



NIH PUBLIC ACCESS

Author Manuscript

Obes Rev. Author manuscript; available in PMC 2013 March 01.

Published in final edited form as:

Obes Rev. 2012 September ; 13(9): 810–821. doi:10.1111/j.1467-789X.2012.01016.x.

The Expanding Burden of Cardiometabolic Risk in China: the China Health and Nutrition Survey

Shengkai Yan^a, Jiang Li^b, Shuang Li^b, Bing Zhang^c, Shufa Du^d, Penny Gordon-Larsen^d, Linda Adair^d, and Barry Popkin^d

^aGuang Zhou Improve Medical Instruments Co., LTD, Guangzhou, China

^bDepartment of Laboratory Medicine, China-Japan Friendship Hospital, No.2 Yinghua Road East, Chaoyang, Beijing, China

^cNational Institute of Nutrition and Food Safety, Chinese Center for Disease Control and Prevention, No 29 Nanwei Road, Beijing, China

^dDepartment of Nutrition, Gillings School of Global Public Health, Carolina Population Center, University of North Carolina, 123 W. Franklin St., Chapel Hill, NC

Abstract

Background—China faces a major increase in cardiovascular disease, yet there is limited population-based data on risk factors, particularly in children.

Methods and Results—Fasting blood samples, anthropometry and blood pressure were collected on 9,244 children and adults aged 7 years in late 2009 as part of the national China Health and Nutrition Survey. Prevalent overweight, elevated blood pressure, and cardiometabolic risk factors: glucose, HbA1c, triglycerides (TG), total cholesterol (TC), high and low density lipoprotein cholesterol (HDL-C and LDL-C), and C-reactive protein (CRP) are presented.

Results—11% of Chinese children and 30% of Chinese adults are overweight. Rates of diabetes, dyslipidemia, hypertension, and inflammation are high and increased with age and were associated with urbanization. Approximately 42% of children have at least one of the following: pre-diabetes or diabetes, hypertension, high TC, LDL-C, TG, and CRP and low HDL-C, as do 70% males and 60% females aged 18–40 years and >86% of males and females 40 years.

Conclusions—HbA1c findings suggest that as many as 29.4 million Chinese children and 415.8 million Chinese adults may be prediabetic or diabetic. The high prevalence in less urban areas and across all income levels suggests that cardiometabolic risk is pervasive across rural and urban China.

Keywords

China; children; diabetes; cardio-metabolic; adults

Corresponding Author: Barry M. Popkin, Carolina Population Center, University of North Carolina, 123 W. Franklin St., Chapel Hill, NC 27516, Phone: 919-966-1732, Fax: 919-966-9159, popkin@unc.edu.

Conflicts of Interest: No author has a conflict of interest

P.G.L. and L.S.A. contributed to study design, P.G.L., B.M.P. and L.S.A. contributed to data analysis, and all authors contributed to writing of the manuscript. P.G.L. and L.S.A. 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. NIH had no role in the design and conduct of the study; collection, management, analysis, and interpretation of the data; and preparation, review, or approval of the manuscript.

Introduction

Obesity rates in China across all age-gender groups have risen greatly in the past two decades with particularly profound increases in BMI in children at the upper portion of the distribution over the past decade^{1, 2}. Concurrently, China has seen a major increase in cardiometabolic risk, including high rates of hypertension, stroke and diabetes³⁻⁵. Underlying factors, such as diet and physical activity have also changed, including increased consumption of refined carbohydrates, saturated fat, and added sugars, and declines in work, leisure and transport-related physical activity^{1, 6}. These changes extend to poor and rural areas, where the most rapid increases in overweight and obesity have recently occurred⁷. With China's rapid economic growth, the burden of obesity has shifted to lower income groups, mirroring what is typically seen in high income countries⁸. Further complicating the disease profile in China is the relatively higher cardiometabolic risk at relatively low body mass index (BMI)⁹ and at younger ages¹⁰ in Asian populations as well as higher insulin resistance relative to Western populations, even at the same level of BMI¹¹.

Using a representative, randomly selected sample from nine provinces across China, with anthropometry, blood pressure measures and fasting blood samples from 9,244 individuals aged 7 and older, we examine age and sex-specific prevalence of overweight, hypertension, and biomarkers of diabetes, dyslipidemia, and inflammation. In addition, we examine differences in cardiometabolic risk factors across levels of urbanization and income. Further, we provide a comparative analysis of cardiometabolic risk relative to patterns in the US and in other Asian countries.

Methods

The China Health and Nutrition Survey (CHNS) is the only large-scale longitudinal, household-based survey in China¹². The CHNS has followed individuals randomly selected from 228 communities and designed to represent a set of large provinces with a range of economic and demographic variation, covering approximately 56% of China's population, including Liaoning, Shandong, Heilongjiang, Henan, Jiangsu, Hubei, Hunan, Guizhou, and Guangxi (from north to south). The CHNS age and gender distribution is slightly older than that of the Chinese census for 2009, since Muslim provinces which have higher fertility rates than other parts of China were not included in the CHNS. Thus, CHNS includes approximately 3% more individuals aged 60 and older and 3% fewer children and adolescents compared to the national China census¹². A multistage, random cluster process was used to draw the sample in each province, with the baseline survey in 1989, and fasting blood collected for the first time in 2009. The analysis sample includes 9,244 individuals aged 7 and older who provided fasting blood and anthropometry in 2009. Unlike other surveys, the number of individuals who provided anthropometry is slightly greater (79 more individuals) than those providing blood samples, likely due to the ordering of data collection. There were no statistically significant differences in overweight and age between individuals providing blood and anthropometry data. The sample of 2009 participants represented 85.6% of the 2006 participants. Survey protocols, instruments, and the process for obtaining informed consent for this study were approved by the institutional review committees of the University of North Carolina at Chapel Hill, the National Institute of Nutrition and Food Safety, Chinese Center for Disease Control and Prevention, and the China-Japan Friendship Hospital, Ministry of Health.

During home visits spanning 3 days, survey data were collected through interviews with each household member, including anthropometrics (height, weight, skinfold thicknesses and body circumferences), detailed individual and household level dietary data, measures of time spent in a wide range of daily activities and detailed individual, household and

community socio-demographic data¹². Individuals older than 7 years visited a neighbourhood clinic to have trained physicians collect fasting blood samples. Individuals unable to attend the clinic had blood samples collected at home. To avoid missing children in boarding schools and migrant workers, special efforts were made to schedule visits in early morning or in the weekend when these participants were at home. All interviewers had 7 days of training provided by the collaborating teams and were overseen via site visits to monitor data collection in each site visit by the University of North Carolina at Chapel Hill, the China Centers for Disease Control and Prevention and the China-Japan Friendship Hospital at selected locations in each province. Measurement of cardiometabolic disease risk factors.

Following an overnight fast, blood was collected by venipuncture and tested immediately for glucose and hemoglobin A1c (HbA1c). Plasma and serum samples were then frozen, and stored at -86°C for later laboratory analysis. All samples were analyzed in a national central lab in Beijing (medical laboratory accreditation certificate ISO 15189:2007) with strict quality control. We focus on blood pressure [systolic and diastolic BP] and biomarkers of cardiometabolic disease risk related to diabetes [fasting glucose measured with the GOD-PAP method (Randox Laboratories Ltd., UK), HbA1c via high-performance liquid chromatography system (model HLC-723 G7; Tosoh Corporation, Tokyo, Japan)]; dyslipidemia [total cholesterol (TC), high and low density lipoprotein cholesterol (HDL-C and LDL-C) all measured using glycerol-phosphate oxidase method and the PEG-modified enzyme method respectively by determiner reagents (Kyowa Medex Co., Ltd, Tokyo, Japan) and triglycerides (TG) using glycerol-phosphate oxidase method and the PEG-modified enzyme method respectively by determiner reagents (Kyowa Medex Co., Ltd, Tokyo, Japan)]. All lipids measures were on the Hitachi 7600 automated analyzer (Hitachi Inc., Tokyo, Japan); inflammation [high sensitivity C-reactive protein (CRP)] via the immunoturbidimetric method with Denka Seiken, Japan reagents (Hitachi 7600 automated analyzer, Hitachi Inc., Tokyo, Japan). Levels of biomarkers were categorized to represent risk using cutoff points recommended by the International Diabetes Federation (IDF)¹³ or in other published literature, with separate cutoff points for adults and children, and for males and females where appropriate (Table 1).

Systolic (SBP) and diastolic (DBP) blood pressure were measured on the right arm, using mercury sphygmomanometers with appropriate cuff sizes. Measures were collected in triplicate after a 10 minute seated rest and the mean of the three measurements was used in analyses. Adult hypertension was determined (according to the IDF cut point (SBP/DBP 140/90 mmHg) or taking blood pressure medication. For youth <18 years, hypertension risk was defined as blood pressure above the 85th age, sex, and height-specific reference¹⁴.

Overweight. Height was measured without shoes to the nearest 0.1 cm using a portable SECA stadiometer; weight was measured without shoes and in light clothing to the nearest 0.1 kg on a calibrated beam balance. For adults, overweight is defined as BMI $\geq 25\text{ kg/m}^2$ and obesity as BMI $\geq 30\text{ kg/m}^2$ ¹⁵. For youth, age- and sex-specific reference data for the BMI $\geq 25\text{ kg/m}^2$ and BMI $\geq 30\text{ kg/m}^2$ equivalent cut-points from the International Obesity Task Force were used to classify overweight and obesity to be consistent with the adult definitions¹⁶.

We also present population-specific BMI percentile distribution data by age to characterize trends in BMI in children and adolescents in China over the past two decades. Using quantile regression^{17, 18}, we characterize the mean BMI value associated with the population-specific 95th percentile in 10 and 15 year olds from 1991 to 2009 based on a regression of BMI data against age and age squared.

Other variables—Age was recorded as the respondent's age on the date of exam and categorized into the following groups: 7–11, 12–17, 18–29, 30–39, 40–49, 50–59, 60–69, and 70 years). Urbanicity is defined using a multidimensional 12 component urbanization index that captures community-level physical, social, cultural, and economic environments and which represents the heterogeneity that would be otherwise missed in a urban/rural measure based only on population density (mean=67.2; SD=19.5)¹⁹. A high urbanization index represents a large population living closely together in a physical environment providing an efficient transport system; a communication network; good-quality health care; higher-level education; water, sewer, and electric lines. Household income is based on detailed measures of all income-earning activities of all household members (mean=37,979, SD=45,584 yuan). Tertiles of the urbanization index are used to define low, medium, and high urbanicity and household income per capita.

Comparative cardiometabolic risk data—For wider comparison with the US, we used most recently published data from the National Health and Nutrition Survey^{20, 21, 22} and from the SEARCH for Diabetes in Youth study.²³ We used nationally representative data from the Korean Health and Nutrition Survey for individuals 10 and older collected in 2009, which we analyzed for this paper. We conducted a systematic other literature search for using the following key words alone and in combination: non-insulin-dependent diabetes mellitus, type 2 diabetes mellitus, inflammation, children, adolescents, youth, and then for adults to obtain nationally representative studies from Asia on this topic. We limited our review only to national or very large-scale studies, which limited the number or included studies.

Statistical Analyses

Statistical analyses were conducted using Stata (Release 11.0 SE, Stata Corporation, College Station, TX). For descriptive analyses, percentages were calculated for categorical variables, while means were calculated for continuous variables. The nonparametric test for trend was used to test for age differences in cardiometabolic risk factors. Logistic regression models were used to test for differences in risk factor prevalence by income or urbanization in age and sex-stratified models.

Results

Approximately 12% of children and adolescents were overweight (Table 2). Obesity was higher among the 7–11 year olds (3%) than among the 12–17 year olds (1%). Among adults, overweight increased with age with highest rates in middle aged adults; approximately 30% were overweight and 4% obese in the full sample.

All cardiometabolic risk factors (except low HDL-C in males) increased with age (p for trend, $p < 0.0001$), Figures 1A–1C. Pre-diabetes (IFG and impaired HbA1c) and diabetes were highest in individuals older than 50 years (Figure 1A). Diabetes prevalence was similar when based on fasting glucose or HbA1c, although pre-diabetes was higher using HbA1c. Markers of dyslipidemia were comparatively high in females than males, with particularly high prevalence of dyslipidemia in individuals over the age of 40 years (Figure 1B). Similarly, elevated blood pressure and inflammation were comparatively higher in individuals over the age of 40 years (Figure 1C). Values for all risk markers are shown in Supplemental Table 1.

Of major public health relevance is the fact that in the pediatric population, 42% of children aged 7–17 had at least one of the following cardiometabolic risk factors: pre-diabetes/diabetes (HbA1c $\geq 5.7\%$), hypertension, high TC, high LDL, low HDL, high TG, high CRP. In contrast, among adults aged 18 to 40 years, 70% of males and 60% of females had at least

one risk factor, whereas approximately 83% of 40–60 year olds and over 90% of those 60 years and older had at least one elevated risk factor (Figure 2).

The observed elevated cardiometabolic risk factors at younger ages are consistent with distribution of overweight in the CHNS pediatric population. Elsewhere, we have shown that the mean BMI at the Chinese BMI 95th population-specific percentile at age 6 in 2006 was 24.8 kg/m², whereas the US 95th population-specific percentile for 6 year olds was 22.2 kg/m².¹ Looking across the past two decades, the increase in mean BMI at the population-specific BMI for China was biggest between 1991 and 2000 and between 2006 and 2009 and that the increase has been comparatively larger in 10 year olds than 13 year olds.

After adjusting for age, higher urbanization was associated with higher levels of most risk factors, although there were some sex differences in which specific markers related to urbanicity, particularly in individuals over the age of 40 years (Table 4). Urbanization was unrelated to impaired HbA1c and prehypertension (males and females). Comparing older (>40 yr) to younger (<40) individuals, some interesting relationships emerged. The positive association of urbanization was stronger or seen only among older individuals for overweight, IFG, diabetes (males and females), and low HDL (males only). The direction of association of urbanization with risk indicators was opposite in older versus younger females for high TG, and prehypertension. For these risk factors, prevalence was lower in the highest urbanization category among younger women, but higher in the highest urbanization category among older women.

Associations of elevated cardiovascular risk factors with income were less consistent. Higher income was associated with higher prevalence of overweight, diabetes and high TG only in males. Income was unrelated to lipids (except LDL in highest tertile), prehypertension, or CRP in males, and unrelated (positively) to all markers except overweight in the highest tertile in females. Higher income was associated with lower risk of high TG and high CRP in younger women, but with lower risk of diabetes and hypertension in older women (Table 4).

Comparison of CHNS findings to findings in the United States and other Asian countries

We compare our CHNS findings with other nationally representative studies from the US and the few published studies in Asia (Table 4). In Table 4A we show that diabetes and inflammation prevalence was higher in CHNS adolescents than in US adolescents. Although in the CHNS study we are unable to distinguish Type 1 and Type 2 diabetes, available evidence for East Asia suggests almost no Type 1 insulin dependent diabetes among youth^{24–26}. Comparisons of diabetes across China, South Korea and Taiwan (the only other Asian countries with nationally representative data), suggests higher rates in China (Table 4B). For adults, on the other hand, diabetes prevalence is higher in South Korea, Hong Kong, Singapore, and Malaysia than in China (Table 4C).

Discussion

The observed findings indicate strong age-related differences in overweight and cardiometabolic risk. These estimates highlight the huge burden that the health care system will need to confront. Of particular concern is the relatively high risk in the pediatric population, important not only because of current impact on child health, but also because risk tends to track into adulthood and even worsen. Based on HbA1c, the currently recommended indicator for diabetes²⁷, we observed rates of diabetes of 0.9% and pre-diabetes of 14.9% in those aged 7–17 years. Although the CHNS data do not match the Chinese census (the CHNS sample has 3% more adults aged 60 and older and 3% fewer under the age of 19), we can approximate the numbers of individuals affected using Chinese

census data from 2009²⁸. Using the census data, our findings suggest that about 1.7 million Chinese children under the age of 18 are diabetic and about 27.7 million are pre-diabetic. The high rates of pre-diabetes as measured by HbA1c in youth (15%) and adults (30%) portend a public health problem of massive proportion. Furthermore, over one-third of children under the age of 18 had high levels of at least one cardiometabolic risk factor and rates increase to 85% in those >40 years. Of additional concern is the high prevalence of cardiometabolic risk in less urban areas and across all income levels.

Of additional concern is the observation that diabetes and inflammation rates are higher in the Chinese pediatric population than in the US pediatric population and in other Asian countries. In contrast, diabetes rates among adults were comparable to those in the US and other Asian countries. Furthermore, obesity and pre-diabetes measured by HbA1c were twice as high in 7–11 versus 12–18 year olds, it is possible that the younger cohorts who are growing up in a very different environmental setting than the adolescents and adults grew up in and are thus experiencing more dramatic health consequences than the adolescent Chinese population. Coupled with the higher BMI values at the upper end of the distribution in Chinese youth, relative to that in the US, UK and Australia¹, the future health of the current pediatric generation of Chinese youth is concerning, particularly given the higher cardiometabolic risk at lower BMI in Asians^{29–31}.

We also observed age-related increases in cardiometabolic risk across all markers (with the exception of low HDL in males), paralleling the increase in overweight in this population. Diabetes, hypertension and inflammation were higher at older ages, despite a drop off in overweight. More than three quarters of adults had at least one elevated cardiometabolic risk factor, which increased to over 90% in individuals aged 60 and older. A recent study of diabetes in a large national study of Chinese adults, ages 20 yr and older⁵, used fasting and 2-hour glucose levels to define impaired fasting glucose and diabetes. We calculated age- and sex-adjusted estimates (not shown) in the same age range as Yang to provide comparison between our results and theirs. Our overall age and sex-adjusted estimate of diabetes for adults aged 20 and older based on HbA1c (6.7%) is lower than the Yang age- and sex-standardized estimate of 9.7% for the same age group. In comparison with Yang et al., we observed an age- and sex-adjusted estimate of pre-diabetes of 30%, whereas Yang et al., found 15.5%. Compared to the fasting glucose cutoff point of 100 mg/dl (5.6 mmol/l), the HbA1c cutoff point of 5.7% is less sensitive but more specific and has a higher positive predictive value to identify people at risk for later development of diabetes³². Consistent with their study, we find similar age distributions and higher rates of diabetes in men compared to women. Our population study of cardiometabolic risk factors in Chinese adults is the most current data available.

In addition to age-related increases in risk and high rates of cardiometabolic risk in youth, we observed high prevalence of risk in less urban areas and in across all income levels, suggesting pervasive risk across all parts of China. This is of major concern as these risk factors were heretofore less common in rural China. In prior work, we have shown that the burden of overweight is shifting from the rich to the poor in China and other countries, and we have found comparatively rapid increases in obesity in rural and lower income populations⁷. In addition, we have shown an increased burden of obesity among women, but an opposite trend in men⁸. In the past in low to middle income countries, higher levels of chronic disease risk were associated with affluence. Our findings suggest that as cardiometabolic risk increases in China, even rural populations will be affected.

There are a few limitations to this analysis. This cross-sectional analysis does not examine temporal changes in cardiometabolic risk factors owing to the fact that biomarker data were only collected in the 2009 round of the CHNS. Despite the overall large sample size, some

age and sex-specific subgroups are small, and thus may provide unstable estimates of prevalence for rarer outcomes. Our main objective was to determine the prevalence of a wide range of cardiometabolic risk factors by age and gender, and across urbanization and income. To provide comparability, we used conventional, anthropometric and cardiometabolic cutpoints. Nevertheless they may not represent true risk in this population if, for example, disease develops at lower BMI in Asian versus other populations, which is possible given higher visceral fat at the same level of BMI in Asians versus Europeans⁹. We used the BMI cutoff points of 25 and 30 kg/m² for adults and the IOTF 25 and 30 kg/m² equivalent for youth to allow for comparison of youth and adults. We did not exclude obese individuals from this analysis as we were interested in estimating risk in the full Chinese population; nonetheless when we exclude the 369 obese individuals in the sample from the analysis we find very similar results that with rounding are almost identical. We are not able to distinguish type 1 from type 2 diabetes in our sample, although no children reported insulin injection use. Whereas it is possible that the comparatively higher rates of pre-diabetes as measured by HbA1c in children versus adolescents could relate to type 1 diabetes, the rate of type 1 diabetes in China is among the lowest in the world, with an estimate of 0.1 per 100,000/year³³. The differences in pre-diabetes and diabetes as measured by HbA1c versus fasting glucose in youth, could possibly relate to differential meaning of these measures in youth or transient increases in insulin resistance with puberty^{34–37}. While we include comparative data from several countries, we were limited in the availability of published data, particularly in South East and South Asian countries.

Important strengths of the study include its population based design, representation of a wide range of urban and rural communities across China, its inclusion of fasting blood data for children down to the age of 7 year, and the wide range of biomarkers collected. The data from this national study suggests a high prevalence of overweight and elevated cardiometabolic risk factors in China that is higher across age groups and is evident even in the pediatric population. Further, rates were markedly high even in less urban and lower income groups within China. Our HbA1c results suggest a very high burden of chronic disease risk starting at a young age, with 1.7 million Chinese children 7–18 having diabetes (HbA1c), another 27.7 million considered pre-diabetic. For adults, the observed findings that suggest 82.1 million adults have diabetes and 334 million are pre-diabetic, are higher than that of Yang⁵. Further we provide evidence that this is just the beginning of high prevalence of elevated cardiometabolic risk factors, given the increase in overweight with age. Elsewhere we have discussed some of the economic burden of this coming crisis⁶. The observed findings foreshadow substantial and increasing chronic disease burden in China over the next several decades.

Acknowledgments

We thank Ms. Frances Dancy, BS, UNC Carolina Population Center for her helpful administrative assistance, Jennifer Poti, BS for helpful research assistance, Jim Terry, AB, Phil Bardsley, PhD, Donna Miles, PhD, and Dan Blanchette, BA for programming and technical support. None of the individuals acknowledged received compensation for any role in the study.

Funding

NIH (R01-HD30880, DK056350, R24 HD050924, R01-HD38700, R01-HL108427, and R21DK089306) with added financial support from the Chinese Center for Disease Control and Prevention National Institute of Nutrition and Food Safety and the China-Japan Friendship Hospital, Ministry of Health of China.

References Cited

1. Popkin BM. Recent dynamics suggest selected countries catching up to US obesity. *Am J Clin Nutr.* 2010; 91:284S–88S. [PubMed: 19906804]

2. Popkin BM, Adair LS, Ng SW. Global nutrition transition and the pandemic of obesity in developing countries. *Nutr Rev.* 2012; 70:3–21. [PubMed: 22221213]
3. Lee CM, Huxley RR, Lam TH, et al. Prevalence of diabetes mellitus and population attributable fractions for coronary heart disease and stroke mortality in the WHO South-East Asia and Western Pacific regions. *Asia Pac J Clin Nutr.* 2007; 16:187–92. [PubMed: 17215197]
4. Gu D, Gupta A, Muntner P, et al. Prevalence of cardiovascular disease risk factor clustering among the adult population of China: results from the International Collaborative Study of Cardiovascular Disease in Asia (InterAsia). *Circulation.* 2005; 112:658–65. [PubMed: 16043645]
5. Yang W, Lu J, Weng J, et al. Prevalence of diabetes among men and women in China. *N Engl J Med.* 2010; 362:1090–1101. [PubMed: 20335585]
6. Popkin BM. Will China's nutrition transition overwhelm its health care system and slow economic growth? *Health Aff (Millwood).* 2008; 27:1064–76. [PubMed: 18607042]
7. Dearth-Wesley T, Wang H, Popkin BM. Obesity dynamics in China: The poor are catching up. *Eur J Clin Nutr.* 2007; 18:1–6.
8. Jones-Smith JC, Gordon-Larsen P, Siddiqi A, Popkin BM. Emerging disparities in overweight by educational attainment in Chinese adults (1989–2006). *Int J Obes.* 2011 10.1038/ijo.2011.134
9. WHO Expert Consultation. Appropriate body-mass index for Asian populations and its implications for policy and intervention strategies. *Lancet.* 2004; 363:157–163. [PubMed: 14726171]
10. Chan JCN, Malik V, Jia W, et al. Diabetes in Asia: epidemiology, risk factors, and pathophysiology. *JAMA.* 2009; 301:2129–2140. [PubMed: 19470990]
11. Deurenberg P, Deurenberg-Yap M, Guricci S. Asians are different from caucasians and from each other in their body mass index/body fat per cent relationship. *Obes Rev.* 2002; 3:141–6. [PubMed: 12164465]
12. Popkin BM, Du S, Zhai F, Zhang B. Cohort Profile: The China Health and Nutrition Survey--monitoring and understanding socio-economic and health change in China, 1989–2011. *Int J Epidemiol.* 2009; 39:1435–1440. [PubMed: 19887509]
13. International Diabetes Federation. 2006. The IDF consensus worldwide definition of the metabolic syndrome http://www.idf.org/webdata/docs/MetS_def_update2006.pdf
14. National High Blood Pressure Education Program. The fourth report on the diagnosis, evaluation, and treatment of high blood pressure in children and adolescents. *Pediatrics.* 2004; 114:555–576. [PubMed: 15286277]
15. World Health Organisation International Association for the Study of Obesity International Obesity TaskForce. The Asia-Pacific perspective: redefining obesity and its treatment. *Health Communications; Sydney.* 2000. p. 1-56.
16. Cole TJ, Bellizzi MC, Flegal KM, Dietz WH. Establishing a standard definition for child overweight and obesity worldwide: international survey. *Br Med J.* 2000; 320:1240–1243. [PubMed: 10797032]
17. Koenker R, Hallock, Kevin F. Quantile Regression. *J Econ Perspect.* 2001; 15:143–156.
18. Cole TJ, Green PJ. Smoothing reference centile curves: the LMS method and penalized likelihood. *Stat Med.* 1992; 11:1305–1319. [PubMed: 1518992]
19. Jones-Smith JC, Popkin BM. Understanding community context and adult health changes in China: development of an urbanicity scale. *Soc Sci Med.* 2011; 71:1436–1446. [PubMed: 20810197]
20. Martinson ML, Teitler JO, Reichman NE. Health across the life span in the United States and England. *Am J Epidemiol.* 2011; 173:858–865. [PubMed: 21389038]
21. Dowd JB, Zajacova A, Aiello AE. Predictors of inflammation in U.S. children aged 3–16 years. *Am J Prev Med.* 2010; 39:314–320. [PubMed: 20837281]
22. Johnson WD, Kroon JJ, Greenway FL, Bouchard C, Ryan D, Katzmarzyk PT. Prevalence of risk factors for metabolic syndrome in adolescents: National Health and Nutrition Examination Survey (NHANES), 2001–2006. *Arch Pediatr Adolesc Med.* 2009; 163:371–377. [PubMed: 19349567]
23. SEARCH for Diabetes in Youth Study Group. The burden of diabetes mellitus among us youth: prevalence estimates from the SEARCH for Diabetes in Youth Study. *Pediatrics.* 2006; 118:1510–1518. [PubMed: 17015542]

24. Kitagawa T, Owada M, Urakami T, Tajima N. Epidemiology of type 1 (insulin-dependent) and type 2 (non-insulin-dependent) diabetes mellitus in Japanese children. *Diabetes Res Clin Practice Suppl.* 1994; 24:S7-13.R.
25. Wei JN, Chuang LM, Lin CC, Chiang CC, Lin RS, Sung FC. Childhood diabetes identified in mass urine screening program in Taiwan, 1993–1999. *Diabetes Res Clin Practice.* 2003; 59:201–206.
26. Yang Z, Wang K, Li T, et al. Childhood diabetes in China: enormous variation by place and ethnic group. *Diabetes Care.* 1998; 21:525–529. [PubMed: 9571336]
27. International Expert Committee report on the role of the A1C assay in the diagnosis of diabetes. *Diabetes Care.* 2009; 32:1327–1334. [PubMed: 19502545]
28. National Bureau of Statistics of China. Beijing: China Statistics Press. 2010. *China Statistical Yearbook 2010.*
29. Nguyen TT, Adair LS, Suchindran CM, He K, Popkin BM. The association between body mass index and hypertension is different between East and Southeast Asians. *Am J Clin Nutr.* 2009; 89:1905–1912. [PubMed: 19369374]
30. Nguyen T, Adair LS, Stevens J, Popkin BM. Prediction of hypertension by different anthropometric indices in adults: the change in estimate approach. *Public Health Nutr.* 2010; 13:639–646. [PubMed: 19758482]
31. World Health Organization Expert Consultation. Appropriate body-mass index for Asian populations and its implications for policy and intervention strategies. *Lancet.* 2004; 363:157–163. [PubMed: 14726171]
32. American Diabetes Association. Diagnosis and classification of diabetes mellitus. *Diabetes Care Suppl.* 2010; 33:62–69.
33. The DIAMOND Project Group. Incidence and trends of childhood Type 1 diabetes worldwide 1990–1999. *Diabetic Med.* 2006; 23:857–866. [PubMed: 16911623]
34. Saaddine JB, Fagot-Campagna A, Rolka D, et al. Distribution of HbA(1c) levels for children and young adults in the U.S: Third National Health and Nutrition Examination Survey. *Diabetes Care.* 2002; 25:1326–1330. [PubMed: 12145229]
35. Moran A, Jacobs DR Jr, Steinberger J, et al. Insulin resistance during puberty: results from clamp studies in 357 children. *Diabetes.* 1999; 48:2039–2044. [PubMed: 10512371]
36. Eldeirawi K, RBL. Predictors of hemoglobin A1c in a national sample of nondiabetic children: the Third National Health and Nutrition Examination Survey, 1988–1994. *Am J Epidemiol.* 2003; 157:624–632. [PubMed: 12672682]
37. Shultis WA, Leary SD, Ness AR, et al. Haemoglobin A1c is not a surrogate for glucose and insulin measures for investigating the early life and childhood determinants of insulin resistance and Type 2 diabetes in healthy children. An analysis from the Avon Longitudinal Study of Parents and Children (ALSPAC). *Diabetic Med.* 2006; 23:1357–1363. [PubMed: 17116188]
38. Wei J-N, Sung F-C, Lin C-C, Lin R-S, Chiang C-C, Chuang L-M. National surveillance for type 2 diabetes mellitus in Taiwanese children. *JAMA.* 2003; 290:1345–1350. [PubMed: 12966126]
39. Chan JCN, Malik V, Jia W, et al. Diabetes in Asia. *JAMA.* 2009; 301:2129–2140. [PubMed: 19470990]
40. Ministry of Health. WHO Global Infobase: National Health Survey 2004. 2005. Singapore: World Health Organization; 2005. Web site. <http://www.who.int/infobase/>
41. Mustafa N, Kamarudin NA, Ismail AA, et al. Prevalence of abnormal glucose tolerance and risk factors in urban and rural Malaysia. *Diabetes Care.* 2011; 34:1362–1364. [PubMed: 21498788]

Figure 1A.

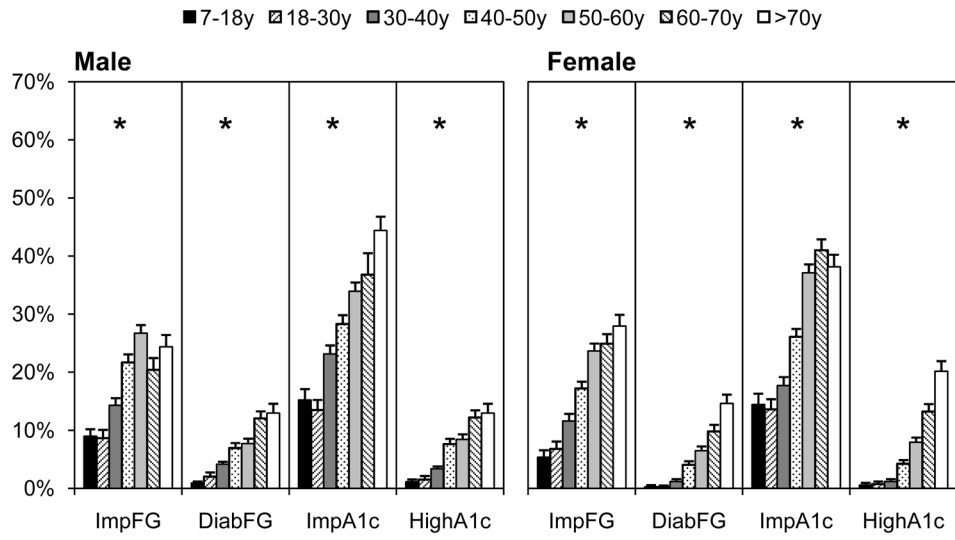


Figure 1B.

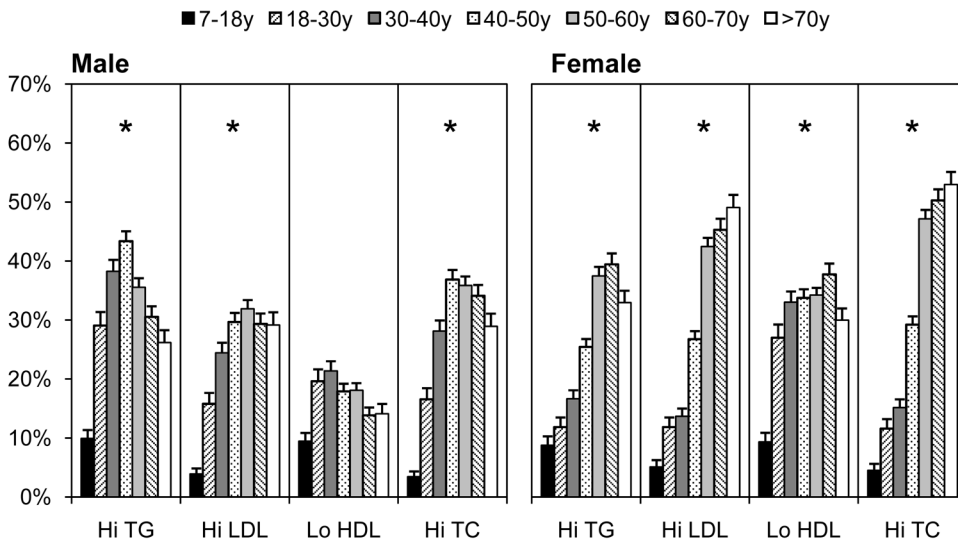


Figure 1C.

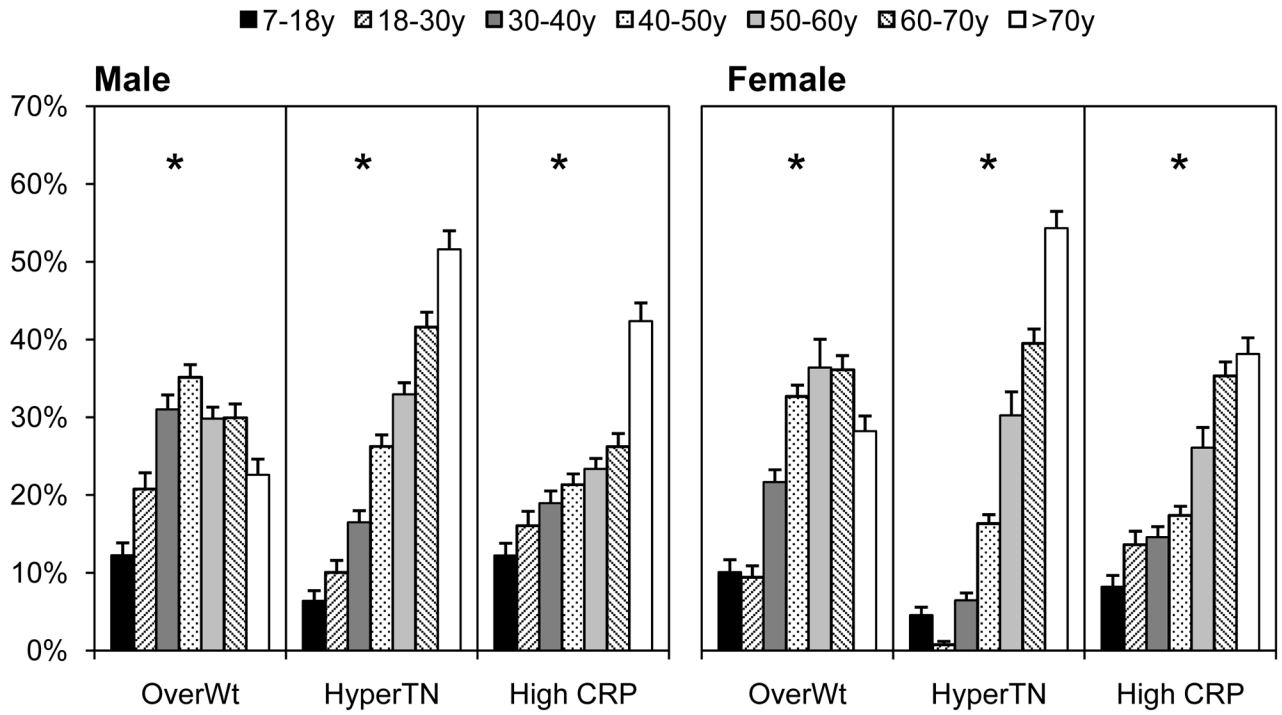


Figure 1.
 A–C. Cardiometabolic Risk Across Several Indicators by Age, 2009 China Health And Nutrition Survey
 Cutpoints shown in Table 1, abbreviations below
 Over WT overweight
 Imp FG impaired fasting glucose
 Diab FG, diabetes based on fasting glucose
 Imp A1c, impaired HbA1c
 High A1c, hemoglobin HbA1c
 High TG, triglycerides
 High LDL, high low density lipoprotein cholesterol
 Low HDL, low high density lipoprotein cholesterol
 High TC, high total cholesterol
 Hyper TN hypertension
 High CRP, Inflammation measured by C-reactive protein

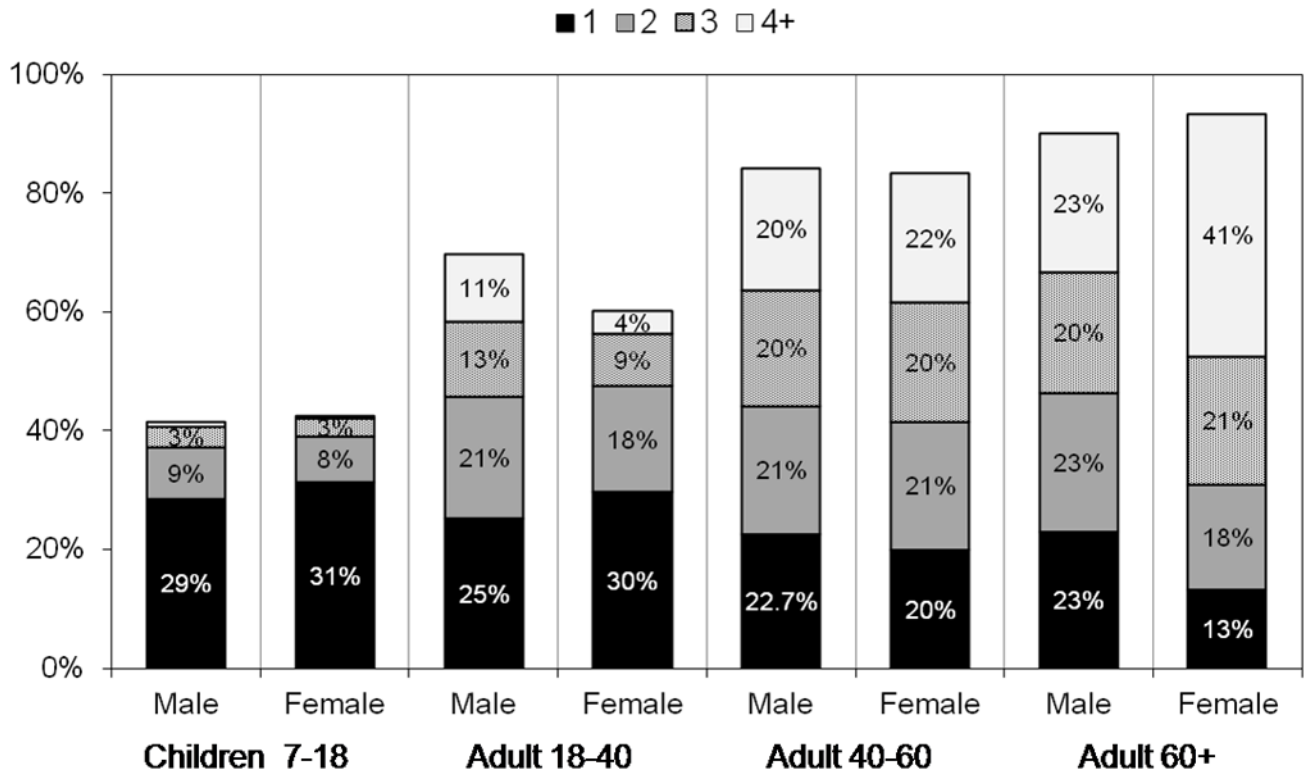


Figure 2. Proportion of The CHNS Sample with at Least one Elevated Cardiometabolic Risk Factor by Age Group (age 7–17 Years; age 18–39 Years, age 40–60 years; and age 60 Years). Proportion of the population with one or more of the following cardiometabolic risk factors: impaired/diabetic (HbA1c $\geq 5.7\%$), hypertension (140/90, or age-sex specific percentiles for pediatric population), high TC (≥ 200 mg/dL), high LDL (>130 mg/dL), low HDL (<40 mg/dL), high TG (≥ 150 mg/dL), high CRP (High CRP (≥ 3)).

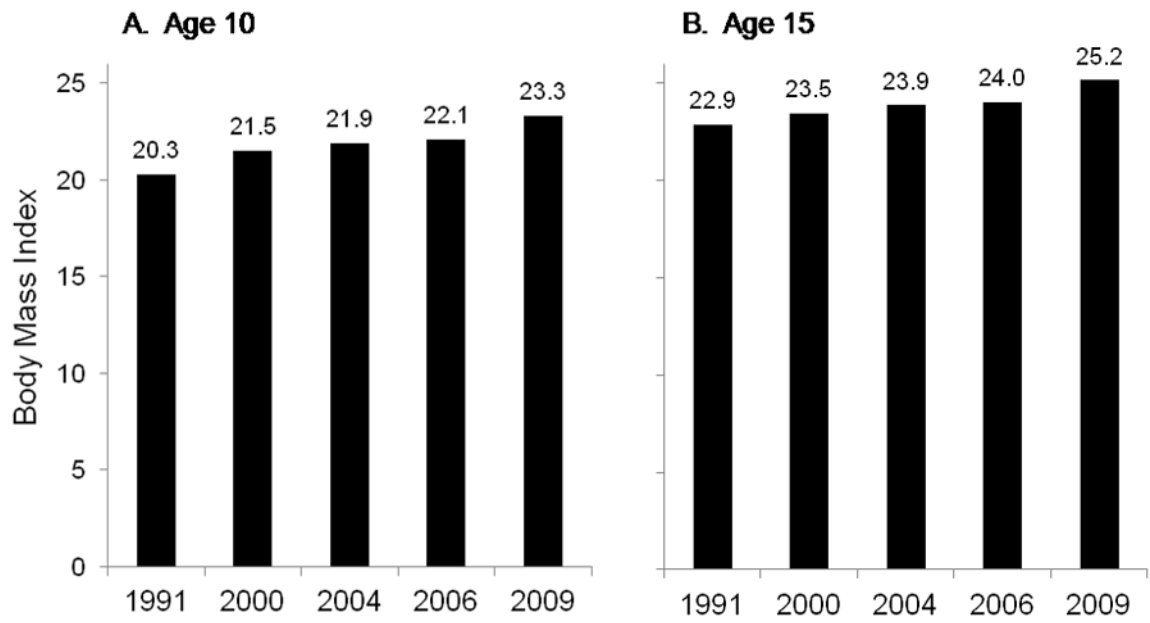


Figure 3.
Mean BMI value associated with the population-specific 95th percentile in 10 and 15 year olds from 1991 to 2009
Based on quantile regression models of BMI data against age and age squared

Table 1

Biomarkers, laboratory analyses methods and definitions of cardiometabolic risk, 2009 China Health and Nutrition Survey fasting blood samples

Biomarker	Risk Indicator	Definition
Glucose	Impaired fasting glucose	Glucose 100 & < 126 mg/dL
	Diabetes	Glucose 126 mg/dL or taking diabetes medication
HbA1C	Impaired glucose control	HbA1c 5.7 & < 6.5%
	Diabetes	HbA1c 6.5% or taking diabetes medication
Total Cholesterol	High TC	200 mg/dL
HDL-C	Low HDL	Male: < 40 mg/dL; female < 50 mg/dL; pediatric < 40 mg/dL
LDL-C	High LDL	> 130 mg/dL
Triglycerides	High TG	150 mg/dL
High Sensitivity CRP	High GRP	3 mg/dL

Table 2

Mean BMI and percent overweight and obese in CHNS participants

Age Group	N 4,417	Mean BMI (SD)	Overweight [BMI 25 (or pediatric equivalent ¹⁸); % (95% CI)]	Obesity [BMI 30 (or pediatric equivalent ¹⁸); % (95% CI)]
MALES				
7–11* y	211	17.0 (2.9)	12.8 (10.5, 15.1)	3.2 (2.0, 4.4)
12–17* y	205	19.0 (3.4)	11.7 (9.5, 14.0)	1.4 (0.6, 2.2)
Total 7–17 y	416	18.0 (3.4)	12.3 (10.6, 13.9)	2.3 (1.6, 3.0)
18–29 y	374	22.2 (3.5)	20.8 (18.7, 22.9)	2.3 (1.5, 3.1)
30–39 y	611	24.0 (3.5)	31.0 (29.1, 32.9)	4.5 (3.7, 5.3)
40–49 y	857	24.0 (3.3)	35.2 (33.5, 36.8)	4.8 (4.1, 5.5)
50–59 y	973	23.5 (3.2)	29.8 (28.4, 31.3)	3.4 (2.8, 3.9)
60–69 y	659	23.2 (3.4)	30.0 (28.2, 31.7)	3.3 (2.6, 4.0)
70 y	431	22.5 (4.0)	22.6 (20.6, 24.6)	4.1 (3.2, 5.0)
Total 18 y	3905	23.3 (3.4)	29.5 (28.8, 30.3)	3.8 (3.5, 4.1)
Age Group	N 4,827	Mean BMI (SD)	Overweight [BMI 25 (or pediatric equivalent ¹⁸); % (95% CI)]	Obesity [BMI 30 (or pediatric equivalent ¹⁸); % (95% CI)]
FEMALES				
7–11 y*	160	16.3 (3.0)	12.2 (9.6, 14.8)	3.0 (1.7, 4.3)
12–17 y*	184	19.0 (3.0)	8.2 (6.2, 10.2)	1.1 (0.3, 1.8)
Total 7–17y	344	18.0 (3.2)	10.1 (8.5, 11.7)	2.0 (1.2, 2.7)
18–29 y	392	21.0 (3.0)	9.4 (8.0, 10.9)	0.8 (0.3, 1.2)
30–39 y	665	23.0 (3.4)	21.7 (20.1, 23.3)	3.9 (3.1, 4.6)
40–49 y	1039	24.0 (3.3)	32.7 (31.2, 34.1)	4.4 (3.8, 5.1)
50–59 y	1098	24.0 (3.3)	36.4 (35.0, 37.9)	4.6 (4.0, 5.2)
60–69 y	697	24.0 (4.0)	36.1 (34.3, 38.0)	5.7 (4.8, 6.6)
70 y	519	23.1 (4.0)	28.2 (26.2, 30.2)	5.6 (4.6, 6.5)
Total 18 y	4410	23.4 (4.0)	29.9 (29.2, 30.6)	4.4 (4.1, 4.7)
Age Group	N	Mean BMI (SD)	Overweight [BMI 25 (or pediatric equivalent ¹⁸); % (95% CI)]	Obesity [BMI 30 (or pediatric equivalent ¹⁸); % (95% CI)]
TOTAL POPULATION				
Pediatric	760	17.7 (3.3)	11.3 (10.1, 12.4)	2.2 (1.6, 2.7)
Adult	8315	23.4 (3.5)	29.7 (29.2, 30.2)	4.1 (3.9, 4.3)

*7–17 year old age group broken into two categories given variation in growth for children and adolescents

Table 3

Association of urbanization (first panel) and income (second panel) with cardiometabolic risk in younger and older adults.

		Prevalence Odds ratios from logistic regression models, stratified by age			
		Males		Females	
Urbanization		Age 18–40 years n=1,014	Age 40 years n=2,970	Age 18–40 years n=1,068	Age 40 years n=3,406
		Overweight	Low	1.00	1.00
Med	1.16 (0.82, 1.64)		1.70 (1.39, 2.08)	0.84 (0.58, 1.24)	1.23 (1.03, 1.47)
Hi	1.22 (0.86, 1.73)		1.86 (1.53, 2.27)	0.67 (0.45, 1.00)	1.20 (1.01, 1.43)
IFG	Low	1.00	1.00	1.00	1.00
	Med	0.84 (0.51, 1.38)	1.45 (1.16, 1.82)	1.31 (0.75, 2.30)	1.55 (1.24, 1.93)
	Hi	1.17 (0.73, 1.87)	1.72 (1.38, 2.14)	1.84 (1.08, 3.15)	1.88 (1.52, 2.33)
Diabetic	Low	1.00	1.00	1.00	1.00
	Med	1.48 (0.62, 3.51)	1.42 (1.03, 1.96)	0.97 (0.14, 6.92)	1.50 (1.06, 2.10)
	Hi	1.53 (0.64, 3.62)	1.75 (1.28, 2.38)	2.41 (0.46, 12.50)	2.01 (1.45, 2.78)
Impaired A1c	Low	1.00	1.00	1.00	1.00
	Med	0.83 (0.57, 1.20)	0.89 (0.73, 1.08)	0.80 (0.54, 1.18)	0.97 (0.81, 1.16)
	Hi	0.75 (0.51, 1.10)	0.87 (0.72, 1.05)	0.62 (0.41, 0.93)	0.96 (0.80, 1.15)
High A1c	Low	1.00	1.00	1.00	1.00
	Med	2.69 (0.95, 7.65)*	1.35 (0.98, 1.86)	2.92 (0.30, 28.26)*	1.86 (1.37, 2.53)
	Hi	2.12 (0.72, 6.27)*	1.77 (1.31, 2.39)	6.80 (0.83, 55.59)*	2.01 (1.49, 2.71)
High TG	Low	1.00	1.00	1.00	1.00
	Med	1.36 (0.98, 1.88)	1.43 (1.18, 1.73)	0.75 (0.50, 1.12)	1.12 (0.94, 1.34)
	Hi	1.96 (1.42, 2.70)	1.89 (1.57, 2.27)	0.64 (0.42, 0.97)	1.24 (1.04, 1.48)
High LDL	Low	1.00	1.00	1.00	1.00
	Med	1.71 (1.16, 2.53)	2.07 (1.70, 2.54)	1.16 (0.74, 1.83)	1.47 (1.23, 1.75)
	Hi	1.99 (1.35, 2.93)	1.82 (1.49, 2.23)	1.38 (0.89, 2.15)	1.61 (1.36, 1.91)
Low HDL	Low	1.00	1.00	1.00	1.00
	Med	1.24 (0.85, 1.81)	1.19 (0.92, 1.56)	0.67 (0.49, 0.92)	0.89 (0.74, 1.06)
	Hi	1.34 (0.92, 1.95)	2.29 (1.80, 2.92)	0.68 (0.49, 0.93)	1.17 (0.99, 1.39)
High TC	Low	1.00	1.00	1.00	1.00
	Med	1.63 (1.12, 2.36)	2.05 (1.69, 2.48)	1.40 (0.90, 2.18)	1.38 (1.17, 1.64)

Prevalence Odds ratios from logistic regression models, stratified by age

Urbanization	Males				Females			
	Age 18-40 years n=1,014	Age 40 years n=2,970	Age 18-40 years n=1,068	Age 40 years n=3,406	Age 18-40 years n=1,068	Age 40 years n=3,406	Age 18-40 years n=1,068	Age 40 years n=3,406
	Hi	Low	Hi	Low	Hi	Low	Hi	Low
Hypertension	1.86 (1.29, 2.68)	1.00	1.79 (1.48, 2.16)	1.00	1.41 (0.91, 2.19)	1.00	1.45 (1.23, 1.72)	1.00
	0.78 (0.55, 1.10)	1.20 (1.01, 1.44)	1.02 (0.64, 1.65)	1.14 (0.96, 1.35)	0.54 (0.31, 0.94)	1.42 (1.21, 1.68)		
Pre-HTN	0.63 (0.41, 0.98)*	1.13 (0.94, 1.36)	0.32 (0.13, 0.77)	1.32 (1.11, 1.57)				
High CRP	1.72 (1.14, 2.59)	1.05 (0.86, 1.28)	1.03 (0.68, 1.56)	1.13 (0.93, 1.37)				
	1.66 (1.10, 2.51)	0.99 (0.81, 1.21)	0.86 (0.56, 1.32)	1.33 (1.11, 1.61)				

Income	Males				Females			
	Age 18-40 years	Age 40 years	Age 18-40 years	Age 40 years	Age 18-40 years	Age 40 years	Age 18-40 years	Age 40 years
	Low	Med	Hi	Low	Med	Hi	Low	Med
Overweight	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00
	1.00 (0.69, 1.44)	1.19 (0.97, 1.45)	0.68 (0.45, 1.02)	1.12 (0.94, 1.34)				
IFG	1.07 (0.75, 1.52)	1.59 (1.31, 1.93)	0.76 (0.51, 1.12)	1.29 (1.08, 1.54)				
	0.98 (0.58, 1.66)	1.16 (0.93, 1.44)	0.88 (0.58, 1.35)	0.88 (0.74, 1.06)				
Diabetic	1.13 (0.69, 1.87)	1.20 (0.96, 1.49)	0.97 (0.65, 1.46)	1.08 (0.91, 1.30)				
	0.59 (0.24, 1.45)	1.06 (0.77, 1.46)	4.05 (0.47, 34.84)	0.67 (0.49, 0.93)				
Impaired A1c	0.86 (0.39, 1.91)	1.51 (1.12, 2.04)	2.16 (0.22, 20.84)	0.88 (0.65, 1.17)				
	0.75 (0.49, 1.12)	0.95 (0.78, 1.14)	0.85 (0.41, 1.18)	1.08 (0.84, 1.36)				
High A1c	0.95 (0.65, 1.41)	1.04 (0.86, 1.26)	0.86 (0.42, 1.78)	0.93 (0.73, 1.20)				
	0.48 (0.17, 1.37)	0.94 (0.69, 1.29)	1.34 (0.32, 5.65)	0.73 (0.55, 0.97)				
High TG	0.92 (0.39, 2.12)	1.45 (1.08, 1.93)	0.71 (0.14, 3.56)	1.04 (0.80, 1.36)				
	1.00	1.00	1.00	1.00				

	Income	Males			Females		
		Age 18–40 years	Age 40 years	Age 18–40 years	Age 18–40 years	Age 40 years	Age 40 years
High LDL	Med	1.27 (0.91, 1.79)	1.23 (1.02, 1.49)	0.77 (0.51, 1.16)	1.10 (0.92, 1.31)		
	Hi	1.04 (0.75, 1.46)	1.33 (1.11, 1.61)	0.64 (0.42, 0.98)	1.16 (0.97, 1.38)		
	Low	1.00	1.00	1.00	1.00		
Low HDL	Med	1.28 (0.85, 1.92)	1.08 (0.89, 1.31)	0.93 (0.59, 1.45)	1.10 (0.93, 1.30)		
	Hi	1.34 (0.90, 1.98)	1.13 (0.93, 1.37)	0.76 (0.48, 1.19)	0.88 (0.75, 1.05)		
	Low	1.00	1.00	1.00	1.00		
High TC	Med	1.14 (0.76, 1.71)	1.21 (0.95, 1.53)	0.77 (0.55, 1.07)	1.01 (0.85, 1.20)		
	Hi	1.12 (0.75, 1.65)	1.17 (0.92, 1.49)	0.85 (0.62, 1.17)	1.08 (0.91, 1.29)		
	Low	1.00	1.00	1.00	1.00		
Hypertension	Med	1.33 (0.90, 1.96)	0.99 (0.82, 1.19)	1.00 (0.65, 1.53)	1.13 (0.96, 1.33)		
	Hi	1.28 (0.88, 1.87)	1.11 (0.92, 1.33)	0.65 (0.41, 1.01)	0.92 (0.77, 1.08)		
	Low	1.00	1.00	1.00	1.00		
Pre-HTN	Med	0.81 (0.57, 1.16)	0.92 (0.77, 1.10)	1.07 (0.64, 1.79)	0.85 (0.73, 1.01)		
	Hi	0.62 (0.44, 0.89)	0.89 (0.74, 1.06)	0.80 (0.47, 1.37)	0.79 (0.67, 0.93)		
	Low	1.00	1.00	1.00	1.00		
High CRP	Med	0.83 (0.53, 1.28)	0.95 (0.79, 1.14)	0.97 (0.47, 2.01)	0.87 (0.73, 1.03)		
	Hi	0.65 (0.42, 1.02)	0.99 (0.82, 1.19)	0.81 (0.39, 1.69)	0.77 (0.64, 0.91)		
	Low	1.00	1.00	1.00	1.00		
	Med	0.92 (0.60, 1.41)	0.91 (0.74, 1.11)	0.79 (0.51, 1.20)	0.96 (0.80, 1.15)		
	Hi	1.11 (0.74, 1.67)	0.85 (0.69, 1.03)	0.62 (0.40, 0.95)	1.03 (0.86, 1.24)		

NOTE: urbanization and income categorized by tertiles, with approximately 300 per tertile in the 18–40 year age group and approximately 1,000 per tertile in the 40 year age group

* Unstable estimates due to n<10 in the reference category

Sex- and age (18–40 years; Age 40 years)-stratified crude logistic regression models including 3 levels each for urbanicity and income

Table 4

Comparison of CHNS findings to findings in the United States and other Asian countries using fasting blood and C-reactive protein

Adolescent Males	CHNS (Aged 12–18 years)	United States Publications (Aged 12–19 years)
A. Adolescent Males only: CHNS and US adolescents		
HbA1c 6.5%	1.9%	0.5% NHANES ²⁰
CRP 3 mg/dL	12.1%	8.5% NHANES ²¹
Glucose 126 mg/dL	0.9%	0.28% NHANES ²² ; 0.22% SEARCH ^{**23}
IFG (100–126 mg/dL)	8.4%	20% NHANES ²²
Country	Glucose 126 mg/dL Plus Medication	IFG (100–126 mg/dL)
B. Adolescents: CHNS and other Asian countries		
CHNS ages 12–17	0.9%	8.4%
South Korea, KHANES 10–17 ^{**}	0.1%	5.3%
Taiwan 10–18 ³⁸	<0.0	--
Country	Glucose 126 mg/dL Plus Medication	IFG (100–126 mg/dL)
C. Adults: CHNS, US, and other Asian countries		
CHNS ages 18 and older	6.9%	20.0%
NHANES 18 and over 2011 ^{22*}	8.8%	--
South Korea, KHANES 10–17 ^{**}	8.1%	17.8%
Hong Kong ³⁹	9.8%	--
Malaysia ³⁹	11.0%	--
Thailand 2004 ³⁹	6.7	--
Singapore 2004 ⁴⁰	8.9 (men); 7.6 (women)	--
Malaysia ⁴¹	12.6%	22.1%

--No data available

* Source: NHANES weighted to be national representative²²

** Use of weighted data from the Korean Health and Nutrition Examination Survey 2009

*** For both males and females