

ORIGINAL ARTICLE

Prevalence of the metabolic syndrome and its components in secondary school student population in the city of Douala, Cameroon

NADINE BILOG^{1,2}, ELYSÉE CLAUDE BIKA LELE^{1,2}, JERSON MEKOULOU NDONGO^{1,2},
YVES JULIEN MBAMA BILOA³, JOSIANE BINDI NGASSE BWEGNE², PEGUY BRICE ASSOMO NDEMBA^{2,4},
NOËL BABAYANA ETAGA^{1,2}, SAMUEL HONORÉ MANDENGUE^{1,2}, LAURENT SERGE ETOUNDI NGOA⁴,
ABDOU TEMFEMO⁵, BIENVENU BONGUE³, JESSICA GUYOT³ and CLARISSE NOEL AYINA AYINA^{1,2}

¹Department of Animal Biology, Faculty of Science, University of Douala; ²Physiology and Medicine of Physical Activities and Sports Unit, University of Douala, Cameroon; ³Sainbiose Inserm U1059 Laboratory, Jean Monnet University, Saint-Étienne, France; ⁴Faculty of Medicine and Biomedical Sciences, University of Yaounde 1 Yaounde; ⁵Faculty of Medicine and Pharmaceutical Sciences, University of Douala, Cameroon

DOI: 10.4081/jphia.2023.2465

Abstract. While the burden of metabolic syndrome (MetS) is still increasing in sub-Saharan Africa, there is a lack of data among young Cameroonian population. The aim of this study was to evaluate the prevalence of MetS and its components among secondary school students in Douala. This was a cross-sectional prospective study carried out on 803 students recruited from February to May 2021 in public and private secondary schools in Douala city, Cameroon. MetS was assessed according to the IDF/AHA/NHLBI 2009 consensus definition. The data collection consisted of a questionnaire on sociodemographic characteristics, measurement of anthropometric parameters (height, weight, body mass index (BMI), waist circumference) and overnight fasting blood sample. Blood pressure (BP), fasting blood glucose, HDL cholesterol and triglycerides were measured using standard methods. The mean age was 18±3 years, 73.3% female. The prevalence of MetS was 27.4%, common among participants aged ≥16 years, and higher in females compare to males (33.7% vs. 11.1%, P<0.0001). The prevalence of MetS components i.e abdominal obesity, high BP, fasting hyperglycemia, low-level HDL cholesterol and hypertriglyceridemia were 14.1, 18.1, 42.8, 51.4 and 38.6% respectively. All MetS components were significantly higher in females

compared to males except for high BP which was similar among the genders. In our study population, the prevalence of MetS is high and this calls for improved monitoring to limit the evolution of associated cardiometabolic complications among young Cameroonians.

Introduction

The metabolic syndrome (MetS) refers to a particular state of morbidity characterized by a constellation of several metabolic abnormalities i.e. glucose intolerance; insulin resistance; dyslipidemia (particularly hypertriglyceridemia and low-level HDL cholesterol); high blood pressure; abdominal obesity (1) There is an ongoing rise of prevalence of the MetS globally, and particularly in developing countries, mainly due to the evolution of obesity linked to poor diet, sedentary lifestyle and the acceleration of economic and demographic transition (2). Globally, the prevalence of MetS is increasing in many countries like USA (3) and China (4) and there is now about one quarter of people having MetS in the world (1). In recent years, the global prevalence of obesity, diabetes, and hypertension has increased significantly (5,6), which contributes to an increase in the prevalence of MetS (7). This syndrome is associated with increase prevalence of cardiovascular diseases and diabetes which are major public health concerns worldwide (8) although there is still ongoing discussion on some MetS component thresholds in black people (9,10) and definition and diagnostic criteria are not yet harmonized in teenager (11).

Increase in sex steroidal hormones and other hormones during puberty may be associated with obesity and insulin resistance which may persist in adulthood and result in increased CVD risk (12). Insulin resistance which is a key mechanism in the development of MetS (13) is almost physiological during pubertal stage. Actually, fuel metabolism is

Correspondence to: Clarisse Noel Ayina Ayina, Department of Animal Biology, Faculty of Science, University of Douala, PO Box 24157 Douala, Cameroon
E-mail: clarisseayina@gmail.com

Key words: metabolic syndrome, prevalence, secondary school, Cameroon

24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58

1 altered during puberty to preserve lean muscle mass and
2 to maximize fat as and alternated fuel source (14). Besides,
3 puberty associated with abdominal obesity is integral part
4 of the mechanisms associated with the development of the
5 MetS (5,15,16) including hypertension, dyslipidemia and
6 hyperglycaemia. Globally, the prevalence of MetS ranges from
7 1.2 to 22.6% in youth and 9.0 to 35.0% in adults (13,17).

8 In Cameroon, the scientific literature does not have
9 sufficient data to establish the national prevalence of MetS.
10 However, some studies estimate the prevalence of MetS to be
11 between 7.4 and 21.4% (18,19), depending on the definition
12 used, the region, the year the study was conducted, the age
13 group of the target population. The main objective of our study
14 was to determine the prevalence of MetS and its components
15 in students in the city of Douala using the 2009 consensus
16 definition of MetS.

17 **Methods**

20 *Design.* This was a cross-sectional prospective study carried
21 out from February through May 2021 in two public and one
22 private secondary schools in the city of Douala (which is
23 the economical capital and one of the most populated city in
24 Cameroon). Participants were students aged 10 to 27 years,
25 regularly registered in the school in all levels.

27 *Ethical considerations.* The study was approved by the insti-
28 tutional ethical board of the University of Douala (ethical
29 clearance N°2508 CEI-Udo/02/2021/M). All the participants
30 included were briefed on the object of the study and were
31 asked to sign informed consent. Data were collected by a
32 trained survey officer and each survey sheet were coded for
33 privacy. All data were stored on a secured computer.

35 *Procedure.* The study procedure consisted of questionnaire
36 administration, anthropometric measurement and blood
37 sample collection. Participants were provided with question-
38 naires and were asked to fast overnight and the blood samples
39 and anthropometric parameters were collected in the morning
40 before 12 PM.

41 Data collection and all measurements were performed
42 in a secured place in the school. Items in the questionnaire
43 included socio-demographic parameters i.e. age, gender,
44 education, physical activity, family and personal history of
45 diabetes, obesity, hypertension and stoke, smoking and alcohol
46 consumption.

48 *Measurements.* Body weight was measured in kilograms (kg)
49 using electronic medical scale. Height was measured to the
50 nearest 0.5 cm with fixed stadiometer. BMI was calculated as
51 weight in kilograms divided by the square of height in meters
52 (kg/m^2). Overweight was defined as a BMI at least $25 \text{ kg}/\text{m}^2$
53 and obesity as a BMI at least $30 \text{ kg}/\text{m}^2$. Waist circumference
54 was measured using a tape measure, halfway between the last
55 rib and the anterior superior iliac spine at the end of expiration.

56 Blood pressure and heart rate measurements were
57 performed after a 15-min rest, in the seated position, following
58 standardized recommendations for blood pressure measure-
59 ment. Three consecutive BP measurements were taken at
60 time intervals of at least 5 min using a validated automated

sphygmomanometer (THUASNE 3W1-A) with the cuff's
width adjusted to the arm's circumference.

Participants were instructed to fast for at least 10 h
overnight and venous blood samples were collected into
vacuum-stoppered heparinized tubes. The samples were then
centrifuged at 3000 rotations per minute for 10 min before
being analyzed using COBAS C111® automated analyzer.
Blood glucose, lipids profile (HDL, LDL, total cholesterol and
triglycerides) were assayed from the samples.

MetS was assessed in participants aged 16 years and
older according to IDF/AHA/NHLBI 2009 collective
consensus (20) which defined MetS for at least three of the
following features: Waist circumference (WC) ≥ 94 cm for
male and ≥ 80 cm for female; Triglycerides ≥ 150 mg/dl; HDL
cholesterol: < 40 mg/dl for males and < 50 mg/dl female; fasting
blood glucose: ≥ 100 mg/dl; Blood pressure $\geq 130/85$ mmHg.
For participants aged less than 16 years, the same criteria were
applied except for WC who was considered high if it is higher
or equal to the 90th percentile for age and sex (21).

Statistical analysis. Data collected were recorded using
Microsoft office Excel 2016 software and analyzed using SPSS
24 (IBM Statistics). Data were presented as mean \pm standard
deviation (SD) for quantitative variables and counts and
percentages for qualitative variables. Student-t test was used
to compare quantitative variables while Chi-square test was
used for qualitative variable comparisons. Differences were
considered significant for $P < 0.05$.

90 **Results**

92 We recruited 803 participants, mean age 18 ± 3 years with 73.3%
93 females ($n=589$). Table I shows the anthropometric, biological
94 and social characteristics of the study population. Mean age
95 was similar between male and female ($P=0.753$), height, SBP,
96 and tobacco consumption levels were significantly higher in
97 male compared to female. On the contrary, hip circumfer-
98 ence, heart rate, BMI, alcohol consumption and physical
99 activity were significantly higher in female compared to male.
100 There were no significant gender differences in weight, waist
101 circumference, DBP, fasting blood glucose, HDL cholesterol,
102 LDL cholesterol, total cholesterol and triglycerides.

103 Table II shows the prevalence of MetS and its components
104 among participants stratified by age groups and compared
105 between male and female. The prevalence of MetS in the
106 whole sample was 27.4%; common in female participants.
107 The common MetS components were low HDL cholesterol
108 (51.4%), hyperglycemia (42.8%) and hypertriglyceridemia
109 (38.6%). All the components' frequencies in the sample were
110 significantly higher in female compared to male; except for
111 high blood pressure that was similar between the two groups.
112 MetS and its components were common among ≥ 16 years
113 group. Hyperglycemia, low-level HDL cholesterol and hyper-
114 triglyceridemia were most represented abnormalities among
115 ≥ 16 years group while only 3 participants had hypertriglyc-
116 eridemia in < 16 years group. In the < 16 years group, only
117 low-level HDL cholesterol was significantly higher in female
118 while in the ≥ 16 years group all the parameters were signifi-
119 cantly higher in female compared to male, except for blood
120 glucose that was similar between the two groups.

Table I. Anthropometric, biological and social characteristics of participants by gender.

	Over all (N=803)	Male (N=214)	Female (N=589)	P-value
Age (years)	18±3	18±3	18±3	0.753
Weight (kg)	59.0±11.3	60.3±11.0	58.6±11.4	0.063
Height (cm)	162±8	167±10	160±7	<0.0001
BMI (kg/m ²)	22.6±6.8	21.6±3.0	23.0±7.7	0.013
Waist circumference (cm)	73±8	72±7	73±8	0.210
Hip circumference (cm)	77±8	75±7	78±9	<0.0001
Systolic BP (mmHg)	118±13	121±11	116±13	<0.0001
Diastolic BP (mmHg)	72±10	71±10	72±11	0.153
Heart rate (beat/min)	82±14	75±13	85±14	<0.0001
Fasting blood glucose (mg/dl)	97±15	96±14	98±16	0.092
HDL cholesterol (mg/dl)	47±12	48±11	47±12	0.277
LDL cholesterol (mg/dl)	136±47	134±42	136±48	0.694
Total cholesterol (mg/dl)	212±49	212±44	212±051	0.996
Triglycerides (mg/dl)	144±56	142±52	145±058	0.523
Physical activity	562 (70)	164 (20.4)	398 (49.6)	0.013
Alcohol consumption	421 (65.8)	133 (73.1)	288 (62.9)	0.014
Tobacco consumption	23 (3.6)	13 (7.1)	10 (2.2)	0.005

Results are presented as mean ± standard deviation and count (percentage); N, Number of participant; BMI, body mass index; BP, blood pressure; HDL, high density lipoprotein; LDL, low density lipoprotein.

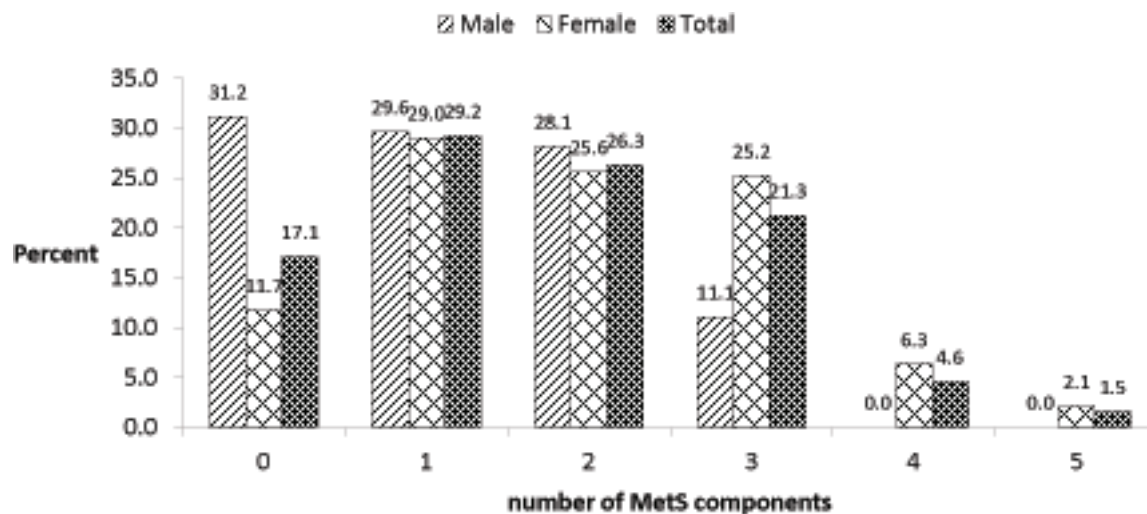


Figure 1. Distribution of number of MetS components by age group.

Fig. 1 shows the number of MetS components in the population and according to gender. Regarding the components of MetS in the general population, 17.1% had no MetS component, 55.5% had one or two components, and 6.1% had four to five components. Only female participants had more than three MetS components and 11.1% male compared to 25.2% female had three components. Differences were significant between the two groups (P<0.0001).

Fig. 2 shows frequencies of different association of MetS components. The most frequent association of MetS components was hyperglycemia + hypertriglyceridemia + low-level HDL cholesterol with 51.6%. Other components associations represented less than 10% of the sample.

Discussion

This was a cross-sectional descriptive study aimed to contribute to the literature around the prevalence of metabolic syndrome and its components among secondary school children in Cameroon. One of the major findings in this study is the 27.4% prevalence of MetS found according the IDF/AHA/NHLBI consensus of 2009 criteria. Low-level HDL cholesterol (51.4%), hyperglycemia (48.8%) and hypertriglyceridemia (38.6%) were the main drivers of that prevalence. MetS was significantly predominant in female (33.7%) compared to male (11.1%) and in participants aged 16 years and older (33%) compared to their younger counterparts (1.6%). Few studies have reported

Table II. Metabolic syndrome and its components stratified by age groups and compared between male and female.				
Under 16 years (n=167)				
	Total (N=167)	Male (N=52)	Female (N=115)	P-value
Waist circumference ($\geq 90^{\text{e}}$ percentile)	0	0	0	-
Blood glucose (≥ 100 mg/dl)	23 (18.0)	7 (14.9)	16 (19.8)	0.652
Blood pressure ($\geq 130/85$ mmHg)	32 (19.2)	9 (17.3)	23 (20.0)	0.844
Triglycerides (≥ 150 mg/dl)	3 (1.8)	0	3 (2.6)	0.585
HDL cholesterol (< 40 mg/dl)	97 (58.1)	13 (25.0)	84 (73.0)	< 0.0001
Metabolic syndrome	2 (1.6)	0	2 (2.5)	0.729
16 years and older (n=636)				
	Total (N=636)	Male (N=162)	Female (N=474)	P-value
Waist circumference (≥ 94 ♂; 80 ♀ cm)	113 (17.8)	1 (0.6)	112 (23.6)	< 0.0001
Blood glucose (≥ 100 mg/dl)	315 (53.3)	80 (52.6)	235 (53.5)	0.923
Blood pressure ($\geq 130/85$ mmHg)	113 (17.8)	36 (22.2)	77 (16.2)	0.11
Triglycerides (≥ 150 mg/dl)	307 (48.3)	68 (42.0)	239 (50.4)	0.077
Cholesterol HDL (< 40 ♂; 50 ♀ mg/dl)	316 (49.7)	31 (19.1)	285 (60.1)	< 0.0001
Metabolic syndrome	195 (33.0)	22 (14.5)	173 (39.4)	< 0.0001
All participants (n=803)				
	Total (N=803)	Male (N=214)	Female (N=589)	P-value
Obesity	113 (14.1)	1 (0.5)	112 (19.0)	< 0.0001
Hyperglycemia	308 (42.8)	89 (23.4)	219 (64.8)	< 0.0001
High blood pressure	145 (18.1)	45 (21.0)	100 (17.0)	0.224
Hypertriglyceridemia	310 (38.6)	68 (31.8)	242 (41.1)	0.021
Low-level HDL cholesterol	413 (51.4)	44 (20.6)	369 (62.6)	< 0.0001
Metabolic syndrome	197 (27.4)	22 (11.1)	175 (33.7)	< 0.0001

Results are presented as number of participant (percentage); HDL, high density lipoprotein.

on MetS in Cameroon in young age as well as in adulthood with various prevalence mainly depending on the diagnostic criteria. We used in this study the IDF/AHA/NHLBI consensus of 2009 criteria which is the latest the most frequently used definition of MetS (8,11,19,21).

The prevalence of MetS found in our study (27.7%) was higher than that found in a population of students aged 16-21 years old in the city of Yaoundé (20.3%) (19). The prevalence of MetS found in that study was also lower than that of the group of 16 years and older in our study (33%). On the contrary, our prevalence was almost similar to the 32.5% prevalence found on older population (mean age 44 ± 17 years) in West region Cameroon (22). Our results confirm the ongoing epidemiological transition faced by Douala and Yaounde which are respectively economic and politic capital in Cameroon. Other studies using the IDF to define MetS had also found a lower prevalence compared to our study i.e. Ethiopia (4.8%) (23), South Africa (3-6%) populations (24), Congo-Brazzaville (15.9%) (25) and USA (9.3%) (26) although higher prevalence

using the IDF criteria were found in Iran (37.4%) and Tunisia (45.5%) (27). Discrepancies in the prevalence of MetS could also been explained by genetic and environmental divergences among populations that could influence the development of MetS components like obesity (28).

Low-level HDL Cholesterol (51.4%), hyperglycemia (42.8%) and hypertriglyceridemia (38.6%) were the main drivers of the MetS in our study. These findings are consistent with some studies (29,30) while other studies found high blood pressure (31,32) and higher WC (33) as main character of MetS. Study on student in Yaounde (19) found similar trend for low-level HDL cholesterol (49.3%) but lower prevalence for hyperglycemia (20%) and hypertriglyceridemia (12.9%) compare to our study. The 13.6% prevalence of abdominal obesity found in our study was higher than that reported in Yaoundé (12.1%) (19), but lower in that reported in Egypt (78%) (34), in Spain (31.4%) (35), and in a meta-analysis involving 14 African and Middle Eastern countries (67.6%) (36). Abdominal obesity, Low-level HDL

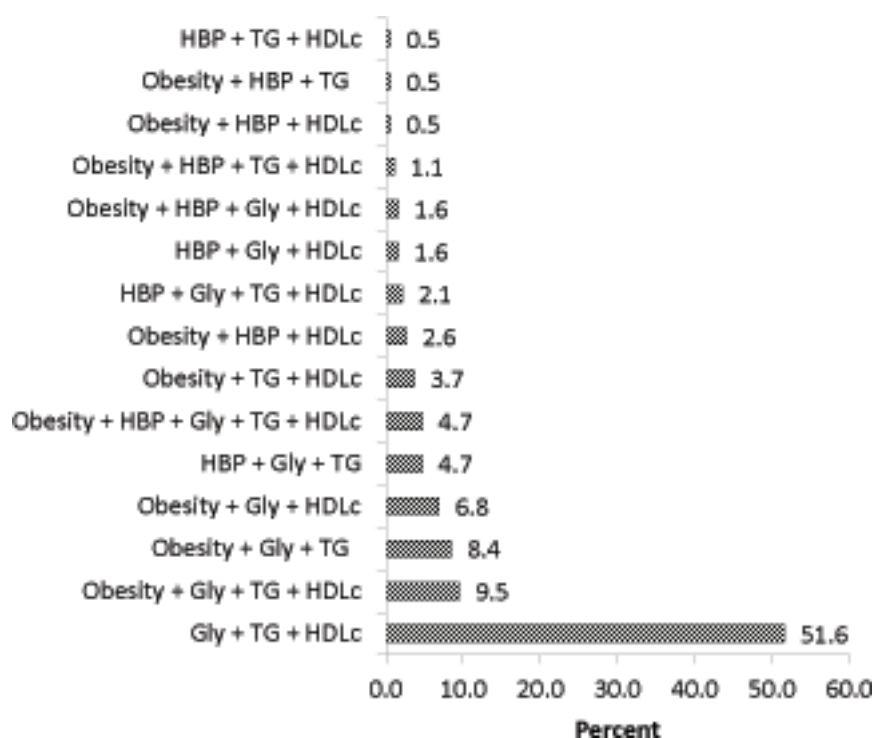


Figure 2. Association of MetS components in individuals with MetS. MetS, metabolic syndrome; HBP, High blood pressure; TG, hypertriglyceridemia; HDLc, low-level High Density Lipoprotein cholesterol; Gly, hyperglycemia.

cholesterol and hypertriglyceridemia in the adolescent have a great impact on the development of cardiovascular disease in adulthood. The high prevalence of this components combined with the large difference found between younger participants (<16 years) and older participants (>=16 years) emphasize the necessity for a rapid diagnosis of MetS in adolescents and targeted interventions to reduce the risk of cardiovascular mortality in adulthood (37).

MetS and all its components (except high blood pressure) were significantly more frequent in female compare to male. This result is consistent with those obtained in Yaoundé students, Cameroon (19) and in other populations (30). Another study on the American population showed a higher proportion in men (38). Apart from gender, age also has a significant influence on the metabolic syndrome, but appears here as a protective factor for younger people because only participant who have more than 16 years old are affected.

Approximately 14.6% of the participants had at-risk blood pressure levels according to the IDF/AHA/NHLBI consensus definition. This lower prevalence of hypertension than those observed in studies conducted in Yaoundé (19,39) is justified by the fact that our study population had a low consumption of cigarette which is a main factor for hypertension (40). This result is in line with the Tunisian study conducted in 2010 (41), and can be explained by the presence of obesity in the MetS, because the prevalence of hypertension increases with the severity of obesity and excess abdominal fat (42). Regarding fasting hyperglycemia, about 42.8% of the participants had a fasting blood glucose level above 100 mg/dl. This result is similar to those found in southwestern Benin in 2015 (43) and could be explained by a diet very rich in sugar or even an abnormality of glucose regulation in the participants.

In this study, various associations of the components of MetS were observed in people with MetS, but the most predominant association was hyperglycemia + hypertriglyceridemia + low-level HDL cholesterol (51.6%). This result corroborates the role of these three components as the main leaders of MetS in our population. Our results however differ from that obtained in a Canadian survey (44), which instead identified the association Obesity + low-level HDL cholesterol + hypertriglyceridemia as the most frequent. These differences could be explained by the diversity of the populations studied in terms of gender, age, lifestyle and environment, which are socio-demographic factors that enable the development of MetS components.

Limitations

The main limitation of our study is that we recruited participants in only three secondary schools of the Douala city. This is however mitigate by the considerable sample size of our study which probably include participants from every part of the city. Other studies using probabilistic sampling procedure are necessary to draw more precise conclusions.

Conclusions

MetS is a major concern in our population with a 27.4% prevalence. More than 80% of the sample had at least one MetS component and the most prevalent components found in our study were Low-level HDL cholesterol, hyperglycemia and hypertriglyceridemia. MetS was strongly related to age and gender. Compared to other studies, our study shows rise of MetS in adolescents and young adults and emphasizes the necessity for public health policies targeting dietary habits and

1 other factors that could help curve the rise of MetS in young
2 people.

3 **Acknowledgments**

4 The authors express their grateful thanks to the survey officers,
5 all the participants of the survey, and to the administrative
6 officers of the school involved in the study for their facilitation
7 and their dedication to this study.

8 **Funding**

9 None.

10 **Contributions**

11 AACN, MSH, designed the study protocol, and wrote the first
12 manuscript draft; BLEC, MBYJ, led the statistical analyses
13 and contributed to the manuscript draft; BN, MBYJ, ENB,
14 BNBJ, contributed to logistic and data collection; MNJ, GJ,
15 ANPB, BB, ENLS, TA, critically contributed to analysis,
16 discussion, and interpretation of the data, and to the writing
17 of the manuscript. All the authors approved the final version
18 to be published.

19 **Ethical approval and consent to participate**

20 The study was approved by the institutional ethical board
21 of the University of Douala (ethical clearance N 2508
22 CEI-Udo/02/2021/M).

23 **Availability of data and material**

24 Data and materials are available by the authors.

25 **Informed consent**

26 All the participants included were briefed on the object of the
27 study and were asked to sign informed consent.

28 **Conflict of interest**

29 The authors declare no potential conflict of interest.

30 Accepted: 24, March 2023; submitted: 13, January 2023.

31 **References**

- 32 1. Saklayen MG: The global epidemic of the metabolic syndrome. *Curr Hypertens Rep* 20: 12, 2018.
- 33 2. Weiss R, Dziura J and Burgert TS: Obesity and the metabolic
34 syndrome in children and adolescents. *N Engl J Med* 350:
35 2362-2374, 2004.
- 36 3. National Center for Health Statistics, Division of Health
37 Interview Statistics. Crude and age-adjusted percentage of
38 civilian, non-institutionalized adults with diagnosed diabetes,
39 United States, 1980-2010. National Center for Chronic Disease
40 Prevention and Health Promotion, Ed. Atlanta, GA, Centers
41 for Disease Control and Prevention, Division of Diabetes
42 Translation, 2012.
- 43 4. Wang Y, Mi J, Shan XY, Wang QJ and Ge KY: Is China facing
44 an obesity epidemic and the consequences? The trends in obesity
45 and chronic disease in China. *Int J Obes (Lond)* 31: 177-188,
46 2007.
- 47 5. Junquero D and Rival Y: Metabolic syndrome: Which definition
48 for what treatment(s)?]. *Med Sci (Paris)* 21: 1045-1053, 2005.
- 49 6. Zimmet P, Alberti KG and Shaw J: Global and societal implica-
50 tions of the diabetes epidemic. *Nature* 414: 782-787, 2001.
- 51 7. Després JP, Poirier P, Bergeron J, Tremblay A, Lemieux I and
52 Alméras N: From individual risk factors and the metabolic
53 syndrome to global cardiometabolic risk. *Eur Heart J* 10: 24-33,
54 2008.
- 55 8. Vaduganathan M, Mensah GA, Turco J, Turco JV, Fuster V and
56 Roth GA: The global burden of cardiovascular diseases and risk:
57 A Compass For Future Health. *J Am Coll Cardiol* 80: 2361-2371,
58 2022.
- 59 9. Sumner AE: Ethnic differences in triglyceride levels and high-density
60 lipoprotein lead to underdiagnosis of the metabolic syndrome in
black children and adults. *J Pediatr* 155: S7.e7-e11, 2009.
- 10 10. Yu SSK, Ramsey NLM, Castillo DC, Ricks M and Sumner AE:
Triglyceride-based screening tests fail to recognize cardiometabolic
disease in African immigrant and African-American men. *Met Synd Rel Dis* 11: 15-20, 2013.
- 11 11. Alberti KG, Zimmet P and Shaw J: Metabolic syndrome-a
12 new worldwide definition. A Consensus Statement from the
13 International Diabetes Federation. *Diabet Med* 23: 469-480,
14 2006.
- 15 12. Reinehr T and Toschke AM: Onset of puberty and cardiovascular
16 risk factors in untreated obese children and adolescents: A 1-year
17 follow-up study. *Arch Pediatr Adolesc Med* 163: 709-715, 2009.
- 18 13. Eckel RH, Grundy SM and Zimmet PZ: The metabolic syndrome.
19 *Lancet* 365: 1415-1428, 2005.
- 20 14. Kelsey MM and Zeitler PS: Insulin resistance of puberty. *Curr
21 Diab Rep* 16: 64, 2016.
- 22 15. Gamila S and Dallongeville J: Épidémiologie du syndrome
23 métabolique en France. *Med Nutr* 39: 89-94, 2003.
- 24 16. Ghedada Y and Beddar L: Historical perspective: Visceral
25 obesity and related comorbidity in Joannes Baptista Morgagni's
26 *De Sedibus et Causis Morborum per Anatomen Indagata*.
27 *Diabetes Metab* 42:57-58, 2016.
- 28 17. Cameron AJ, Shaw JE and Zimmet PZ: The metabolic syndrome:
29 Prevalence in worldwide populations. *Endocrinol Metab Clin
30 North Am* 33: 351-375, 2004.
- 31 18. Fezeu LK, Assah FK, Balkau B, Mbanya DS, Kengne AP,
32 Awah PK and Mbanya JC: Ten-years changes in central obesity
33 and BMI in rural and urban Cameroon. *Obesity* 16: 1144-1147,
34 2008.
- 35 19. Ngo-Song MC, Kingue A, Fouejou-Wamba BG, Abega-Ebene PJ,
36 Ngondi JL and Oben JE: Prevalence of metabolic syndrome
37 among 16-21 years urban Cameroonian using NCEP ATPIII and
38 IDF criteria. *BMJ* 4: 2483-2493, 2014.
- 39 20. Alberti KG, Robert RH, Grundy SM, Zimmet PZ, Cleeman JI,
40 Donato KA, Fruchart JC, James WP, Loria CM, Smith SC Jr, *et al*:
41 Harmonizing the metabolic syndrome: A joint interim state-
42 ment of the International Diabetes Federation Task Force on
43 Epidemiology and Prevention; National Heart, Lung, and Blood
44 Institute; American Heart Association; World Heart Federation;
45 International Atherosclerosis Society; and International
46 Association for the Study of Obesity. *Circulation* 120: 1640-1645,
47 2009.
- 48 21. Zimmet P, Alberti KG, Kaufman F, Tajima N, Silink M,
49 Arslanian S, Wong G, Bennett P, Shaw J and Caprio S; IDF
50 Consensus Group: The metabolic syndrome in children and
51 adolescents-an IDF consensus report. *Pediatr Diabetes* 8:
52 299-306, 2007.
- 53 22. Marbou WJT and Kuete V: Prevalence of metabolic syndrome
54 and its components in Bamboutos Division's adults, west region
55 of Cameroon. *Biomed Res Int* 2019: 9676984, 2019.
- 56 23. Tran A, Gelaye B, Girma B, Lemma S, Berhane Y, Bekele T,
57 Khali A and Williams MA: Prevalence of metabolic syndrome
58 among working adults in Ethiopia. *Int J Hypertens* 2011: 193719,
59 2011.
- 60 24. Sekokotla MA, Goswami N, Sewani-Rusike CR, Iputo JE and
Nkeh-Chungag BN: Prevalence of metabolic syndrome in
adolescents living in Mthatha, South Africa. *Ther Clin Risk
Manag* 13: 131-137, 2017.
- 25 25. Gombet T, Longo-Mbenza B, Ellenga-Mbolla B, Ikama MS,
Mokondjimobe E, Kimbally-Kaky G and Nkoua JL: Aging,
female sex, migration, elevated HDL-C, and inflammation
are associated with prevalence of metabolic syndrome among
African bank employees. *Int J Gen Med* 5: 495-503, 2012.
- 26 26. Topè AM and Rogers PF: Metabolic syndrome among students
attending a historically black college: Prevalence and gender
differences. *Diabetol Metab Syndr* 5: 2, 2013.

- 1 27. Delvari A, Forouanafar MH, Alikhani S, Sharifian A and
2 Kelishadi R: First nationwide study of the prevalence of the
3 metabolic syndrome and optimal cutoff points of waist circum-
4 ference in the Middle East: The national survey of risk factors for
5 noncommunicable diseases of Iran. *Diabetes Care* 32: 1092-1097,
6 2009.
- 7 28. Colombet Z, Perignon M, Salanave B, Landais E, Martin-Prével Y,
8 Allès B, Drogé S, Amiot-Carlin MJ and Méjean C: La qualité
9 de l'alimentation contribue-t-elle à expliquer les différences
10 socioéconomiques de la prévalence du syndrome métabolique
11 dans les Antilles Françaises? *Nutr Clin Met* 33: 110, 2019.
- 12 29. Huang TT, Shimel A, Lee RE, Delancey W and Strother ML:
13 Metabolic risks among college students: Prevalence and gender
14 differences. *Metab Syndr Relat Disord* 5: 365-372, 2007.
- 15 30. Fernandes J and Lofgren IE: Prevalence of metabolic syndrome
16 and individual criteria in college students. *J Am Coll Hlth* 59:
17 313-321, 2011.
- 18 31. Yen SL, Chiu TY, Lin YC, Lee YC, Lee LT and Huang KC:
19 Obesity and hepatitis B infection are associated with increased
20 risk of metabolic syndrome in university freshmen. *Int J Obes*
21 (Lond) 32: 474-480, 2008.
- 22 32. Burke JD, Reilly RA, Morrell JS and Lofgren IE: The University
23 of New Hampshire's young adult health risk screening initiative.
24 *J Am Diet Assoc* 1751-1758, 2009.
- 25 33. Park J, Mendoza JA, O'Neil CE, Hilmers DC, Liu Y and
26 Nicklas TA: A comparison of the prevalence of the metabolic
27 syndrome in the United States (US) and Korea in young adults
28 aged 20 to 39 years. *Asia Pac J Clin Nutr* 17: 471-482, 2008.
- 29 34. Assaad-Khalil SH, Mikhail MM, Aati TA, Zaki A, Helmy MA,
30 Megallaa MH, Hassanien R and Rohoma KH: Optimal waist
31 circumference cutoff points for the determination of abdominal
32 obesity and detection of cardiovascular risk factors among adult
33 Egyptian population. *Indian J Endocrinol Metab* 19: 804-810,
34 2015.
- 35 35. Lopez-Sobaler AM, Aparicio A, Aranceta-Bartrina J, Gil Á,
36 González-Gross M, Serra-Majem L, Varela-Moreiras G and
37 Ortega RM: Overweight and general and abdominal obesity in
38 a representative sample of Spanish adults: Findings from the
39 ANIBES Study. *Biomed Res Int* 2016: 8341487, 2016.
- 40 36. Alsheikh-Ali AA, Omar MI, Raal FJ, Rashed W, Hamoui O,
41 Kane A, Alami M, Abreu P and Mashhoud WM: Cardiovascular
42 risk factor burden in Africa and the middle east: The Africa
43 middle east cardiovascular epidemiological (ACE) study. *PLoS*
44 *One* 9: e102830, 2014.
- 45 37. Berenson GS, Srinivasan SR, Bao W, Newman WP III, Tracy RE
46 and Wattigney WA: Association between multiple cardiovascular
47 risk factors and atherosclerosis in children and young adults. *The*
48 *Bogalusa Heart Study*. *N Engl J Med* 338: 1650-1656, 1998.
- 49 38. Ford ES, Giles WH and Dietz WH: Prevalence of the metabolic
50 syndrome among US adults: Finding from the third National Health
51 and Nutrition Examination Survey. *JAMA* 287: 356-359, 2002.
- 52 39. Enyegue Mandob D, Messi Z and Noa Ndoua CC: Facteurs de
53 risque sociodémographiques de syndrome métabolique dans une
54 population de Yaoundé-Cameroun. *Int J Adv Res* 6: 596-605,
55 2018.
- 56 40. Benowitz NL, Kuyt F and Jacob P III: Influence of nicotine on
57 cardiovascular and hormonal effects of cigarette smoking. *Clin*
58 *Pharmacol Ther* 36: 74-81, 1984.
- 59 41. Allal-Elasmi M, Haj Taieb S, Hsairi M, Zayani Y, Omar S,
60 Sanhaji H, Jemaa R, Feki M, Elati J, Mebazaa A and Kaabachi N:
The metabolic syndrome: Prevalence, main characteristics and
association with socio-economic status in adults living in Great
Tunis. *Diabetes Metab* 36: 204-208, 2010.
- Rein, hypertension artérielle et
syndrome métabolique. *Revue Générale*. 2008. [https://www.
realites-cardiologiques.com/2007/01/31/rein-hypertension-arterielle-
et-syndrome-metabolique/](https://www.realites-cardiologiques.com/2007/01/31/rein-hypertension-arterielle-et-syndrome-metabolique/) Accessed on 27/07/2022.
- Aspects épidé-
miologiques du syndrome métabolique au sein de la population
obèse de la commune Ouidah au Sud-ouest du Bénin. *Antropo* 33:
111-116, 2015.
- Macpherson M, De Groh M, Loukine L, Prud'homme D and
Dubois L: Prévalence du syndrome métabolique et ses facteurs
de risque chez les enfants et les adolescents canadiens: Enquête
Canadienne sur les mesures de la santé, cycle 1 (2007-2009)
et cycle 2 (2007-2011). *Promotion de la santé et prévention
des maladies chroniques au canada Recherche, politiques et
pratiques* 36: 37-45, 2016.