

# Metabolic Syndrome and Incident Asthma in Chinese Adults: An Open Cohort Study

This article was published in the following Dove Press journal:  
*Diabetes, Metabolic Syndrome and Obesity: Targets and Therapy*

Haixia Wang  
Chenxiao Bai  
Mo Yi  
Yuanmin Jia  
Yizhang Li  
Di Jiang  
Ou Chen

School of Nursing, Cheeloo College of  
Medicine, Shandong University, Jinan  
250012, Shandong, People's Republic of  
China

**Background:** Although metabolic syndrome is a well-known risk factor for many non-communicable diseases, its contribution to asthma remains controversial.

**Objective:** The aim of this study was to explore the associations of metabolic syndrome and its components with incident asthma in Chinese adults.

**Methods:** We conducted an open cohort study of participants who were asthma-free at baseline ( $n=42,304$ ) in the Shandong multi-center health check-up longitudinal study from 2004 to 2015. Participants aged  $\geq 20$  years and had regular physical examination (once a year) more than three times during follow-up.

**Results:** Ninety subjects (38 women and 52 men) developed incident asthma over 12 years of follow-up. Our study suggested that metabolic syndrome itself was not significantly associated with incident asthma in either women or men ( $P>0.050$ ). Interestingly, we found that overweight and/or obesity was a risk factor for incident asthma among women but not men in the Cox proportional hazards model after adjusting covariates (adjusted incidence rate ratio (IRR)= 2.940, 95% confidence interval (CI): 1.467–5.894,  $P=0.002$ ). The result was consistent with the Poisson regression model (hazard ratio (HR)= 2.241, 95% CI: 1.135–4.988,  $P=0.026$ ). After stratifying according to overweight and/or obesity, we found that female subjects with overweight and obesity were associated with the occurrence of incident asthma ( $P<0.050$ ). However, we did not find this result among men.

**Conclusion:** Metabolic syndrome was not significantly associated with incident asthma in both women and men; however, overweight and/or obesity was shown to be a significant risk factor for incident asthma but only in women, not in men.

**Keywords:** overweight and/or obesity, metabolic syndrome, risk factor, incident asthma

## Introduction

Asthma is a chronic inflammatory disease caused by gene–environment interactions, with symptoms of wheezing, coughing, chest tightness, and shortness of breath.<sup>1</sup> Prevalence rates for asthma have been increasing in many countries over the past few decades.<sup>2–4</sup> Approximately 235 million people worldwide suffer from asthma.<sup>5</sup> Asthma has become one of the most common diseases in the world, resulting in a substantial disease burden.<sup>6</sup> Therefore, it is particularly important to prevent asthma by identifying risk factors for asthma, such as smoking, allergic rhinitis, eczema, psychosocial factors, rhinitis, sinusitis, lung infections, alcohol consumption, air pollution and occupational exposure.<sup>7–10</sup>

Overweight and/or obesity is a vast public health problem and a major risk factor for asthma in adults.<sup>11,12</sup> Overweight and/or obesity is considered part of the metabolic syndrome in China.<sup>13</sup> Metabolic syndrome is a well-known cluster of

Correspondence: Ou Chen  
School of Nursing, Cheeloo College of  
Medicine, Shandong University, Jinan  
250012, Shandong, People's Republic of  
China  
Tel +8615990991181  
Email chenou@sdu.edu.cn

cardiometabolic risk factors associated with an increased risk of multiple chronic diseases, including cancer and cardiovascular disease, and is associated with an approximately 2-fold higher risk of cardiovascular disease and a 5-fold or more increase in the risk of type 2 diabetes mellitus.<sup>14–16</sup> In addition, metabolic syndrome is associated with lung function impairment.<sup>17–19</sup>

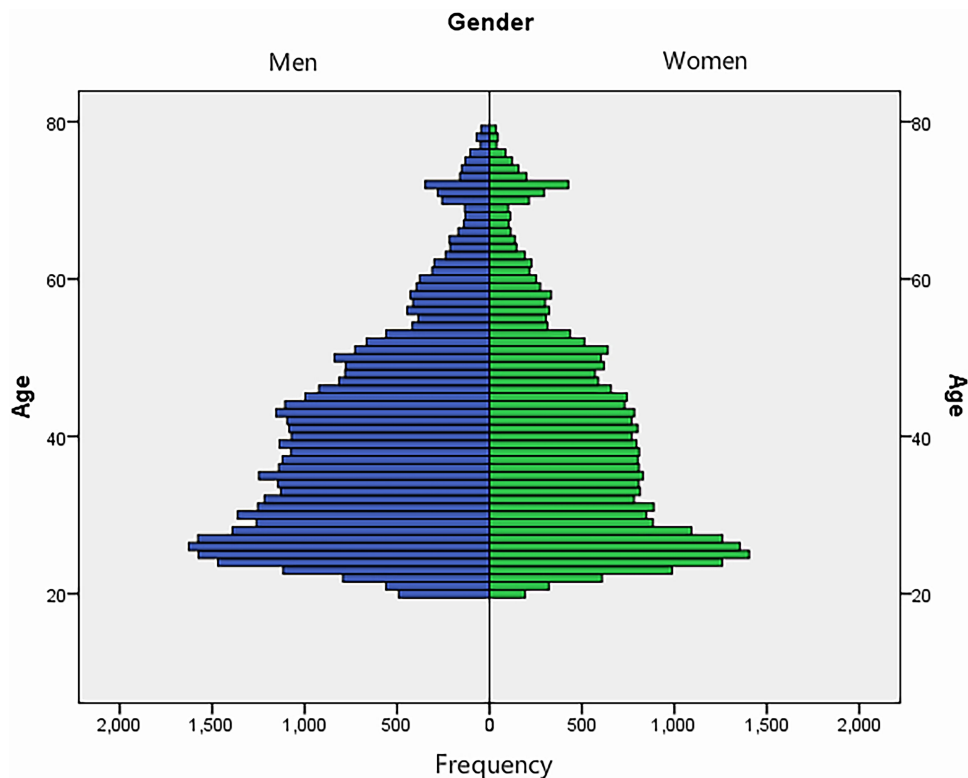
More recently, literature has emerged that offers contradictory findings of the epidemiological link between metabolic syndrome and asthma.<sup>20–23</sup> Prior studies that have noted that metabolic syndrome and its components (high waist circumference (WC), lower high-density lipoprotein cholesterol (HDL-C) and elevated glucose, insulin resistance (IR) or diabetes) were associated with an increased risk of asthma.<sup>22,23</sup> However, some evidence suggested that metabolic syndrome predicted incident asthma among women but not men.<sup>20</sup> In addition, surveys such as that conducted by Tomita et al<sup>21</sup> have shown that metabolic syndrome was not a risk factor for the incidence of late-onset asthma. However, whether metabolic syndrome was a risk factor for asthma was not known in the Chinese population. Based on this, our study aimed to explore the relationship between metabolic syndrome and its components and incident asthma in Chinese adults.

## Methods

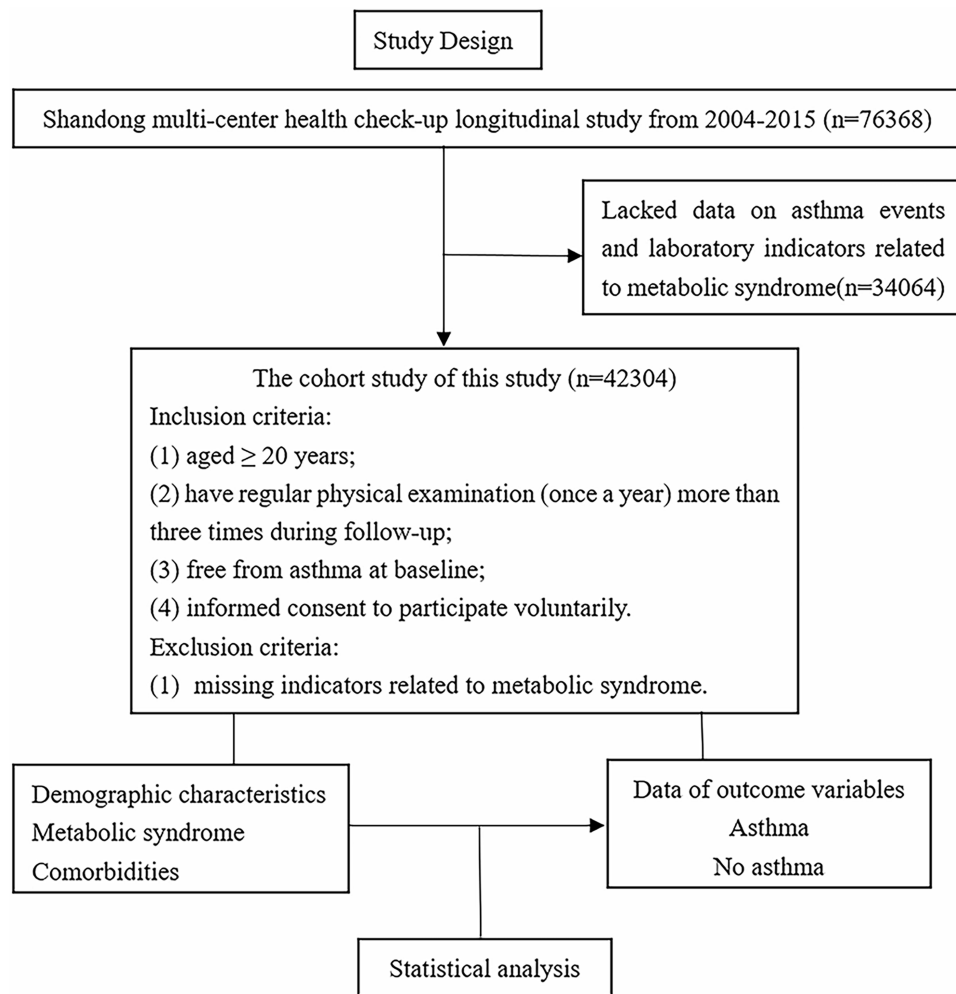
### Study Design

This was an open cohort study that the original participants exited continuously and new objects joined at any time, covering more than 20 health management centres in Shandong Province from January 2004 to December 2015. The longitudinal observational study included 76,368 participants who were free from asthma at baseline, covering nearly all of Shandong Province.<sup>24</sup> The study design and age distribution of this population are shown in Figures 1 and 2. The study was approved by the Institutional Ethical Committee, School of Nursing, Cheeloo College of Medicine, Shandong University (2003-R-036). The study was in compliance with the Declaration of Helsinki for clinical research. All participants provided written informed consent before participating in the study.

The research data were obtained through questionnaire surveys, healthy physical measurements and laboratory examinations of the participants. The questionnaire survey included age, gender, disease history, drinking status and smoking status. Physical measurement indicators included height, weight, systolic blood pressure (SBP) and diastolic blood pressure (DBP). Laboratory indicators included



**Figure 1** Age distribution of the population.



**Figure 2** Study design.

fasting plasma glucose (FPG), triglycerides (TGs), and HDL-C, etc. Height and weight were measured with light clothing without shoes; BP was measured using Omron HEM-907 by the cuff-oscillometric method on the right arm in sitting position after a 5-min rest, and the mean SBP and DBP values of two measurements were recorded, respectively. In addition, participants needed fasting for more than 12 h.

In the follow-up, we collected asthma-related diseases (rhinitis, nasal polyps, pneumonia, bronchiectasis, pulmonary infection, bronchitis, and Chronic obstructive pulmonary disease (COPD)), these diseases were self-reported by participants, including nasal polyps. The doctor at the medical center also confirmed the diagnosis based on the patient's symptoms and related laboratory tests, because nasal polyps coexist with sinusitis.<sup>25</sup> In this study, nasal polyps were only nasal polyps without accompanying sinusitis.

## Inclusion and Exclusion Criteria

The inclusion criteria were as follows: 1) participants aged  $\geq 20$  years; 2) participants had regular physical examination (once a year) more than three times during follow-up; 3) participants in the study who did not have asthma at baseline diagnosed by doctors at the Health Management Center; and 4) individuals who provided informed consent to participate voluntarily. The exclusion criteria were as follows: missing indicators related to metabolic syndrome at baseline (height, weight, SBP, DBP, FPG, TGs, and HDL-C). Finally, this study included 42,304 subjects. The median follow-up time was 3.1 years.

## Dependent and Independent Variables

The primary dependent variable was physician-diagnosed asthma during the annual physical examination. The diagnostic criteria for asthma were according to the guidelines

for Asthma Section, Respiratory Disease Branch, Chinese Medical Association<sup>26</sup> ([Supplementary](#)).

The primary independent variable was metabolic syndrome, including four components. The diagnosis of metabolic syndrome was made using the criteria recommended by the Chinese Medical Association Diabetes Division (CDS) in 2004:<sup>27</sup> 1) overweight and/or obesity: BMI  $\geq 25.0$  kg/m<sup>2</sup>; 2) hyperglycaemia: FPG  $\geq 6.1$  mmol/L (110mg/dL), 2hPG  $\geq 7.8$  mmol/L (140mg/dL), and/or diagnosis of and treatment for diabetes; 3) hypertension: SBP/DBP  $\geq 140/90$  mmHg and/or confirmation and treatment of hypertension; 4) TGs  $\geq 1.7$  mmol/L (150mg/dL) and/or HDL-C  $< 0.9$  mmol/L (35mg/dL) (male) or  $< 1.0$  mmol/L (39mg/dL) (female). Metabolic syndrome was diagnosed when 3 or more of the above 4 components met the criteria.

## Statistical Analysis

Considering that studies have shown that the relationship between metabolic syndrome and asthma is different between men and women, we divided the participants into two groups of men and women for analysis.<sup>20</sup> A comparison of baseline characteristics (age, sex, BMI, smoking status, drinking status and some respiratory diseases) between participants included and excluded from this cohort study was performed by the chi-square test for categorical variables and the Wilcoxon rank sum test for continuous variables. Multiple analyses were performed using the Cox proportional hazards model to determine the association between metabolic syndrome and its components and incident asthma. Considering the low incidence of asthma in this study, we also performed Poisson regression model to verify the reliability of the results. We included metabolic syndrome and the components of metabolic syndrome in model 1 and model 2, respectively, and adjusted the three covariates of age, drinking and smoking status. Then, statistically significant variables such as overweight and/or obesity and other statistically significant variables in univariate analysis were simultaneously entered into the statistical models to compare the impact of those factors on the outcome; Considering that significant lung diseases might affect our findings, we further excluded participants who diagnosed COPD at follow-up (n=184). The excluded data were also included in the two models for analysis. In addition, we also included participants with BMI data but not included in this study into the Poisson regression and Cox proportional hazards model for statistical analysis. We further included

stratification of overweight and obesity into the model for analysis. The statistical significance threshold of IRRs and 95% CIs was  $P < 0.05$ . All analyses were performed using IBM SPSS Statistics 24.

## Results

### Identification and Characteristics of the Participants

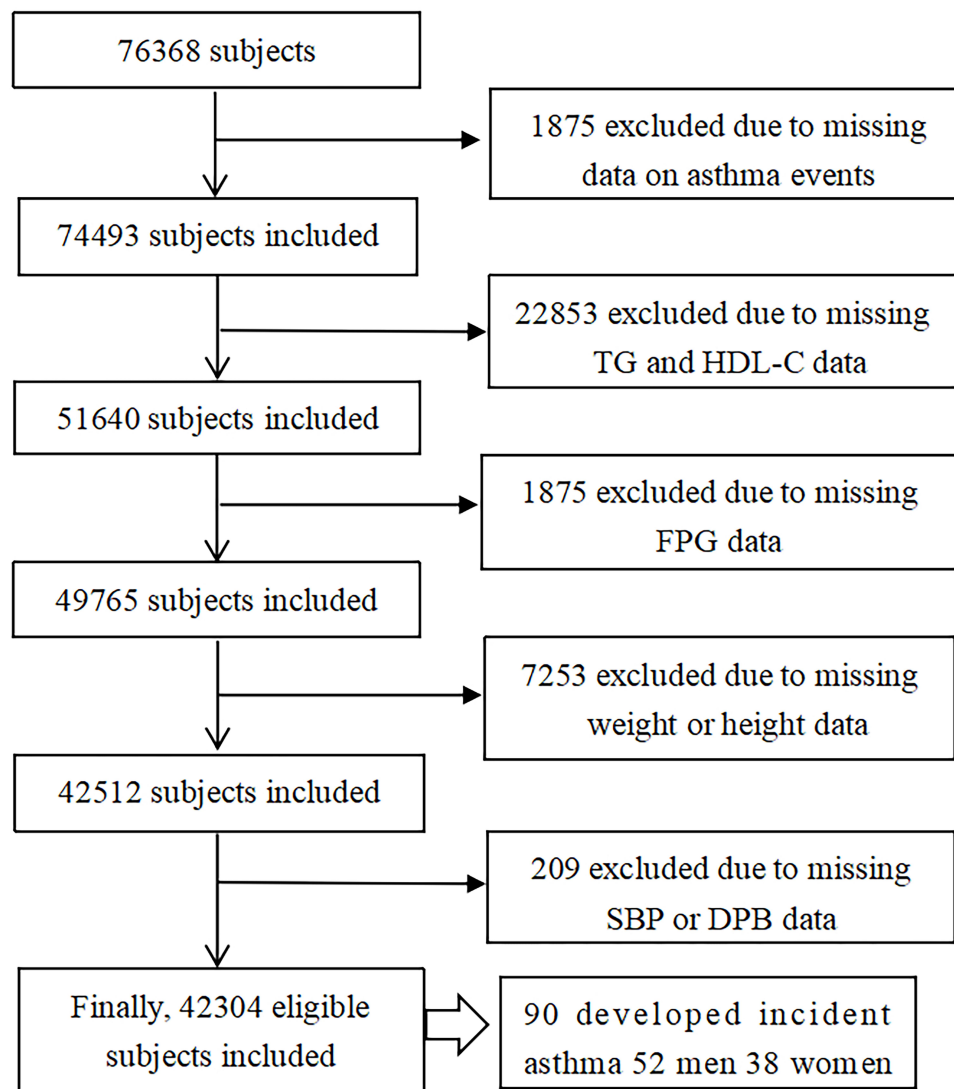
Among the 76,368 original participants at the baseline examination in the open cohort, 42,304 eligible subjects were included in the study ([Figure 3](#)). Ninety eligible subjects developed incident asthma during the follow-up, including 38 women (42.22%) and 52 men (57.78%). Compared to the excluded participants, included participants exhibited different demographic characteristics, generally lower age, more women, less smoking, less drinking and less asthma-like respiratory diseases ([Supplementary Table 1](#)).

### Characteristics of the Participants

As shown in [Table 1](#), we found that older individuals, drinker (people who are drinking or used to drink), smokers (people who are smoking or have smoked), people with higher DBP or those who had respiratory diseases (nasal polyps, pneumonia, bronchitis, COPD, bronchiectasis) were more likely to develop incident asthma among men. In addition, the results showed that people who were older, had higher TGs, were overweight and/or obese, and had incident respiratory diseases (pneumonia, bronchitis, COPD, bronchiectasis) were more likely to develop incident asthma among women.

### Relationship Between Metabolic Syndrome and Its Components and Incident Asthma

The associations of metabolic syndrome and its components with incident asthma are shown in [Table 2](#). Metabolic syndrome could not predict incident asthma in men and women after adjusting for age, drinking and smoking status ( $P > 0.05$ ) in model 1. Of the four metabolic syndrome components, only the associations for overweight or/and obesity remained relatively stable after adjustment for all covariates among women ( $P < 0.05$ ) in model 2. However, this result was not found among men. We also used the Poisson regression model for analysis, and the analysis results were similar in the Cox proportional hazard model analysis ([Supplementary Table 2](#)). The



**Figure 3** Overview of study participant.

**Abbreviations:** SBP, systolic blood pressure; DBP, diastolic blood pressure; FPG, fasting plasma glucose; TG, triglyceride; HDL-C, high-density lipoprotein cholesterol.

relationship between overweight and/or obesity and incident asthma was statistically significant after adjusting for and unadjusted covariates (age, smoking and drinking status) (Table 3, Supplementary Table 3).

After stratifying by overweight and obesity, we found that overweight and obesity were associated with the occurrence of incident asthma among women ( $P < 0.05$ ). However, we did not find this result among men (Table 4). We used the Poisson regression model for analysis, and the analysis results were similar in the Cox proportional hazard model analysis (Supplementary Table 4). The Characteristics of participants with BMI data but excluded in the study in supplementary Table 5. We also verified this result in excluded population including women (Supplementary Table 6) and men (Supplementary Table 7).

Considering that significant lung diseases might affect our findings, we further excluded participants who self-reported COPD and bronchiectasis at follow-up ( $n=184$ ). The excluded data were also included in the Poisson regression model and the Cox proportional hazards model for analysis. We found that the results were not affected after excluding patients' self-reported COPD and bronchiectasis in women (Supplementary Table 8) and men (Supplementary Table 9).

## Discussion

This prospective, open cohort study was conducted in China using data from multiple health management centres. The results of the specific health examinations suggested that overweight and/or obesity was a risk factor for incident

**Table 1** Characteristics of 42,304 Participants Included in the Study

Characteristics	Men (26,009)			Women (16,295)		
	Asthma (n=52)	Control (n=25,957)	Pvalue	Asthma (n=38)	Control (n=16,257)	Pvalue
Age, years	46.67(11.43)	42.25(12.57)	0.00*	47.71(10.95)	43.22(13.32)	0.01*
BMI, kg/m <sup>2</sup>	25.49(3.48)	25.78(3.51)	0.55	23.77(3.24)	23.47(3.57)	0.55
SBP	133.21(21.23)	130.44(17.08)	0.31	120.82(17.05)	120.68(18.92)	0.96
DBP	83.94(10.13)	81.29(12.32)	0.04*	74.66(10.67)	72.84(11.56)	0.26
HDL-C	52.12(13.14)	49.69(11.25)	0.13	57.27(11.95)	56.57(11.79)	0.71
TGs	160.09(100.93)	158.06(147.73)	0.88	116.78(66.13)	102.05(80.82)	0.04*
FPG	100.24(16.66)	99.08(25.39)	0.58	93.47(10.30)	94.44(20.26)	0.60
BMI						
<25kg/m <sup>2</sup>	26(50.00)	10,760(41.45)		18(47.37)	10,498(64.58)	
≥25kg/m <sup>2</sup>	26(50.00)	15,197(58.55)	0.21	20(52.63)	5759(35.42)	0.03*
HDL-C						
≥35mg/dl(male)/≥39mg/dl(female)	49(94.23)	24,454(94.21)		36(94.94)	15,534 (95.55)	
<35mg/dl(male)/<39mg/dl(female)	3(5.77)	1503(5.79)	1.00	2(5.26)	723(4.55)	0.81
TGs						
<150mg/dl	30(57.69)	16,653(64.16)		28(73.68)	13,760(84.64)	
≥150mg/dl	22(42.31)	9340(35.98)	0.33	10(26.32)	2497 (15.36)	0.07
FPG						
<110mg/dl	41(78.85)	22,114(85.19)		36(94.94)	15,534 (95.55)	
≥110mg/dl	11 (21.15)	3843 (14.81)	0.20	2(5.26)	723(4.55)	0.38
BP						
<140/90mmHg	32 (61.54)	17,587 (67.75)		32(84.21)	13,565(83.44)	
≥140/90mmHg	20 (38.46)	8370 (32.25)	0.34	6(15.79)	2692(16.56)	0.90
History of hypertension	7 (13.46)	2258 (8.70)	0.23	3 (7.89)	995 (6.12)	0.65
History of diabetes	1 (1.92)	830 (3.20)	0.61	0(0.00)	395 (100.00)	
Metabolic Syndrome	10 (19.23)	4830 (18.61)	0.91	2 (5.26)	1177 (7.24)	0.64
Drinking status	22(42.31)	6681(25.74)	0.01*	0(0.00)	41(0.25)	/
Smoking status	19 (36.54)	5349 (20.61)	0.01*	0(0.00)	0 (0.00)	/
Componentsofmetabolicsyndrome						
Overweight or obesity	26(50.00)	15,197(58.55)	0.21	20(52.63)	5759(35.42)	0.03*
Hyperglycemia	12 (23.08)	3943 (15.19)	0.12	2(5.26)	1609 (9.90)	0.35
Hypertension	20 (38.46)	8832 (34.03)	0.50	7(18.42)	2963(18.23)	0.98
Dyslipidemia	22 (42.31)	9783 (37.69)	0.49	11(28.95)	2837(17.45)	0.07
Respiratory diseases						
Rhinitis	1 (1.92)	97 (0.37)	0.10	0(0.00)	28(0.00)	/
Nasal polyps	1 (1.92)	41 (0.16)	0.01*	0(0.00)	11(0.00)	/
Pulmonary infection	1 (1.92)	119 (0.46)	0.15	2(5.26)	82(0.50)	0.00*
Pneumonia	7 (13.46)	308 (1.19)	0.00*	7(18.42)	198(0.01)	0.00*
Bronchitis	8 (15.38)	313 (1.21)	0.00*	10(26.32)	236(0.02)	0.00*
COPD	7 (13.46)	95 (0.37)	0.00*	3(7.89)	23(0.00)	0.00*
Bronchiectasis	1 (1.92)	39(0.15)	0.01*	1(2.63)	23(0.001)	0.00*

Note: \*P<0.05.

Abbreviations: BMI, body mass index; SBP, systolic blood pressure; DBP, diastolic blood pressure; BP, blood pressure; HDL-C, high-density lipoprotein cholesterol; TGs, triglycerides; FPG, fasting plasma glucose; COPD, chronic obstructive pulmonary disease; IRR, incident rate ratios.

asthma among only women, not men. However, there was no significant correlation between metabolic syndrome and incident asthma in either men or women. These findings, based

on longitudinal analysis of a large adult population data set, confirm the importance of overweight and/or obesity as a risk factor for incident asthma in adult women.



**Table 2** Longitudinal Associations Between the Metabolic Syndrome and Incident Asthma

Statistical Models	Men		Women	
	HR (95% CI)	P	HR (95% CI)	P
Model 1				
Metabolic syndrome+	0.81(0.40–1.63)	0.56	0.45(0.10–1.95)	0.29
Metabolic syndrome-	0.92(0.46–1.83)	0.81	0.82(0.20–3.40)	0.78
Model 2				
Hypertension+	0.87(0.48–1.60)	0.66	0.55(0.22–1.35)	0.19
Dyslipidemia+	1.06(0.59–1.90)	0.84	1.23(0.58–2.58)	0.59
Hyperglycemia+	1.42(0.73–2.79)	0.31	0.37(0.09–1.61)	0.19
Overweight and/or obesity+	0.77(0.43–1.37)	0.37	2.94(1.47–5.89)	0.00*
Hypertension-	1.03(0.57–1.85)	0.92	0.76(0.32–1.80)	0.53
Dyslipidemia-	1.04(0.58–1.86)	0.90	1.36(0.65–2.86)	0.42
Hyperglycemia-	1.70(0.87–3.30)	0.12	0.45(0.11–1.94)	0.29
Overweight and/or obesity-	0.76(0.42–1.35)	0.35	3.54(1.79–7.01)	0.00*

**Notes:** +Adjusted age, smoking and drinking status; -unadjusted age, smoking and drinking status. \* P<0.05.

**Abbreviations:** HR, hazard ratio; IRR, incident rate ratios.

**Table 3** Longitudinal Associations Between Overweight and/or Obesity and Incident Asthma

Statistical Models	Men		Women	
	HR (95% CI)	P	HR (95% CI)	P
BMI-				
<25kg/m <sup>2</sup>				
≥25kg/m <sup>2</sup>	1.61(0.33–7.74)	0.56	6.86(1.84–25.58)	0.00*
BMI+				
<25kg/m <sup>2</sup>				
≥25kg/m <sup>2</sup>	0.96(0.18–5.15)	0.97	5.05(1.26–20.22)	0.02*

**Notes:** +Adjusted age, smoking and drinking status; -unadjusted age, smoking and drinking status. \*P<0.05.

**Abbreviations:** HR, hazard ratio; IRR, incident rate ratios; BMI, body mass index.

In recent years, findings on metabolic syndrome and asthma have been inconsistent. Studies in the US, Norway, and South Korea have shown that metabolic syndrome is

associated with incident asthma.<sup>20,22,23</sup> However, among these studies, the large retrospective cohort study from Norway showed that high WC (adjusted OR 1.62, 95% CI 1.36–1.94) and elevated glucose or diabetes (adjusted OR 1.43, 95% CI 1.01–2.04) were associated with an increased risk of incident asthma in adults. In addition, a cohort followed for over 25 years in the USA reported that after adjusting for BMI, the correlation between metabolic syndrome and asthma attacks was no longer statistically significant among women ( $P=0.44$ ). Interestingly, a retrospective cohort study in Japan showed that obesity, not metabolic syndrome, was a significant risk factor for the incidence of late-onset asthma but only in middle-aged Japanese women,<sup>21</sup> which was similar to the result of our open cohort study. Our observations suggest that overweight and/or obesity, not metabolic syndrome, is a significant risk factor for incident asthma, but only in Chinese women, not in men. The inconsistency between the findings of previous studies and our current study may be due to the differences in study designs, populations and diagnostic criteria for metabolic syndrome.

**Table 4** Longitudinal Associations Between BMI and Incident Asthma

Variables	Men		Women	
	HR (95% CI)	P	HR (95% CI)	P
BMI 25.0–29.9kg/m <sup>2</sup> -	0.813(0.461–1.435)	0.48	3.194(1.593–6.404)	0.001*
BMI≥30kg/m <sup>2</sup> -	0.783(0.276–2.275)	0.67	8.134(3.126–21.168)	0.000*
BMI 25.0–29.9kg/m <sup>2</sup> +	0.784(0.443–1.387)	0.40	2.711(1.310–5.610)	0.007*
BMI≥30kg/m <sup>2</sup> +	0.769(0.268–2.211)	0.63	6.405(2.339–17.541)	0.000*

**Notes:** +Adjusted age, smoking and drinking status; -unadjusted age, smoking and drinking status. \*P<0.05.

**Abbreviations:** HR, hazard ratio; IRR, incident rate ratios; BMI, body mass index.

We need to carefully interpret the results for the components, as these components in the definition of metabolic syndrome are dichotomous and do not allow for adjustment.<sup>13</sup> In our study, TGs were statistically significant in univariate analysis among women ( $P=0.037$ ). However, the association was no longer statistically significant after adjustment for covariates ( $P>0.050$ ). Further studies should address the individual component in more detail; for example, repeated measurements of consecutive serum levels in a cohort study should be performed to eliminate confounding factors and help identify the associations of the components of metabolic syndrome with asthma.

There are several possible mechanisms connecting overweight and/or obesity and incident asthma. We observed that obesity may change pulmonary physiology and lung function in adults.<sup>28,29</sup> Excessive accumulation of fat in the chest and abdomen might cause compression of the lungs, accompanied by reduced lung volume. In fact, we have shown that obesity increased the folding of the surrounding airways and parenchyma, especially in asthmatic patients with late-onset disease.<sup>30</sup> In addition, adipocytokines (leptin and resistin) produced by adipose tissue have been suggested to promote systemic inflammation, which may induce asthma.<sup>31,32</sup> Patients with  $\text{aBMI} \geq 25 \text{ kg/m}^2$  showed higher levels of leptin, and this cytokine difference was also found in female patients with asthma ( $P < 0.05$ ).<sup>33</sup> On the other hand, asthma and obesity have considerable genetic components. The obesity–asthma association may also be due to common genes shared by both obesity and asthma. Candidate gene studies have reported a few genes associated with asthma and obesity, such as PRKCA and NR3C1.<sup>34,35</sup> The specific mechanism of the relationship between overweight and/or obesity and asthma still needs more research to explore.

There were some limitations to our study. First, this study lacks demographic characteristics such as birthplace, residence, occupation, marriage, income, etc. This may be due to the incomplete basic information filled out by the research subjects in various medical examination centers, which may have an impact on the results. Second, the measurement of various indicators of metabolic syndrome over time was not included, as some study subjects did not participate in regular physical examinations. Third, the doctor diagnosed asthma based on the patient's lung function indicators and clinical manifestations, but we did not collect data on

the lung function indicators, which may affect the research results to a certain extent. Fourth, metabolic syndrome was diagnosed using the diagnostic criteria recommended by the CDS in 2004; however, it did not include WC, which may not reflect overweight and/or obesity, as many studies have shown that WC was closely related to asthma.<sup>36–38</sup> Therefore, we hope that the diagnostic criteria for metabolic syndrome will be updated in China.

In conclusion, the study confirmed overweight and/or obesity as a risk factor for incident asthma only among women, not men. However, our study suggested that metabolic syndrome was not significantly associated with incident asthma in both women and men. The findings of this study warrant further research, particularly addressing the associations between the components of metabolic syndrome and incident asthma in prospective studies.

## Abbreviations

HR, hazard ratio; IRR, incident rate ratios; CI, confidence interval; WHO, World Health Organization; SBP, systolic blood pressure; DBP, diastolic blood pressure; FPG, fasting plasma glucose; TGs, triglycerides; HDL-C, high-density lipoprotein cholesterol; IR, insulin resistance; CDS, Chinese Medical Association Diabetes Division; BMI, body mass index; COPD, chronic obstructive pulmonary disease; WC, waist circumference.

## Funding

This work was supported by the National Natural Science Foundation of China (Grant 81400072), the Natural Science Foundation of Shandong Province (Grant 2013HQ047), the Medicine and health development plan of Shandong Province (Grant 2017WS113), the humanities and social sciences youth team project of Shandong University (Grant IFYT18036). The funding bodies were not involved in the design of the study, the collection, analysis, and interpretation of data and the preparation of the manuscript.

## Disclosure

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper. The authors report no conflicts of interest in this work.



## References

- Mims JW. Asthma: definitions and pathophysiology. *Int Forum Allergy RH*. 2015;5(Suppl 1):S2–S6.
- Moraes TJ, Sears MR, Subbarao P. Epidemiology of asthma and influence of ethnicity. *Semin Respir Crit Care Med*. 2018;39(1):3–11. doi:10.1055/s-0037-1618568
- Aaron SD, Boulet LP, Reddel HK, Gershon AS. Underdiagnosis and overdiagnosis of asthma. *Am J Respir Crit Care*. 2018;198(8):1012–1020. doi:10.1164/rccm.201804-0682CI
- Collaborators GCRD. Global, regional, and national deaths, prevalence, disability-adjusted life years, and years lived with disability for chronic obstructive pulmonary disease and asthma, 1990–2015: asystematic analysis for the global burden of disease study 2015. *Lancet Respir Med*. 2017;5(9):691–706.
- Organization WH. *Chronic Respiratory Diseases: Scope: Asthma*. Vol. 2020; 2020:235
- GINA. *Main Report - Global Initiative for Asthma - GINA*. Vol. 2020; 2019:2019
- Beasley R, Semprini A, Mitchell EA. Risk factors for asthma: is prevention possible? *Lancet (London)*. 2015;386(9998):1075–1085. doi:10.1016/S0140-6736(15)00156-7
- Ilmarinen P, Tuomisto LE, Kankaanranta H. Phenotypes, risk factors, and mechanisms of adult-onset asthma. *Mediat Inflamm*. 2015;2015:514868. doi:10.1155/2015/514868
- Lin J, Wang W, Chen P, et al. Prevalence and risk factors of asthma in mainland China: the CARE study. *Respir Med*. 2018;137:48–54. doi:10.1016/j.rmed.2018.02.010
- Lieberoth S, Backer V, Kyvik K, et al. Intake of alcohol and risk of adult-onset asthma. *Respir Med*. 2012;106(2):184–188. doi:10.1016/j.rmed.2011.11.004
- Peters U, Dixon AE, Forno E. Obesity and asthma. *J Allergy Clin Immunol*. 2018;141(4):1169–1179. doi:10.1016/j.jaci.2018.02.004
- Ciprandi G, Schiavetti I, Fontana R, Sorbello V, Ricciardolo F. Overweight and obesity as risk factors for impaired lung function in patients with asthma: a real-life experience. *Allergy Asthma Proc*. 2014;35(4):E62–E71. doi:10.2500/aap.2014.35.3773
- Alberti KGMM, Eckel RH, Grundy SM, et al. Harmonizing the metabolic syndrome: a joint interim statement of the International diabetes federation task force on epidemiology and prevention; national heart, lung, and blood institute; American heart association; world heart federation; international atherosclerosis society; and international association for the study of obesity. *Circulation*. 2009;120:1640–1645.
- Sookoian S, Pirola CJ. Metabolic syndrome: from the genetics to the pathophysiology. *Curr Hypertens Rep*. 2011;13(2):149–157. doi:10.1007/s11906-010-0164-9
- Samson SL, Garber AJ. Metabolic syndrome. *Endocrinol Metab Clin*. 2014;43(1):1–23. doi:10.1016/j.ecl.2013.09.009
- Ali Z, Ulrik CS, Agner T, Thomsen SF. Association between atopic dermatitis and the metabolic syndrome: a systematic review. *Dermatology (Basel)*. 2018;234(3–4):79–85. doi:10.1159/000491593
- Baffi CW, Wood L, Winnica D, et al. Metabolic syndrome and the lung. *Chest*. 2016;149(6):1525–1534. doi:10.1016/j.chest.2015.12.034
- Leone N, Courbon D, Thomas F, et al. Lung function impairment and metabolic syndrome: the critical role of abdominal obesity. *Am J Respir Crit Care*. 2009;179(6):509–516. doi:10.1164/rccm.200807-1195OC
- Forno E, Han Y, Mullen J, Celedon JC. Overweight, obesity, and lung function in children and adults—a meta-analysis. *J Allergy Clin Immunol*. 2018;6(2):570–581.
- Assad N, Qualls C, Smith LJ, et al. Body mass index is a stronger predictor than the metabolic syndrome for future asthma in women. The longitudinal CARDIA study. *Am J Respir Crit Care Med*. 2013;188:319–326. doi:10.1164/rccm.201303-0457OC
- Tomita Y, Fukutomi Y, Irie M, et al. Obesity, but not metabolic syndrome, as a risk factor for late-onset asthma in Japanese women. *Allergol Int*. 2019;68(2):240–246. doi:10.1016/j.alit.2018.10.003
- Park S, Choi N, Kim S, Lee C. The relationship between metabolic syndrome and asthma in the elderly. *Sci Rep*. 2018;8(1):9378.
- Brumpton BM, Camargo CAJ, Romundstad PR, Langhammer A, Chen Y, Mai X. Metabolic syndrome and incidence of asthma in adults: the HUNT study. *Eur Respir J*. 2013;42(6):1495–1502. doi:10.1183/09031936.00046013
- Liu Y, Xing P, Xu X, et al. Shandong multi-center health management longitudinal cohort. *J Shandong Univ*. 2017;55(06):30–36.
- Fokkens V, Lund V, Mullol J, et al. European position paper on rhinosinusitis and nasal polyps 2012. *Rhinology*. 2012;50(1):1–298. doi:10.4193/Rhino50E2
- Asthma Section RDBC. Guidelines for the prevention and treatment of bronchial asthma (definition, diagnosis, treatment, education and management of bronchial asthma). *Chin J Int Med*. 2003;(11):68–73.
- Group CMAD. Recommendations of the Chinese medical association diabetes division on metabolic syndrome. *Clin J Diabetes*. 2004;(03):5–10.
- Watson RA, Pride NB, Thomas EL, et al. Reduction of total lung capacity in obese men: comparison of total intrathoracic and gas volumes. *J Appl Physiol*. 2010;108(6):1605–1612. doi:10.1152/jappphysiol.01267.2009
- Dixon AE, Peters U. The effect of obesity on lung function. *Expert Rev Respir Med*. 2018;12(9):755–767. doi:10.1080/17476348.2018.1506331
- Al-Alwan A, Bates JHT, Chapman DG, Kaminsky DA, DeSarno MJ, Irvin CG. The nonallergic asthma of obesity. A matter of distal lung compliance. *Am J Respir Crit Care*. 2014;189(12):1494–1502. doi:10.1164/rccm.201401-0178OC
- Sood A, Shore SA. Adiponectin, leptin, and resistin in asthma: basic mechanisms through population studies. *J Allergy (Cairo)*. 2013;2013:785835. doi:10.1155/2013/785835
- Kattan M, Kumar R, Bloomberg GR, et al. Asthma control, adiposity, and adipokines among inner-city adolescents. *J Allergy Clin Immunol*. 2010;125(3):584–592. doi:10.1016/j.jaci.2010.01.053
- Muc M, Todo-Bom A, Mota-Pinto A, Vale-Pereira S, Loureiro C. Leptin and resistin in overweight patients with and without asthma. *Allergol Immunopath*. 2014;42(5):415–421. doi:10.1016/j.aller.2013.03.004
- Murphy A, Tantisira KG, Soto-Quiros ME, et al. A positional candidate gene for body mass index and asthma. *Am J Hum Genet*. 2009;85(1):87–96. doi:10.1016/j.ajhg.2009.06.011
- Beuther DA, Weiss ST, Sutherland ER. Obesity and asthma. *Am J Respir Crit Care*. 2006;174(2):112–119. doi:10.1164/rccm.200602-231PP
- Kronander U, Falkenberg M, Zetterström O. Prevalence and incidence of asthma related to waist circumference and BMI in a Swedish community sample. *Respir Med*. 2004;98(11):1108–1116. doi:10.1016/j.rmed.2004.03.022
- Jeong A, Imboden M, Hansen S, et al. Heterogeneity of obesity-asthma association disentangled by latent class analysis, the SAPALDIA cohort. *Respir Med*. 2017;125:25–32. doi:10.1016/j.rmed.2017.02.014
- Vangeepuram N, Teitelbaum SL, Galvez MP, Brenner B, Doucette J. Measures of obesity associated with asthma diagnosis in ethnic minority children. *J Obes*. 2011;2011.

## Diabetes, Metabolic Syndrome and Obesity: Targets and Therapy

Dovepress

### Publish your work in this journal

Diabetes, Metabolic Syndrome and Obesity: Targets and Therapy is an international, peer-reviewed open-access journal committed to the rapid publication of the latest laboratory and clinical findings in the fields of diabetes, metabolic syndrome and obesity research. Original research, review, case reports, hypothesis formation, expert opinion

and commentaries are all considered for publication. The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit <http://www.dovepress.com/testimonials.php> to read real quotes from published authors.

Submit your manuscript here: <https://www.dovepress.com/diabetes-metabolic-syndrome-and-obesity-targets-and-therapy-journal>