

Lipid Accumulation Product is Associated with Urinary Albumin-creatinine Ratio in Chinese Prediabetic Population: A Report from the REACTION Study

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Background: Lipid accumulation product (LAP) as a powerful marker of visceral obesity is an independent risk factor of chronic kidney disease. The present study attempted to explore the association between LAP and albuminuria in prediabetic individuals.

Methods: We conducted a cross-sectional study and enrolled 26,529 participants with prediabetes over 40 years old with prediabetes from seven provinces in China. LAP was calculated from waist circumference and fasting triglycerides. Elevated albuminuria was defined by urinary albumin-creatinine ratio (uACR) ≥ 30 mg/g. Propensity score matching was applied to reduce bias, comparison between LAP and other traditional visceral obesity indices was performed and multiple logistic regression models were conducted to assess the association between LAP and albuminuria in the prediabetic population.

Results: Individuals with uACR ≥ 30 mg/g were older and had higher BP, BMI, WC, TG, fasting insulin, glycohemoglobin and LAP, as well as lower eGFR and HDL level. Multiple logistic regression analysis showed elevated LAP was associated with increased odds of albuminuria (OR [95%CI]_{Q2 vs Q1} 1.09 [0.94, 1.27], OR [95%CI]_{Q3 vs Q1} 1.13 [0.97, 1.31], OR [95%CI]_{Q4 vs Q1} 1.42 [1.21, 1.67], *P* for trend=0.018), and superior over waist-to-hip ratio or waist-to-height ratio. Stratification indicated that the prediabetic population with higher LAP level and characterized by female gender, middle age, being overweight, and rise in blood pressure were more likely to have increased uACR.

Conclusion: Elevated level of LAP was associated with increased albuminuria in the prediabetic population in China.

Keywords: albuminuria, lipid accumulation product, prediabetes, visceral obesity

Plain Language Summary

1. This is the first multicenter, large-sampled cross-sectional study of association between Lipid accumulation product (LAP) and urinary albumin to creatinine ratio (uACR) in the Chinese prediabetic population.

2. LAP is positively correlated with increased uACR. Prediabetic individuals with higher LAP level and characterized by female gender, middle age, being overweight and rise in blood pressure were more likely to have increased uACR.

Introduction

Data from the International Diabetes Federal Diabetes Atlas 2019 indicated that prediabetes is an increasingly common global condition as 1 in 13 adults

have impaired glucose tolerance.¹ The estimated standardized prevalence of prediabetes among the Chinese adult population was 35.7%.² Compared with nondiabetic population, individuals with prediabetes had a 15% higher risk of prevalent chronic kidney disease (CKD) (OR: 1.15, 95%CI: 1.02–1.32) according to the REACTION study.³ Besides, dramatically increased prevalence of obesity also substantially affected the patterns of CKD. Adjusted hazard ratios for declined estimated glomerular filtration rate (eGFR) were 2.02 (95%CI: 1.80–2.27) with body mass indices (BMI) 40 kg/m² comparing to BMI of 25 kg/m².⁴ A recent UK Biobank study that enrolled 0.4 million people proved that each 5 kg/m² ascending in BMI was associated with a 47% (46–49%) increase in the odds of higher urinary albumin-to-creatinine ratio (uACR) category.⁵

To further evaluate obesity-associated risks, an alternative continuous index of lipid over accumulation, the lipid accumulation product (LAP), was calculated from waist circumference (WC, cm) and fasting triglycerides (TGs) (mmol/L): (WC: 65)×TG (men) and (WC: 58)×TG (women), initially for better identifying visceral obesity and insulin resistance, elevated glucose, and diabetes than BMI did.⁶ Recent studies found that LAP also acquitted itself well in predicting cardiovascular risks^{7,8} and identifying CKD^{9,10} in the healthy population. However, evidence in association between LAP and renal dysfunction in the prediabetic population is lacking, thus the aim of our study was to explore the association between LAP and albuminuria in the prediabetic population.

Methods

Participants and Study Design

Cross-sectional data from seven provinces across China was collected in this study, as a part of the ongoing risk evaluation of cancers in Chinese diabetic individuals a longitudinal (REACTION) study. Baseline assessments included a standardized questionnaire supplemented by interviews, standardized physical and functional measurements and the collection of biological samples. A total of 47,807 participants over 40 years old were recruited from May to December 2011. The protocol was approved by the Clinical Research Ethics Review Board of Rui-Jin Hospital (No. 2014–25), which was consistent with the Declaration of Helsinki. All participants gave written informed consent.

Participants with self-reported diabetes, kidney diseases, or undergoing lipid-lowering, antihypertensive (including angiotensin converting enzyme inhibitors or angiotensin receptor blocker) or hypoglycemic treatment, and those without data on adiposity measures, blood pressure, glycosylated hemoglobin (HbA1c), fasting blood glucose (FBG), 2 h postload blood glucose (PBG), urinary albumin or urinary creatinine were excluded. In total 26,529 participants with prediabetes were enrolled. Further propensity matching was used to reduce potential bias. Eventually, 16,631 eligible participants were ultimately included in the final analysis (Figure 1).

Data Collection and Measurements

The REACTION study consists of a standardized interview, followed by a physical examination and blood and urine collection at authorized examination centers. Self-reported information on demographics (age, gender), health conditions (diagnosed hypertension, cardiovascular diseases, and family history of diabetes), smoking and drinking habits were obtained during the interview. The prescription medication information was recorded by the interviewer of the medication names from the package provided by the participant. Height and weight were measured during the examination. BMI was computed by dividing weight in kilograms by height in meters squared (kg/m²). Waist circumference (WC) and hip circumference (HC) were measured in centimeters. LAP, waist-to-hip ratio (WHR) and waist-to-height (WhtR) Ratio were computed. Blood pressure (BP) was measured three times with intervals of one minute in a seated position after a resting period of five minutes.

After 8–12 h overnight fasting, venous blood samples were collected followed by second venipuncture sampling in 2 h after oral glucose tolerance test (OGTT): consumption of a 300 mL glucose solution containing 75 g glucose within 5 min. Fasting insulin was tested via photochemiluminescence; FBG, PBG, HbA1c, high-density lipoprotein cholesterol (HDL), low-density lipoprotein cholesterol (LDL), total cholesterol (TC), TG, alanine transferase (ALT), aspartate transferase (AST) and serum creatinine were measured via enzymatic methods.

Participants answering “no” to self-report diagnosed diabetes underwent OGTT. Prediabetes is defined as having either impaired glucose tolerance (IGT) with 2-h postload plasma glucose (PPG) ≥ 7.8 mmol/L but < 11.1 mmol/L and/or impaired fasting glucose (IFG) with fasting plasma glucose (FPG) ≥ 6.0 mmol/L but < 7.0

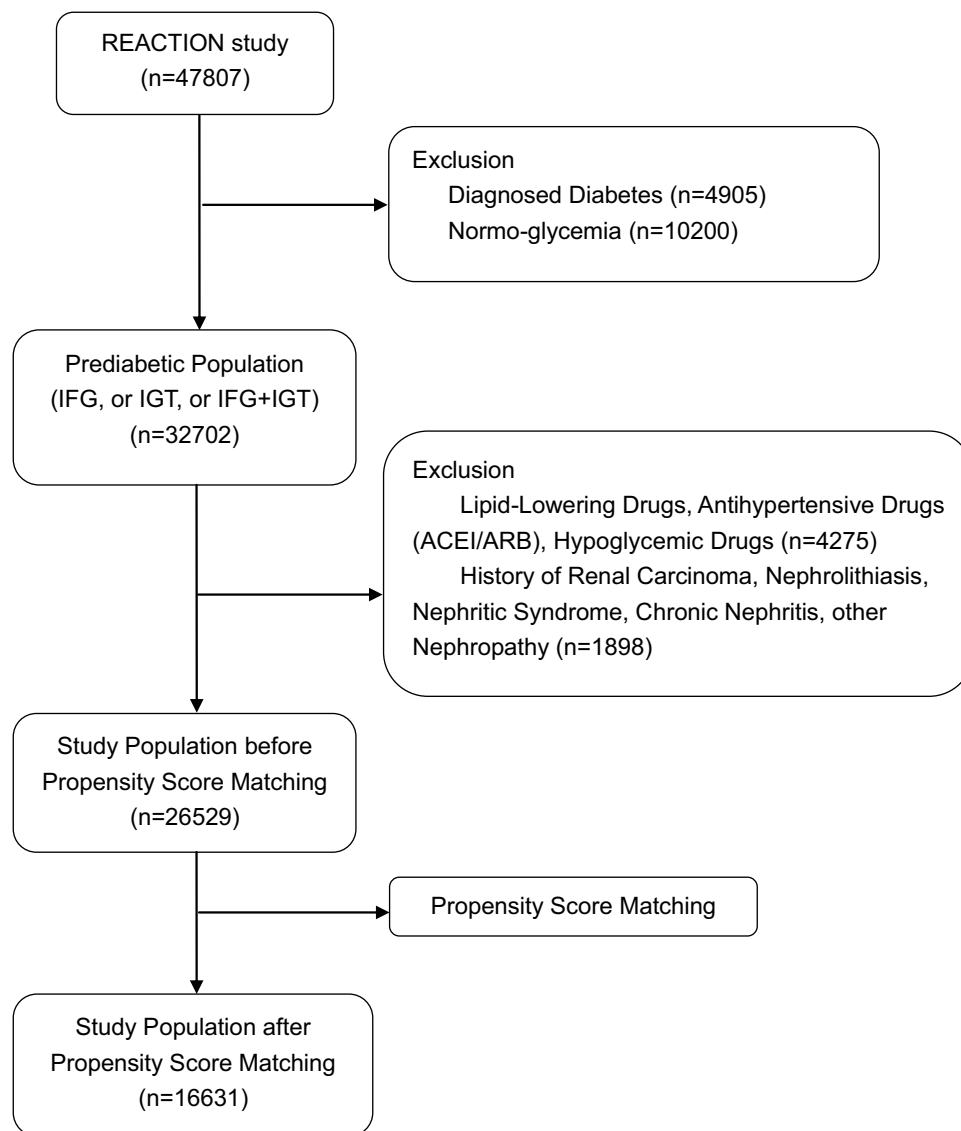


Figure 1 Flow chart of study population.

mmol/L.¹¹ Prediabetes was defined as IFG in absence of IGT or a combination of both.¹²

The Chronic Kidney Disease Epidemiology Collaboration formula fitting Asian females was applied to calculate eGFR (mL/min/1.73 m²).¹³ Random spot urine samples were obtained, and urine albumin and creatinine were measured using frozen specimens. Urine albumin was measured using solid-phase fluorescence immunoassay; urine creatinine was measured using the modified Jaffe kinetic method in the same laboratory. The uACRs were calculated using formula as: urinary albumin (mg)/urinary creatinine (g). Each authorized examination center used the same range and units of measurement. The uACR was divided into two groups: normo-albuminuria: uACR

<30 mg/g and increased albuminuria: uACR ≥30 mg/g, indicating kidney damage according to KDIGO-CKD guidelines.¹⁴ LAP was divided into quartiles for the analysis.

Based on daily cigarette consumption, smokers were classified into regular smokers (at least one cigarette per day) and occasional smokers (less than one cigarette per day or seven cigarettes per week). Based on drinking frequency, regular drinkers were defined as drinking at least once a week for over six months and occasional drinkers drank no more than once a week.

Statistical Analysis

Statistical analysis was performed via Empower (R) version 2.2 (www.Empowerstats.com, X&Y Solutions Inc.,

Boston, MA, USA), R (<http://www.Rproject.org>) and SPSS version 26.0 (IBM Corp., Armonk, NY, USA). Propensity score matching was applied to control the bias of baseline characteristics between the two groups of uACRs. Matching was performed using a 1:4 ratio protocol to match all covariates, with a calliper width equal to 0.05 of the standard deviation of the logit of the propensity score. Continuous variables were presented as mean \pm SD or median (IQR) for skewed variables. Categorical variables are presented as percentages (%). The Kruskal–Wallis test was applied for comparison of continuous variables between the uACR groups; chi-squared test was used to analyze categorical variable. Logistic regression analysis with unadjusted and multivariate-adjusted models was conducted to determine the association between LAP quartiles and the risk of increased albuminuria. Conventional risk factors were adjusted. Confounding factors that changed the matched ORs by at least 10% were also selected for adjustment. Model 1 was unadjusted. Model 2 was adjusted for age, sex and BMI. Model 3 was additionally adjusted for smoking, drinking, family history of diabetes, and CVD history, based on Model 2. Model 4 was further adjusted for SBP, DBP, HDL, FBG, PBG, and HbA1c based on Model 3. In all models, OR (95%CI) were calculated. To further examine the association between LAP quartiles and the risk of increased albuminuria, stratified analyses were performed with gender, age, BP (normal: systolic blood pressure (SBP) <120 and diastolic blood pressure (DBP) <80 mmHg, borderline BP: SBP: 120–139 and/or DBP 80–89 mmHg, hypertension: SBP \geq 140 or DBP \geq 90 mmHg); BMI: underweight (<18.5 kg/m²), normal weight (18.5–23.9 kg/m²), overweight (24–27.9 kg/m²), and obesity (\geq 28.0 kg/m²).¹⁵ Potential interactions between LAP quartiles and strata variables were assessed in the logistic regression analysis after fully adjustment. *P*-values <0.05 (two-sided) were considered statistically significant.

Results

Selected Characteristics by uACR Category

After propensity score matching, individuals with uACR \geq 30 mg/g were older and had higher BP, BMI, WC, TG, fasting insulin, glycohemoglobin and LAP, as well as lower eGFR and HDL level than those with normo-albuminuria (*P* \leq 0.001) (Table 1). The percentage of participants with extreme obesity (defined as BMI \geq 30 kg/m²) in uACR

<30 mg/g group was 2.01% (n=281), and 5.64% (n=149) in uACR \geq 30 mg/g group (Table S1). A slight but significant difference of both WHR (*P*=0.040) and WHtR (*P*=0.035) between uACRs groups was observed. In pairwise comparisons, those with increased albuminuria were more likely to be female (*P* \leq 0.001). Participants with increased albuminuria were more likely to be smokers or drinkers. The prevalence of CVD events was greater in individuals with increased uACR (2.73% vs 1.39%, *P* \leq 0.001). However, neither TC nor LDL presented significant difference among groups. Equilibrium was also seen in the prevalence of stroke (0.45% vs 0.64%, *P*=0.253) and myocardial infarction (0.15% vs 0.15%, *P*=0.988) in pairwise comparisons.

Comparison of Central Adiposity Associated with Increased uACR

Multiple logistic regression models for odds of increased uACR with central adiposity indices including WHR, WHtR and LAP (Table S2) were constructed in the total population in unadjusted and fully adjusted models. Each index, both in its quartile and continuous form was significantly associated with increased uACR in the nonadjusted model. However, after further adjustments, only the fourth quartiles of LAP (OR: 1.42, 95%CI: 1.21–1.67, *P*<0.001) and the fourth quartile of WHtR (OR: 1.36, 95%CI: 1.18–1.57, *P*<0.001) remained significantly associated with increased uACR, whereas WHR was not. The association between uACR \geq 30 mg/g and continuous values of the WHR, WHtR, and LAP was consistent with the finding delivered by categorical values: each unit elevation of LAP was associated with 34% higher odds of increased uACR (OR: 1.34; 95%CI: 1.00–1.68, *P*<0.001). No effect of interaction in WHR or WHtR with uACR was observed after adjustment (also Table S2). Compared to those classic adiposity indices, LAP was more likely to be closely associated with higher odds of increased uACR.

Associations Between Lipid Parameters and uACR

As a component of LAP, TG and other lipid parameters (TC, HDL-C and LDL-C) were further analyzed for their association with increased uACR after adjusting for multiple confounders in order to learn which lipid parameter was more closely related to increased albuminuria. Compared with TC, HDL-C, and LDL-C, although attenuated after full adjustment in Model 4, only the TG \geq 2.3 mmol/L category was significantly associated with higher

Table 1 Characteristics of Study Population by uACR Category After Propensity Score Matching

Variables	uACR <30 mg/g	uACR ≥30 mg/g	P-value
n	13,987	2644	
Age, years	54.86±8.84	59.41±9.88	<0.001
Men, %	4592 (32.83%)	553 (20.92%)	<0.001
BMI, kg/m ²	22.99±3.25	24.18±3.72	<0.001
SBP, mmHg	121.00 (110.00–133.00)	130.00 (116.00–146.00)	<0.001
DBP, mmHg	74.00 (68.00–81.00)	77.00 (70.00–85.00)	<0.001
TC, mmol/L	4.83±1.09	4.88±1.13	0.051
TG, mmol/L	1.44±1.02	1.65±1.14	<0.001
HDL, mmol/L	1.35±0.36	1.30±0.33	<0.001
LDL, mmol/L	2.79±0.85	2.80±0.87	0.671
FBG, mmol/L	5.76±0.59	5.78±0.59	0.031
PBG, mmol/L	8.59±1.32	8.72±1.25	0.014
HbA1c, %	5.93±0.44	6.01±0.48	<0.001
Fasting insulin, mU/L	7.60 (5.50–10.30)	8.10 (5.80–11.20)	<0.001
ALT, U/L	14.00 (10.00–20.00)	14.00 (11.00–20.00)	0.051
AST, U/L	20.00 (16.00–24.00)	21.00 (17.00–25.00)	0.022
GGT, U/L	18.00 (13.00–27.00)	19.00 (14.00–29.00)	0.024
eGFR, mL/min/1.73m ²	118.46 (106.04–132.31)	113.38 (100.04–127.47)	<0.001
WC, cm	82.00 (75.00–88.00)	84.05 (78.00–92.00)	<0.001
WHR	0.87 (0.82–0.91)	0.88 (0.83–0.92)	0.040
WHtR	0.51 (0.47–0.54)	0.53 (0.49–0.58)	0.035
LAP	25.02 (14.36–41.72)	34.02 (19.80–55.67)	<0.001
MI history, %			0.988
Yes	21 (0.15%)	5 (0.15%)	
No	13,966 (99.85%)	2639 (99.85%)	
Stroke history, %			0.253
Yes	91 (0.64%)	13 (0.45%)	
No	13,896 (99.36%)	2631 (99.55%)	
CVD history, %			<0.001
Yes	196 (1.39%)	73 (2.73%)	
No	13,791 (98.61%)	2571 (97.27%)	
Smoking habits, %			<0.001
No	11,537 (82.47%)	2354 (89.05%)	
Occasional	356 (2.55%)	43 (1.64%)	
Regular	2094 (14.97%)	247 (9.31%)	
Drinking habits, %			<0.001
No	9850 (70.42%)	2116 (80.03%)	
Occasional	3096 (22.14%)	407 (15.40%)	
Regular	1041 (7.44%)	121 (4.57%)	

(Continued)

Table I (Continued).

Variables	uACR <30 mg/g	uACR ≥30 mg/g	P-value
DM family history, %			0.007
Yes	2173 (15.54%)	357 (13.49%)	
No	11,814 (84.46%)	2287 (86.51%)	
LAP quartiles			<0.001
Q1	3737 (26.72%)	412 (15.58%)	
Q2	3595 (25.70%)	561 (21.22%)	
Q3	3473 (24.83%)	695 (26.29%)	
Q4	3182 (22.75%)	976 (36.91%)	

Notes: Data expressed as mean ±SD for continuous variables or median (IQR) for skewed variables and percentage (%) for categorical variables.

Abbreviations: uACR, urinary albumin to creatinine ratio; BMI, body mass index; TC, total cholesterol; TG, triglyceride; HDL, high-density lipoprotein cholesterol; LDL, low-density lipoprotein cholesterol; FBG, fasting blood glucose; PBG, 2-h postload blood glucose; HbA1c, glycosylated hemoglobin; ALT, alanine transferase; AST, aspartate transferase; GGT, gamma-glutamyl transferase; eGFR, estimated glomerular filtration rate; WC, waist circumference; WHR, waist circumference to hip circumference ratio; WHtR, waist circumference to height ratio; LAP, lipid accumulation product index; MI, myocardial infarction; CVD, cardiovascular disease; DM, diabetes mellitus.

odds of increased uACR for both genders in all models (OR: 1.12, 95%CI: 1.03–1.51, $P=0.042$ in men; OR: 1.18, 95%CI: 1.00–1.35, $P=0.026$ in women; OR: 1.15, 95%CI: 1.01–1.31, $P=0.040$ in total population)(Table S3). HDL-C <1.0 mmol/L was significantly associated with increased uACR only in women in the fully adjusted model (OR: 1.14, 95%CI: 1.06–1.45, $P=0.026$) (also Table S3). There was no significant and independent association between TC or LDL-C and increased uACR.

Association of LAP Quartiles with Increased uACR

To investigate the stability of the association between LAP quartiles and increased uACR, multiple logistic regression analysis for the association between LAP quartiles and increased uACR in the prediabetic population were carried out (Table 2). Compared to participants with the first quartile of LAP, those with a higher level of LAP had higher odds for increased uACR among Models 1–3 (P for trend <0.001). After further adjustment for SBP, DBP, HDL, FBG, PBG, HbA1c, AST and GGT, the association was still significant in Model 4 (OR [95%CI]_{Q2 vs Q1} 1.09 [0.94, 1.27], OR [95%CI]_{Q3 vs Q1} 1.13 [0.97, 1.31], OR [95%CI]_{Q4 vs Q1} 1.42 [1.21, 1.67], P for trend=0.018).

Associations Between the LAP Index and uACR for Stratified Groups of Gender, Age, and BMI

Stratified analysis was performed to verify the stability of the relationship between LAP and uACR. In gender stratification (Table 3) (P for interaction=0.511), after full

adjustment for BMI, family history of DM, drinking and smoking habit, SBP, DBP, HDL, FBG, PBG, and HbA1c, prediabetic women with elevated LAP was associated with increased odds of uACR ≥30 mg/g (OR [95%CI]_{Q2 vs Q1} 1.12 [0.94, 1.33], OR [95%CI]_{Q3 vs Q1} 1.29 [1.08, 1.53], OR [95%CI]_{Q4 vs Q1} 1.48 [1.23, 1.79], P for trend <0.001); however, no significant association between LAP and uACR was observed in men (P for trend=0.052).

In age stratification (also Table 3) (P for interaction=0.395), middle aged (40–59 years old) participants with higher LAP, were more likely to have increased uACR compared to their lower counterpart (OR [95%CI]_{Q2 vs Q1} 1.06 [0.88, 1.28], OR [95%CI]_{Q3 vs Q1} 1.21 [1.01, 1.46], OR [95%CI]_{Q4 vs Q1} 1.47 [1.20, 1.79], P for trend <0.001); while in the older group (age ≥60 years old), the trend vanished.

Although significant associations of elevated LAP with increased uACR were observed in participants with all range of BP, a rise in BP indicating increased odds of increased uACR (Table 4). In participants with normal BP (SBP <120 and DBP <80 mmHg), elevated LAP was associated with increased uACR (OR [95%CI]_{Q2 vs Q1} 1.15 [0.94, 1.41], OR [95%CI]_{Q3 vs Q1} 1.27 [1.02, 1.57], OR [95%CI]_{Q4 vs Q1} 1.36 [1.06, 1.74], P for trend=0.010); furthermore, those high LAP level (Q4 quartile) participants with borderline BP (≤120 SBP <140 and/or ≤80 DBP <90 mmHg) or hypertension (SBP ≥140 or DBP ≥90 mmHg) were 1.45 (95%CI: 1.09–1.94, P for trend=0.048) or 2.08 (95%CI: 1.40–3.09, P for trend=0.001) times to have increased uACR compared to its lowest counterpart, indicating that ascending BP was related to increased albuminuria (P for interaction=0.012).

Table 2 Odds Ratio of uACR ≥ 30 mg/g by LAP Quartiles in Prediabetic Population

Variables	Odds Ratio (95%CI) of Increased uACR by LAPI				P-value for Trend
	LAPI Quartiles				
	Q1	Q2	Q3	Q4	
	≤ 15.11	15.12–26.40	26.41–43.94	≥ 43.95	
uACR category					
Model 1	1.00	1.42 (1.24–1.62)	1.82 (1.59–2.07)	2.78 (2.46–3.15)	<0.001
P-value		<0.001	<0.001	<0.001	
Model 2	1.00	1.12 (0.98–1.29)	1.21 (1.05–1.39)	1.56 (1.34–1.80)	<0.001
P-value		0.099	0.008	<0.001	
Model 3	1.00	1.12 (0.97–1.29)	1.20 (1.04–1.39)	1.55 (1.33–1.79)	<0.001
P-value		0.120	0.010	<0.001	
Model 4	1.00	1.09 (0.94–1.27)	1.13 (0.97–1.31)	1.42 (1.21–1.67)	0.018
P-value		0.238	0.111	<0.001	

Notes: Model 1 adjusted for: none. Model 2 adjusted for: age; sex; BMI. Model 3 adjusted for: age; sex; BMI; DM family history; drinking; smoking; CVD history. Model 4 adjusted for: age; sex; BMI; DM family history; drinking; smoking; CVD history; SBP; DBP; HDL; FBG; PBG; HbA1c.

Abbreviations: uACR, urinary albumin to creatinine ratio; eGFR, estimated glomerular filtration rate; LAP, lipid accumulation product index; BMI, body mass index; DM, diabetes mellitus; CVD, cardiovascular disease; HDL, high-density lipoprotein cholesterol; FBG, fasting blood glucose; PBG, 2-h postload blood glucose; HbA1c, glycosylated hemoglobin.

Table 3 Odds Ratio of uACR ≥ 30 mg/g by LAP Quartiles in Gender and Age Stratified Groups with Prediabetes

Variables	Odds Ratio (95%CI) of Increased uACR by LAPI LAPI Quartiles				P-value for Trend	P for Interaction
	Q1	Q2	Q3	Q4		
UACR ≥ 30 mg/g						
Gender						0.511
Men	1.00	1.03 (0.77–1.38)	0.69 (0.58–1.05)	1.31 (0.95–1.79)	0.052	
Women	1.00	1.12 (0.94–1.33)	1.29 (1.08–1.53)*	1.48 (1.23–1.79)*	<0.001	
Age, years						0.395
<60	1.00	1.06 (0.88–1.28)	1.21 (1.01–1.46)*	1.47 (1.20–1.79)*	<0.001	
≥ 60	1.00	1.02 (0.78–1.34)	0.89 (0.68–1.17)	1.09 (0.82–1.45)	0.227	

Notes: *Significance of Pvalue ($P < 0.05$). Adjusted for BMI; DM family; drinking; smoking; SBP; DBP; HDL; FBG; PBG; HbA1c.

Abbreviations: uACR, urinary albumin to creatinine ratio; eGFR, estimated glomerular filtration rate; LAP, lipid accumulation product index; eGFR BMI, body mass index; DM, diabetes mellitus; CVD, cardiovascular disease; HDL, high-density lipoprotein cholesterol; FBG, fasting blood glucose; PBG, 2-h postload blood glucose; HbA1c, glycosylated hemoglobin.

In accordance with BMI (P for interaction=0.256), participants who were overweight ($24 \leq \text{BMI} < 28 \text{ kg/m}^2$) were more likely to have increased uACR (OR [95%CI] $_{\text{Q2 vs Q1}}$ 2.15 [1.19, 3.87], OR [95%CI] $_{\text{Q3 vs Q1}}$ 1.89 [1.06, 3.36], OR [95%CI] $_{\text{Q4 vs Q1}}$ 2.34 [1.32, 4.16], P for trend <0.001). However, there was no significant relationship between LAP quartiles and uACR among participants with obesity, normal weight or underweight (also Table 4).

Discussion

Main Findings

In the present study, we found that an ascending level of LAP was significantly associated with increased uACR in

prediabetic women rather than men after adjustment for a wide spectrum of biochemical and lifestyle confounders. The multiple regression models adjustment for age, sex, BMI, DM family history, drinking and smoking habit, CVD history, BP, HDL, glucose, and HbA1c, witnessed an attenuated trend. Further stratification revealed that the prediabetic population with higher LAP level and characterized by middle age (40–59 years old) and overweight (BMI 24–27.9 kg/m^2) were more likely to have increased uACR. Besides, the incidence of albuminuria was positively correlated with elevated BP in the higher LAP group. To the best of our knowledge, the present study is the first multicenter, large sample clinical survey on the

Table 4 Odds Ratio of uACR \geq 30 mg/g by LAP Quartiles in Metabolic Stratified Groups with Prediabetes

Variables	Odds Ratio (95%CI) of Increased uACR by LAPI LAPI Quartiles				P-value for Trend	P-value for Interaction
	Q1	Q2	Q3	Q4		
uACR \geq30 mg/g						0.012
BP, mmHg						
SBP <120 and DBP <80	1.00	1.15 (0.94–1.41)	1.27 (1.02–1.57)*	1.36 (1.06–1.74)*	0.010	
\leq 120 SBP <140 and/or \leq 80 DBP <90	1.00	1.02 (0.77–1.35)	1.21 (0.92–1.60)	1.45 (1.09–1.94)*	0.048	
SBP \geq 140 or DBP \geq 90	1.00	1.44 (0.97–2.13)	1.29 (0.87–1.91)	2.08 (1.40–3.09)*	0.001	
BMI, kg/m²						0.256
<18.5	1.00	0.87 (0.45–1.68)	0.71 (0.26–1.94)	0.70 (0.14–3.52)	0.224	
18.5–23.9	1.00	0.99 (0.83–1.18)	1.05 (0.87–1.27)	1.10 (0.88–1.39)	0.121	
24–27.9	1.00	2.15 (1.19–3.87)*	1.89 (1.06–3.36)*	2.34 (1.32–4.16)*	<0.001	
\geq 28	1.00	2.13 (0.48–9.41)	1.79 (0.45–7.18)	1.96 (0.50–7.68)	0.463	

Notes: *Significance of P-value ($P < 0.05$). Adjusted for age; sex; BMI; DM family; drinking; smoking; SBP; DBP; HDL; FBG; PBG; HbA1c.

Abbreviations: uACR, urinary albumin to creatinine ratio; LAPI, lipid accumulation product index; BMI, body mass index; DM, diabetes mellitus; CVD, cardiovascular disease; HDL, high-density lipoprotein cholesterol; FBG, fasting blood glucose; PBG, 2-h postload blood glucose; HbA1c, glycosylated hemoglobin.

relationship between LAP and uACR in a middle-aged Chinese prediabetic population. Therefore, such people should be vigilant about the detection, avoidance, and intervention of risk factors to prevent the proteinuria development.

LAP presented as a surrogate marker of visceral adiposity, was reportedly a powerful predictor of prediabetes and CKD, respectively.^{9,16,17} A prospective study in Europe enrolled 148 women with polycystic ovary syndrome proved that elevated LAP was associated with prediabetes.¹⁸ Another community-based cross-sectional survey involving 1777 Chinese participants indicated that LAP showed a better performance in discriminating IFG risk than BMI.¹⁷ Meanwhile, a cross-sectional study of 4658 diabetic participants conducted in China confirmed that in women, increased LAP level was significantly associated with a greater prevalence of DKD; furthermore, elevated LAP was significantly associated with higher uACR in both genders.¹⁹ In the current study, LAP was proven to be more closely associated with uACR in prediabetes, than WHR and WHtR, and independent of measurement of general adiposity is consistent with previous data.⁵

Prediabetes was described as a precursor of chronic kidney disease, and vice versa, microvascular dysfunction preceded and contributed to hyperglycemia through impairment of insulin-mediated glucose disposal and insulin secretion.^{20–22} Meanwhile, visceral obesity was also associated with microvascular abnormality, including insulin-induced microvascular dilation and recruitment, impairments in functional capillary density, and endothelium-dependent vasodilation in the absence of general obesity,^{23–25} supporting the notion that

visceral adiposity was an important driver of DKD.²⁰ In-depth research found lipid metabolism dysregulation occurring systemically and specifically in the kidneys in diabetic individuals.^{26–28} High fructose feeding rats resulted in alterations in lipid metabolism, significantly elevated plasma and tissue lipid peroxides.²⁹ Other in vivo studies have also described lipid accumulation in the kidney cortex in DKD mouse models, demonstrating renal lipids accumulation and lipids peroxide are correlating with the development of glomerulosclerosis in diabetic setting.^{26,27,30}

More previous data have reported lipids crucial roles in the development of CKD.³¹ A study involving 15,362 diabetic patients in Italy found low HDL-C and high TG levels were independent risk factors for the development of DKD over four years.³² Data from the UKPDS study with a median 15 years of follow-up also showed independent risk factors for albuminuria including increased waist circumference, TG, DLD-C in diabetic population.³³ Our study has also described lipids parameter associations with albuminuria in Chinese prediabetic population to show that: after full adjustment of confounding factors (1) elevated TG was significantly associated with increased uACR in both genders, (2) descending HDL-C was associated with uACR in women alone, which was consistent with previous evidence and might explain LAP, as an inexpensive tool to monitor total lipid accumulation was superior over other anthropometrical indices by combining two efficient and independent risk factors in identifying DKD.

In the current study, the association between elevated LAP level and increased albuminuria was attenuated after further

adjustment for BP, HDL-C, FBG, and PBG in Model 4 compared with Models 1 to 3, suggesting that metabolic abnormalities added to the risk of albuminuria. Evidence showed modifiable factors including poor glycemic control, hypertension,³⁴ dyslipidemia,^{35,36} and obesity^{37–39} contribute to the development of CKD in patients with diabetes. Besides, further stratification analysis demonstrated that female gender, middle age (40–59 years old) rather than the older group (≥ 60 years old), rise in blood pressure and weight increase the odds of proteinuria. (1) Sex hormones influenced body fat distribution and CKD progression, which subsequently affect the association of visceral adiposity with CKD in different genders. In premenopausal women, subcutaneous adipose tissue fat accumulated, whereas in postmenopausal women and men, due to the rapid decline in estrogens, more abdominal visceral adipose tissue fat accumulated.⁴⁰ Thus we speculated that the gender-specific discrepancy in the relationship between LAP and uACR is due to sexual hormone-induced fat distribution. (2) Advanced age was believed to be a nonmodifiable risk factor of CKD,⁴¹ we speculated that the senior residents take a more healthy diet and lifestyle into consideration to prevent the development of microvascular complications as reported before.⁴² Thus, in the older group we could not observe the significant relationship on LAP and uACR. (3) Much evidence proved that better blood pressure^{41,43,44} and weight control delay the development of diabetic renal dysfunction.⁴⁵ Thus the early detection and intervention on those modifiable and metabolic risk factors can reduce the additive risk of albuminuria in the prediabetic population.

In place of sophisticated imaging methods for estimating the lipid burden in an isolated tissue, LAP as an inexpensive tool to monitor total lipid accumulation and a surrogate marker of visceral adiposity was reportedly a better predictor of diabetes, a risk factor of CKD and all-cause mortality, compared with central adiposity such as BMI. To date, despite the extensive research on LAP as visceral obesity and CKD, previous studies have failed to investigate the visceral obesity and uACR in the prediabetic population at high-risk of developing diabetes. Thus the current study provides evidence on elevated level of LAP can be applied as an effective indicator of diabetic nephropathy. We believe in taking LAP into consideration as part of the management strategy to reduce risk of diabetic nephropathy in further clinical practice.

Limitation

There are several limitations to this study. First, this is a retrospective single-center cross-sectional study; the

analysis of the causative effect of LAP on DKD is limited. The causal relationship between LAP and uACR requires further prospective investigation. Second, other than bringing the existing adjustment variables into analysis, the absence of other confounders may interact with dependent and independent variables. Third, as the participants enrolled in the REACTION study were aged ≥ 40 years old, a considerable number of middle-aged men were occupied with full-time jobs, while more women were available for the informed consent process and the following recruitment, which might also partly cause the insignificance in relationship of LAP and uACR in men.

Conclusion

In this study, we observed that the elevated level of LAP was associated with increased albuminuria in the prediabetic population in China. Individuals with higher LAP who are female, middle-aged (40–59 years old), hypertensive and overweight have increased odds of proteinuria. Thus, the marker of visceral adiposity, LAP might be a useful clinical indicator of diabetic nephropathy in the prediabetic Chinese population. Our study sheds light onto the perspective for LAP as a reliable and convenient tool identifying diabetic microvascular disease and a therapeutic marker to refine the distribution and deposition of visceral fat and lipid accumulation other than just weight loss.

Abbreviations

ALT, alanine transferase; AST, aspartate transferase; BMI, body mass index; BP, blood pressure; CKD, chronic kidney disease; CVD, cardiovascular disease; DKD, diabetic kidney disease; DM, diabetes mellitus; eGFR, estimated glomerular filtration rate; FBG, fasting blood glucose; GGT, gamma-glutamyl transferase; HbA1c, glycosylated hemoglobin; HC, hip circumference; HDL, high-density lipoprotein cholesterol; IFG, impaired fasting glucose; IGT, impaired glucose tolerance; LDL, low-density lipoprotein cholesterol; MI, myocardial infarction; OGTT, oral glucose tolerance test; PBG, 2-h postload blood glucose; TC, total cholesterol; TG, triglyceride; uACR, urinary albumin to creatinine ratio; WC, waist circumference; WHR, waist circumference to hip circumference ratio; WHtR, waist circumference to height ratio; LAP, lipid accumulation product index.

Data Sharing Statement

All data used to support the findings of this study are available from the corresponding author upon request.

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Disclosure

The authors report no conflicts of interest in this work.

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