

Research Article

Cite this article: Chen M, Shi S, Wang S, Huang Y, Zhou F, and Zhong VW (2024) Prevalence of cardiometabolic diseases in underweight: a nationwide cross-sectional study. *British Journal of Nutrition* **132**: 1654–1662. doi: [10.1017/S0007114524002885](https://doi.org/10.1017/S0007114524002885)

Received: 14 June 2024
Revised: 11 October 2024
Accepted: 7 November 2024
First published online: 11 November 2024

Keywords:

Underweight; Cardiometabolic diseases; NHANES; Prevalence; BMI

Abbreviations:

CMD, cardiometabolic disease; NHANES, National Health and Nutrition Examination Survey; PR, prevalence ratio

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Prevalence of cardiometabolic diseases in underweight: a nationwide cross-sectional study

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Abstract

This study aimed to estimate the nationwide prevalence of cardiometabolic diseases (CMD) among adults with underweight in the US general population. Using data from the National Health and Nutrition Examination Survey (1999–2020), we estimated the age-standardised prevalence of dyslipidemia, hypertension, diabetes, chronic kidney disease, CVD and the presence of zero or at least two CMD. Multivariable Poisson regressions were used to compare CMD prevalence between subgroups, adjusting for age, sex and race/ethnicity. Among the 855 adults with underweight included, the weighted mean age was 40.8 years, with 68.1 % being women and 70.4 % non-Hispanic White. The estimated prevalence rates were 23.4 % for dyslipidemia (95 % CI 19.4 %, 27.5 %), 15.6 % for hypertension (95 % CI 13.3 %, 17.8 %), 2.5 % for diabetes (95 % CI 1.5 %, 3.5 %), 7.9 % for chronic kidney disease (95 % CI 6.9 %, 8.8 %) and 6.1 % for CVD (95 % CI 4.3 %, 7.9 %). The prevalence of having zero and at least two CMD was 50.6 % (95 % CI 44.1 %, 57.0 %) and 12.3 % (95 % CI 8.1 %, 16.4 %), respectively. Non-Hispanic Black adults had significantly higher prevalence of diabetes (adjusted prevalence ratio, 3.35; 95 % CI 1.35, 8.30) compared with non-Hispanic White adults. In conclusion, approximately half of the underweight adults had at least one CMD, and 12.3 % had at least two CMD. Prevention and management of CMD in underweight adults are critical yet neglected public health challenges.

Cardiometabolic diseases (CMD) are a group of common but often preventable diseases encompassing CVD and metabolic disorders, which are a major public health concern worldwide⁽¹⁾. Existing studies have focused on quantifying burden of CMD in general populations or people with obesity^(2,3). Data regarding the burden of CMD specifically among people with underweight are scarce. However, underweight has been associated with increased risk of CVD and mortality, according to repeatedly reported J- or U-shaped associations of BMI with CVD and mortality in both general and diseased populations^(4–7). In the USA, about 1.6 % of adults aged 20 years or older were underweight in 2017–2018, equivalent to about 4 million individuals⁽⁸⁾. However, the nationwide prevalence of CMD among US adults with underweight is unclear.

Using nationally representative data from the National Health and Nutrition Examination Survey (NHANES) in 1999–2020, we sought to estimate the prevalence of a range of CMD among US adults with underweight.

Methods*Data source and study design*

NHANES is a continuous, multistage, nationally representative survey of the non-institutionalised civilian resident US population. The survey has been conducted periodically in 2-year cycles since 1999, collecting data through in-home interviews and study visits at mobile examination centres. However, the NHANES programme suspended field operations in March 2020 due to the pandemic of coronavirus disease 2019. Therefore, data collected from 2019 to March 2020 were combined with the data from the 2017–2018 cycle to form a nationally representative sample. This study included ten cycles between 1999–2000 and 2017–2020. The overall response rate ranged from 51 % to 84 % for the interview component and from 46.9 % to 80 % for the examination component. Participants aged 20 years or older were included. Pregnant women were excluded.

The National Center for Health Statistics Research Ethics Review Board approved NHANES. This study was conducted according to the guidelines laid down in the Declaration of Helsinki, and all procedures involving human subjects/patients were approved by the Shanghai Jiao Tong University School of Medicine Public Health and Nursing Research Ethics Review Committee (ethics number: SJUPN-202102-B). Written informed consent was obtained from all participants.

Underweight and covariates

Height and weight were collected by trained health technicians. BMI was computed by dividing an individual's weight in kilograms by their height in metres squared. BMI < 18.5 kg/m² defined underweight and the inclusion criterion of this study. Self-reported information on age, sex, race/ethnicity, education and medical conditions was collected during the household interview. Race/ethnicity was self-reported according to fixed-category questions. Alcohol consumption, leisure-time physical activity, sleep duration, smoking status and dietary intake were self-reported. Diet quality was assessed by Healthy Eating Index-2015 (HEI-2015)⁽⁹⁾.

Definition of cardiometabolic diseases

CMD included dyslipidemia, hypertension, prediabetes, diabetes, chronic kidney disease and CVD. Dyslipidemia was defined as having a total cholesterol level ≥ 240 mg/dl or a HDL-cholesterol level < 40 mg/dl for men or < 50 mg/dl for women or self-reported current use of lipid-lowering drugs⁽¹⁰⁾. Hypertension was defined as having blood pressure $\geq 130/80$ mm Hg or self-reported current use of antihypertensive drugs. Blood pressure was based on the average of all available measurements. Diabetes was defined as having a self-reported diagnosis of diabetes by a physician or other health professional, a fasting plasma glucose level ≥ 126 mg/dl or a Hb A1c level ≥ 6.5 %. Fasting plasma glucose was measured among those who were fasted for 8 to < 24 h. Plasma glucose data between 2005–2006 and 2017–2020 were calibrated according to the recommended equation from NHANES⁽¹¹⁾. Among participants without being diagnosed with diabetes before, a Hb A1c level of 5.7–6.4 % or a fasting plasma glucose level of 100–125 mg/dl defined prediabetes. Chronic kidney disease was defined as having a urine albumin-to-creatinine ratio ≥ 30 mg/g or an estimated glomerular filtration rate < 60 ml/min/1.73 m²⁽¹²⁾. Urine and serum creatinine levels were calibrated⁽¹³⁾. Estimated glomerular filtration rate was computed according to the Chronic Kidney Disease Epidemiology Collaboration equation⁽¹⁴⁾. CVD was a composite endpoint of self-reported congestive heart failure, CHD, heart attack and stroke. Two or more CMD commonly co-occur within an individual⁽²⁾. Therefore, having zero and at least two of the following diseases were studied: dyslipidemia, hypertension, diabetes, chronic kidney disease and CVD.

Statistical analysis

The characteristics of the study participants were described using weighted percentages or weighted mean (se). The prevalence of CMD alone and in combination was estimated in the total sample and by age (20–39, 40–59 and ≥ 60 years), sex (men and women), race/ethnicity (non-Hispanic White, non-Hispanic Black, Hispanic and other) and education (less than high school, high school graduate, some college and college graduate or higher). Estimates were age-standardised to the 1999–2020 NHANES non-pregnant adult population with underweight. Multivariable

Poisson regressions were used to estimate the prevalence ratios (PR) for comparing the prevalence of CMD between subgroups, adjusting for age, sex and race/ethnicity. Subgroup differences in the PR of CMD were obtained from weighted Poisson regression models. *P* values were adjusted using false discovery rate (FDR) corrections.

Post hoc analyses were conducted to help interpret the prevalence of CMD in adults with underweight. First, the prevalence of CMD in the general population was compared by weight category (underweight, normal weight, overweight and obese) using multivariable Poisson regressions, adjusting for age, sex and race/ethnicity. Using underweight as the reference, PR and 95 % CI were derived for other weight categories. Second, the prevalence of CMD among adults with underweight was estimated by different lifestyle factors: non-excessive drinking (≥ 4 –5 drinks/d, or ≤ 14 drinks/week for men or ≤ 7 drinks/week for women; yes/no)⁽¹⁵⁾, meeting physical activity guidelines (≥ 150 min per week of moderate-intensity or ≥ 75 min per week of vigorous-intensity leisure-time activity; yes/no), meeting recommended sleep duration (sleep duration of 7–9 h per d; yes/no), smoking status (self-reported and grouped into three categories: current, former and never) and low HEI-2015 score (< 50 or ≥ 50). Subgroup differences were compared using the *F* tests.

Appropriate sampling weights and design variables were considered to account for the stratified, multistage probability cluster sampling method. Complete case analysis was implemented for primary analysis unless missing data for specific analysis exceeded 10 % according to the NHANES analytical guidelines⁽¹⁶⁾. When missing data exceeded 10 %, the original sampling weights of the respondent sample were adjusted using a weight factor that accounted for the differences between respondents and non-respondents, based on the Lohr's method⁽¹⁷⁾. A *post hoc* analysis revealed sex and racial/ethnic differences between individuals with missing data and those without; age and education attainment did not differ between the two groups (online Supplementary eTable 1). Participants were therefore classified into eight subgroups defined by sex (men and women) and race/ethnicity (non-Hispanic Black, non-Hispanic White, Hispanic and other)⁽¹⁸⁾. The weight factor for each subgroup was calculated as the sum of the weights for all eligible individuals (including those with missing data) divided by the sum of the weights for those with complete data. The adjusted sampling weight for each respondent in a subgroup was then multiplied by the subgroup weight factor. A sensitivity analysis using multiple imputations was also conducted to assess the robustness of the prevalence estimates in the presence of missing data ≥ 10 %. We employed PROC MI procedure in SAS (v 9.4) and used the fully conditional specification method⁽¹⁹⁾. The following covariates were included: age, sex, race/ethnicity, education level, estimated glomerular filtration rate, urine albumin-to-creatinine ratio and chronic kidney disease status. All missing values were assumed to be missing at random. All analyses were conducted using SAS statistical software, version 9.4 (SAS Institute Inc.).

Results

A total of 855 participants with underweight were analysed. Specific sample size for each outcome varied (Fig. 1). The weighted mean age was 40.8 years (SE, 0.6), 68.1 % were women and 70.4 % were non-Hispanic White (Table 1). Missing data were found for dyslipidemia (*n* 79 (9.2 %)), hypertension (*n* 39 (4.6 %)), chronic kidney disease (*n* 86 (10.0 %)) and CVD (*n* 5 (0.6 %)). For

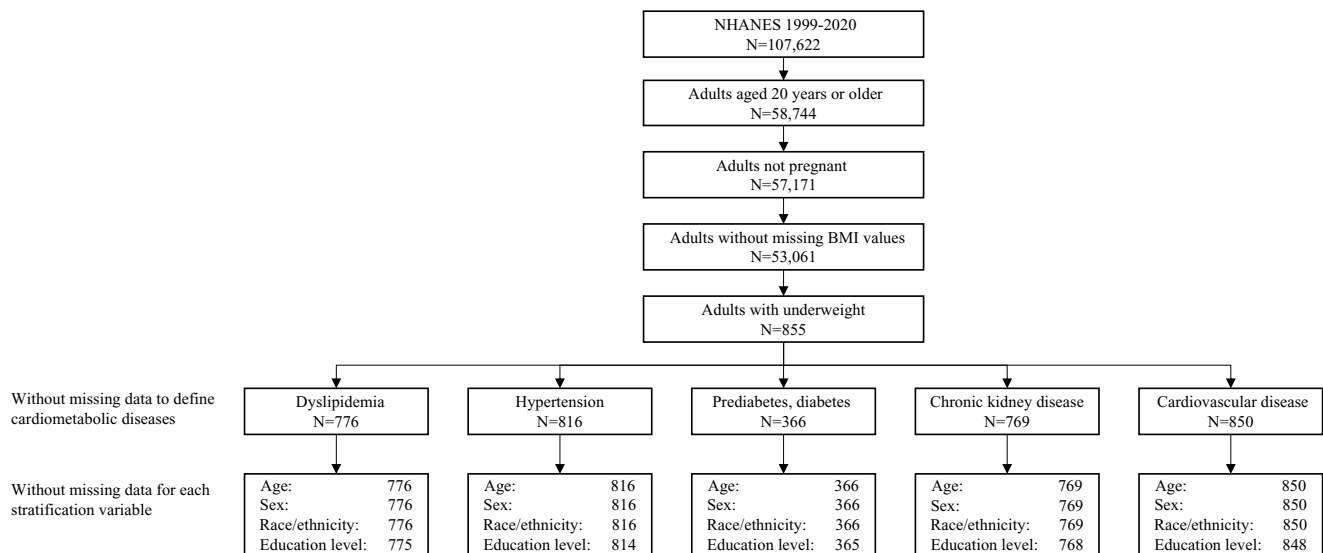


Figure 1. Flow chart of the study sample.

Table 1. Characteristics for adults with underweight*

Characteristics	No. of participants	Weighted %
Total	855	100.0
Age group (years)		
20–39	407	54.7
40–59	209	27.8
≥ 60	239	17.5
Sex		
Men	334	31.9
Women	521	68.1
Race/ethnicity [†]		
Non-Hispanic White	412	70.4
Non-Hispanic Black	209	11.7
Hispanic	92	6.8
Other	142	11.1
Education level [‡]		
Less than high school	228	20.4
High school graduate	197	24.7
Some college	242	28.6
College graduate or higher	186	26.3

*Underweight was defined as having BMI < 18.5 kg/m².

[†]Race/ethnicity was self-reported. The 'other' group included other non-Hispanic races or multiple races.

[‡]Two participants refused to report or did not know their education level.

stratification variables, missing data were found only for education (n 2 (0.2 %)).

The estimated prevalence of each CMD among adults with underweight was 23.4 % (95 % CI 19.4 %, 27.5 %) for dyslipidemia, 15.6 % (95 % CI 13.3 %, 17.8 %) for hypertension, 21.2 % (95 % CI 16.6 %, 25.8 %) for prediabetes, 2.5 % (95 % CI 1.5 %, 3.5 %) for diabetes, 7.9 % (95 % CI 6.9 %, 8.8 %) for chronic kidney disease

and 6.1 % (95 % CI 4.3 %, 7.9 %) for CVD (Table 2). The estimated prevalence of having zero and at least two CMD was 50.6 % (95 % CI 44.1 %, 57.0 %) and 12.3 % (95 % CI 8.1 %, 16.4 %), respectively. Regarding the subgroup results, the prevalence of all CMD was significantly higher in older adults aged at least 60 years than young adults aged 20–39 years, except for dyslipidemia (PR 1.81 (1.04, 3.16), FDR-adjusted P = 0.08) (Table 3). No sex difference in the prevalence of CMD was identified. Racial/ethnic difference in the prevalence of CMD was only found for diabetes. Non-Hispanic Black adults had a significantly higher prevalence of diabetes (PR 3.35 (95 % CI 1.35, 8.30), FDR-adjusted P = 0.045) than non-Hispanic White adults. The prevalence of having zero CMD was significantly lower in older and middle-aged adults than young adults. The prevalence of having at least two CMD was significantly higher in older and middle-aged adults than young adults. No significant difference in the prevalence of composite CMD outcomes by sex, race or education level was identified.

Post hoc analyses showed that adults with underweight had the highest prevalence of chronic kidney disease among all weight categories (all P < 0.01). No significant difference in the prevalence of CVD was found across weight categories (all P > 0.05) (online Supplementary eTable 2). Among adults with underweight, the prevalence of all CMD was significantly lower in those who met the physical activity guideline recommendation than those who did not meet. The prevalence of all CMD, except for diabetes, was significantly lower in never smokers than former or current smokers (online Supplementary eTable 3). Adults with healthier lifestyle behaviours (e.g. non-excessive drinking, meeting physical activity guidelines, never smoking and not low diet quality) had a significantly higher prevalence of having zero CMD than that in their counterparts (all P < 0.05). No significant difference in the prevalence of all CMD, except for CVD, was found between those who met the recommended sleep duration and those who did not.

The prevalence of chronic kidney disease among adults with underweight using different missing data handling methods was similar. Compared with complete case analysis, only small differences in the point estimates were found using adjusted weights (\leq 0.3 %) and multiple imputations (\leq 1.4 %) (online Supplementary eTable 4).

Table 2. Prevalence of cardiometabolic diseases among adults with underweight* (Percentages and 95 % confidence intervals)

Characteristics	Dyslipidemia [†]			Hypertension [†]			Prediabetes [‡]			Diabetes		
	No.	Prevalence, % ^{††}	95 % CI	No.	Prevalence, % ^{††}	95 % CI	No.	Prevalence, % ^{††}	95 % CI	No.	Prevalence, % ^{††}	95 % CI
Total	776	23.4	19.4, 27.5	816	15.6	13.3, 17.8	366	21.2	16.6, 25.8	366	2.5	1.5, 3.5
Age group (years)												
20-39	376	19.4	14.4, 24.5	385	10.2	6.2, 14.3	182	10.6	5.4, 15.7	182	0.4	-0.2, 0.9 ^{¶¶}
40-59	183	23.4	15.2, 31.7	202	33.6	24.3, 42.9	87	27.1	17.5, 36.6	87	2.5	0.3, 4.7 ^{¶¶}
≥ 60	217	36.3	27.0, 45.6	229	69.3	62.2, 76.3	97	46.8	31.7, 62.0	97	8.7	4.0, 13.5
Sex												
Men	292	17.9	12.4, 23.5	324	32.4	26.5, 38.4	142	30.5	21.0, 40.1	142	3.2	1.2, 5.3 ^{¶¶}
Women	484	25.9	20.6, 31.2	492	24.3	19.9, 28.6	224	17.1	12.0, 22.3	224	2.0	0.8, 3.1
Race/ethnicity ^{§§}												
Non-Hispanic White	377	24.6	19.3, 29.9	396	25.5	20.7, 30.3	171	19.8	13.8, 25.8	171	1.6	0.5, 2.6 ^{¶¶}
Non-Hispanic Black	181	23.9	17.1, 30.7	199	39.7	33.5, 45.9	74	23.3	13.6, 33.1	74	4.9	1.6, 8.1 ^{¶¶}
Hispanic	83	16.1	8.7, 23.6	85	21.8	14.4, 29.1	40	29.2	15.5, 42.9	40	3.7	-0.4, 7.5 ^{¶¶}
Other	135	20.1	11.6, 28.6	136	25.5	18.0, 33.0	81	25.1	13.8, 36.3	81	4.1	0.8, 7.5 ^{¶¶}
Education level												
Less than high school	205	27.7	19.2, 36.3	222	34.2	25.3, 43.2	104	24.4	15.3, 33.5	104	3.4	1.1, 6.0 ^{¶¶}
High school graduate	178	27.0	18.3, 35.6	184	34.6	24.5, 44.6	75	29.7	19.4, 40.0	75	2.8	0.3, 5.3 ^{¶¶}
Some college	218	22.9	16.1, 29.6	228	22.8	17.4, 28.1	103	20.6	12.1, 29.2	103	2.7	0.9, 4.5 ^{¶¶}
College graduate or higher	174	17.6	9.1, 26.2	180	18.8	12.8, 24.8	83	13.2	4.8, 21.6 ^{¶¶}	83	0.8	-0.8, 2.4 ^{¶¶}
Chronic kidney disease ^{†††}												
Characteristics	No.	Prevalence, %	95 % CI ^{††}	CVD ^{**}			Having 0 CMD ^{††}			Having at least 2 CMD ^{††}		
				No.	Prevalence, %	95 % CI ^{††}	No.	Prevalence, %	95 % CI ^{††}	No.	Prevalence, %	95 % CI ^{††}
Total	769	7.9	6.9, 8.8	850	6.1	4.3, 7.9	352	50.6	44.1, 57.0	353	12.3	8.1, 16.4
Age group (years)												
20-39	374	10.1	8.5, 11.7	407	1.5	1.4, 1.5	173	65.4	56.7, 74.1	177	8.8	2.3, 15.4 ^{¶¶}
40-59	185	19.6	18.4, 20.7	208	8.6	7.2, 10.1	85	37.6	24.0, 51.2	82	28.4	15.0, 41.9
≥ 60	210	41.0	38.3, 43.8	235	17.0	13.8, 20.2	94	18.7	8.0, 29.3	94	55.0	41.1, 69.0
Sex												
Men	285	14.7	13.9, 15.5	330	4.4	4.0, 4.8	136	44.8	34.4, 55.3	135	17.5	11.2, 23.8
Women	484	19.6	18.1, 21.1	520	7.0	6.0, 8.0	216	51.7	43.7, 59.7	218	24.4	17.0, 31.9
Race/ethnicity ^{§§}												
Non-Hispanic White	371	18.0	16.6, 19.5	410	6.0	5.0, 6.9	169	48.7	40.4, 57.0	166	23.0	15.8, 30.2
Non-Hispanic Black	175	24.6	23.5, 25.8	208	7.5	6.8, 8.1	67	40.2	28.4, 51.9	69	25.9	16.9, 34.8

(Continued)

Table 2. (Continued)

Characteristics	Chronic kidney disease ^a			CVD ^{**}			Having 0 CMD ^{††}			Having at least 2 CMD ^{††}		
	No.	Prevalence, %	95 % CI ^{††}	No.	Prevalence, %	95 % CI ^{††}	No.	Prevalence, %	95 % CI ^{††}	No.	Prevalence, %	95 % CI ^{††}
Hispanic	85	14.5	13.9, 15.1	91	8.6	8.2, 8.9	36	47.5	31.7, 63.3	39	14.6	7.6, 21.7
Other	138	13.6	13.4, 13.8	141	4.4	4.3, 4.5	80	60.5	48.7, 72.2	79	19.7	11.1, 28.3
Education level ^{‡‡‡}												
Less than high school	199	24.8	19.9, 29.8	225	10.1	9.3, 10.9	98	40.9	26.6, 55.1	97	20.9	12.9, 28.9
High school graduate	177	20.8	18.8, 22.9	196	8.1	5.6, 10.6	73	40.8	28.0, 53.6	74	34.5	18.1, 50.9
Some college	219	15.1	14.1, 16.2	241	6.6	6.1, 7.1	98	50.8	38.7, 62.9	102	21.9	13.1, 30.6
College graduate or higher	173	13.6	12.7, 14.5	186	0.9	0.7, 1.1	82	62.8	50.1, 75.5	79	13.0	5.0, 21.0

CMD, cardiometabolic diseases.

^aUnderweight was defined as BMI < 18.5 kg/m².

^bDyslipidemia was defined as having a total cholesterol level ≥ 240 mg/dl or a HDL-cholesterol level < 40 mg/dl for men or < 50 mg/dl for women or self-reported current use of lipid-lowering drugs.

^cHypertension had blood pressure ≥ 130/80 mm Hg or self-reported current use of antihypertensive drugs.

^dPrediabetes had a Hb A1c level of 5.7–6.4 % or a fasting plasma glucose level of 100–125 mg/dl among people without self-reported diabetes.

^eDiabetes was defined as having a self-reported diagnosis of diabetes by a physician or other health professional, a fasting plasma glucose level ≥ 126 mg/dl or a Hb A1c level ≥ 6.5 %.

^fChronic kidney disease was defined as having a urine albumin-to-creatinine ratio ≥ 30 mg/g or an estimated glomerular filtration rate < 60 ml/min/1.73 m².

^gCVD was a composite endpoint of self-reported congestive heart failure, CHD, heart attack and stroke.

^hCMD included dyslipidemia, hypertension, diabetes, chronic kidney disease and CVD.

ⁱEstimates by age group were unadjusted. Other estimates were age-standardised to the 1999–2020 National Health and Nutrition Examination Survey non-pregnant adult population with underweight, using the age groups 20–39 years, 40–59 years and 60 years or older.

^jRace/ethnicity was self-reported. The 'other' group included other non-Hispanic races or multiple races.

Discussion

Among adults with underweight in the USA, only 50.6 % had absence of CMD and 12.3 % lived with at least two CMD. Diabetes and hypertension disproportionately affected non-Hispanic Blacks. The prevalence of CMD was higher in older than younger adults but did not vary by education level. No sex difference was observed.

Adults with underweight are commonly perceived as having a low burden of CMD, but our results did not support this. In our study, only half of adults with underweight had no CMD. Of all the CMD examined, the prevalence of dyslipidemia was the highest achieving about 23 %, followed by hypertension of about 16 %, while the prevalence of diabetes was relatively low. Published studies that reported the prevalence of CMD often merged the underweight adults into the normal-weight group^(20,21) and rarely estimated the prevalence of CMD in underweight separately^(22,23). Only one study conducted specifically among US adults with underweight was identified based solely on self-reported data, reporting a prevalence of CVD of 7.3 % in 2013, similar to the estimate from our study⁽²⁴⁾. The high prevalence of CMD in the underweight population implies that being underweight does not necessarily mean being cardiometabolically healthy⁽²³⁾. A cross-sectional study found that nearly 20 % of the underweight population were classified as metabolically abnormal, defined as having two or more criteria of metabolic syndrome⁽²⁵⁾. Ectopic fat deposition in the liver and pancreas may confer a large role on the development of CMD in the underweight population⁽²⁶⁾.

Similar to the findings from overweight and obese populations, multimorbidity of CMD was also common in underweight based on our study. Unhealthy lifestyle behaviours are well-established risk factors for CMD. Evidence has shown that unhealthy lifestyle behaviours may even cause more severe health problems in non-obese individuals, including those who were underweight, than obese adults⁽²⁷⁾. In our study, the prevalence of CMD was highly prevalent among underweight adults with unhealthy lifestyle behaviours. Of all the lifestyle factors except for sleep duration, compared with underweight adults with a healthier lifestyle, those with a less healthier lifestyle had a 6–16 % higher prevalence of having at least two CMD. Despite the differential influences of each lifestyle factor, the findings suggest that targeting multiple unhealthy lifestyle behaviours may be needed for the prevention and management of CMD among adults with underweight.

Subgroup differences by demographic variables and socio-economic status in the prevalence of CMD were not as widely present in the underweight population as previously reported in general adult populations^(28,29), suggesting that the entire underweight group was at risk of developing CMD. This distinction suggests possibilities of different aetiologies and risk factor profiles for CMD between adults with underweight and those with overweight/obesity, which require future investigations to elucidate. Nonetheless, the disproportionate burden of diabetes in non-Hispanic Black adults with underweight was in line with the racial/ethnic disparities in diabetes well-described in general adult populations^(30,31). Racial/ethnic disparities in the prevalence of CMD found in previous studies were largely due to racial/ethnic disparities in the prevalence of overweight and obesity, social risk factors and lifestyle behaviours^(32,33). The NHANES data did not allow accurate classification of diabetes type, but it is possible that type 1 diabetes accounted for a substantial proportion. To understand pathophysiological mechanisms leading to CMD in underweight, it is critical to understand the causes of underweight

Table 3. Subgroup differences in the prevalence of CMD among adults with underweight* (Prevalence ratio and 95 % confidence intervals)

Characteristics	Dyslipidemia†		FDR-adjusted P value		Hypertension§		FDR-adjusted P value		Prediabetes		FDR-adjusted P value		Diabetes*		FDR-adjusted P value		
	Prevalence ratio†	95 % CI	Prevalence ratio†	95 % CI	Prevalence ratio†	95 % CI	Prevalence ratio†	95 % CI	Prevalence ratio†	95 % CI	Prevalence ratio†	95 % CI	Prevalence ratio†	95 % CI	Prevalence ratio†	95 % CI	
Age group (years)																	
20-39	1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		
40-59	1.15	0.66, 1.98	0.63	1.65, 6.48	3.27	1.65, 6.48	0.001	0.001	2.76	1.41, 5.42	0.009	0.009	8.32	2.12, 32.55	0.009	0.009	
≥ 60	1.81	1.04, 3.16	0.08	3.67, 12.20	6.69	3.67, 12.20	<.001	<.001	4.47	2.46, 8.13	<.001	<.001	25.49	6.07, 106.92	<.001	<.001	
Sex																	
Men	1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		
Women	1.44	0.89, 2.32	0.14	0.57, 1.07	0.78	0.57, 1.07	0.13	0.13	0.55	0.32, 0.96	0.049	0.049	0.72	0.31, 1.64	0.47	0.47	
Race/ethnicity§§																	
Non-Hispanic White	1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		
Non-Hispanic Black	1.04	0.68, 1.60	0.86	1.03, 1.98	1.43	1.03, 1.98	0.12	0.12	0.97	0.43, 2.19	0.95	0.95	3.35	1.35, 8.30	0.045	0.045	
Hispanic	0.65	0.27, 1.58	0.86	0.39, 1.45	0.75	0.39, 1.45	0.60	0.60	1.49	0.78, 2.83	0.72	0.72	3.04	0.63, 14.57	0.21	0.21	
Other	0.85	0.47, 1.54	0.86	0.58, 1.55	0.95	0.58, 1.55	0.83	0.83	1.15	0.61, 2.17	0.95	0.95	2.92	1.18, 7.22	0.045	0.045	
Education level																	
Less than high school	1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		
High school graduate	0.93	0.49, 1.75	0.82	0.67, 1.79	1.09	0.67, 1.79	0.73	0.73	1.38	0.65, 2.92	0.62	0.62	0.91	0.20, 4.06	0.96	0.96	
Some college	0.79	0.44, 1.43	0.66	0.43, 1.15	0.70	0.43, 1.15	0.24	0.24	0.89	0.45, 1.78	0.75	0.75	1.09	0.30, 3.96	0.96	0.96	
College graduate or higher	0.58	0.34, 0.99	0.15	0.37, 1.10	0.64	0.37, 1.10	0.24	0.24	0.59	0.25, 1.40	0.62	0.62	0.40	0.13, 1.23	0.36	0.36	
Chronic kidney disease**																	
Prevalence ratio†		95 % CI		FDR-adjusted P value		CVD††		FDR-adjusted P value		Having 0 CMD##		FDR-adjusted P value		Having at least 2 CMD##		FDR-adjusted P value	
Prevalence ratio†		95 % CI		Prevalence ratio†		95 % CI		Prevalence ratio†		95 % CI		Prevalence ratio†		95 % CI		Prevalence ratio†	
Age group (years)																	
20-39	1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		1.00 (reference)		
40-59	1.87	0.93, 3.78	0.08	1.82, 25.81	6.86	1.82, 25.81	0.005	0.005	0.58	0.37, 0.89	0.02	0.02	3.06	1.47, 6.35	0.007	0.007	

Table 3. (Continued)

Characteristics	Chronic kidney disease**		FDR-adjusted P value		CVD ^{††}		FDR-adjusted P value		Having 0 CMD ^{‡‡}		FDR-adjusted P value		Having at least 2 CMD ^{‡‡}		FDR-adjusted P value
	Prevalence ratio [†]	95% CI	Prevalence ratio [†]	95% CI	Prevalence ratio [†]	95% CI	Prevalence ratio [†]	95% CI	Prevalence ratio [†]	95% CI	Prevalence ratio [†]	95% CI	Prevalence ratio [†]	95% CI	
≥ 60	3.99	2.20, 7.23	<.001	4.12, 40.83	<.001	0.29	0.16, 0.51	<.001	6.20	3.44, 11.19	<.001			<.001	
Sex															
Men	1.00	(reference)			1.00	(reference)			1.00	(reference)			1.00	(reference)	
Women	1.39	0.78, 2.47	0.27	0.82, 3.64	0.15	1.16	0.88, 1.52	0.30	1.44	0.91, 2.29	0.14			0.14	
Race/ethnicity ^{§§}															
Non-Hispanic White	1.00	(reference)			1.00	(reference)			1.00	(reference)			1.00	(reference)	
Non-Hispanic Black	1.49	0.86, 2.57	0.48	0.49, 3.20	0.63	0.86	0.63, 1.18	0.54	1.26	0.76, 2.09	0.56			0.56	
Hispanic	0.79	0.32, 1.96	0.61	0.49, 3.20	0.63	0.99	0.63, 1.56	0.97	0.53	0.31, 0.91	0.09			0.09	
Other	0.78	0.39, 1.57	0.61	0.21, 2.12	0.63	1.26	1.00, 1.58	0.18	0.89	0.55, 1.46	0.66			0.66	
Education level															
Less than high school	1.00	(reference)			1.00	(reference)			1.00	(reference)			1.00	(reference)	
High school graduate	0.83	0.40, 1.72	0.62	0.33, 1.90	0.62	1.00	0.58, 1.71	1.00	1.57	0.83, 2.98	0.36			0.36	
Some college	0.58	0.30, 1.09	0.15	0.19, 1.93	0.59	1.23	0.77, 1.96	0.60	0.99	0.50, 1.98	0.98			0.98	
College graduate or higher	0.56	0.28, 1.10	0.15	0.02, 0.89	0.12	1.49	0.91, 2.43	0.39	0.62	0.29, 1.33	0.36			0.36	

CMD, cardiometabolic diseases; FDR, false discovery rate.

*Underweight was defined as BMI < 18.5 kg/m².

[†]Prevalence ratio was estimated using Poisson regressions, adjusting for age, sex and race/ethnicity when appropriate.

[‡]Dyslipidemia was defined as having a total cholesterol level ≥ 240 mg/dl or a HDL-cholesterol level < 40 mg/dl for men or < 50 mg/dl for women or self-reported current use of lipid-lowering drugs.

[§]Hypertension had blood pressure ≥ 130/80 mm Hg or self-reported current use of antihypertensive drugs.

^{||}Pre-diabetes had a Hb A1c level of 5.7–6.4% or a fasting plasma glucose level of 100–125 mg/dl among people without self-reported diabetes.

^{**}Chronic kidney disease was defined as having a self-reported diagnosis of diabetes by a physician or other health professional, a fasting plasma glucose level ≥ 126 mg/dl or a Hb A1c level ≥ 6.5%.

^{††}CVD was a composite endpoint of self-reported congestive heart failure, CHD, heart attack and stroke.

^{‡‡}CMD included dyslipidemia, hypertension, diabetes, chronic kidney disease and CVD.

^{§§}Race/ethnicity was self-reported. The 'other' group included other non-Hispanic races or multiple races.

^{|||}P values were adjusted using FDR method to control the type I error for multiple comparisons in subgroup differences.

itself. Unlike underweight mainly resulting from inadequate nutrition in many low- and middle-income countries, underweight in the USA may be multifactorial, including malnutrition, chronic diseases and a personal choice due to body image dissatisfaction, among others⁽³⁴⁾. Effective prevention and management of CMD are not possible without correctly understanding the underlying causes of underweight.

Although the underlying contributors to the high prevalence of CMD in underweight populations are poorly understood, improving cardiometabolic health among underweight people is clearly an urgent public health need. Published data on characterising distributions of lifestyle risk factors in underweight are scarce. Our study found that underweight people who had a healthier lifestyle, including non-excessive drinking, more physical activity, never smoking and higher-quality diet, had a lower prevalence of various CMD. These findings suggest that improving lifestyle may be critical to improving cardiometabolic health in people with underweight as in people with overweight or obesity⁽³⁵⁾. Furthermore, evidence has shown that underweight individuals tended to have lower cardiorespiratory fitness compared with those with normal weight^(36,37). Lower cardiorespiratory fitness is known to be associated with higher risks of CMD and mortality^(38,39). Cardiorespiratory fitness can possibly be improved through reducing alcohol intake, eating healthy diet and increasing physical activity, especially resistance training^(39–43), but these data may not be equally applicable to underweight people. Whether such lifestyle modifications would result in a similar improvement in cardiorespiratory fitness specifically in people with underweight requires further investigation.

Strengths and limitations

To our knowledge, this is the first study to comprehensively characterise the landscape of CMD in underweight, using both self-reported and laboratory data from a large nationally representative sample. However, this study has several limitations. First, misdiagnosis of CMD was possible due to the use of self-reported data and one-time laboratory measurements. Second, this was a cross-sectional study; thus, the causal relationship between underweight and CMD cannot be inferred. Third, because of the small sample size, several subgroup estimates had a relative standard error greater than 30 % and thus should be interpreted with caution. Fourth, missing data may have caused bias in some estimates, but we used both multiple imputations and weight adjustment approach to address missing data. Results from these two approaches were similar.

Conclusions

Contrary to the commonly assumed low burden of CMD in the underweight population, nearly half of adults with underweight had at least one CMD, and nearly one-eighth had at least two CMD. Screening of CMD in underweight population may be considered. More resources should be allocated to prevention and management of CMD in this understudied group.

Supplementary material. For supplementary material/s referred to in this article, please visit <https://doi.org/10.1017/S0007114524002885>

Acknowledgements. The authors would like to express our deepest appreciation to all who contributed to this study.

This study was supported by the Innovative Research Team of High-Level Local Universities in Shanghai.

Drs F. Z. and V. W. Z. had full access to all the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis. Concept and design: M. C., F. Z. and V. W. Z. Acquisition, analysis or interpretation of data: all authors. Drafting of the manuscript: M. C. and V. W. Z. Critical revision of the manuscript for important intellectual content: all authors. Statistical analysis: M. C. and V. W. Z. Administrative, technical or material support: F. Z. and V. W. Z.

All authors have completed the ICMJE uniform disclosure form at www.icmje.org/doi_disclosure.pdf and declare: support from the Innovative Research Team of High-Level Local Universities in Shanghai for the submitted work; no financial relationships with any organisations that might have an interest in the submitted work in the previous 3 years; no other relationships or activities that could appear to have influenced the submitted work.

The National Center for Health Statistics Research Ethics Review Board approved NHANES. All participants signed informed consent. Shanghai Jiao Tong University School of Medicine Public Health and Nursing Research Ethics Review Committee approved this study, approval number (SJUPN-202102-B).

The data that support the findings of this study are openly available at <https://www.cdc.gov/nchs/nhanes/>.

References

- Sattar N, Gill JMR & Alazawi W (2020) Improving prevention strategies for cardiometabolic disease. *Nat Med* **26**, 320–325.
- Cheng X, Ma T, Ouyang F, *et al.* (2022) Trends in the prevalence of cardiometabolic multimorbidity in the United States, 1999–2018. *Int J Environ Res Public Health* **19**, 4726.
- Powell-Wiley TM, Poirier P, Burke LE, *et al.* (2021) Obesity and cardiovascular disease: a scientific statement from the American Heart Association. *Circulation* **143**, e984–e1010.
- Milajerdi A, Djafarian K, Shab-Bidar S, *et al.* (2018) Pre- and post-diagnosis body mass index and heart failure mortality: a dose-response meta-analysis of observational studies reveals greater risk of being underweight than being overweight. *Obes Rev* **20**, 252–261.
- Bhaskaran K, Dos-Santos-Silva I, Leon DA, *et al.* (2018) Association of BMI with overall and cause-specific mortality: a population-based cohort study of 3.6 million adults in the UK. *Lancet Diabetes Endocrinol* **6**, 944–953.
- Bhaskaran K, dos-Santos-Silva I, Leon DA, *et al.* (2018) Association of BMI with overall and cause-specific mortality: a population-based cohort study of 3.6 million adults in the UK. *Lancet Diabetes Endocrinol* **6**, 944–953.
- Aune D, Sen A, Prasad M, *et al.* (2016) BMI and all cause mortality: systematic review and non-linear dose-response meta-analysis of 230 cohort studies with 3.74 million deaths among 30.3 million participants. *BMJ* **353**, i2156.
- Fryar C, Carroll M & Afful J (2020) Prevalence of Underweight among Adults Aged 20 and Over: United States, 1960–1962 through 2017–2018. NCHS Health E-Stats. <https://www.cdc.gov/nchs/data/hestat/underweight-adult-17-18/underweight-adult.htm> (accessed June 2024).
- Reedy J, Lerman JL, Krebs-Smith SM, *et al.* (2018) Evaluation of the Healthy Eating Index-2015. *J Acad Nutr Diet* **118**, 1622–1633.
- Shin J-I, Bautista LE, Walsh MC, *et al.* (2015) Food insecurity and dyslipidemia in a representative population-based sample in the US. *Prev Med* **77**, 186–190.
- National Center for Health Statistics (2024) NHANES survey methods and analytic guidelines. Centers for Disease Control and Prevention. <https://www.cdc.gov/nchs/nhanes/analyticguidelines.aspx> (accessed June 2024)
- Afkarian M, Zelnick LR, Hall YN, *et al.* (2016) Clinical manifestations of kidney disease among US adults with diabetes, 1988–2014. *JAMA* **316**, 602–610.
- US Centers for Disease Control and Prevention & National Center for Health Statistics (2024) National Health and Nutrition Examination Survey 1999–2000, 2001–2002, 2003–2004, 2005–2006, and 2017–2020 Documentation Files. <http://www.cdc.gov/nchs/nhanes.htm> (accessed June 2024).

14. Levey AS, Stevens LA, Schmid CH, *et al.* (2009) A new equation to estimate glomerular filtration rate. *Ann Intern Med* **150**, 604–612.
15. Taylor AL, Denniston MM, Klevens RM, *et al.* (2016) Association of hepatitis C virus with alcohol use among U.S. adults: NHANES 2003–2010. *Am J Prev Med* **51**, 206–215.
16. Johnson CL, Paulose-Ram R, Ogden CL, *et al.* (2013) National health and nutrition examination survey: analytic guidelines, 1999–2010. *Vital Health Stat* **2** **161**, 1–24.
17. Särndal C-E, Swensson B & Wretman J (1999) *Sampling Design and Analysis*. Albany, NY: Duxbury Press.
18. Gregg EW, Sorlie P, Paulose-Ram R, *et al.* (2004) Prevalence of lower-extremity disease in the US adult population ≥ 40 years of age with and without diabetes: 1999–2000 national health and nutrition examination survey. *Diabetes Care* **27**, 1591–1597.
19. Lee KJ & Carlin JB (2010) Multiple imputation for missing data: fully conditional specification *v.* multivariate normal imputation. *Am J Epidemiol* **171**, 624–632.
20. Gujral UP, Weber MB, Staimez LR, *et al.* (2018) Diabetes among non-overweight individuals: an emerging public health challenge. *Curr Diab Rep* **18**, 60.
21. Brown CD, Higgins M, Donato KA, *et al.* (2000) Body mass index and the prevalence of hypertension and dyslipidemia. *Obes Res* **8**, 605–619.
22. Holmes L, Hossain J, Ward D, *et al.* (2013) Racial/Ethnic variability in hypertension prevalence and risk factors in National Health Interview Survey. *ISRN Hypertens* **2013**, 257842.
23. Khan SS, Ning H, Wilkins JT, *et al.* (2018) Association of body mass index with lifetime risk of cardiovascular disease and compression of morbidity. *JAMA Cardiol* **3**, 280–287.
24. Park D, Lee J-H & Han S (2017) Underweight: another risk factor for cardiovascular disease?: A cross-sectional 2013 Behavioral Risk Factor Surveillance System (BRFSS) study of 491 773 individuals in the USA. *Medicine (Baltimore)* **96**, e8769.
25. Gao B, Zhang L & Zhao M (2016) Underweight but metabolically abnormal phenotype: metabolic features and its association with cardiovascular disease. *Eur J Intern Med* **29**, 46–51.
26. Thomas EL, Parkinson JR, Frost GS, *et al.* (2012) The missing risk: MRI and MRS phenotyping of abdominal adiposity and ectopic fat. *Obesity (Silver Spring)* **20**, 76–87.
27. Kikuchi A, Monma T, Ozawa S, *et al.* (2021) Risk factors for multiple metabolic syndrome components in obese and non-obese Japanese individuals. *Prev Med* **153**, 106855.
28. He J, Zhu Z, Bundy JD, *et al.* (2021) Trends in cardiovascular risk factors in US adults by race and ethnicity and socioeconomic status, 1999–2018. *JAMA* **326**, 1286–1298.
29. Gerdtz E & Regitz-Zagrosek V (2019) Sex differences in cardiometabolic disorders. *Nat Med* **25**, 1657–1666.
30. Ngo-Metzger Q (2022) Diabetes screening: different thresholds for different racial/ethnic groups. *Ann Intern Med* **175**, 895–896.
31. Kim EJ, Kim T, Conigliaro J, *et al.* (2018) Racial and ethnic disparities in diagnosis of chronic medical conditions in the USA. *J Gen Intern Med* **33**, 1116–1123.
32. Min J, Goodale H, Xue H, *et al.* (2021) Racial-ethnic disparities in obesity and biological, behavioral, and sociocultural influences in the united states: a systematic review. *Adv Nutr* **12**, 1137–1148.
33. Maraboto C & Ferdinand KC (2020) Update on hypertension in African-Americans. *Prog Cardiovasc Dis* **63**, 33–39.
34. Furnham A, Badmin N & Sneade I (2002) Body image dissatisfaction: gender differences in eating attitudes, self-esteem, and reasons for exercise. *J Psychol* **136**, 581–596.
35. Kaminsky LA, German C, Imboden M, *et al.* (2022) The importance of healthy lifestyle behaviors in the prevention of cardiovascular disease. *Prog Cardiovasc Dis* **70**, 8–15.
36. Nikolakaros G, Vahlberg T, Auranen K, *et al.* (2017) Obesity, underweight, and smoking are associated with worse cardiorespiratory fitness in Finnish healthy young men: a population-based study. *Front Public Health* **5**, 206.
37. Lee I & Kim B (2020) Association between estimated cardiorespiratory fitness and all-cause mortality in underweight older adults. *Exerc Sci* **29**, 146–153.
38. Lang JJ, Prince SA, Merucci K, *et al.* (2024) Cardiorespiratory fitness is a strong and consistent predictor of morbidity and mortality among adults: an overview of meta-analyses representing over 20·9 million observations from 199 unique cohort studies. *Br J Sports Med* **58**, 556–566.
39. Ross R, Blair SN, Arena R, *et al.* (2016) Importance of assessing cardiorespiratory fitness in clinical practice: a case for fitness as a clinical vital sign: a scientific statement from the American Heart Association. *Circulation* **134**, e653–e99.
40. Perissiou M, Borkoles E, Kobayashi K, *et al.* (2020) The effect of an 8 week prescribed exercise and low-carbohydrate diet on cardiorespiratory fitness, body composition and cardiometabolic risk factors in obese individuals: a randomised controlled trial. *Nutrients* **12**, 482.
41. Baumeister SE, Finger JD, Gläser S, *et al.* (2018) Alcohol consumption and cardiorespiratory fitness in five population-based studies. *Eur J Prev Cardiol* **25**, 164–172.
42. Kaminsky LA, Myers J, Brubaker PH, *et al.* (2024) 2023 update: the importance of cardiorespiratory fitness in the United States. *Prog Cardiovasc Dis* **83**, 3–9.
43. Kaminsky LA, Lavie CJ, Flint K, *et al.* (2022) Working toward optimal exercise prescription: strength training should not be overlooked. *J Cardiopulm Rehabil Prev* **42**, E32–E33.