Dr. Ajay V Kaduskar, Dr. Ajay Yadav, Dr. Ajeet Singh Kothari, Dr. Ajit Kumthekar, Dr. Ajit Sawhney, Dr. Ajoy Kumar Tewari, Dr. Akula Vidya Sagar, Dr. Alam Nawaz, Dr. Almas Talib, Dr. Ambairam Viswanathan, Dr. Amish V Shah, Dr. Amit Maheswari, Dr. Amit Rastogi, Dr. Amitesh Kumar Chatterjee, Dr. Amrut Sultane, Dr. Anand Sudhakar Bhave, Dr. Anand Swaroop Menawat, Dr. Anantha Raman, Dr. Anil Bhargava, Dr. Anil Kumar Agarwal, Dr. Anil Kumar Bhatt, Dr. Anil Maruti Shewale, Dr. Anil Saraswat, Dr. Anilkumar A Kustagi, Dr. Anuj Kumar Saha, Dr. Anurag Bajpai, Dr. Anurag Srivastava, Dr. Arjun Baidya, Dr. Arun Kamble, Dr. Arun Kumar, Dr. Arun Singhal, Dr. Arvind K Mishra, Dr. Arvind Kumar, Dr. Arvind Sharma, Dr. Ashok Appasaheb Jadhav, Dr. Ashok B Malipatil, Dr. Ashok H Sancheti, Dr. Ashok Kumar Yadav, Dr. Ashok M Parekh, Dr. Ashok Solanki, Dr. Ashok Varma, Dr. Ashwin K Vaghani, Dr. Ashwini Joshi, Dr. Asish Kumar Basu, Dr. B A Rudrawadi, Dr. B Ashok Kumar, Dr. B Bosco, Dr. B K Kakkad, Dr. B K Srinivasa Murthy, Dr. B Karthik Rao, Dr. B R K Reddy, Dr. B Rajaganesan, Dr. B Rama Krishna Reddy, Dr. B Ramulu, Dr. B Siva SubramanyamDr. B Sivasita Rammiah, Dr. Bala Raju Gundam, Dr. Balbhadra Kumar Agrawal, Dr. Balraj Bosco, Dr. Bandaru Shiva Subrahmanyam, Dr. Basavana Gouda, Dr. Bharat Bhushan Mittal, Dr. Bhaskar M Patil, Dr. Bhubeneswar Dutta, Dr. Binay Prasad, Dr. Binoda Nand Jha, Dr. Biplab Bandhopadhyay, Dr. Birju S Mori, Dr. Biswanath Biswas, Dr. Bobby K Mathew, Dr. Bongu Karthik Rao, Dr. C Somanathan, Dr. C H Naveen Kumar, Dr. C Jagadeesh, Dr. C Somanathan, Dr. C T Patil, Dr. Chaitanya N Buva, Dr. Chandan Kumar Patra, Dr. Chandan Sarmah, Dr. Chandrakant Gudage, Dr. Chandrakant Patil, Dr. Chandrashekhar Desai, Dr. Chetan I Velani, Dr. D Ganekal Prashanth, Dr. D Ragothaman, Dr. D S Prasad, Dr. Dante Ruskin, Dr. Dayanidhi N Meher, Dr. Debaprasad Chakraborty, Dr. Debarchan C Jena, Dr. Debasis Giri, Dr. Debotosh Sen Purkayastha, Dr. Deepak Kumar Dasmohapatra, Dr. Deepak S Bhosle, Dr. Deepal K Parekh, Dr. Devang M Desai, Dr. Dhanaraj Singh Chongtham, Dr. Dharmesh Jain, Dr. Dharmesh Solanki, Dr. Dhruv K Singh, Dr. Dileep Nagesh Mane, Dr. Dilip Joshi, Dr. Dilip Kumar Gupta, Dr. Dinesh Agarwal, Dr. Dinesh K Garg, Dr. Dinesh Kansal, Dr. Dinesh Kumar, Dr. Dnyanoba Bhaskar, Dr. Dosapati Ramesh, Dr. Durga Prasad Bhimala, Dr. Dwarakanath C S, Dr. E M Surendra, Dr. Faiz Ahmad, Dr. G Aravindan, Dr. G Bala Raju, Dr. G Kiran, Dr. G Mohan, Dr. G Pavan Kumar Reddy, Dr. G Raja Gopal, Dr. G S Gupta, Dr. G S Mahishale, Dr. G Sathish Kumar, Dr. G Satya Sreenivasa Rao, Dr. G T Rane, Dr. G V Siva Reddy, Dr. G Vijavakumar, Dr. Gajendra S Mahishale, Dr. Ganesh H K, Dr. Ganesha Moorthy S N, Dr. Ganga Kiran, Dr. Gangadhara T K, Dr. George Paulose, Dr. Girija Nandan Singh, Dr. Gopala Venkata Giridhar, Dr. H S Paul, Dr. Hari Har Trivedi, Dr. Hari Vinaya Kumar, Dr. Harish Basera, Dr. Himanshu Mehta, Dr. Hiranmoy Bhattacharyya, Dr. Hiranmoy Paul, Dr. Ibrahim Mohamed Rowther, Dr. J Aman Kumar, Dr. J Girithara Gopala Krishnan, Dr. J Nagaraju, Dr. J Rajesh Kumar, Dr. J Sangumani, Dr. Jagdish Gotur , Dr. Jagulal Agrawal, Dr. Javedahmed A Shaikh, Dr. Jayashri P Shembalkar, Dr. Jayesh Patel, Dr. Jayesh Shah, Dr. Jayshree Swain, Dr. Jibesh Kumar Sarkar, Dr. Jilse George, Dr. Jitendra Singh Kushwaha, Dr. K Arumugam, Dr. K Balakrishnan, Dr. K Bhuvanesh, Dr. K C Jindal, Dr. K D Purohit, Dr. K Eswaran, Dr. K G Suresh, Dr. K Hari Babu, Dr. K Jayarami reddy, Dr. K M Jeyabalaji, Dr. K Nagesh, Dr. K R Ravish, Dr. K S R Swany, Dr. K Sarath Chandra Babu, Dr. K Seetharamaiah, Dr. KV Asaithambi, Dr. Kailash Chand Lohani, Dr. Kalinga B E, Dr. Kalpana S Mehta, Dr. Kalyan Mitra, Dr. Kalyan Mullick, Dr. Kamlesh Thakker, Dr. Kampa Sarat Chandra Babu, Dr. Kanakesa Thevar Balakrishnan, Dr. Kandula Sai, Dr. Kapil S Borawake, Dr. Kasargaon Paten Vitthal Rao, Dr. Kedar Chand Jindal, Dr. Kiran Belsare, Dr. Kishan M Jani, Dr. Konatham Haribabu, Dr. Krishna Kumar, Dr. Kshitij Shankhdhar, Dr. Kuchatu Sreedharan Udayabhanu, Dr. Kummaraganti Sitarama Swamy, Dr. Laxman Kumar Sahu, Dr. Liladhar S Vora, Dr. Loganathan M, Dr. Lokesh Abrol, Dr. M A Karmur, Dr. M Bheemasena Achar, Dr. M Durairaj, Dr. M E Ramakrishnan, Dr. M J Stanley Ambroise, Dr. M K Guha, Dr. M Manju Bhargavi, Dr. M Nagesh Prabhu, Dr. M P Patel, Dr. M R A Rafique, Dr. M Ramesh Kumar, Dr. M S Suri, Dr. M Viyay Mohan, Dr. Madan Pal Singh, Dr. Madhav K Mandal, Dr. Madhav Pavuluri, Dr. Madhukar Rai, Dr. Mahendra Kumar Jaiswal, Dr. Mahesh Omprakash Baheti, Dr. Mallikarjuna Rao, Dr. Mamtesh Gupta, Dr. Manish Sachdev, Dr. Manmohan Sharma, Dr. Manohar K N, Dr. Manoj Hari Naik, Dr. Manoj Kabra, Dr. Manoj Kumar Dash, Dr. Manoj Kumar Rawat, Dr. Manoj Kumar Shrivastav, Dr. Manoj S Chawla, Dr. Metta Madhu, Dr. Mohaideen Pitchai, Dr. Mohammad Jameel, Dr. Mohammed Parvez Syed, Dr. Mohan Magdum, Dr. Mohammad Jameel, Dr. Mohammad d Parvez, Dr. Mohit Kumar Santra, Dr. Monika Mahajan, Dr. Mritunjay Behera, Dr. Mritunjay Kumar Sinha, Dr. Munibalakrishna S, Dr. Murali Madhav, Dr. Murthy S, Dr. N B Srivastava, Dr. N Ethiraj, Dr. N M D Athaullah, Dr. N Muthukumarasamy, Dr. N S Ramesh, Dr. N Sudha, Dr. Nagaraj S, Dr. Nandakumar S, Dr. Narayan P Gherwara, Dr. Narottam Kumar Singh, Dr. Nasimullah Attaullah Mohd, Dr. Natarajan Selvarajan, Dr. Navneet Agarwal, Dr. Neelambar Bhatt, Dr. Neeraj Chaudhary, Dr. Neeresh Jain, Dr. Nellaiappan Muthukumarasamy, Dr. Nihar Ranjan Samal, Dr. Nilesh M Detroja, Dr. Nipun Gupta, Dr. Niranjan Tripathy, Dr. Nirmal N Nehete, Dr. Nitin Vijay Gore, Dr. Noor Khan, Dr. Om Prakash Prasad, Dr. Omer Mustafa Hasan, Dr. P Amareshwar, Dr. P Chakradhar, Dr. P L Saravanan, Dr. P Murali Madhav, Dr. P Naga Bhushanam, Dr. P Omprakash, Dr. P Ranga Prasad, Dr. P Ravi Kaladhar Reddy, Dr. P Ravi Kumar, Dr. P Satish Kumar, Dr. P Subramanyam Reddy, Dr. P V B Sridhar, Dr. Palaniappan Saravanan, Dr. Pankaj Desai, Dr. Pankaj Jain, Dr. Pankaj Kumar Agarwal, Dr. Pankaj Kumar Gupta, Dr. Paramesh S, Dr. Paras Jain Dr. Parimal Swamy, Dr. Patnala Chakradhar, Dr. Pawan Agarwal, Dr. Phillips Routray, Dr. Pinakin Soni, Dr. Piyush Desai, Dr. Pothamsetty Ravikaladhar Reddy, Dr. Pothiraju Naicker Subbiah, Dr. Prabal Kumar Ghosh, Dr. Prabhakar R, Dr. Prabhat Agarwal, Dr. Prabhat Pandey, Dr. Prabhu Rami Reddy, Dr. Pradeep N Shantagiri, Dr. Prahlad R Chebbi, Dr. Prakash Chandra Patra, Dr. Prakash N, Dr. Prakash S Shende, Dr. Prakash T V, Dr. Pramod D Paritekar, Dr. Pranabes Ray, Dr. Prasanta Kumar Mishra, Dr. Prasanth Sankar, Dr. Pratap Kumar Mishra, Dr. Preamkumar Asokumar, Dr. Preeti Gholap, Dr. Prem Kumar Madkan, Dr. Pruthvi B C, Dr. Punit Arora, Dr. Puttaraj K R, Dr. R Balamurugan, Dr. R K Khinsvera, Dr. R K Ramesh, Dr. R Mehetre, Dr. R P Chaubey, Dr. R Prabakar, Dr. R Rajapaul, Dr. R Rajarathinam, Dr. R Rajendra Prasad, Dr. R Rajendran, Dr. R Ravi Chandra Reddy, Dr. R Srinivasan, Dr. Rachamallu Ravichandra Reddy, Dr. Rahamuddin Ahmed, Dr. Rahul Jain, Dr. Rahul Kapur, Dr. Rahul Mathur, Dr. Raj Kumar Lalwani Dr. Rajapaul Ramasamy, Dr. Rajarathinam, Dr. Rajeev Agarwal, Dr. Rajeev Kasliwal, Dr. Rajeev Tyagi, Dr. Rajendra B Auti, Dr. Rajendra Kumar Khinvasara, Dr. Rajesh Nanda, Dr. Rajesh Regonda, Dr. Rajeshwar Kant Chandra Mishra, Dr. Rajeshwar Singh, Dr. Rajnish Kumar Dahuja, Dr. Rajnish Saxena, Dr. Rakesh Agarwal, Dr. Rakesh Kumar Aran, Dr. Rakesh Kumar Singh, Dr. Rakesh Prasad, Dr. Ram Kumar Bhardwaj, Dr. Raman Boddula, Dr. Ramesh Prasad Srivastava, Dr. Ranabir Bose, Dr. Ranasing D Wabale, Dr. Ranga P Prasad, Dr. Rantu Barman, Dr. Rasheed Ali, Dr. Ravi Kant Mishra, Dr. Regonda Rajesh, Dr. Ritesh Kumar Agrawala, Dr. Riyaj Umar Mujawar, Dr. Rohan V Ainchwar, Dr. Roshan M, Dr. S Azhagam Perumal, Dr. S Chandrashekar, Dr. S D M Sekhar, Dr. S Iqbaluddin Ahmed, Dr. S Jaidev, Dr. S N Ganesha Moorthy, Dr. S Nagaraj, Dr. S P Sathish Kumar, Dr. S P Singh, Dr. S Pothiraju, Dr. S Rajkumar, Dr. S Sahubar Sadique, Dr. S Sathyanarayana Murthy, Dr. S Vanchilingam, Dr. S Velavutham, Dr. Sabeer T K, Dr. Sachin Mohan Lakade, Dr. Sailesh Lodha, Dr. Sandeep Bhatnagar, Dr. Sandeep Kumar Gupta, Dr. Sanjay Garg, Dr. Sanjay Gujrati, Dr. Sanjay Gupta, Dr. Sanjay Kishor, Dr. Sanjay Kumar Giri, Dr. Sanjay Kumar Jain, Dr. Sanjay Mahajan, Dr. Sanjay More, Dr. Sanjay S More, Dr. Sanjeev R Phatak, Dr. Sanjeev Saxena, Dr. Sankalingam Azhagam perumal, Dr. Sankar Joyti Parashar, Dr. Sarat Chandra Choudhury, Dr. Sarat Chandra Mohanty, Dr. Sarat Keot, Dr. Satish Raikar, Dr. Satish Sutradhar, Dr. Savita Agarwal, Dr. Seema A Bagri, Dr. Shah Abrar, Dr. Shahul Hameed, Dr. Shailesh Narayan Palekar, Dr. Shailesh Srivastava, Dr. Shamsudeen M, Dr. Shamsudheen S, Dr. Shankar Jyoti Parashar, Dr. Shantharaman D, Dr. Sharad Kumar, Dr. Sharan R Pawad shetter, Dr. Shehla Shaikh, Dr. Shivaprasad C, Dr. Shrikant Shankarrao Wasavade, Dr. Snehal R Tanna, Dr. Sri Krishna Mundhra, Dr. Sri Prasad Mohanty, Dr. Srinath Vittal Kasal, Dr. Stanley Ambroise, Dr. Subhash Chandra Javaram Balleke, Dr. Subodh Banzal, Dr. Subodh Chandra, Dr. Subrata Bhattacharya, Dr. Sudha N Sathiyan, Dr. Sudharshan Murthy Karkala Achutha, Dr. Sudhir Ranjan Pattnaik, Dr. Sukhen Kumar Saha, Dr. Suman Kirti, Dr. Sumit Mukherjee, Dr. Sunil Bansal, Dr. Sunil Dhand, Dr. Surekha B Shetty, Dr. Suren G Chavan, Dr. Surendra Kumar Bhatter, Dr. Suresh Agrawal, Dr. Suresh Mittal, Dr. Suresh S Sawardekar, Dr. Sushil Jhawar, Dr. Sushil Kumar, Dr. Sushil Shukla, Dr. Sushil Upadhyay, Dr. Suyog Jawahar Doshi, Dr. Swayamsidha Mangaraj, Dr. Syed Mohammad Razi, Dr. Syed Sultan Ibrahim, Dr. T K Sabeer, Dr. T M Venkateswara Rao, Dr. Taruni Devi N G, Dr. Tejpal Shah, Dr. U Rajanikanth, Dr. Uday Pundalik Nayak, Dr. Udaya Bhanu K S, Dr. Umesh Akkalkotkar, Dr. Umesh Masand, Dr. Umesh Singh, Dr. Usmangani I Khatri, Dr. V Channaraya, Dr. V Dineshkumar, Dr. V Mahadevan, Dr. V Manakavala Perumal, Dr. V Rajendran, Dr. V Sandeep Reddy, Dr. V Saravanan, Dr. V Sathiyamoorthy, Dr. V Vigneshwaran, Dr. V Y Manakavala Perumal, Dr. Varsha Jagtap, Dr. Venkataraman S, Dr. Vijay Gurung, Dr. Vijay Kumar Aggarwal, Dr. Vijay Narayan Tiwari, Dr. Vikas Namdeo Desale, Dr. Vikas S Patil, Dr. Vikram Jain, Dr. VIkrant B Ghatnatti, Dr. Vinay Kumar Dhandhania, Dr. Vinay Kumar Singh, Dr. Vinay Sinha, Dr. Vinayak S Kubal, Dr. Vineet Agrawal, Dr. Vineet Sabharwal, Dr. Vipin Srivastava, Dr. Virendra C Chauhan, Dr. Virendrasinh C Chauhan, Dr. Vishal Chopra, Dr. Vishal Kastwar, Dr. Vishwanath Parsewar, Dr. Vivek Mehta, Dr. Vivek Narasinh Annachhatre, Dr. Y S Ravi Kumar, Dr. Y S Tomar, Dr. Yalamanchi Sadasiva Rao, Dr. Yash Bahulikar, Dr. Yogesh Mehrotra, and Dr. Zia Ul Haque.

Ethics approval and consent to participate

The study protocol was approved by an Independent Ethics Committee. As this was a retrospective study, consent was not obtained.

Financial support and sponsorship

The study was funded by USV Pvt Ltd.

Conflicts of interest

Dr. Mahesh V. Abhyankar and Dr. Santosh Revankar are employees of USV Pvt Ltd. All other authors have no conflicts of interest.

References

- 1. Saeedi P, Petersohn I, Salpea P, Malanda B, Karuranga S, Unwin N, *et al.* Global and regional diabetes prevalence estimates for 2019 and projections for 2030 and 2045: Results from the international diabetes federation diabetes atlas, 9th ed.ition. Diabetes Res Clin Pract 2019;157:107843. doi: 10.1016/j.diabres. 2019.107843.
- 2. Anjana RM, Deepa M, Pradeepa R, Mahanta J, Narain K, Das HK, *et al.* Prevalence of diabetes and prediabetes in 15 states of India: Results from the ICMR-INDIAB population-based cross-sectional study. Lancet Diabetes Endocrinol 2017;5:585-96.
- 3. Anjana RM, Shanthi Rani CS, Deepa M, Pradeepa R, Sudha V, Divya Nair H, *et al.* Incidence of diabetes and prediabetes and predictors of progression among Asian Indians: 10-year follow-up of the Chennai urban rural epidemiology study (CURES). Diabetes Care 2015;38:1441-8.
- 4. Dutta D, Mondal SA, Kumar M, Hasanoor Reza AH, Biswas D, Singh P, *et al.* Serum fetuin-A concentration predicts glycaemic outcomes in people with prediabetes: A prospective study from Eastern India. Diabet Med 2014;31:1594-9.
- 5. Dutta D, Mukhopadhyay S. Intervening at prediabetes stage is critical to controlling the diabetes epidemic among Asian Indians. Indian J Med Res 2016;143:401-4.
- 6. Sahoo SK, Zaidi G, Vipin VP, Chapla A, Thomas N, Yu L, *et al.* Heterogeneity in the aetiology of diabetes mellitus in young adults: A prospective study from north India. Indian J Med Res 2019;149:479-88.
- 7. Thanabalasingham G, Pal A, Selwood MP, Dudley C, Fisher K, Bingley PJ, *et al.* Systematic assessment of aetiology in adults with a clinical diagnosis of young-onset type 2 diabetes is a successful strategy for identifying maturity-onset diabetes of the young. Diabetes Care 2012;35:1206-12.
- 8. Oommen AM, Abraham VJ, George K, Jose VJ. Prevalence of risk factors for non-communicable diseases in rural & urban Tamil Nadu. Indian J Med Res 2016;144:460-71.
- 9. Ashok P, Kharche JS, Joshi AR. Evaluation of risk for type 2 diabetes mellitus in medical students using Indian Diabetes Risk Score. Indian J Med Sci 2011;65:1-6.
- 10. Mizokami-Stout K, Cree-Green M, Nadeau KJ. Insulin resistance in type 2 diabetic youth. Curr Opin Endocrinol Diabetes Obes 2012;19:255-62.
- 11. Würtz P, Mäkinen VP, Soininen P, Kangas AJ, Tukiainen T,

Kettunen J, *et al.* Metabolic signatures of insulin resistance in 7,098 young adults. Diabetes 2012;61:1372-80.

- 12. Ormazabal V, Nair S, Elfeky O, Aguayo C, Salomon C, Zuñiga FA. Association between insulin resistance and the development of cardiovascular disease. Cardiovasc Diabetol 2018;17:122.
- 13. American Diabetes Association. 2. Classification and diagnosis of diabetes: Standards of medical care in diabetes-2020. Diabetes Care 2020;43:S14-31.
- 14. Misra A, Chowbey P, Makkar BM, Vikram NK, Wasir JS, Chadha D, *et al.* Consensus statement for diagnosis of obesity, abdominal obesity and the metabolic syndrome for Asian Indians and recommendations for physical activity, medical and surgical management. J Assoc Physicians India 2009;57:163-70.
- 15. Tripathy JP, Thakur JS, Jeet G, Jain S. Prevalence and determinants of comorbid diabetes and hypertension: Evidence from non communicable disease risk factor STEPS survey, India. Diabetes Metab Syndr 2017;11(Suppl 1):S459-65.
- 16. India State-Level Disease Burden Initiative Diabetes Collaborators. The increasing burden of diabetes and variations among the states of India: The Global Burden of Disease Study 1990-2016. Lancet Glob Health 2018;6:e1352-62.
- 17. Patel M, Patel IM, Patel YM, Rathi SK. A hospital-based

observational study of type 2 diabetic subjects from Gujarat, India. J Health Popul Nutr 2011;29:265-72.

- Geetha A, Gopalakrishnan S, Umadevi R. Study on the impact of family history of diabetes among type 2 diabetes mellitus patients in an urban area of Kancheepuram district, Tamil Nadu. Int J Community Med Public Health 2017;4:4151-6.
- 19. Borgharkar SS, Das SS. Real-world evidence of glycemic control among patients with type 2 diabetes mellitus in India: The TIGHT study. BMJ Open Diabetes Res Care 2019;7:e000654.
- 20. Veera RB, Alekhya K, Kavitha D, Annapurna A. Assessment of pre-diabetes in young adults of age 18-40yrs in Andhra University College of Pharmacy, Visakhapatnam. Indian J Pharm Sci 2012;5:53-6.
- 21. Asiimwe D, Mauti GO, Kiconcol R. Prevalence and risk factors associated with type 2 diabetes in elderly patients aged 45-80 years at Kanungu district. J Diabetes Res 2020;1:1-5.
- 22. Bellou V, Belbasis L, Tzoulaki I, Evangelou E. Risk factors for type 2 diabetes mellitus: An exposure-wide umbrella review of meta-analyses. PLoS One 2018;13:e0194127.
- 23. Selvin E, Steffes MW, Zhu H, Matsushita K, Wagenknecht L, Pankow J, *et al.* Glycated hemoglobin, diabetes, and cardiovascular risk in nondiabetic adults. N Engl J Med 2010;362:800–11.