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SYSTEMATIC REVIEW

Type 2 Diabetes Mellitus in Nepal from 2000 to 2020: A

systematic review and meta-analysis [version 1; peer review: 3

approved with reservations]

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Abstract

Aims: To evaluate the prevalence and risk factors of type 2 diabetes mellitus (T2DM) from 2000-2020 in various parts of Nepal. Methods: PubMed, Embase, Scopus, and Google Scholar were searched using the appropriate keywords. All Nepalese studies mentioning the prevalence of T2DM and/or details such as risk factors were included. Studies were screened using Covidence. Two reviewers independently selected studies based on the inclusion criteria. Meta-analysis was conducted using Comprehensive Meta-Analysis Software v.3. Results: Total 15 studies met the inclusion criteria. The prevalence of T2DM, pre-diabetes, and impaired glucose tolerance in Nepal in the last two decades was 10% (CI, 7.1%- 13.9%), 19.4% (CI, 11.2%- 31.3%), and 11.0% (CI, 4.3%- 25.4%) respectively. The prevalence of T2DM in the year 2010-15 was 7.75% (CI, 3.67-15.61), and it increased to 11.24% between 2015-2020 (CI, 7.89-15.77). There were 2.19 times higher odds of having T2DM if the body mass index was \geq 24.9 kg/m². Analysis showed normal waist circumference, normal blood pressure, and no history of T2DM in a family has 64.1%, 62.1%, and 67.3% lower odds of having T2DM, respectively. Conclusion: The prevalence of T2DM, pre-diabetes, and impaired glucose tolerance in Nepal was estimated to be 10%, 19.4%, and 11% respectively.

Keywords

Blood Pressure, Body Mass Index, Diabetes Mellitus Type 2, Nepal

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Introduction

In 2019, the International Diabetes Federation (IDF) estimated that 463 million adults worldwide had diabetes¹. The statistics showed that these individuals were in the age range of 20 to 79, have diabetes, and 79.4% were from low- and middle-income countries1. Additionally, IDF estimates that the global prevalence of diabetes will be 578.4 million by 2030, with this rising to 700.2 million by 2045 among adults aged between 20 to 79 years¹ In the region of Southeast Asia, the prevalence of diabetes was 8.8% in 2019, and this is projected to increase to 9.7% by 2030². Type 2 diabetes mellitus (T2DM) still remains a major cause of worldwide morbidity and mortality, which leads to complications such as neuropathy, nephropathy, stroke, and coronary artery disease³. In 2017, over 10, 000 individuals died due to T2DM or diabetes-related complications in Nepal, which is the 11th most common cause of disability in terms of disability-adjusted life years (DALYs) (1226 DALYs per 10,000 population)⁴. In 2020, the prevalence of T2DM in Nepal was 8.5% (95% CI 6.9-10.4%), which was higher than that of 8.4% (95% CI 6.2-10.5%) in 2014^{5,6}. Similarly, in 2020 the prevalence of pre-diabetes was 9.2% (95% CI 6.6 - 12.6%) compared to 2014, which was 10.3% (95% CI 6.1-14.4%)^{5,6}. In the advent of growing non-communicable diseases, a Multi-Sectoral Action Plan has been adopted by the government of Nepal to prevent and control non-communicable diseases including T2DM7. However, there have not been many studies that evaluate the risk factors of T2DM in Nepal, which can be helpful for the prevention and control of this disease. We conducted this review to evaluate the prevalence and risk factors of pre-diabetes and T2DM in Nepal over the past 20 years, by pooling the studies done in various parts of the country.

Methods

Protocol registration

The systematic review is registered in PROSPERO (CRD42020215247). It is documented as per the guidelines of the Meta-Analysis of Observational Studies in Epidemiology (MOOSE)^{8,9}.

Information sources and search strategy

Electronic databases such as PubMed, PubMed Central, Google Scholar, Scopus, and Embase were used to search relevant articles (Extended data file 1¹⁰). Published articles from 2000 to 2020 were searched with the use of the appropriate keywords such as "diabetes mellitus", "high blood sugar", "type 2 diabetes", "prevalence", "risk factor" and "Nepal" along with relevant Boolean operators.

Eligibility criteria

All published studies that took place in Nepal from 2000–2020 were included in this review. These studies comprised of cross-sectional studies, case series that reported on more than 50 patients, cohort study, randomized control trial (RCTs) that were based on prevalence of T2DM and/or its related issues such as risk factors, outcome, and outcome predictors.

Editorials, commentaries, viewpoint articles without adequate data on T2DM and its related issues were excluded. Furthermore,

studies that took place before 1999, outside of Nepal, as well as those that were on Type 1 and gestational diabetes were excluded.

Study selection

The studies were selected with the use of Covidence¹¹. The title and abstract were screened based on the inclusion criteria independently by two authors (SL, SN). Discrepancies were resolved by consensus obtained from the third author (AM). Further full-text review (SN, AM) was done independently, and discrepancies (SL) were resolved.

Data collection process and data extraction

Three authors (SL, AM, and SN) were independently involved in the data extraction and adding that to a standardized form in Excel. The accuracy and completion of each other's work was verified by all the reviewers. The characteristics extracted for each selected study included, first author, year, study design, sample size, study location, prevalence rate, and risk factors of T2DM such as Body Mass Index (BMI), exercise (moderate to high level of exercise (\geq 30 minutes/days) is taken as adequate), waist circumference (\geq 85 cm in females, and \geq 90 cm in males were defined as high), family history, fruit and vegetable serving per day, alcohol, smoking/tobacco, literacy, and increased blood pressure (BP) (\geq 140/90 mmHg is taken as hypertensive) (Please see *Underlying data*¹²).

Data analysis

Comprehensive Meta-Analysis Software (CMA) v.3 was used to analyze the extracted data.

Definition of the condition

T2DM was defined as a fasting blood glucose (FBG) of \geq 126 mg/dl (7.0 mmol/l) or a 2-h oral glucose tolerance test (OGTT) blood glucose level of \geq 200 mg/dl (11.1 mmol/l). Prediabetes was defined as FBG level between 100 (5.6 mmol/l) and 125 mg/dL (< 7 mmol/l) or a 2-h OGTT blood glucose level between 140 (7.8 mmol/l) and 199 mg/dl (11 mmol/l). Impaired glucose tolerance (IGT) was defined as two-hour glucose levels of 140 to 199 mg per dL (7.8 to 11.0 mmol) on the 75-g oral glucose tolerance test¹³.

Bias assessment

Bias assessment of the included studies was done by the Joanna Briggs Institute (JBI) tool (Table 1)¹⁴.

Assessment of heterogeneity

The heterogeneity in the included studies was assessed based on the Cochrane Handbook for Systematic reviews by the I² statistics (I²>50%)¹⁵. Thus, a random-effects model with the inverse variance heterogeneity model was performed. If I²>50% significant heterogeneity random effect model was preferred. If I²<50% then fixed effect model was preferred.

Sensitivity analysis

The sensitivity analysis was performed by excluding studies that did not show any significant difference in the prevalence of T2DM.

Author/year	Was the sample frame appropriate to address the target population?	Were study participants sampled in an appropriate way?	Was the sample size adequate?	Were the study subjects and the setting described in detail?	Was the data analysis conducted with sufficient coverage of the identified sample?	Were valid methods used for the identification of the condition?	Was the condition measured in a standard, reliable way for all participants?	Was there appropriate statistical analysis?	Was the response rate adequate, and if not, was the low response rate managed appropriately?
Sharma B ¹⁶ <i>et al.</i> 2019	yes	yes	No	Yes	Yes	yes	yes	yes	yes
Gyawali B ¹⁷ <i>et al.</i> 2018	yes	yes	Yes	Yes	Yes	yes	yes	yes	yes
Sharma SK ¹⁸ et al. 2011	yes	yes	Yes	Yes	Yes	yes	yes	yes	yes
Sharma SK ¹⁹ et al. 2013	yes	yes	Yes	Yes	yes	yes	yes	yes	yes
Chhetri MR ²⁰ et al. 2009	yes	yes	Yes	Yes	yes	yes	yes	yes	yes
Paudyal G ²¹ <i>et al.</i> 2008	yes	yes	Yes	Yes	yes	yes	yes	yes	yes
Bhandari GP ²² et al. 2014	yes	yes	Yes	Yes	yes	yes	yes	no	yes
Karki P ²³ <i>et al</i> . 2000	yes	yes	Yes	Yes	yes	yes	yes	no	yes
Paudel S ²⁴ et al. 2020	yes	yes	Yes	Yes	yes	yes	yes	yes	yes
Koirala S ²⁵ et al. 2018	yes	yes	yes	Yes	yes	yes	yes	yes	yes
Ranabhat K ²⁶ et al. 2016	ou	yes	ou	Yes	yes	yes	yes	yes	yes
Mehta KD ²⁷ et al. 2011	yes	yes	yes	Yes	yes	yes	yes	yes	yes
Shrestha UK ²⁸ et al. 2006	yes	yes	yes	Yes	yes	yes	yes	yes	yes
Dhimal M ²⁹ et al. 2019	yes	no	yes	Yes	yes	yes	yes	yes	yes
Kushwaha A ³⁰ <i>et al.</i> 2020	no	yes	no	Yes	yes	yes	yes	yes	yes

Table 1. JBI checklist for bias assessment.

A total of 15 studies were included in the quantitative analysis.

The random effects meta-analysis assessment of 15 studies

indicated T2DM prevalence at 10% (95% CI, 7.1%- 13.9%)

(Figure 2). Sensitivity analysis was performed with the exclusion

of individual studies which resulted in no significant differences

The assessment of T2DM prevalence between 2010-2015 with

the use of random-effects meta-analysis was 7.75% (Propor-

tion, 0.0775; 95% CI, 0.0367-0.1561; studies: 4; I²:99.62),

in the prevalence of T2DM (*Extended data* file 2, Figure 1¹⁰)

Quantitative synthesis

Prevalence of T2DM

Result

A total of 4651 studies were analyzed after thorough database search, of which 736 were identified as duplicates and removed. Title and abstracts of 3915 studies were screened and 3822 studies were excluded. The full-text eligibility of 92 studies was assessed and 77 studies were excluded for definite reasons. A total of 15 studies were included in the qualitative and quantitative analysis. The following information is depicted in the PRISMA flow diagram (Figure 1).

Qualitative summary

A qualitative summary of the individual study is presented in (Table 2).



PRISMA 2009 Flow Diagram

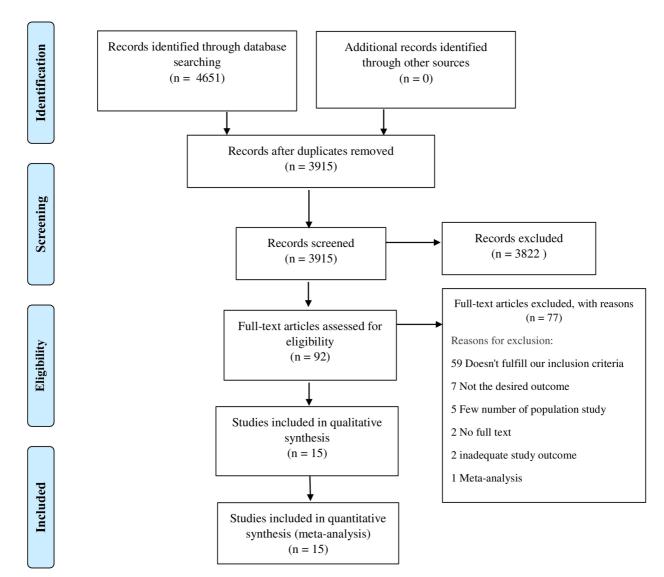


Figure 1. PRISMA flow diagram.

Author/s	Study Year	Study Design	Sample Size	Study Area	Pre-diabetes	T2DM	IGT
Dhimal M ²⁹ <i>et al.</i> 2019	2019	Cross-sectional study	12557	72 districts (all provinces)		1067/12557	
Shrestha UK ²⁸ et al. 2006	2006	2006 Cross-sectional study 1012	1012	Seven wards of metropolitan and sub-metropolitan of Nepal		192/1012	107/1012
Kushwaha A ³⁰ et al. 2020	2020	2020 Cross-sectional study 114	114	Community Hospital		5/114	
Sharma B ¹⁶ et al. 2019	2019	Cross-sectional study	320	Morang	55/320	38/320	57/320
Gyawali B ¹⁷ <i>et al.</i> 2018	2018	Cross-sectional study	2310	Lekhnath municipality	302/2310	271/2310	
Sharma SK ¹⁸ <i>et al.</i> 2011	2011	Cross-sectional study	14425	Eastern Nepal		889/14008	
Sharma SK ¹⁹ <i>et al.</i> 2013	2013	Cross-sectional study	3218	Dharan		242/3218	
Chhetri MR ²⁰ et al. 2009	2009	2009 Cross-sectional study 1633	1633	Kathmandu valley		422/1633	
Paudyal G ²¹ et al. 2008	2008	2008 Cross-sectional study 1475	1475	Mulpani ,Gothar Kathmandu valley		60/1475	34/1475
Bhandari G ²² <i>et al.</i> 2014	2014	2014 Cross-sectional study 11901	11901	31 selected hospital institutions (28 non-speciality)		391/11901	
Karki P ²³ <i>et al.</i> 2000	2000	Cross-sectional Study	1,840	Outpatient clinic of BPKIHS		116/1840	
Paudel S ²⁴ <i>et al.</i> 2020	2020	Secondary analysis of 1977 the data	1977	Across Nepal		179/1977	
Koirala S ²⁵ et al. 2018	2018	Cross-sectional study	188(85M/103F)	Mustang district	59/188	9/188	
Ranabhat K ²⁶ <i>et al.</i> 2016	2016	2016 Cross-sectional study 154 (80M/74F)	154 (80M/74F)	Tribhuwan University Teaching Hospital of Nepal		66/154	
Mehta KD ²⁷ et al. 2011	2011	2011 Cross-sectional study 2006(1096M/910F) Sunsari , Eastern Nepal	2006(1096M/910F)	Sunsari , Eastern Nepal		422/2006	80/289
BPKIHS, B.P. Koirala Institute of Health Sciences; F, female; IGT, Impaired Glucose Test; M, Male.	Health Sc	ciences; F, female; IGT, Impai	red Glucose Test; M, Mal	ai			

Table 2. Qualitative summary.

while this value increased to 11.24%, between 2015–2020 (Proportion, 0.1124; 95% CI, 0.0789-0.1577; 1²: 96.74) (Figure 3).

In relation to the study setting, the re-analysis of the data with the use of the random-effects model showed that 10.4% among surveyed adult population based on community-based studies had T2DM (Proportion, 0.1040; 95% CI, 0.0668-0.1596) (*Extended data* file 2, Figure 2), while 9.23% among hospital/Directly

observed, treatment short-course (DOTS) center-based studies have this disease (Proportion, 0.0923; 95% CI, 0.0509-0.1617) (*Extended data* file 2, Figure 3^{10}).

Pre-diabetes was present in 19.4% (Proportion, 0.194; 95% CI, 11.2%- 31.3%) (*Extended data* file 2, Figure 4) and IGT in 11.0% (Proportion, 0.110; 95% CI, 4.3%- 25.4%) (*Extended data* file 2, Figure 5^{10}).

	FIEVE				3	
Study name					Event rate and 95%	CI
	E vent rate	Lower limit	Upper limit	Total		
Bhandari GP, et al. 2014	0.033	0.030	0.036	391/11901		
Paudyal G, et al. 2008	0.041	0.032	0.052	60/1475		
Kushuwaha A, et al. 2020	0.044	0.018	0.101	5/114		
Koirala S, et al. 2018	0.048	0.025	0.089	9/188		
Karki P, et al. 2000	0.063	0.053	0.075	116/1840		
Sharma SK, et al. 2011	0.063	0.060	0.068	889 / 14008		
Sharma SK, et al. 2013	0.075	0.067	0.085	242/3218		
Dhimal M, et al. 2019	0.085	0.080	0.090	1067 / 12557		
Paudel S, et al. 2020	0.091	0.079	0.104	179/1977		
Gyawali B, et al. 2018	0.117	0.105	0.131	271/2310		
Sharma B, et al. 2019	0.119	0.088	0.159	38 / 320		
Shrestha UK, et al. 2006	0.190	0.167	0.215	192/1012		
Mehta KD, et al. 2011	0.210	0.193	0.229	422 / 2006		
Chhetri MR, et al. 2009	0.258	0.238	0.280	422 / 1633		
Ranabhat K, et al. 2016	0.429	0.353	0.508	66 / 154		
	0.100	0.071	0.139			
					-1.00 -0.50 0.00 0.50	1

Prevalence of Diabetes Mellitus

Figure 2. Prevalence of T2DM in Nepal.

Prevalence of T2DM in Nepal in last decade

Froup by	Study name	Stati	stics for each	study		Event rate and 95%
ime		Event rate	Lower limit	Upper limit	Total	
010-15	Bhandari GP, et al. 2014	0.033	0.030	0.036	391 / 11901	1 🗰
010-15	Sharma SK, et al. 2011	0.063	0.060	0.068	889 / 14008	🖬
010-15	Sharma SK, et al. 2013	0.075	0.067	0.085	242 / 3218	
010-15	Mehta KD, et al. 2011	0.210	0.193	0.229	422 / 2006	
010-15		0.078	0.037	0.156		
015-20	Kushuwaha A, et al. 2020	0.044	0.018	0.101	5/114	
015-20	Koirala S, et al. 2018	0.048	0.025	0.089	9/188	
015-20	Dhimal M, et al. 2019	0.085	0.080	0.090	1067 / 12557	
015-20	Paudel S, et al. 2020	0.091	0.079	0.104	179 / 1977	
015-20	Gyawali B, et al. 2018	0.117	0.105	0.131	271 / 2310	
015-20	Sharma B, et al. 2019	0.119	0.088	0.159	38/320	
015-20	Ranabhat K, et al. 2016	0.429	0.353	0.508	66 / 154	📫
015-20		0.112	0.079	0.158		

Figure 3. Prevalence of T2DM in Nepal taking consideration of time frame from 2010-2020.

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-1.00 -0.50 0.00 0.50 1.00

Risk factors of T2DM

Exercise. Random-effects model that incorporated data from six studies on exercise showed that the difference in T2DM status between adequate and inadequate exercise groups were not statically significant (OR, 0.75, 95% CI, 0.49-1.16; I², 67.85%) (Figure 4).

BMI. Fixed-effect meta-analysis of five studies that reported on the BMI indicated that with a BMI \geq 24.9 kg/m² the odds of having T2DM is 2.19 times higher than with BMI <24.9 kg/m² (OR, 2.197; 95% CI, 1.799-2.683) (Figure 5).

Waist circumference. Individuals with healthy waist circumference had 64.1% lower odds of having T2DM compared with those with high waist circumference (OR, 0.361; 95% CI, 0.284-0.460; I², 0%) (*Extended data* file 2, Figure 4¹⁰).

Smoking status. The random-effects meta-analysis of four T2DM studies based on smoking status indicated that the differences in T2DM status among smokers and non-smoker were not significant (OR, 0.752; 95% CI, 0.366-1.546; I^2 ; 87.2%) (Figure 6).

Alcohol consumption. T2DM status in relation with alcohol consumption was assessed by four studies with the use of random-effects model. The results showed that T2DM status among alcoholic and non-alcoholic groups were not statistically significant (OR, 0.750; 95% CI, 0.439-1.281 I²; 37.72%) (Figure 7).

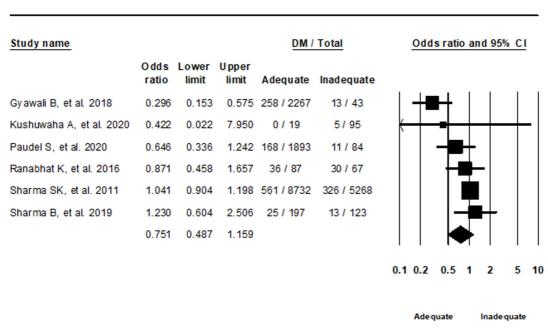
BP. Fixed-effect meta-analysis of three studies that have reported on T2DM status in relation with BP has indicated that the odds of individuals with normal BP having T2DM is 62.1% lower than those with high BP (OR, 0.379; 95% CI, 0.290-0.495) (Figure 8).

Literacy. The assessment of four studies that reported on T2DM based on literacy status did not show any significant differences in T2DM between literate and illiterate groups (OR, 1.165; 95% CI, 0.664-2.045; I², 93.61%) (Figure 9).

Family history. The random-effects meta-analysis of three studies indicated that the odds of T2DM in individuals without a family history of T2DM were 67.3% lower in comparison to those with a family history (OR, 0.327; 95% CI, 0.202-0.529; I², 56.62%) (Figure 10).

Fruits and vegetables intake. The data assessment of the two studies that had reported on T2DM status in relation to fruits and vegetable intake did not reach a significant difference (OR, 0.933; 95% CI, 0.441-1.976; I², 78.72%). (*Extended data* file 2, Figure 7¹⁰).

Publication bias. Publication bias among the included studies were tested with the use of Egger's test and was presented in a Funnel plot. The prevalence of T2DM in the Funnel plot showed an asymmetric distribution of studies, which suggested publication bias (*Extended data* file 2, Figure 8¹⁰).



Exercise and occurence of T2DM

Figure 4. Forest plot showing exercise status and T2DM in Nepal.

Study name	Sta	tistics fo	or each s	study	DM /	Total
	Odds ratio	Lower limit		p-Value	BMI>/= 24.9	BMI<24.9
Sharma B, et al. 2019	1.227	0.446	3.375	0.692	5/36	33/284
Ranabhat K, et al. 2016	1.671	808.0	3.459	0.166	51/110	15/44
Gyawali B, et al. 2018	2.145	1.601	2.874	0.000	206 / 1422	65 / 888
audel S, et al. 2020	2.455	1.799	3.350	0.000	86 / 578	93 / 1399
shuwaha A, et al. 202	013.948	0.753	258.473	0.077	5/53	0/61
	2.197	1.799	2.683	0.000		

BMI and occurence of T2DM

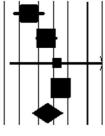
0.10.2 0.5 1 2 5 10 BMI>/=24.9 BMI<24.9

Figure 5. Forest plot showing BMI category and T2DM in Nepal.

Smoking and T2DM

Sta	atistics fo	r each s	tudy	DM / T	otal	Odds ratio
Odds ratio	Lower limit	Upper limit	p-Value	Non-smoker	Smokers	and 95% C
0.317	0.152	0.660	0.002	24 / 262	14 / 58	
0.718	0.433	1.188	0.197	91 / 224	41 / 84	
1.190	0.127	11.142	0.879	4 / 88	1/26	│┽┼╞
1.403	0.957	2.056	0.083	238 / 1945	33 / 365	
0.752	0.366	1.546	0.439			
	Odds ratio 0.317 0.718 1.190 1.403	Odds ratio Lower limit 0.317 0.152 0.718 0.433 1.190 0.127 1.403 0.957	Odds ratioLower limitUpper limit0.3170.1520.6600.7180.4331.1881.1900.12711.1421.4030.9572.056	ratiolimitlimitp-Value0.3170.1520.6600.0020.7180.4331.1880.1971.1900.12711.1420.8791.4030.9572.0560.083	Odds ratio Lower limit Upper limit p-Value Non-smoker 0.317 0.152 0.660 0.002 24 / 262 0.718 0.433 1.188 0.197 91 / 224 1.190 0.127 11.142 0.879 4 / 88 1.403 0.957 2.056 0.083 238 / 1945	Odds ratio Lower limit Upper limit p-Value Non-smoker Smokers 0.317 0.152 0.660 0.002 24 / 262 14 / 58 0.718 0.433 1.188 0.197 91 / 224 41 / 84 1.190 0.127 11.142 0.879 4 / 88 1 / 26 1.403 0.957 2.056 0.083 238 / 1945 33 / 365

io CI



0.10.2 0.5 1 2 5 10

Non-smoker Smoker

Figure 6. Forest plot showing smoking status and T2DM in Nepal.

Study name				DM / T	otal	Odds ratio and 95% Cl
	Odds ratio	Lower limit	Upper limit	Non-alcoholic	Alcoholic	
Kushuwaha A, et al. 2020	0.055	0.003	1.022	0 / 68	5/46	
Ranabhat K, et al. 2016	0.581	0.150	2.253	61 / 145	5/9	┤┼╼╪┼┤┃│
Sharma B, et al. 2019	0.679	0.344	1.340	18 / 177	20/140	+∰+
Gyawali B, et al. 2018	1.001	0.689	1.453	235 / 2003	36/307	
	0.750	0.439	1.281			

Alcochol consumption and T2DM

Non-alcoholic Alcoholic

0.10.20.51 2 510

Figure 7. Forest plot showing alcohol consumption status and T2DM in Nepal.

Blood pressure and T2DM

Study name	Sta	tistics fo	or each s	study	DM/	Total	Odds ratio
	Odds ratio	Lower limit	Upper limit	p-Value	Normal BP	High BP	and 95% CI
Gyawali B, et al. 2018	0.332	0.256	0.430	0.000	114 / 1513	157 / 797	
Paudel S, et al. 2020	0.386	0.281	0.531	0.000	65/1137	114/840	
Sharma B, et al. 2019	0.694	0.310	1.558	0.376	29/261	9/59	
	0.367	0.302	0.446	0.000			

0.10.20.51 2 510

Normal BP High BP

Figure 8. Forest plot showing blood pressure status and T2DM in Nepal.

Study name				DM /	Total
	Odds ratio	Lower limit	Upper limit	Literate	Illiterate
Gyawali B, et al. 2018	0.770	0.596	0.995	113 / 1095	158 / 1215
Sharma SK, et al. 2011	0.770	0.668	0.888	575 / 9805	314 / 4197
Paudel S, et al. 2020	2.496	1.819	3.424	111/822	68/1155
Ranabhat K, et al. 2016	1.359	0.626	2.950	53/119	13/35
	1.165	0.664	2.045		

Literacy and T2DM

0.10.2 0.5 1 2 5 10

Literate Illiterate

Figure 9. Forest plot showing literacy status and T2DM in Nepal.

Family history of T2DM

Study name	<u>Sta</u>	tistics fo	or each :	study	DM/	Total	Odds ratio and 95% Cl
	Odd s ratio	Lower limit	Upper limit	p-Value	No	Yes	
Sharma B, et al. 2019	0.196	0.096	0.400	0.000	14 / 225	24/95	╞╋╋╾┤╎╎╎╎
Gyawali B, et al. 2018	0.315	0.241	0.413	0.000	164 / 1855	107 / 455	
Ranabhat K, et al. 2016	0.588	0.288	1.202	0.145	44 / 112	22/42	││→╪═┼│││
	0.327	0.202	0.529	0.000			

0.10.20.51 2 510

No Yes

Figure 10. Forest plot showing the family history of T2DM and Diabetes status in patients in Nepal.

Discussion

The prevalence of T2DM, pre-diabetes, and IGT in Nepal was found to be 10%, 19.4%, and 11% respectively. Our results show that in Nepal obesity is the highest risk factor for T2DM, while individuals with normal waist circumference and lack of family history of T2DM had lower risk of T2DM.

The estimated prevalence of T2DM was higher than that reported in WHO STEP wise approach to Surveillance (STEPS) survey in 2013 (3.6%), and previous meta-analyses (8.4%) and 8.5%)^{5,6,31}. Similarly, the estimated prevalence of prediabetes in our study was almost double than what has been reported in other studies^{5,6}. One explanation for this finding can be the rapid urbanization, and migration from rural to urban areas which has promoted a sedentary lifestyle among individuals, along with consumption of unhealthy foods³². As per our study, high BMI was the main cause of T2DM in Nepal. In South Asia, lifestyle factors such as poor diet, and increased sedentary behaviors with limited physical activities have contributed to the rise of overweight and obesity among children and adolescents³³. Rapid development of the economic situation in developing countries like Nepal has resulted in a change of diet rich in cereal and vegetables to one with animal products and processed food with high fat and sugar content³⁴. In a study by Hills et al. the prevalence of overweight in Nepal was estimated to be 16.7%, with a higher prevalence in women (19.6%) compared to men (13.6%)³⁴. Obesity is closely linked with premature onset of T2DM and cardiovascular disease³⁵. A similar increasing trend of T2DM led by obesity is seen in Africa as well³⁶. It is important to target T2DM risk factors in order to take control of this disease in Nepal. Our finds highlight the importance of exercise and a healthy diet to prevent the increased morbidity among individuals with T2DM in this country. Shrestha et al. found that the T2DM awareness to be low, with nearly half of the population unaware of the fact that they had this disease⁶. Increasing public awareness about noncommunicable diseases like T2DM and hypertension, and the need to implement a healthy lifestyle is of paramount importance given that our results indicated that individuals with normal blood pressure had less chance of developing T2DM compared to those with hypertension. Increased intake of oily foods, reusing cooking oils which can cause increased conversion of unsaturated fats to trans fats, and low consumption of fruits and vegetables have been found throughout South Asia^{37,38}. These unhealthy dietary habits lead to increased risks of non-communicable diseases like T2DM and hypertension. Thus, interventions are needed to better manage the overweight and obesity epidemic. This can be achieved through various measures such as opening public parks in the cities for exercise, educating the population about what a healthy lifestyle entails such as decreasing the intake of oily foods, increasing the intake of fruits and vegetable, as well as improving the quality of food. Our study has several strengths. Firstly, we performed comprehensive literature search to pool the results of fifteen studies over the last twenty years to evaluate the prevalence of T2DM in Nepal. In addition, no prior meta-analysis has evaluated the risk factors for T2DM, specifically IGT in Nepal, prior to our study. We also analyzed data based on a time frame, where

significant increase in T2DM prevalence was observed in Nepal when comparing 2010–2015 with 2015–2020. Our study had some limitations. There was heterogeneity in the studies due to variation in the T2DM diagnostic criteria, different demographics of the population, etc. Most of the included studies were based on specific areas such as province 1 and 3, and not enough studies have been done on a national scale. Finally, risk factors for T2DM were not reported in all the studies that were included.

Conclusion

The prevalence of T2DM, pre-diabetes and IGT in Nepal was estimated to be 10%, 19.4% and 11% respectively. Obesity is the major risk factor of T2DM in Nepal and people with normal waist circumference, normal blood pressure and lack of family history of T2DM had lower odds of developing this disease.

Data availability statement Underlying data

Figshare: Diabetes Mellitus in Nepal from 2000 to 2020: A systematic review and meta-analysis

https://doi.org/10.6084/m9.figshare.14706648.v112

The project contains the following underlying data:

Dataset: Quantitative data, glycemic control, socio-economic status, BMI, exercise, T2DM prevalence, waist circumference, family history, fruit and vegetable serving per day, alcohol consumption, smoking, education, and BP)

Extended data

Figshare: Diabetes Mellitus in Nepal from 2000 to 2020: A systematic review and meta-analysis

https://doi.org/10.6084/m9.figshare.14854065.v110

The project contains the following underlying data:

- Data file 1: Electronic search details
- Data file 2: Additional analysis
- Data file 3: PRISMA checklist

Data are available under the terms of the Creative Commons Zero "No rights reserved" data waiver (CC0 1.0 Public domain dedication).

Authors' contributions

DBS, PB, and YRS contributed to the concept and design, analysis, and interpretation of data. DBS, PB, AM, SL, SN, AA, and AP contributed to the literature search, data extraction, review, and initial manuscript drafting. YRS, SN, and AA interpretation of data, revising the manuscript for important intellectual content, and approval of the final manuscript.

All authors were involved in drafting and revising the manuscript and approved the final version.

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Version 1

Reviewer Report 06 September 2021

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Prajwal Gyawali 匝

School of Health and Wellbeing, Faculty of Health, Engineering and Sciences, University of Southern Queensland, Toowoomba, QLD, Australia

A) Background

1. Please check for sentence structure;

"In 2019, the International Diabetes Federation (IDF) estimated that 463 million adults worldwide had diabetes1. The statistics showed that these individuals were in the age range of 20 to 79, **have diabetes**, and 79.4% were from low- and middle-income countries."

- 2. Reference number 4 used in the background does not match with the authors' claim. This reference is more focused on foetal programming contributing to the prevalence of obesity in Nepal.
- 3. There are already two systematic reviews published in this sector, references 5 and 6:

Gyawali B, Sharma R, Neupane D, *et al.*: Prevalence of type 2 diabetes in Nepal: a systematic review and meta-analysis from 2000 to 2014¹.

Shrestha N, Mishra SR, Ghimire S, *et al.*: Burden of Diabetes and Prediabetes in Nepal: A Systematic Review and Meta-Analysis².

4. Authors need to provide a valid reason for conducting this systematic review in the same area in the background section. Paper by Shrestha N. *et al.*² was published less than 1 year ago and they have included 14 papers for the final analysis.

B) Methods

1. The authors have reported the prevalence of impaired glucose tolerance and prediabetes. Why were these keywords not used in the search strategy? Likewise, have the authors used the terminology 'glucose' as a keyword in a search strategy?

C) Discussion

1. The metanalysis published in 2020 by Shrestha N. *et al.*² included 14 papers in the final

analysis. All papers were after 2000. They reported the prevalence of pre-diabetes is much less than what the current paper has reported. The reason provided by the authors for this large difference is not properly addressed in the discussion. This is very important to address.

2. The papers should be discussed in sub-sections.

References

1. Gyawali B, Sharma R, Neupane D, Mishra SR, et al.: Prevalence of type 2 diabetes in Nepal: a systematic review and meta-analysis from 2000 to 2014.*Glob Health Action*. 2015; **8**: 29088 PubMed Abstract | Publisher Full Text

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Are the rationale for, and objectives of, the Systematic Review clearly stated?

No

Are sufficient details of the methods and analysis provided to allow replication by others? $\ensuremath{\mathbb{No}}$

Is the statistical analysis and its interpretation appropriate?

I cannot comment. A qualified statistician is required.

Are the conclusions drawn adequately supported by the results presented in the review? Partly

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Stroke

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Reviewer Report 13 August 2021

https://doi.org/10.5256/f1000research.57408.r91625

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? Chudchawal Juntarawijit 匝

Department of Natural Resources and Environment, Faculty of Agriculture, Natural Resources, and Environment, Naresuan University, Phitsanulok, Thailand

Background:

- 1. In the introduction, more information about the prevalence and risk factors of diabetes should be presented and reorganized for easy reading. The research problem and hypothesis also need to be clearly articulated.
- 2. The rationale for the review was also not clearly described in the context of what is already known. As there have been few studies in Nepal, it was not a good reason for conducting meta-analysis. Doing a meta-analysis using a small number of studies might cause bias.

Methods:

- 1. Study methods were poorly described and hard to read (see a good example study by Miller *et al.*¹). To help readers replicate the study, the authors should provide more detailed information on electronic search strategy, methods of results synthesis, and sensitivity analysis.
- 2. Using only the global search engine, e.g. PubMed, Embase, Scopus, and Google Scholar might cause publication bias if there were some researches papers published locally or in other databases. In point had not been mention in the study limitation.

Results:

1. Results were inadequately described. In the qualitative summary, the authors should talk about the important features of the information in Table 1. In quantitative synthesis, information on how the pooled prevalence was calculated is also needed. In every figure, the total events should be indicated, and the figure legend should contain more information.

Discussion:

- 1. In the Discussion, more information should be provided and reorganized for an easy read.
- 2. In the study limitation, the authors mention T2DM diagnostic criteria and demographic data as a problem. However, the authors did not mention how the problems affect the results and how the study handled them.
- 3. The statement: "Our finds highlight the importance of exercise and a healthy diet to prevent the increased morbidity among individuals with T2DM in this country", needs more justification. Was the information from the results of this study or from the literature?
- 4. In an attempt to identify diabetes risk factors, the study limited only those studies from Nepal. How this will affect the study results?

Conclusion:

 In the conclusion statement, only obesity was claimed to cause diabetes. This conclusion might need more justification since obesity and other risk factors were not extensively discussed. This result was also in contrast to that observed by Jayawardena *et al.*², a similar study using data from all countries in South Asia. In that study, family history, urban residency, age, higher BMI, sedentary lifestyle, hypertension, and waist-hip ratio were found to be associated with an increased risk of diabetes. 2. The conclusion statement should also provide study limitations and recommendations for public health use.

References

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Are the rationale for, and objectives of, the Systematic Review clearly stated? Partly

Are sufficient details of the methods and analysis provided to allow replication by others? Partly

Is the statistical analysis and its interpretation appropriate?

Partly

Are the conclusions drawn adequately supported by the results presented in the review? Partly

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Environmental health science

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Author Response 29 Aug 2021

Alok Atreya, Lumbini Medical College, Palpa, Nepal

Background: In the introduction, more information about the prevalence and risk factors of diabetes should be presented and reorganized for easy reading. The research problem and hypothesis also need to be clearly articulated.

Reply: Thank you for the comment. We have presented the situation of T2DM in a funnel pattern in the background giving an overview of global, regional, and national scenarios. We have rephrased and edited as per requirement by the reviewer's comment to further clarify the rationale.

The rationale for the review was also not clearly described in the context of what is already

known. As there have been few studies in Nepal, it was not a good reason for conducting meta-analysis. Doing a meta-analysis using a small number of studies might cause bias.

Reply: Thank you for the comment. We have edited the rationale as per requirement by the reviewer's comment to further clarify the rationale. We included 15 prevalencebased studies with justifiable sample sizes and assessed the quality of those individual studies using the JBI bias tool to reduce the bias of including studies. Due to the urge of the evidence-based practice, we authors think pooling of the data from available individual studies to give pooled prevalence is justifiable.

Methods: Study methods were poorly described and hard to read (see a good example study by Miller et al.1). To help readers replicate the study, the authors should provide more detailed information on electronic search strategy, methods of results synthesis, and sensitivity analysis.

Reply: Thank you for the comment. We have conducted our study as per the standard guideline and abide by the MOOSE checklist for the meta-analysis of observational studies. Our electronic search details including the link, and additional sensitivity analysis are available in the extended data file and shared publicly.

Using only the global search engine, e.g. PubMed, Embase, Scopus, and Google Scholar might cause publication bias if there were some researches papers published locally or in other databases. In point had not been mention in the study limitation.

Reply: Thank you for the comment. We have conducted our study as per the standard guideline and abide by the MOOSE checklist for the meta-analysis of observational studies. In Nepal, almost all Nepalese local journals are listed in NEPJOL (Nepalese Journal Online), which is the common platform and all the published papers are available through Google scholar so we included the Google scholar database in our review, which is missed by some prior reviews. So, we authors believe including Google scholar and other standard global databases won't miss published literature.

Results: Results were inadequately described. In the qualitative summary, the authors should talk about the important features of the information in Table 1. In quantitative synthesis, information on how the pooled prevalence was calculated is also needed. In every figure, the total events should be indicated, and the figure legend should contain more information.

Reply: Thank you for the comment. We elaborated qualitative summary as "A qualitative summary of all 15 individual studies is presented in (Table 2). Nine studies were done in a community setting while the rest six were done in the hospital setting. Two community-based studies and one hospital-based study included samples from different parts of Nepal to represent the country, while the rest were loco-regional studies." For transparency of our work we made our data available in public data repository Figshare; link: https://doi.org/10.6084/m9.figshare.14706648.v1. Regarding data pooling methods, based on heterogeneity we used a random or fixed-effect model. And we mentioned the model used in every section before the

description of the individual results and forest plot. Regarding total events, I agree with the reviewer, however, due to the default setting of the software CMA-3, we could not edit the figure.

Discussion:

In the Discussion, more information should be provided and reorganized for an easy read. In the study limitation, the authors mention T2DM diagnostic criteria and demographic data as a problem. However, the authors did not mention how the problems affect the results and how the study handled them. The statement: "Our finds highlight the importance of exercise and a healthy diet to prevent the increased morbidity among individuals with T2DM in this country", needs more justification. Was the information from the results of this study or from the literature?

Reply: Thank you for the comment. There are perceived barriers for the noncompliance for physician's advice to a special diet and regular exercise as shown in a Nepalese study. We have added a reference to justify this statement.

In an attempt to identify diabetes risk factors, the study limited only those studies from Nepal. How this will affect the study results?

Reply: Thank you for the comment. Diabetes prevalence and risk differ across different societies and regions across the globe. This study is solely aimed to estimate prevalence and risk factors in the context of Nepal to build foundation evidence to make evidence-based practice in Nepal so may be different than other regions across the globe.

Conclusion: In the conclusion statement, only obesity was claimed to cause diabetes. This conclusion might need more justification since obesity and other risk factors were not extensively discussed. This result was also in contrast to that observed by Jayawardena et al. 2, a similar study using data from all countries in South Asia. In that study, family history, urban residency, age, higher BMI, sedentary lifestyle, hypertension, and waist-hip ratio were found to be associated with an increased risk of diabetes. The conclusion statement should also provide study limitations and recommendations for public health use.

Reply: Thank you for the comment. The conclusion statement states obesity as one of the many major risk factors for diabetes. We kindly apologize, but we have not claimed only obesity causes diabetes. The limitations have already been provided at the end of the discussion section, so due to fear of duplication, we did not include them in the conclusion section.

Competing Interests: No competing interests were disclosed.

Reviewer Report 29 July 2021

https://doi.org/10.5256/f1000research.57408.r89249

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Sijan Basnet 匝

Department of Internal Medicine, Reading Hospital and Medical Centre, West Reading, PA, USA

This is a very well-written manuscript with robust statistical analysis.

1. Please mention the limitations of your study if any.

2. Most of the studies cited are from the 2010s. There is one from 2000 with 1840 subjects from an outpatient clinic in a city that may not be representative of the country's demographics and one from 2006 with 1012 subjects from seven wards that are being used to study the trend. Does this skew your results in any way? Also, the authors justify the increasing prevalence of rapid urbanization but the above studies were done in cities.

Are the rationale for, and objectives of, the Systematic Review clearly stated? Yes

Are sufficient details of the methods and analysis provided to allow replication by others? $\ensuremath{\mathsf{Yes}}$

Is the statistical analysis and its interpretation appropriate?

Yes

Are the conclusions drawn adequately supported by the results presented in the review? $\ensuremath{\mathsf{Yes}}$

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Internal Medicine

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Author Response 29 Aug 2021

Alok Atreya, Lumbini Medical College, Palpa, Nepal

Thank you for the comment. Due to the relatively few studies from 2000-2010. We did an analysis based on a published study between 2010-2015, and 2015-2020 as "The assessment of T2DM prevalence between 2010–2015 with the use of random-effects meta-analysis was 7.75% (Proportion, 0.0775; 95% CI, 0.0367-0.1561; studies: 4; I ²:99.62), while this value increased to 11.24%, between 2015–2020 (Proportion, 0.1124; 95% CI, 0.0789-0.1577; I ²: 96.74)". Analysis showed some increasing trends of T2DM in Nepal which is of concern.

Being our meta-analysis is a secondary analysis of the published literature, we do have some limitations due to primary studies variation. We have included our study limitations as "Our study had some limitations. There was heterogeneity in the studies due to variation in the T2DM diagnostic criteria, different demographics of the population, etc. Most of the included studies were based on specific areas such as provinces 1 and 3, and not enough studies have been done on a national scale. Finally, risk factors for T2DM were not reported in all the studies that were included."

Competing Interests: No competing interests were disclosed.

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