ORIGINAL RESEARCH Association of Hyper-Triglyceridemic Waist Phenotype and Diabetic Vascular Complication in the Chinese Population

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Background: Diabetic vascular complications are the leading cause of crippling and death of diabetic patients and seriously affect patients' quality of life. It is essential to control the risk factors contributing to vascular complications in patients with Type 2 diabetes (T2DM). This study aimed to examine the association between hyper-triglyceridemic waist phenotype (HWP) and the risk of vascular complication index of diabetes in T2DM patients.

Methods: The participants with type 2 diabetes mellitus in this study registered at the National Metabolic Management Center (MMC) of Beijing Luhe Hospital from June 2017 to June 2021. Data were collected by trained personnel according to the protocol. The questionnaire containing information on demographic characteristics and lifestyle factors (including alcohol drinking and cigarette smoking et al) was administered by trained interviewers. Logistic regression analysis assessing the associations between the hypertriglyceridemic waist phenotype and vascular complication index of diabetes. In addition, the subgroup analysis was performed by age, sex, HbA1c, hypertension or not, and education level.

Results: After data cleaning, a total of 3221 participants with T2DM were enrolled. The median (IQR) duration of diabetes was 47.00 (3.00, 125.00) months. Compared to the participants in the Normal triglycerides level and Normal waist circumference group (NTNW), those in the Elevated triglycerides level and Enlarged waist circumference group (HTGW) have a higher risk of CKD-related vascular complications; the OR of decreased estimated glomerular filtration rate (GFR) and elevated urinary albumin creatinine ratio (UACR) were 2.21 (95% CI:1.32-3.82) and 2.18 (95% CI:1.69–2.81), respectively. Moreover, compared to the participants in the NTNW group, the ORs of the decreased anklebrachial index (ABI) and elevated Brachial-ankle pulse wave velocity (baPWV) were 2.24 (95% CI:1.38-3.80) and 1.63 (95% CI:1.28-2.06) in the HTGW group.

Conclusion: In summary, there was an association between hyper-triglyceridemic waist phenotype and diabetic vascular complications in the Chinese population.

Keywords: hyper-triglyceridemic waist phenotype, type 2 diabetes mellitus, vascular complication

Introduction

Type 2 diabetes mellitus (T2DM) is a significant health problem, especially in low and middle-income countries.¹ The new epidemiological investigation showed that 537 million people had diabetes in 2021.² Diabetic vascular complications are the leading cause of crippling and death of diabetic patients and seriously affect patients' quality of life, including diabetic nephropathy (DN), cardiovascular (CVD), and diabetic retinopathy (DR) et al^{3-5} Therefore, it is essential to control the risk factors contributing to vascular complications in patients with T2DM.

In the new decade, several studies have concentrated on finding metabolism-related indexes to predict chronic disease complications. For example, Wu et al found a relationship between triglyceride glucose (TyG) index and arterial stiffness progression.⁶ Moreover, there was an association between Visceral Adiposity Index (VAI) and artery disease in normal-weight adults with hypertension.⁷

The hyper-triglyceridemic waist phenotype (HWP) is a physical characterized classified by triglycerides (TG) and waist circumference, which is characterized by enlarged waist circumference and elevated TG levels.⁸ As a composite index-related metabolism, HWP was used to predict hypertension.⁹ In addition, several studies have shown that HWP was associated with coronary artery disease and metabolic syndrome.^{10–12} Long-term obesity leads to abnormal lipid metabolism.¹³ When TG level is present at a high level, heparin activates lipoprotein lipase, which increases intravascular lipolysis of circulating TG, thereby increasing tissue exposure to free fatty acids, ultimately leading to insulin resistance and impaired B-cell function.¹⁴ As a result, there is an association between HWP and T2DM. However, the relationship between the risk of vascular complications and HWP in diabetic patients is unclear.

Therefore, this study aimed to investigate the association between HWP and the risk of vascular complication index of diabetes in T2DM patients.

Method

Study Participants

The participants with type 2 diabetes mellitus in this study registered at the National Metabolic Management Center (MMC) of Beijing Luhe Hospital from June 2017 to June 2021. The MMC is a national project to manage metabolic patients according to the same standard. All the patients accepted systematic physical examination, blood sample collection, and oral questionnaire interviews. The protocol of this project was published previously.¹⁵ The MMC is a long-term follow-up program. Considering the life expectancy of patients, the present study only included participants under 80 years.

Participants were excluded according to the following criteria: (1) Pregnant or nursing women. (2) Malignant tumor or life expectancy <5 years, (3) Acute complications of diabetes, (4) Presence of severe heart, kidney, and liver disease.

The protocol was approved by the Medical Ethics Committee of Beijing Luhe Hospital, Capital Medical University. This study was performed by the Declaration of Helsinki. All participants provided written informed consent.

Data Collection

Data were collected by trained personnel according to the protocol. The questionnaire containing information on demographic characteristics, and lifestyle factors such as alcohol drinking and cigarette smoking, was administered by trained interviewers. For the participants who smoked daily or almost daily, smoking status was defined as "yes". And for the participants who drank weekly or nearly weekly, drinking status was described as "yes". Education level was categorized as less than high school and high school or more.

Weight and height were measured with a standard protocol at 8:00–10:00 AM, and body mass index (BMI) was calculated as weight divided by height squared. Peripheral venous blood was withdrawn from all subjects in the morning after fasting for 8 hours or more. Glycated hemoglobin (HbA1c) levels were assayed using the method of high-performance liquid chromatography (HPLC) with a D10 set (Bio-RAD, Hercules, CA, USA). An auto biochemical analyzer measured LDL, HDL, and triglyceride (AU5800, Beckman Coulter, USA).

Brachial-ankle pulse wave velocity (baPWV) and ankle-brachial index (ABI) were measured by an automated recording apparatus with participants in the supine position after at least 5 minutes of rest (BP-203RPE III, form PWV/ABI, Omron Healthcare Co.). In this study, baPWV was calculated as La–Lb/ Δ T (La and Lb are the distance from the heart to the brachium, respectively, Δ T is the time between the wavefront of the brachial waveform and that of the ankle waveform).

Diabetic retinopathy (DR) severity was classified according to the International Clinical Diabetic Retinopathy and Diabetic Macular Oedema Disease Severity Scales.¹⁶ DR of the present study was defined as severe non-proliferative diabetic retinopathy and proliferative diabetic retinopathy, also called vision-threatening DR (VTDR).

Definitions

The 1999 World Organization criteria were used to diagnose T2DM. Patients who had fasting plasma glucose \geq 7.0 mmol/L or 2-h plasma glucose \geq 11.1 mmol/L or a self-reported physician diagnosis were diagnosed with T2DM.

Triglyceride waist circumference phenotypes were divided into four groups. (1) Normal TG level and Normal waist circumference, NTNW (TG \leq 1.7 mmol/L and males' waist <90cm or females' waist <85cm). (2) Normal TG level and Enlarged waist circumference, NTEW (TG \leq 1.7 mmol/L and males' waist \geq 90 cm or females' waist \geq 85cm). (3) Elevated TG level and Normal waist circumference, ETNW (TG \geq 1.7 mmol/L and males' waist < 90 cm or females' waist < 85cm). (4) Elevated TG level and Enlarged waist circumference, HTGW (TG> 1.7 mmol/L and males' waist \geq 90 cm or females' waist \geq 90 cm or females' waist < 85cm).

ABI and baPWV were the arterial stiffness-related index; decreased ABI and elevated baPWV were defined as <0.9 and >1400 m/s, respectively.^{17–19} Decreased estimated glomerular filtration rate (GFR) and elevated urinary albumin creatinine ratio (UACR) were defined as <60mL/min/1.73 m² and \geq 30mg/g.²⁰ GFR was calculated by the aCKD-EPI formula.²¹

Statistical Analysis

Continuous variables were described as mean \pm standard deviation (SD) or median [interquartile range (IQR)], and categorical variables were described as frequency (%). Data should be evaluated for normal distribution and, if necessary, the logarithmic transformation would be performed for statistical analysis. The differences between groups were assessed using Pearson's χ^2 test for categorical variables and the *t*-test or Mann–Whitney *U*-test for continuous variables.

The associations between the hyper-triglyceridemic waist phenotype and vascular complication index of diabetes were assessed by the Logistic regression model. Two models were constructed with different covariables: model 1 inputs major variables of diabetes, including sex, duration, and HbA1c. History of hypertension, smoking, drinking, education, LDL, and HDL were primary factors in the vascular research and associated with the vascular complication index of diabetes.^{9,22–24} To avoid these confounding effects, model 2 adjusted for a history of hypertension, smoking, drinking, education, LDL, HDL, plus the covariable of model 1.

In addition, the subgroups analysis was performed by age (<60, ≥60 years), sex (male, female), HbA1c (<7.0%, $\ge7.0\%$), hypertension or not, an education level (less than high school, high school education or higher level).

R statistical software (version 4.1.2) was used for statistical analysis. Two-tailed P <0.05 was considered statistically significant.

Results

Demographic and Clinical Characteristics

The data cleaning procedure can be seen in Figure 1. After excluding participants aged not 20–80 years, missing value, and using antilipemic agents, there were 3221 participants including the current analysis.

Table 1 shows the demographic and clinical characteristics of this study. A total of 1914 (59.4%) participants were males. The median (IQR) duration of diabetes was 47.00 (3.00, 125.00) months. Compared to the participants in the NTNW group, those in the HTGW group were younger with a shorter duration of diabetes, higher fasting blood glucose, HbA1c, baPWV, UACR, and ABI (all P < 0.01). Moreover, the participants in the HTGW tend to be obese, which has a larger value of physical measurement index including waist circumference, body weight, and BMI.

The Associations Between the HWP and Vascular Complication Index of Diabetes

The HWP was associated with the vascular complication index of diabetes in T2DM patients. Table 2 gives the model information on the vascular complication index of T2DM, which includes decreased ABI, decreased GFR, elevated

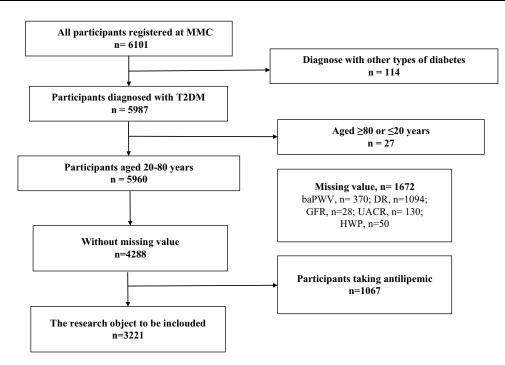


Figure I Date cleaning procedure. After excluding participants aged not 20–80 years, missing value, and using antilipemic agents, there were 3221 participants including the current analysis.

UACR, elevated baPWV, and DR. The effect of HWP on the vascular complication index was described as odds ratios (ORs) and 95% confidence interval (CI).

Compared to the participants of NTNW, those in the HTGW group have a higher risk of CKD-related vascular complications; the OR of decreased GFR and elevated UACR were 2.21 (95% CI:1.32–3.82) and 2.18 (95% CI:1.69–2.81), respectively. Moreover, compared to the participants in the NTNW group, the ORs of decreased ABI and elevated baPWV were 2.24 (95% CI:1.38–3.80) and 1.63 (95% CI:1.28–2.06) in the HTGW group. However, there was no association between DR and HWP.

Stratified Analysis

We further examined the association between vascular complication index in subgroups of age (<60 years, \geq 60 years), sex, HbA1c (<7.0%, \geq 7.0%), history of hypertension, and education (less than high school, high school education or higher level), the results were shown in forest plots (Figure 2).

In groups of HbA1c<7.0%, both the 95% CI of OR of deceased ABI and GFR crossed the invalid line (P>0.05), and the OR was 0.95 (95% CI:0.37–2.48) and 2.38 (95% CI:0.95–6.45), respectively. The stratified analysis results for each vascular complication index show differences in patients with a history of hypertension or not, except for UACR.

Discussion

In this study, we found that HWP was associated with vascular complications of diabetes: Compared with the participants of NTNW, those patients of HTGW have a higher risk of CKD-related vascular complications (elevated UACR and decreased GFR) and diabetic macrovascular complications (elevated baPWV and decreased ABI). This implies that HWP could be a screening tool for diabetic vascular complications and may help to reduce the incidence of serious diabetic vascular complications.

Several studies suggested that HWP was associated with coronary artery disease and metabolic syndrome in community crowds. Compared with Isabelle's study, our results adjusted for HbA1c in diabetes patients, demonstrating the consistency of the conclusions in the insulin-resistant states. In addition, the present study focuses on

Table	Demographic	Information and	l Clinical	Character	of the	Present Study
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	Total	NTNW	ETNW	NTEW	HTGW	P*
n	3221	660	275	1209	1077	-
Males, n (%)	1914 (59.4)	374 (56.7)	174 (63.3)	673 (55.7)	693 (64.3)	<0.001
Age, years, mean (SD)	50.00 (12.27)	51.53 (11.92)	47.19 (10.90)	52.52 (12.00)	46.94 (12.32)	<0.001
Course, months, median	47.00	61.00	50.00	61.00	34.00	<0.001
(IQR)	(3.00, 125.00)	(10.00,135.25)	(3.00,115.00)	(5.00,136.00)	(1.00, 98.00)	
Height, cm, mean (SD)	166.53 (8.62)	164.68 (8.34)	165.64 (8.05)	166.12 (8.61)	168.36 (8.61)	<0.001
Weight, kg, mean (SD)	74.57 (14.52)	62.22 (8.74)	65.97 (8.73)	76.66 (13.12)	82.00 (14.10)	<0.001
BMI, kg/m2, mean (SD)	26.76 (4.03)	22.90 (2.38)	23.98 (2.15)	27.68 (3.60)	28.81 (3.64)	<0.001
Waist, cm, mean (SD)	94.28 (10.54)	82.04 (5.64)	84.35 (4.53)	97.95 (7.68)	100.21 (8.61)	<0.001
High school education, n (%)	1947 (60.6)	388 (59.1)	185 (67.3)	650 (53.9)	724 (67.3)	<0.001
Hypertension, n (%)	1104 (34.3)	145 (22.0)	75 (27.3)	477 (39.5)	407 (37.8)	<0.001
Drinking, n (%)	1327 (41.2)	233 (35.4)	125 (45.5)	461 (38.2)	508 (47.2)	<0.001
Smoking, n (%)	1093 (33.9)	192 (29.1)	107 (38.9)	377 (31.2)	417 (38.7)	<0.001
LDL-c mmol/l, mean (SD)	3.24 (0.91)	3.02 (0.83)	3.55 (0.93)	3.05 (0.84)	3.50 (0.94)	<0.001
HDL-c, mmol/l, mean (SD)	1.20 (0.29)	1.31 (0.32)	1.15 (0.26)	1.22 (0.28)	1.11 (0.27)	<0.001
Fasting glucose, mmol/L,	9.50 (4.05)	8.93 (4.20)	10.15 (4.19)	9.06 (3.79)	10.18 (4.07)	<0.001
mean (SD)						
Triglyceride, mmol/L,	1.51 (1.05, 2.32)	0.96 (0.75, 1.28)	2.51 (2.00, 3.71)	1.19 (0.94, 1.43)	2.57 (2.05, 3.68)	<0.001
median (IQR)						
HbAlc,%, mean (SD)	8.48 (2.30)	8.24 (2.53)	8.97 (2.40)	8.32 (2.16)	8.68 (2.25)	<0.001
Decreased GFR, n (%)	177 (5.5)	23 (3.5)	16 (5.8)	66 (5.5)	72 (6.7)	0.006*
Decreased ABI, n (%)	188 (5.8)	21 (3.2)	12 (4.4)	61 (5.0)	94 (8.7)	<0.001*
Elevated baPWV, n (%)	2065 (64.1)	381 (57.7)	162 (58.9)	840 (69.5)	682 (63.3)	0.003*
Elevated UACR, n (%)	914 (28.4)	119 (18.0)	100 (36.4)	312 (25.8)	383 (35.6)	<0.001*
DR, n (%)	125 (3.9)	22 (3.3)	17 (6.2)	50 (4.1)	36 (3.3)	0.685*

Note: *P for trend was calculated by Cochran Armitage Test.

Abbreviations: BMI, Body mass index; HDL-C, High-density lipoprotein cholesterol; LDL-C, Low-density lipoprotein cholesterol; TG, Triglyceride; HbA1c, Glycated hemoglobin; NTNW, Normal Triglyceride and Waist; ETNW, Elevated Triglyceride and normal Waist; NTEW, Normal Triglyceride and enlarged Waist; HTGW, Hypertriglyceridemic-waist; DR, Diabetic retinopathy; UACR, Urinary Albumin Creatinine Ratio; GFR, Glomerular filtration rate; ABI, ankle brachial index; baPWV, Brachial-ankle pulse wave velocity.

analyzing the association between arterial stiffness index (ABI and baPWV) and HWP.¹⁰ The result of CKD-related vascular complications was consistent with Yan's study.²⁵ However, we did not find an association between HWP and DR.

Although the underlying mechanism of HWP with the vascular complication of diabetes has not been fully understood, there are several possible explanations. HWP is a particular biomarker that comprises both enlarged waist circumference and elevated TG.⁸ First, Elevated triglyceride levels cause heparin to activate lipoprotein lipase and increase intravascular lipolysis of circulating triglycerides, thereby increasing tissue exposure to free fatty acids, ultimately leading to insulin resistance.¹³ Second, the sedentary lifestyle contributes to the formation of the HTGW state, and the sedentary life was associated with atherosis in diabetes. The mechanism between HWP and vascular complication may be similar to the association between VAI and CVD, while both VAI and HWP are metabolic indicators.^{26,27} Visceral fat cells stimulate the secretion of inflammatory factors such as interleukin-6 (IL-6), tumor necrosis factor α (TNF- α), and adipokines, which lead to the production of reactive oxygen in arterioles.²⁵ Therefore, pro-inflammatory cytokines and reactive oxygen species produced by obesity can directly affect endothelial cells, leading to endothelial dysfunction.²⁸

In our study, there was an age difference in the result between HWP and vascular complications of diabetes. This phenomenon may be contributing to the prevalence of HWP in different age groups.²⁹ A similar result was also seen in Huelgas's study.¹¹ Elderly population has a lower prevalence of HTGW, and the survival bias should be considered. Moreover, the association between HWP and vascular complications of diabetes was more significant in the population

	Model	I	Model 2		
	OR	Р	OR	Р	
Decreased ABI					
NTNW	Reference	_	Reference	-	
ETNW	1.32[0.62, 2.69]	0.460	1.13 [0.53, 2.33]	0.740	
NTEW	1.57 [0.96, 2.66]	0.082	1.49 [0.91, 2.55]	0.124	
HTGW	2.68 [1.68, 4.48]	<0.001	2.24 [1.38, 3.80]	0.002	
Decreased GFR					
NTNW	Reference	-	Reference	-	
ETNW	2.64 [1.32, 5.21]	0.005	2.25 [1.10, 4.54]	0.024	
NTEW	1.63 [1.00, 2.73]	0.056	1.38 [0.84, 2.34]	0.222	
HTGW	2.78 [1.71, 4.70]	<0.001	2.21 [1.32, 3.82]	0.003	
Elevated UACR					
NTNW	Reference	-	Reference	-	
ETNW	2.52 [1.82, 3.48]	<0.001	2.34 [1.68, 3.26]	<0.001	
NTEW	1.52 [1.20, 1.94]	0.001	1.33 [1.04, 1.71]	0.023	
HTGW	2.53 [1.99, 3.23]	0.001	2.18 [1.69, 2.81]	<0.001	
Elevated baPWV					
NTNW	Reference	-	Reference	-	
ETNW	1.39 [1.02, 1.90]	0.038	1.37 [0.99, 1.89]	0.056	
NTEW	1.69 [1.36, 2.11]	<0.001	1.49 [1.19, 1.88]	0.001	
HTGW	1.73 [1.39, 2.16]	<0.001	1.63 [1.28, 2.06]	<0.001	
DR					
NTNW	Reference	-	Reference	-	
ETNW	1.96 [0.99, 3.82]	0.048	2.02 [1.00, 4.02]	0.046	
NTEW	1.25 [0.75, 2.15]	0.402	1.19 [0.70, 2.06]	0.533	
HTGW	1.07 [0.62, 1.90]	0.807	1.04 [0.58, 1.90]	0.889	

Table 2Logistic Regression Analysis Assessing the Associations Between theHypertriglyceridemic-Waist Phenotype and Vascular Complication Index of Diabetes

Notes: Model I: adjust for sex, duration, HbA1c. Molde 2: adjust for history of hypertension, smoking, drinking, education, LDL, HDL, and plus the covariable in model I.

Abbreviations: NTNW, Normal Triglyceride and Waist; ETNW, Elevated Triglyceride and normal Waist; NTEW, Normal Triglyceride and enlarged Waist; HTGW, Hypertriglyceridemic-waist; DR, Diabetic retinopathy; GFR, Glomerular filtration rate; ABI, ankle brachial index; baPWV, Brachial-ankle pulse wave velocity.

of high academic level, which may be illustrated by a sedentary lifestyle in these people. In addition, the stratified analysis results for ABI and baPWA show differences in patients with a history of hypertension or not, which may contribute to the interaction between d blood pressure and arterial stiffness.³⁰

This study also has several limitations. Firstly, due to the cross-sectional study design, we cannot obtain an accurate causal relationship and cannot rule out reverse causality. More cohort studies should be designed to clarify the association between HWP and vascular complications in type 2 diabetes. In future research, we will analyze the association of HWP and vascular complications in a cohort using causal inference methods. Moreover, although a series of sensitivity analyses adjusted the result of the association between HWP and vascular complications, residual confounders still existed, such as the usage of hormones and other drugs. Last, we could not include the patients with type 1 diabetes and gestational diabetes mellitus who have younger ages and morbid states, which should be a reappraisal of the association between HWP and vascular complications.

Therefore, this study found an association between HWP and vascular complications of diabetes. All these results suggest that HWP is a cost-effective screening tool that can be used in clinical practice and health management. Patients of T2DM should monitor the level and trajectory of HWP, which may help to reduce the incidence of serious diabetic vascular complications. For medical researchers, more simple, accurate, and reliable biomarkers should be developed to assess the risk of chronic disease complications.

А				В			
Item	OR (95%CI)	Р		Item	OR (95%CI)	Р	
Age		·····		Age		•••••	
<60 vear	3.85 [1.97, 8.45]	<0.001		<60 year	2.17 [1.01, 5.24]	0.061	
>=60 year	1.12 [0.48, 2.67]	0.786		>=60 year	2.31 [1.14, 4.82]	0.022	
Sex	1.12 [0.40, 2.07]	0.700		Sex	2.01 [1.11, 1.02]	0.0LL	
Male	3.62 [1.44, 11.04]	0.011		Male	0.56 [0.18, 1.82]	0.317	
Female	1.97 [1.11, 3.67]	0.025		Female	3.14 [1.74, 5.95]	<0.001	
HbA1c	1.07 [1.11, 0.07]	0.020		HbA1c	0.14[1.14, 0.00]	-0.001	
<7.0 %	0.95 [0.37, 2.48]	0.917		<7.0 %	2.38 [0.95, 6.45]	0.074	
>=7.0 %	3.23 [1.74, 6.58]	<0.001		>=7.0 %	2.21 [1.18, 4.35]	0.014	
Hypertension	5.25 [1.74, 0.50]	~0.001		Hypertension	2.21 [1.10, 4.55]	0.010	
No	2.80 [1.53, 5.49]	0.001		No	4.15 [1.91, 9.57]	0.001	
Yes	1.45 [0.65, 3.59]	0.383		Yes	1.52 [0.75, 3.32]	0.001	
Education	1.45 [0.05, 5.59]	0.303			1.52 [0.75, 3.32]	0.27	
	4 70 10 00 0 001	0.158		Education	4 00 10 00 0 001	0 470	
Less then high school	1.73 [0.83, 3.86]	0.156		Less then high school	1.62 [0.82, 3.32]	0.173	· · · ·
High school or more	2.74 [1.42, 5.83]	0.005		High school or more	3.29 [1.48, 8.22]	0.006	
C			0.5 1 1.5 2 2.5 3				0 0.5 1 1.5 2 2.5
C	OR (95%CI)	P	0.5 1 1.5 2 2.5 3	D	OR (95%CI)	P	0 0.5 1 1.5 2 2.5
ltem	OR (95%Cl)	P		Item	OR (95%Cl)	P.	0 0.5 1 1.5 2 2.5
ltem Age				ltem Age			
Item Age <60 year	2.73 [2.01, 3.75]	<0.001		Item Age <60 year	1.67 [1.30, 2.16]	<0.001	
Item Age <60 year >=60 year				Item Age <60 year >=60 year			
Item Age <60 year >=60 year Sex	2.73 [2.01, 3.75] 1.16 [0.72, 1.86]	<0.001 0.542		Item Age <60 year >=60 year Sex	1.67 [1.30, 2.16] 1.57 [0.73, 3.52]	<0.001 0.258	
ltem Age <60 year >=60 year Sex Male	2.73 [2.01, 3.75] 1.16 [0.72, 1.86] 1.58 [1.07, 2.35]	<0.001 0.542 0.022		Item Age <60 year >=60 year Sex Male	1.67 [1.30, 2.16] 1.57 [0.73, 3.52] 1.39 [0.95, 2.06]	<0.001 0.258 0.094	
Item Age <60 year >=60 year Sex Male Female	2.73 [2.01, 3.75] 1.16 [0.72, 1.86]	<0.001 0.542		Item Age <60 year >=60 year Sex Male Female	1.67 [1.30, 2.16] 1.57 [0.73, 3.52]	<0.001 0.258	
Item Age <60 year >=60 year Sex Male Female HbA1c	2.73 [2.01, 3.75] 1.16 [0.72, 1.86] 1.58 [1.07, 2.35] 2.68 [1.92, 3.79]	<0.001 0.542 0.022 <0.001		Item Age <60 year >=60 year Sex Male Female HbA1c	1.67 [1.30, 2.16] 1.57 [0.73, 3.52] 1.39 [0.95, 2.06] 1.73 [1.28, 2.36]	<0.001 0.258 0.094 <0.001	
Item Age <60 year >=60 year Sex Male Female HbA1c <7.0 %	2.73 [2.01, 3.75] 1.16 [0.72, 1.86] 1.58 [1.07, 2.35] 2.68 [1.92, 3.79] 1.96 [1.11, 3.53]	<0.001 0.542 0.022 <0.001 0.023		Item Age <60 year	1.67 [1.30, 2.16] 1.57 [0.73, 3.52] 1.39 [0.95, 2.06] 1.73 [1.28, 2.36] 1.74 [1.10, 2.76]	<0.001 0.258 0.094 <0.001 0.018	
Item Age <60 year	2.73 [2.01, 3.75] 1.16 [0.72, 1.86] 1.58 [1.07, 2.35] 2.68 [1.92, 3.79]	<0.001 0.542 0.022 <0.001		Item Age <60 year	1.67 [1.30, 2.16] 1.57 [0.73, 3.52] 1.39 [0.95, 2.06] 1.73 [1.28, 2.36]	<0.001 0.258 0.094 <0.001	
Item Age <60 year	2.73 [2.01, 3.75] 1.16 [0.72, 1.86] 1.58 [1.07, 2.35] 2.68 [1.92, 3.79] 1.96 [1.11, 3.53] 2.18 [1.65, 2.91]	<0.001 0.542 0.022 <0.001 0.023 <0.001		Item Age <60 year	1.67 [1.30, 2.16] 1.57 [0.73, 3.52] 1.39 [0.95, 2.06] 1.73 [1.28, 2.36] 1.74 [1.10, 2.76] 1.56 [1.18, 2.07]	<0.001 0.258 0.094 <0.001 0.018 0.002	
Item Age <60 year	2.73 [2.01, 3.75] 1.16 [0.72, 1.86] 1.58 [1.07, 2.35] 2.68 [1.92, 3.79] 1.96 [1.11, 3.53] 2.18 [1.65, 2.91] 2.17 [1.59, 3.00]	<0.001 0.542 0.022 <0.001 0.023 <0.001 <0.001		Item Age <60 year	1.67 [1.30, 2.16] 1.57 [0.73, 3.52] 1.39 [0.95, 2.06] 1.73 [1.28, 2.36] 1.74 [1.10, 2.76] 1.56 [1.18, 2.07] 1.84 [1.40, 2.42]	<0.001 0.258 0.094 <0.001 0.018 0.002 <0.001	
Item Age <60 year	2.73 [2.01, 3.75] 1.16 [0.72, 1.86] 1.58 [1.07, 2.35] 2.68 [1.92, 3.79] 1.96 [1.11, 3.53] 2.18 [1.65, 2.91]	<0.001 0.542 0.022 <0.001 0.023 <0.001		Item Age <60 year	1.67 [1.30, 2.16] 1.57 [0.73, 3.52] 1.39 [0.95, 2.06] 1.73 [1.28, 2.36] 1.74 [1.10, 2.76] 1.56 [1.18, 2.07]	<0.001 0.258 0.094 <0.001 0.018 0.002	
Item Age <60 year	2.73 [2.01, 3.75] 1.16 [0.72, 1.86] 1.58 [1.07, 2.35] 2.68 [1.92, 3.79] 1.96 [1.11, 3.53] 2.18 [1.65, 2.91] 2.17 [1.59, 3.00] 2.21 [1.44, 3.43]	<0.001 0.542 0.022 <0.001 0.023 <0.001 <0.001 <0.001		Item Age <60 year	1.67 [1.30, 2.16] 1.57 [0.73, 3.52] 1.39 [0.95, 2.06] 1.73 [1.28, 2.36] 1.74 [1.10, 2.76] 1.56 [1.18, 2.07] 1.84 [1.40, 2.42] 0.76 [0.41, 1.36]	<0.001 0.258 0.094 <0.001 0.018 0.002 <0.001 0.374	
Item Age <60 year	2.73 [2.01, 3.75] 1.16 [0.72, 1.86] 1.58 [1.07, 2.35] 2.68 [1.92, 3.79] 1.96 [1.11, 3.53] 2.18 [1.65, 2.91] 2.17 [1.59, 3.00]	<0.001 0.542 0.022 <0.001 0.023 <0.001 <0.001		Item Age <60 year	1.67 [1.30, 2.16] 1.57 [0.73, 3.52] 1.39 [0.95, 2.06] 1.73 [1.28, 2.36] 1.74 [1.10, 2.76] 1.56 [1.18, 2.07] 1.84 [1.40, 2.42]	<0.001 0.258 0.094 <0.001 0.018 0.002 <0.001	

Figure 2 Stratified analysis of associations between the hypertriglyceridemic-waist phenotype and vascular complication index of diabetes. (A) ABI (B) GFR (C) UACR (D) baPWV.

Abbreviations: DR, Diabetic retinopathy; GFR, Glomerular filtration rate; ABI, ankle-brachial index; baPWV, Brachial-ankle pulse wave velocity.

Conclusion

In summary, there was an association between HWP and diabetic vascular complications in the Chinese population.

Data Sharing Statement

The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

Ethics Approval and Consent to Participate

The Medical Ethics Committee of Beijing Luhe Hospital, Capital Medical University approved the study protocol. This study was performed by the Declaration of Helsinki, and all participants provided written informed consent.

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Disclosure

The authors report no conflicts of interest in this work.

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