RESEARCH

BMC Public Health



Risk of diabetes and expected years in life without diabetes among adults from an urban community in India: findings from a retrospective cohort

Palak Sharma^{1*}, T.R. Dilip¹, Anjali Kulkarni², Udaya Shankar Mishra³ and Yogesh Shejul²

Abstract

Background Diabetes prevalence has increased over the past few decades, and the shift of the burden of diabetes from the older population to the younger population has increased the exposure of longer durations in a morbid state. The study aimed at ascertaining the likelihood of progression to diabetes and to estimate the onset of diabetes within the urban community of Mumbai.

Methods This study utilized an observational retrospective non-diabetic cohort comprising 1629 individuals enrolled in a health security scheme. Ten years of data were extracted from electronic medical records, and the life table approach was employed to assess the probability of advancing to diabetes and estimate the expected number of years lived without a diabetes diagnosis.

Results The study revealed a 42% overall probability of diabetes progression, with age and gender variations. Males (44%) show higher probabilities than females (40%) of developing diabetes. Diabetes likelihood rises with age, peaking in males aged 55–59 and females aged 65–69. Males aged 30–34 exhibit a faster progression (10.6 years to diagnosis) compared to females (12.3 years).

Conclusion The study's outcomes have significant implications for the importance of early diabetes detection. Progression patterns suggest that younger cohorts exhibit a comparatively slower rate of progression compared to older cohorts.

Highlights

- **High Diabetes Progression Rates**: A striking finding is that approximately 2 out of every 5 individuals aged 30 years and older advanced to diabetes within a mere decade.
- Another intriguing observation is the gender disparity in diabetes progression. Males, in particular, exhibited a notably higher likelihood of transitioning to diabetes during this timeframe.
- When we delve into age-specific data, it was found that among women aged 30–34 years, there's an average of 12.3 years expected to pass before a diabetes diagnosis is made. However, for their male counterparts,

*Correspondence: Palak Sharma pal3193@gmail.com

Full list of author information is available at the end of the article



© The Author(s) 2024. **Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicate of the original autory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/. The Creative Commons Dedication waiver (http://creativecommons.org/publicdomain/zero/1.0/) applies to the data made available in this article, unless otherwise stated in a credit line to the data.

this waiting time shrinks significantly to just 10.6 years. This discrepancy indicates a faster progression toward diabetes among males.

Keywords Cohort study, Type 2 diabetes, Disease progression, Electronic medical records (EMR), Life table

Background

The non-communicable diseases (NCD) are the leading cause of premature morbidity and mortality. Globally, there are around 537 million adults aged 20–79 years currently living with diabetes and India alone accommodates around 77 million diabetic people [1]. The prevalence of diabetes has increased many fold in recent decades worldwide, yet among major NCDs, it has usually been underestimated in terms of the cause of death. Diabetes is considered to have little impact on mortality directly when compared to morbidity as it mainly expedites the risk of microvascular complications [2] and various other severe chronic conditions [3]. Only a minority of patients uniquely die due to diabetes [3, 4].

The evolving demographics, epidemiology, and longer lifespans in recent decades have raised concerns about the quality of life for individuals with diabetes. Hence, understanding disease progression and a life-course perspective is vital for designing studies that evaluate intervention impacts in terms of preventing and delaying diabetes [5, 6].

Observing diabetes progression across different age groups and genders reveals the significant impact of demographic variables on both the development and duration of morbidity. It is known that blood glucose concentrations tend to rise with age [7], and previous national surveys reported that 54% people of those who develop diabetes were in the most productive years of their lives (<50 years) with a higher risk of developing chronic complications of diabetes [8, 9]. Most studies, both in India and internationally, indicate a higher prevalence and risk of diabetes among men [10–19], while a few studies reported otherwise [4, 20]. The shift in diabetes onset among the young and increased life expectancy for those with diabetes has prolonged the duration of time spent with diabetes and related sufferings [21, 22].

Determining the onset of diseases, particularly for asymptomatic conditions like diabetes with delayed symptom manifestation, poses a significant challenge. Hence, some studies diverge by reporting the age at diabetes detection rather than onset, introducing variations [22]. Also, numerous studies have employed the life table approach to explore the transition from non-diabetic states to diabetes, assessing the proportion of remaining life with and without diabetes [21, 23, 24]. However, these studies couldn't estimate diabetes onset and progression patterns in real-time cohorts with continuous follow-up and clinical diagnosis using laboratory tests, as many longitudinal studies in India lack consistent follow-up and were primarily conducted cross-sectionally with an interval of eight to ten years [12, 25-27]. With current technological advancements, early detection and screening for diabetes is crucial for preventing or delaying its onset [28]. Hence, this study's main objective is to assess the probability of diabetes progression in a diabetes-free cohort and estimate the remaining diabetes-free life years in the population.

Methods

This retrospective cohort study examines diabetes progression among beneficiaries of a Contributory Health Service Scheme (CHSS), using Electronic Medical Records (EMRs) from a Government hospital in urban Mumbai. The beneficiary population features uniform and universal access to healthcare under CHSS, with moderate to high educational attainment and economic stability.

In 2010-12, about 30,463 CHSS beneficiaries were registered with the hospital, and clinical tests revealed thatof these, 835 beneficiaries aged 30 years and above were newly diagnosed with diabetes. Hence, a sample double the size of these cases i.e. 1669 beneficiaries, matched for age and sex, without diabetes diagnosis during this period, was followed up for next ten years (January 2012 to December 2021) to investigate diabetes progression and its patterns. Detailed population characteristics are available elsewhere [29]. Further, 40 beneficiaries were lost to follow-up in the first year (2012) and were excluded from the analysis due to the unavailability of data on the last visited date. Hence, the analysis in this study is restricted to a total of 1629 beneficiaries.

The main clinical endpoint was the transition from non-diabetic status to clinically diagnosed diabetes. The study adopted the American Diabetes Association 2021 definition to categorize individuals with diabetes [30]. An individual is considered to have diabetes if the fasting plasma glucose (FPG) \geq 126 mg/dl (7.0 mmol/l), or postprandial glucose \geq 200 mg/dl (11.1 mmol/l), or HbA1c \geq 6.5% or that individual is on anti-diabetes medication. Table 1 presents the parameters and the cut off values used for the diagnosis of diabetes in a tabular format (Table 1).

A single decrement life table was constructed to analyze the progression of diabetes within the cohort. In this table, individuals are considered as having exited the study when they are diagnosed with diabetes or when they are lost to follow-up. Loss to follow-up, although an

Table 1 Definition of diabetes and pre-diabetes as per the American Diabetes Association, 2021 [30]

Category	Fasting plasma glucose (FPG)	Postprandial glucose (PPG)	HbA1c
Normoglycemia	FPG < 100 mg/ dL	PPPG < 140 mg/ dL	HbA1c<5.7%
Pre-diabetes	100 < = FPG < 126 mg/ dL	140<=PPPG<200 mg/ dL	5.7<=HbA1c<6.5%
Diabetes	FPG > = 126 mg/ dL	PPPG>=200 mg/ dL	HbA1c>=6.5%

Table 2 Characteristics of sampled beneficiaries that were selected as the synthetic cohort with normoglycemia for the year 2011-2012 (N=1629)

	Overall		Male		Female	
Age group	Frequency (N)	Percentage (%)	Frequency (N)	Percentage (%)	Frequency (N)	Percentage (%)
30–34	50	3.1	8	1.1	42	4.6
35–39	111	6.8	32	4.4	79	8.7
40–44	198	12.2	50	6.9	148	16.3
45-49	213	13.1	85	11.8	128	14.1
50-54	250	15.3	113	15.7	137	15.1
55–59	221	13.6	111	15.4	110	12.1
60–64	209	12.8	115	16.0	94	10.3
65–69	142	8.7	66	9.2	76	8.4
70+	235	14.4	140	19.4	95	10.5
Grand Total	1629	100	720	100	909	100
Mean age (years)	55.2		58.2		52.8	
% total sampled				44.2		55.8

event that leads to exit from the cohort, is not considered a separate decrement in this particular life table.

The probability of an individual being diagnosed with diabetes is given by the formula:

$$Probability of turning diabetic in ten years (nqx) = \frac{ndx}{lx}$$

Where ndx is the number of beneficiaries in the age group x to x+n who turned diabetic in ten years and lx is the number of person years contributed over the period by all individuals at risk in age group x to x+n.

Further, using the age and sex-specific probabilities of progressing to diabetes determined over the ten-year follow-up period, the study estimated the average number of years an individual is expected to live before being diagnosed with diabetes based on the life table columns and formulas. The mathematical formulas used to estimate each column of the life table have been published elsewhere [31]. This analysis provides insights into the expected age of onset of diabetes in the synthetic cohort of beneficiaries with normoglycemia across various age groups for both males and females.

Results

Sample characteristics

Table 2 presents the age-sex distribution of the selected synthetic cohort of beneficiaries who were followed up for a period of ten years. Of the total sample of 1629, 56% were female, and the mean age of the beneficiaries included in the study was 55.2 years. Additionally, more

than one-third of the sample (36%) was aged 60 years and above, implying the population under consideration is slightly an aged population. Notably, the proportion of males aged 60+years was higher (45%) than that of their female counterparts (29.2%).

Ten-year aggregate probability of getting diabetes

Table 3 shows the total number of individuals at the beginning of the year 2012 for each of the age groups, the number and proportion of individuals who turned diabetic at the end of ten years of follow-up with the probability of getting diagnosed with diabetes. As expected with an increase in age, the probability of getting diabetes increases over the period of ten years. There is around an 18% chance that an individual aged 30-34 years will progress to diabetes by the time they turn 40-44 years. The chances of an individual aged 55–59 years progressing to diabetes is as high as 59% (Table 3). The probability of getting diabetes is found to be higher among younger males aged 30-44 years as compared to females of the same age. However, the chi-square value from the log rank test indicated that there is no statistical difference in the probability of getting diabetes among males and females across age groups at any given point in time (Table 3).

Annual progression over the period of ten years

The probability of progressing to diabetes at the end of each year of follow-up for all age groups is presented in Table 4. This analysis reveals a progression pattern to diabetes over the years where 42% of the non-diabetic

Age Number of group beneficiaries		Number of benefi- ciaries diagnosed	Censored	Proportion diagnosed with	Overall Probability of getting diagnosed	Probability of getting diagnosed with diabetes (95% Cls)				
	at baseline'	with Diabetes		diabetes in ten years	with diabetes (95% Cis)	Males	Females			
30-34	50	5	45	10.0	0.18 (0.08,0.38)	0.22 (0.03,0.83)	0.17 (0.07,0.40)			
35-39	111	24	87	21.6	0.36 (0.25,0.48)	0.40 (0.22,0.64)	0.34 (0.22,0.49)			
40-44	198	59	139	29.8	0.46 (0.38,0.55)	0.57 (0.42,0.74)	0.42 (0.32,0.52)			
45–49	213	75	138	35.2	0.52 (0.44,0.6)	0.50 (0.37,0.63)	0.54 (0.44,0.64)			
50-54	250	93	157	37.2	0.54 (0.47,0.62)	0.53 (0.43,0.65)	0.55 (0.45,0.65)			
55-59	221	93	128	42.1	0.59 (0.52,0.67)	0.60 (0.50,0.71)	0.58 (0.47,0.69)			
60–64	209	70	139	33.5	0.50 (0.42,0.59)	0.47 (0.36,0.59)	0.54 (0.43,0.67)			
65–69	142	64	78	45.1	0.62 (0.53,0.71)	0.57 (0.43,0.71)	0.67 (0.54,0.78)			
70+	235	69	166	29.4	0.45 (0.38,0.54)	0.50 (0.41,0.61)	0.38 (0.27,0.51)			

Table 3 Overall proportion of beneficiaries diagnosed with diabetes and the probability of getting diagnosed with diabetes (with 95% Cls) over ten years of follow-up for each age group and among males and females (overall N = 1629)

Table 4 Probability of progressing to diabetes at the end of each of the consecutive year over ten years of follow-up among nondiabetic individuals belonging to various age-group (2012–2021)

Age group	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021
30-34	≈0.0	≈0.0	≈0.0	≈0.0	0.02	0.02	0.07	0.07	0.07	0.16
35–39	≈0.0	0.05	0.07	0.10	0.10	0.13	0.18	0.20	0.22	0.24
40-44	≈0.0	0.03	0.06	0.09	0.14	0.19	0.24	0.25	0.30	0.36
45–49	≈0.0	0.05	0.12	0.15	0.19	0.23	0.30	0.32	0.34	0.42
50-54	≈0.0	0.06	0.10	0.16	0.22	0.26	0.32	0.35	0.38	0.44
55-59	≈0.0	0.10	0.14	0.22	0.26	0.31	0.35	0.39	0.42	0.52
60–64	≈0.0	0.05	0.11	0.16	0.21	0.25	0.29	0.33	0.36	0.41
65–69	≈0.0	0.07	0.13	0.22	0.28	0.36	0.43	0.46	0.49	0.53
70+	≈0.0	0.07	0.11	0.18	0.24	0.28	0.30	0.33	0.36	0.43
Overall	≈0.0	0.06	0.10	0.16	0.20	0.25	0.29	0.32	0.35	0.42

subjects at the onset of the study move into the diabetic state in a span of ten years. This progression pattern indicates that the shift into the diabetic state increases with age. For instance, the younger group aged 30-34 have a relatively slower progression to diabetes (16%) than those who were aged 55 at the start of the study (52%). Subjects aged between 30 and 34 years experienced their first encounter with diabetes after four years (2016) with a very low probability of 0.02, whereas older age groups developed diabetes within one year of the follow-up. Those aged 55-59 years had the highest probability of progression, i.e., a 10% chance of progression to diabetes within a year. The most revealing of this is perhaps the probability of entering the diabetic state at the end of the study period over time ages remains cumulatively greater till the age of 60 following which the systematic pattern gets violated as a major share of the older cohorts become diabetic in the middle of the study period. It was observed that if an individual has not been diagnosed with diabetes at ages 50-59 years, the probability of having diabetes at age 60-64 is almost equal to that for subjects aged 45-49 years.

The annual progression pattern among males and females is presented in Table 5. Overall, 44% of males and 40% of the females progressed to diabetes in ten

years. The results show that the probability of a male aged 35–39 years of developing diabetes at the end of five years (in 2016) is 16% whereas among females of the same age it is only 8%. The fastest progression over ten years was observed among males aged 55–59 years (54%) and females 65–69 years (60%). The same analysis was repeated for a few selected ages and it was found that individuals aged 30 years have nearly zero probability of developing diabetes in the next ten years, while the overall probability ranged from 15% at age 35 years to 61% at age 65 years (Fig. 1).

A life table for a synthetic cohort of the non-diabetic population

Using the ten-year probability of progressing to diabetes (from Table 3), an abridged life table was constructed to estimate the expected number of years remaining without being diagnosed with diabetes for individuals free from diabetes before that age. It was found that an individual aged 30–34 years has 11.9 years without being diagnosed with diabetes, implying that the probability of having diabetes is almost negligible till he/she turns 40–44 years. The estimated age at onset is therefore around 42 years in this urban community. An individual aged 50–54 years is expected to have around 6.3 years before progressing to

Fig. 1 Probability of progression to diabetes over the ten years of follow-up among non-diabetic individuals at a few selected terminal ages (N=270)

2016

2017

50 _____55

2018

2019

60

2020

65

0.70

0.60

0.40

0.30

0.20

0.10

0.00 2012

2013

30

2014

35

2015

40 -45

diabetes, whereas an individual aged 65–69 years is only expected to live for 3.5 years more without being diagnosed with diabetes (Table 6).

Females aged 30–34 years have around 12.3 years more without diabetes diagnosis whereas males of the same age are expected to live around 10.6 more years before progressing to diabetes. These results reveal that males have a shorter waiting time and faster progression to diabetes in comparison to females. The number of years before progressing to diabetes is higher among females up to the age of 45 years only, and later, the pattern changes though the difference in the number of years before being diagnosed with diabetes among males and females isn't statistically significant (Table 7).

Discussion

0.15

2021

Diabetes is a chronic disease that significantly increases the risk of developing other potentially life-threatening conditions. Due to shifts in urbanization and lifestyle factors, there's been a noted change in the onset of diabetes, with the condition now affecting a younger demographic [22, 32]. In light of the limited available evidence regarding the progression and risk of diabetes onset, this study aimed to estimate the likelihood of transitioning to diabetes within a ten-year period among a non-diabetic synthetic cohort. Additionally, the research sought to examine variations in this progression among different age groups and between males and females, employing a life table approach for analysis. The study revealed that, over a ten-year period, 42% of the beneficiaries in the study population were diagnosed with diabetes. Females had a lower likelihood of developing diabetes compared to males. In a recent publication authored by Sharma and colleagues, an investigation within the same study population revealed a lifetime diabetes risk estimated at 40% [19]. This observation underscores the considerable susceptibility of the community to diabetes, indicating a high rate of diabetes incidence within this demographic. Additionally, a parallel study on the lifetime risk of diabetes conducted by Luhar et al. demonstrated a decline in

Table 5 Probability of progressing to diabetes fat the end of each of the consecutive year over ten years of follow-up among males and females

	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021
Males		2015	2011	2015	2010	2017	2010	2015		2021
	0.0	0.0	0.0	0.0	0.0	0.0	0.15	0.15	0.15	0.15
30-34	≈0.0	≈0.0	≈0.0	≈0.0	≈0.0	≈0.0	0.15	0.15	0.15	0.15
35–39	≈0.0	0.09	0.13	0.16	0.16	0.22	0.22	0.25	0.25	0.25
40-44	≈0.0	0.06	0.12	0.16	0.25	0.29	0.34	0.38	0.38	0.46
45–49	≈0.0	0.02	0.07	0.12	0.20	0.24	0.28	0.29	0.30	0.42
50-54	≈0.0	0.06	0.11	0.16	0.22	0.27	0.32	0.36	0.37	0.42
55-59	≈0.0	0.06	0.10	0.16	0.23	0.28	0.32	0.37	0.41	0.54
60-64	≈0.0	0.04	0.08	0.15	0.21	0.22	0.24	0.30	0.32	0.37
65–69	≈0.0	0.05	0.10	0.16	0.24	0.33	0.36	0.40	0.42	0.45
70+	≈0.0	0.06	0.11	0.18	0.24	0.29	0.31	0.37	0.42	0.49
Overall	≈0.0	0.05	0.10	0.16	0.22	0.27	0.30	0.34	0.36	0.44
Females										
Age group	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021
30-34	≈0.0	≈0.0	≈0.0	≈0.0	0.03	0.03	0.05	0.05	0.05	0.16
35-39	≈0.0	0.03	0.05	0.08	0.08	0.09	0.16	0.17	0.20	0.23
40-44	≈0.0	0.02	0.04	0.06	0.10	0.16	0.20	0.21	0.27	0.32
45–49	≈0.0	0.06	0.15	0.17	0.18	0.22	0.31	0.33	0.36	0.42
50-54	≈0.0	0.06	0.09	0.17	0.21	0.26	0.32	0.34	0.38	0.47
55-59	≈0.0	0.14	0.18	0.29	0.29	0.34	0.38	0.41	0.42	0.50
60-64	≈0.0	0.14	0.18	0.29	0.29	0.34	0.38	0.41	0.42	0.50
65–69	≈0.0	0.10	0.16	0.27	0.32	0.38	0.49	0.51	0.57	0.60
70+	≈0.0	0.07	0.11	0.19	0.25	0.26	0.28	0.28	0.28	0.32
Overall	~00	0.06	0.11	0.16	0.10	0.23	0.20	0.31	0.34	0.40

Table 6	Expected number of	vears lived before n	progressing to a	diabetes by	non-diabetic individuals at various ages

Age group as of Ten-year Probabilit year 2012 of of remaining free from diabetes		Probability of getting diagnosed with Diabetes in next 10 years	Number of beneficia- ries surviving at the start of year	Person years		Expected num- ber of remaining years without Diabetes
	рх	qx	lx	Lx	Тх	ex
30–34	0.82	0.18	1000	4545.5	11854.3	11.9
35–39	0.64	0.36	818	3363.6	7308.8	8.9
40-44	0.54	0.46	527	2031.1	3945.2	7.5
45-49	0.48	0.52	285	1054.6	1914.1	6.7
50–54	0.46	0.54	137	498.0	859.5	6.3
55–59	0.41	0.59	63	220.1	361.5	5.8
60–64	0.50	0.50	25	95.5	141.4	5.5
65–69	0.38	0.62	13	43.8	45.9	3.6
70+	0.55	0.45	5	2.1	2.1	0.4

Note nqx: probability of progressing to diabetes between age x to x+n

Ix: number of individuals surviving without diabetes at the beginning of age x

nLx: person years lived without diabetes by lx individuals in the age group x to x+n

Tx: person years lived without diabetes by the individuals after exact age x

ex: Average number of expected years a person aged x is expected to live free of diabetes/before progressing to diabetes

	_				/			/	•												•			
		montor	1 0 1 1 00	boro	+ \ / O O KC	Lin road	60	toro prov	aroccip		abotoc	611	000	d10	bot	$\sim \sim \sim$	<u>~</u>	tom2	200	t	V 10 KIC	2110 2	200	
14010			1 1 1 1 1 1 1 1 1		I VEAIS	111/10/1	110		THESTIN	1 1 () (1)		1 11/	1 1 (1) 1 - (1)	11.4		11111	11 1	PULL		- AL	VALIC		יםיוא	κ.
IUDIC		1			1 V((11.)		1.7.					1.7.V					 11.1	1. 1110	111			111.16		
									-,								 							-

	Males				Female								
Age group	qx	lx	Lx	Tx	ex	qx	lx	Lx	Tx	ex			
30–34	0.22	1000	4444.5	10618.9	10.6	0.17	1000	4565.3	12253.5	12.3			
35–39	0.40	778	3111.2	6174.4	7.9	0.34	826	3434.9	7688.2	9.3			
40-44	0.57	467	1666.7	3063.2	6.6	0.42	548	2168.1	4253.3	7.8			
45–49	0.50	200	752.3	1396.5	7.0	0.54	319	1168.0	2085.3	6.5			
50-54	0.53	101	370.1	644.2	6.4	0.55	148	535.8	917.3	6.2			
55-59	0.60	47	164.6	274.1	5.8	0.58	66	235.9	381.5	5.7			
60–64	0.47	19	71.6	109.4	5.9	0.54	28	101.6	145.6	5.2			
65–69	0.57	10	35.7	37.8	3.8	0.67	13	42.5	44.0	3.5			
70+	0.50	4	2.1	2.1	0.5	0.38	4	1.5	1.5	0.4			

Note nqx: probability of progressing to diabetes between age x to x+n

lx: number of individuals surviving without diabetes at the beginning of age x

nLx: person years lived without diabetes by lx individuals in the age group x to x+n

Tx: person years lived without diabetes by the individuals after exact age x

ex: Average number of years a person aged x is expected to live free of diabetes/before progressing to diabetes

the remaining lifetime risk (95% confidence interval) with age, reaching 37.7% at the age of 60 for women and 27.5% for men [33]. In a previous investigation involving Asian Indians, it was discovered that over a ten-year period, the incidence of individuals progressing from impaired glucose tolerance (IGT) to diabetes varied between 13% and 52% [9]. Another study conducted in India, employing the life table methodology, revealed that the likelihood of maintaining a diabetes-free life diminishes with age. Specifically, only 30% of the overall participants remained free of diabetes beyond the age of 77 [23].

The current study observed that males had a slightly higher likelihood of developing diabetes compared to females. However, it's important to note that these differences in diabetes risk between males and females were not found statistically significant across various age groups at any point during the follow-up period of ten years. Consistent with our findings, several previous studies have also noted an elevated diabetes risk among males [10–18]. An Indian study specifically highlighted that the likelihood of remaining diabetes-free is notably lower for men compared to women, particularly for urban residents and those belonging to higher socioeconomic classes [23]. This investigation utilized data from the India Human Development Survey (IHDS) and discovered that only 26% of urban males and 33% of urban females remained diabetes-free until the age of 73 in India [23]. It has been argued that the observed discrepancy in survival probabilities can possibly be attributed to the high male mortality in the non-diabetic population [34]. It has also been found that the western parts of India (including Maharashtra and Goa), in comparison to

other regions of India, had an overall higher probability of diabetes progression, which was around 71% for males and 79% for females [23].

As age increased, the likelihood of developing diabetes also rose, peaking at around 59 years of age. This agerelated pattern aligns with findings from prior research studies. Notably, the highest probability of diabetes progression was observed among older men (55-59 years) and women (65-69 years), consistent with earlier investigations [10, 13]. While some studies have reported variations in the precise age peak for diabetes incidence, most have identified the highest incidence of type II diabetes in men aged 55-64 years and women aged 65-69 years, with some indicating an overall peak occurring at 55-59 years [10, 13, 15]. For instance, a study led by Singh and colleagues identified the highest likelihood of developing diabetes between the ages of 38 and 58, with an 8% probability [23]. Another ten-year study found that the risk of diabetes increased until the age of 75, after which it declined. Importantly, this study underscored that the rate of progression from pre-diabetes to diabetes among older adults was one-third of that seen in middle-aged individuals. This suggests that, at later stages of life, the probability of transitioning to diabetes is lower when compared to the higher risk observed in middle-aged individuals [35].

Diabetes is typically a condition that doesn't show noticeable symptoms, and tends to develop slowly, sometimes taking several years to manifest, making it easy to overlook. Consequently, this study utilized a ten-year age and sex stratified approach to estimate the probability of progressing to diabetes. Also, it then constructed a synthetic cohort to estimate the additional years a person can expect to live before a diabetes diagnosis. The study found that the population aged 30-34 years is expected to live around 11.9 years before progressing to diabetes, yielding the age at onset to be around 42 years. The onset of diabetes in our study was observed to occur at an age that is relatively early compared to some other Indian studies. For instance, a ten-year prospective study conducted in Kerala, India, found that the average age at which individuals developed diabetes was 47.3 years [12]. The Chennai Urban-Rural Epidemiological study (CURES study) reported the mean age at diagnosis of diabetes among the incident diabetes cases to be 50.9 ± 12.8 years [26]. Studies have also reported that the onset of diabetes in urban India is about a decade earlier than among their western counterparts [36]. The study found a faster progression to diabetes among males, which is in line with previous studies. It has been reported that males are at a higher risk of developing diabetes at an earlier age than females, i.e. age at onset of newly diagnosed diabetes is earlier among males [13, 15].

The observed variations in the estimates can be predominantly attributed to disparities in the age and sex demographics of the current study population compared to those in previous studies as a majority of the mentioned studies have focused on populations aged 20 years and above. Also, research has demonstrated that the principal factors associated with the progression to diabetes in India are urbanization and elevated economic status [23], a trend that is also apparent in the distribution pattern of diabetes-free survival within our study. Furthermore, these elevated estimates in our study may also be attributed to distinctive factors, including an urban-dwelling population, moderate to high levels of educational attainment, economic stability, and the presence of universal and uniform access to healthcare. Moreover, the study's population benefits from a preventative and universal access medical scheme with rigorous screening and diagnostic evaluations. This unique characteristic partly explains the faster diabetes progression observed in this urban community.

Disease progression models often rely on long-term patient observations, but in India, such longitudinal studies are limited due to extended follow-up periods and high associated costs [37-39]. This study stands out in that it utilizes data from an Electronic Medical Record system within a hospital, leveraging clinically confirmed incident diabetes cases to estimate diabetes progression over a decade of follow-up. The study's findings have significant implications for early diabetes detection and the optimal age to commence diabetes screening in the population. While the American Diabetes Association (ADA) currently recommends regular screening for individuals aged 45 and above, evidence from this urban community suggests a need to reassess the starting age for diabetes screening in India. The study corroborates the need to rigorously implement the national guideline to screen all adults aged 30 years and above for diabetes. The screening capacity of health care systems needs to be strengthened exponentially to reach out to these agegroups in the populations [40]. This approach can aid in identifying high-risk or pre-diabetic individuals and may contribute to delaying diabetes onset through prescribed interventions. These data and its outcomes hold substantial importance for implementing preventive strategies against diabetes among a large patient population, with far-reaching economic, social, and psychological implications. Furthermore, such real-time data-driven decisions can serve as compelling case studies in the Indian context, illustrating potential strategies for early detection and diabetes prevention based on estimated onset age and observed progression patterns.

Limitations of the study

The study has a few limitations that warrant acknowledgment. Firstly, it's important to recognize that the study population consisted of individuals who were beneficiaries of the Contributory Health Service Scheme and received healthcare services under a uniform system. Hence, it's essential to exercise caution when attempting to generalize the findings of this study to other regions of the country, as the incidence and progression of diabetes can differ across different settings. Secondly, when comparing our results with those of previous studies, caution is advised due to variations in the age and sex distributions of the populations under study, as well as differences in the definitions used for diabetes. Lastly, the study was unable to account for other factors that can influence the development and progression of diabetes, such as diet, obesity, physical inactivity, and the presence of various comorbid conditions. Consequently, establishing a causal relationship between these variables and diabetes was not feasible within the scope of this study.

Abbreviations

	American Diabetes Association
BMI	Body mass index
CHSS	Contributory Health Service Scheme
CURES	Chennai urban-rural epidemiological study
EMR	Electronic medical records
FPG	Fasting plasma glucose
HbA1c	Glycated hemoglobin
IFG	Impaired fasting glucose
IGT	Impaired glucose tolerance
IHDS	India Human Development survey
NCDs	Non-communicable diseases
PPG	Postprandial glucose

Acknowledgements

This research is part of the CHIPS cohort study jointly undertaken by the Bhabha Atomic Research Centre (BARC), Mumbai, and International Institute for Population Sciences (IIPS), Mumbai. The authors are thankful to the Head of the Medical Division, Bhabha Atomic Research Centre, Mumbai, and the BARC research ethics committee, for all their support for this research study.

Author contributions

All authors have contributed significantly. PS, TRD and USM conceptualized the idea. PS did all the literature review and analysis, and wrote the first draft of the manuscript. PS, TRD, USM, AK and YS reviewed and revised the manuscript to its present form. The manuscript represents honest work. It is original and has not been submitted to any other journal. The manuscript has been read and approved by all the authors.

Funding

This study received no funding from any of the government, nongovernmental organizations or any other institution.

Data availability

The datasets generated and/or analysed during the current study are not publicly available due to data privacy norms of the hospital under consideration. Data are however available from the corresponding author upon reasonable request to pal3193@gmail.com and with permission of Medical Division, Bhabha Atomic Research Center, Mumbai, after due approvals.

Declarations

Ethics approval and consent to participate

The need for consent to participate was waived by an Institutional Review Board (IRB) and was deemed unnecessary according to national regulations by the "**Bhaba Atomic Research Center's Hospital Medical Ethics Committee**" on 7th July, 2021 in the 47th Medical Ethics Committee Meeting. All methods carried out in the study were in accordance with relevant guidelines and regulations. The study used the masked ID of the patients and the authors had no direct contact with the patients.

Consent for publication

Not Applicable.

Competing interests

The authors declare no competing interests.

Author details

¹Department of Family and Generations, International Institute for Population Sciences, Mumbai 400088, India ²Medical Division, Bhabha Atomic Research Center, Mumbai 400088, India ³Department of Bio-statistics and Epidemiology, International Institute for Population Sciences, Mumbai 400088, India

Received: 15 January 2024 / Accepted: 27 March 2024 Published online: 15 April 2024

References

- 1. International Diabetes Federation. IDF Diabetes Atlas 10th edition [Internet]. 2021 [cited 2022 Dec 27]. Available from: www.diabetesatlas.org.
- Pelluri R, Srikanth K, Chimakurthy J, Nagasubramanian VR. Microvascular complications and their Associated risk factors among rural type 2 Diabetic Population: a cross-sectional study. SN Compr Clin Med. 2021;3(2):625–31.
- Sikdar KC, Wang PP, MacDonald D, Gadag VG. Diabetes and its impact on health-related quality of life: a life table analysis. Qual Life Res. 2010;19(6):781–7.
- van Dieren S, Beulens JWJ, van der Schouw YT, Grobbee DE, Neal B. The global burden of diabetes and its complications: an emerging pandemic. 17, Eur J Cardiovasc Prev Rehabilitation. 2010.
- 5. World Health Organization. Global report on diabetes. 2016.
- Jewell NP. Natural history of diseases: statistical designs and issues. Clin Pharmacol Ther. 2016;353–61.
- Alanazi NH, Alsharif MM, Rasool G, Bin A, Alruwaili H, Matrouk A et al. Prevalence of Diabetes and its relation with age and sex in Turaif city, northern Saudi Arabia in 2016–2017. Electron Physician [Internet]. 2017 [cited 2022 Dec 15];9(9):5294–7. http://www.ephysician.irhttphttps://doi. org/10.19082/5294.
- Ramachandran A, Dharmaraj D, Vlswanathan M. Prevalence of Glucose Intolerance in Asian Indians Urban-rural difference and significance of upper body adiposity. Diabetes Care [Internet]. 1992;15(10). http://diabetesjournals. org/care/article-pdf/15/10/1348/441362/15-10-1348.pdf.
- Ramaiya K, Kodali V, Alberti K. Epidemiology of diabetes in asians of the Indian Sub-continent* [Internet]. 11. 1991. Available from: www.pdf4u.com.
- Huebschmann AG, Huxley RR, Kohrt WM, Zeitler P, Regensteiner JG, Reusch JEB. Sex differences in the burden of type 2 diabetes and cardiovascular risk across the life course. Volume 62. Diabetologia: Springer; 2019. pp. 1761–72.
- Rojo-Martínez G, Valdés S, Soriguer F, Vendrell J, Urrutia I, Pérez V et al. Incidence of diabetes mellitus in Spain as results of the nation-wide cohort di@ bet.es study. Sci Rep. 2020;10(1).
- Vijayakumar G, Manghat S, Vijayakumar R, Simon L, Scaria LM, Vijayakumar A et al. Incidence of type 2 diabetes mellitus and prediabetes in Kerala, India: results from a 10-year prospective cohort. BMC Public Health. 2019;19(1).
- 13. Vega T, Gil M, Lozano J. Age and sex differences in the incidence of diabetes mellitus in a population-based Spanish cohort. J Diabetes. 2015;7(3):411–7.
- Urrutia I, Martín-Nieto A, Martínez R, Casanovas-Marsal JO, Aguayo A, del Olmo J et al. Incidence of diabetes mellitus and associated risk factors in the adult population of the basque country, Spain. Sci Rep. 2021;11(1).
- Khan MAB, Hashim MJ, King JK, Govender RD, Mustafa H, Kaabi JA. Epidemiology of type 2 diabetes - global burden of disease and forecasted trends. J Epidemiol Glob Health. 2020;10(1):107–11.

- International Institute for Population Sciences (IIPS), MoHFW NPHCE, Harvard TH. Chan School of Public Health (HSPH), The University of Southern California (USC). Longitudinal Ageing Study in India (LASI). 2020.
- Kutty Vr, Dilip T, Archana A, Gopinathan S, Ramanathan M. Shifting pattern of diabetes among the elderly in India: evidence from the national sample survey organization's data, 2004–2014. Int J Noncommun Dis. 2018;3(2):67.
- Sharma P, Dilip TR, Mishra US, Kulkarni A. The lifetime risk of developing type II diabetes in an urban community in Mumbai: findings from a ten-year retrospective cohort study. BMC Public Health. 2023;23(1).
- Arnetz L, Ekberg NR, Alvarsson M. Sex differences in type 2 diabetes: focus on disease course and outcomes. Diabetes, metabolic syndrome and obesity: targets and therapy. Volume 7. Dove Medical Press Ltd.; 2014. pp. 409–20.
- Bardenheier BH, Lin J, Zhuo X, Ali MK, Thompson TJ, Cheng YJ, et al. Disabilityfree life-years lost among adults aged ≥ 50 years with and without diabetes. Diabetes Care. 2016;39(7):1222–9.
- 22. Mohanty SK, Rodgers J, Singh RR, Mishra RS, Kim R, Khan J, et al. Morbidity compression or expansion? A temporal analysis of the age at onset of non-communicable diseases in India. Geroscience. 2021;43(1):409–22.
- 23. Singh SK, Pedgaonkar S, Puri P, Gupta J. Diabetes-free life in India: application of survival function analysis. J Soc Health Diabetes [Internet]. 2018;6:48–55. Available from:www.joshd.net.
- 24. Tönnies T, Baumert J, Heidemann C, von der Lippe E, Brinks R, Hoyer A. Diabetes free life expectancy and years of life lost associated with type 2 diabetes: projected trends in Germany between 2015 and 2040. Popul Health Metr. 2021;19(1).
- Mohan V, Deepa M, Anjana RM, Lanthorn H, Deepa R. Incidence of Diabetes and Pre-diabetes in a Selected Urban South Indian Population (CUPS-19) [Internet]. 2007. www.japi.org.
- Anjana RM, Rani CSS, Deepa M, Pradeepa R, Sudha V, Nair HD, 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(8):1441–8.
- Khan R, Singh S, Surya J, Sharma T, Kulothunga V, Raman R. Age of onset of diabetes and its comparison with prevalence and risk factors for diabetic retinopathy in a rural population of India. Ophthalmic Res. 2019;61(4):236–41.
- Edlitz Y, Segal E. Prediction of type 2 diabetes mellitus onset using logistic regression-based scorecards. Elife. 2022;11.
- Kulkarni A, Dilip T, Shejul YK, Bhandarkar P, Goswami P, Sharma P et al. Progression and management of diabetes in Indian settings with universal access to health care: Protocol and plans for CHIPS cohort study. medRxiv

[Internet]. 2023 Jan 28 [cited 2023 Jan 30];2023.01.24.23284975. https://www. medrxiv.org/content/https://doi.org/10.1101/2023.01.24.23284975v1.

- American Diabetes Association. Classification and diagnosis of diabetes: standards of medical care in diabetes-2021. Diabetes Care. 2021;44:S15–33.
- Islam A, Mondal MNI, Islam MR, Islam MR, Islam MS, Khan MN, et al. Techniques of life table construction: a review. Int J Humanit Educ Res. 2019;1(2):10–4.
- Nagarathna R, Bali P, Anand A, Srivastava V, Patil S, Sharma G et al. Prevalence of diabetes and its determinants in the young adults Indian Population-Call for yoga intervention. Front Endocrinol (Lausanne). 2020;11.
- Luhar S, Kondal D, Jones R, Anjana RM, Patel SA, Kinra S et al. Lifetime risk of diabetes in metropolitan cities in India Low-to middle-income country SAGE Study on global AGEing and adult health SRS Sample Registration System. Diabetologia [Internet]. 2021;521–9. https://doi.org/10.1007/ s00125-020-05330-1.
- Tachkov K, Mitov K, Koleva Y, Mitkova Z, Kamusheva M, Dimitrova M et al. Life expectancy and survival analysis of patients with diabetes compared to the non diabetic population in Bulgaria. PLoS ONE. 2020;15(5).
- 35. van Herpt TTW, Ligthart S, Leening MJG, Van Hoek M, Lieverse AG, Ikram MA et al. Lifetime risk to progress from pre-diabetes to type 2 diabetes among women and men: comparison between American Diabetes Association and World Health Organization diagnostic criteria. BMJ Open Diabetes Res Care. 2020;8(2).
- Mehta SR, Kashyap AS, Das S. Diabetes mellitus in India: the modern scourge. 65, Medical Journal Armed Forces India. Med J Armed Forces India; 2009. p. 50–4.
- Patel V, Chatterji S, Chisholm D, Ebrahim S, Gopalakrishna G, Mathers C, et al. Chronic diseases and injuries in India. Comment Lancet [Internet]. 2011;377:181. Available from:http://www.thelancet.com.
- Unnikrishnan R, Mohan V. Challenges in estimation of glycated hemoglobin in India. Diabetes Technol Ther. 2013;15(10):897–9.
- Bennasar-Veny M, Fresneda S, López-González A, Busquets-Cortés C, Aguiló A, Yañez AM. Lifestyle and progression to type 2 diabetes in a cohort of workers with prediabetes. Nutrients. 2020;12(5).
- National Health Mission. Module for Multi-Purpose Workers (MPW)-Female/ Male on Prevention, Screening and Control of Common Non-Communicable Diseases. 2018.

Publisher's Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.