Research Article

Prevalence of Metabolic Syndrome and Its Components in Bamboutos Division's Adults, West Region of Cameroon

Wiliane J. T. Marbou 💿 and Victor Kuete 💿

Department of Biochemistry, Faculty of Science, University of Dschang, Dschang, Cameroon

Correspondence should be addressed to Victor Kuete; kuetevictor@yahoo.fr

Received 14 January 2019; Revised 10 April 2019; Accepted 15 April 2019; Published 30 April 2019

Academic Editor: Cristiano Capurso

Copyright © 2019 Wiliane J. T. Marbou and Victor Kuete. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

The prevalence of metabolic syndrome (MetS) and its associated risks remain unappreciated in Bamboutos Division, west region of Cameroon. This study aimed to evaluate the prevalence of MetS, its individual components, and associated risk factors among Bamboutos Division's adults population using a Joint Interim Statement of the International Diabetes Federation (IDF) Task Force on Epidemiology and Prevention definitions parameters. A cross-sectional study was conducted from May 2016 to May 2018 in Mbouda ADLUCEM Hospital and Mbouda District Hospital, two reference hospitals in Bamboutos Division, west region of Cameroon. Interview, physical and clinical examinations, and lipid and fasting blood glucose measurements were conducted for 604 adults. The definition of MetS proposed by IDF was used. The prevalence of MetS was 32.45% with highly significant female predominance (46.11% for females and 14.01 % for males). In the entire participants, the most common abnormalities were low-HDL (82.78%) and hypertriglyceridemia (53.97%) [p<0.001]. Participants with obesity (OR: 16.34; 95% CI: 9.21-28.96), overweight (OR: 7.45; 95% CI: 4.17-13.30), and highest hs-CRP (hs-CRP >11 mg/l) had a higher risk of developing MetS. The most common MetS component was abdominal obesity (OR: 353.13; 95% CI: 136.16-915.81). MetS is prevalent among Bamboutos Division's adults in west region of Cameroon and abdominal obesity is the most common MetS component. This study highlights the need for evidence-based prevention, diagnosis, and management of MetS and its associated factors among Bamboutos Division's adults in Cameroon.

1. Introduction

Noncommunicable diseases are increasingly important causes of morbidity and mortality in Africa due to the rapid demographic transition and changing in lifestyle. In African countries, the prevalence of diabetes in 2017 was 3.3%, the overall prevalence of hypertension was 55.2% in 2017, and overweight and obesity were estimated to cause 3.4 million deaths in 2017 [1-3]. Concern about the health risks associated with rising noncommunicable diseases has become nearly universal in Africans countries. Cardiovascular diseases and type 2 diabetes are recognized as a determining factor in the development of metabolic syndrome (MetS) [4, 5]. Metabolic disorders known as the "metabolic syndrome" are defined as a set of biological and anthropomorphic disturbances whose most easily observable clinical marker is overweight, especially abdominal obesity [4]. According to the available data, foodstuffs rich in fatty and/or sugary foods, combined with low energy outgo, constitute one of the main aetiologies of the MetS. The prevalence of the MetS varies between different populations in Africa. This prevalence also varies according to the criteria used to define MetS, the type of population recruited, and the type and age of the subjects. In addition, studies show an increase in prevalence over time, as a result of changes in diets and lifestyle in developing countries [6]. Thus, the MetS is an emerging public health problem in developing countries.

The prevalence of MetS in Africa ranges from 0% to 50% or even higher depending on the population and the criteria used in defining it [7]. The prevalence of MetS was found to be 35.1% (ATP III) in northwestern Nigeria, 21.8% (IDF) in adults in South Africa attending healthcare facilities in Eastern Cape, 35.73% (IDF) among adults in Morocco, 25.6% among urban Kenyan population, and 38.98% (IDF) in adult men in the Dschang Health District in the west

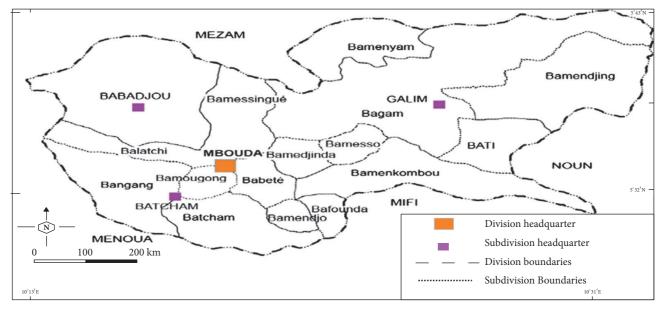


FIGURE 1: Map of Bamboutos Division with subdivision headquarters in west region of Cameroon [14].

region of Cameroon [8–12]. In Cameroon, studies on the distribution of the individual components of MetS are very limited. This study aimed to evaluate the prevalence of MetS, its individual components, and associated risk factors among Bamboutos Division's adults as defined by the Joint Interim Statement of the International Diabetes Federation Task Force on Epidemiology and Prevention [13].

2. Materials and Methods

2.1. Study Type and Population. This was a cross-sectional study conducted from May 2016 to May 2018 in Mbouda ADLUCEM Hospital and District Hospital, two reference hospitals in Bamboutos Division, west region of Cameroon (Figure 1). Participants were randomly selected and included in the study if they were 20 years old and older, attending the two reference hospitals. The human immunodeficiency virus-(HIV-) positive patients, pregnant women, and participants with positive serology for hepatitis B and C were not included in this study.

2.2. Sociodemographic and Anthropometric Data Collection. A structured questionnaire was used to gather information on sociodemographic variables (gender, age, and level of education), smoking status, and physical activity. Smoking was categorized as current, former, and never. Former smokers were those who reported that they had smoked cigarettes during their lifetime but are not currently smoking cigarettes. Drinking was categorized as never, moderate, and heavy. Never drinkers were those who did not drink beer, wine, or hard liquor during the past month. Moderate drinkers had an alcoholic beverage (beer, wine, or hard liquor) less than once per day during the past month. Heavy drinkers were defined as those who ever drank 4 or more alcoholic beverages per day or who drank beer, wine, or hard liquor 1 time per

day during the past month. Physical activity was categorized as low (walking $\leq 60 \text{ min/day}$), moderate (walking $\geq 60 \text{ min/day}$), high (vigorous activity for $\leq 30 \text{ min/day}$), and very high (vigorous activity for $\geq 30 \text{ min/day}$) physical activities.

Trained staff measured anthropometric measurements including weight and height. Body mass index (BMI) was tricked out by dividing the weight in kilograms by the square of the height in meters. BMI was categorized as underweight, <18.5 kg/m², normal, 18.5-24.9 kg/m², overweight, 25-29.9 kg/m², and obesity, \geq 30 kg/m² [15]. Blood pressure (BP) was measured using standardized sphygmomanometer. Trained personnel performed the procedures while the subject was in a sitting position with the arm at the level of the heart and after 5 minutes' rest. Two blood pressure readings were taken from each participant and the average reading of both was used in this study. The participant was labelled as having hypertension if systolic blood pressure was \geq 130 mm Hg or diastolic blood pressure was ≥ 85 mm Hg or if the patient was on antihypertensive medications [13]. At the level of iliac crest, precisely at minimal respiration to the nearest 0.1 cm, the waist circumference was measured.

2.3. Laboratory Measurements. Venous blood was collected after overnight fasting to determine fasting plasma glucose (FPG) using Accu-Chek[®] Active reader, as described by the manufacturer (Roche Diabetes Care GmbH, Sandhofer Strasse 116, 68305 Mannheim, Germany). Plasma concentrations of total cholesterol and triglycerides (TG) were evaluated by enzymatic methods. High-density lipoprotein (HDL) cholesterol was evaluated by enzymatic colorimetric method. Low-density lipoprotein (LDL) cholesterol was calculated using the Friedewald formula if the triglycerides are less than 400 mg/dl (4.6 mmol/l) [16]. Biochemical tests were performed using a MaestroNano[®] Pro Spectrophotometer (Maestrogen, 8275 South Eastern, Avenue #200, Las Vegas,

NV 89123, USA) and the reference commercial kits (Sigma-Aldrich Co., 3050 Spruce Street, St. Louis, MO 63103, USA). We analyzed high sensitivity C-reactive protein (hs-CRP) using ELISA solid phase direct sandwich method (Sigma-Aldrich, St. Louis, USA) with ELx808[™] Microplate reader (BioTek Instruments, Winooski, USA).

2.4. Definitions of Metabolic Syndrome and Dyslipidemia. The criteria used for the definition of metabolic syndrome in adults specified by IDF were applied. Therefore, subjects were considered to have MetS if they had central obesity that was defined by a waist circumference \geq 94 cm in men and \geq 80 cm in women, along with two or more of the following criteria, as per the Joint Interim Statement of the International Diabetes Federation Task Force on Epidemiology and Prevention [13]: high fasting glucose level $\geq 100 \text{ mg/dL}$ (5.6 mmol/L) or patients known to have diabetes mellitus and/or on treatment for diabetes; hypertriglyceridemia-serum triglyceride level \geq 150 mg/dL (1.7 mmol/L); low HDL cholesterol-serum; HDL cholesterol < 40 mg/dL (1.0 mmol/L) in men and < 50 mg/dL (1.3 mmol/L) in women or patients known to have dyslipidemia; high blood pressure [systolic blood pressure $(SBP) \ge 130 \text{ mmHg and/or diastolic blood pressure (DBP)}$ \geq 85 mmHg or patients known to have hypertension and/or on treatment for hypertension]. Dyslipidemia was defined as specified by American Association of Clinical Endocrinologists and American College of Endocrinology [17].

2.5. Assessment of Dietary Intake Frequency. In the present study, dietary intake over the past month was assessed with a structure questionnaire. The questionnaire includes 23 food items highly consumed in the area. Participants were asked about the frequency of each food consumed during the month of their participation in the study. According to the frequency of food intake, each food item was classified into four intervals ([0-4], [4-13], [13-25], and [25-90] times/month).

2.6. Ethical Consideration. Ethical approval was obtained from the Cameroon National Ethics Committee (CNEC), Ministry of Public Health (reference number, 2018/06/ 1054/CE/CNERSH). Prior to data collection, permission was obtained from each manager of the Mbouda ADLUCEM and District Hospital. Information sheets detailing the purpose and process of the study were provided to each participant. Each participant gave written, informed consent for his/her voluntary participation.

2.7. Statistical Analysis. All statistical tests were two-tailed and statistical significance was set at p<0.05. Continuous variables were expressed as mean values \pm Standard Deviation (SD) and compared by using Welch *t*-test. Categorical data were expressed as frequency and compared by using Chisquare tests. We calculated the prevalence of MetS according to the IDF criteria. Logistic Regression Model was used to evaluate the association between MetS and associated risk factors. Epi InfoTM version 7.2.2.6 (CDC, 1600 Clifton Road, Atlanta, GA 30329-4027, USA) was used for statistical analyses.

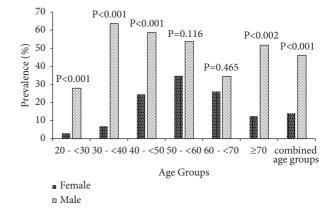


FIGURE 2: Variation in the prevalence of metabolic syndrome by sex and age group in the entire population (n=604). P value (between males and females); MetS, metabolic syndrome.

3. Results

The features of Mbouda's adults by sex who participated in this study are presented in Table 1. 604 [57.45% (n=257) males and 42.55% (n=347) females] participated in this study with the average age of 43.74 \pm 17.21 years. There was a significant difference between the two groups in terms of educational level, drinking and smoking status, physical activities, abdominal obesity (resp., p<0.001), and serum hs-CRP (p=0.008). Female participants (45.87 \pm 16.87 years) were more likely than male participants (40.87 \pm 17.29 years) to be older, with lower levels of systolic blood pressure (Table 1).

The features of the participants according to MetS status are presented in Table 2. Participants with MetS (48.19 ± 15.48 years) were more likely (p<0.001) than the normal participants (41.60 ± 17.61 years) to be older, with higher (p<0.001) levels of total cholesterol, glycaemia, DBP, waist circumference, BMI, and serum hs-CRP (resp., 175.75 ± 85.83 mg/dl; 110.14 ± 49.85 mg/dl; 83.04 ± 12.94 mmHg; 96.56 ± 14.40 cm; 31.56 ± 6.46 Kg/m²; 13.72 ± 26.96 mg/l for participant with MetS compared to 152.25 ± 70.01 mg/dl; 93.50 ± 22.38 mg/dl; 79.11 ± 12.17 mmHg; 77.50 ± 13.89 cm; 25.48 ± 4.91 Kg/m²; 2.12 ± 3.94 mg/l for normal participants). MetS participants were more likely to have abdominal obesity (95.41%), obesity (56.12%), diabetes (44.90%), and hypertension (58.16%) (Table 2).

The variation in the prevalence of MetS by sex and age in the study participants is shown in Figure 2. The overall prevalence of MetS was 32.45 % with highly significant females predominance (46.11% for females and 14.01 % for males; p<0.001). The data show an increase of MetS prevalence with age up to 59 years. A small decline in this prevalence was observed in patients aged 60 years and above (Figure 2). The highest prevalence of MetS was found in male (34.78%) participants aged 50-<60 years and female (63.64 %) participants aged 30-<40 years [also see Table S1].

The variation in the frequency of risk factors associated with MetS by sex and age group in the population is presented in Figure 3. According to these results, in the entire participants, the most common abnormalities were

	IABLE I: Features	LABLE 1: Features of persons who participated in the study according to sex in total participants.	in the study according to se	x in total participants.	
		Total participants	Males	Females	A under (hatering male and famala)
Total, n (%)		604	257 (57.45)	347 (42.55)	<i>p value</i> (detween male and lemale)
Age (years) ^a		43.74±17.21	40.87 ± 17.29	45.87 ± 16.87	< 0.001
Educational level, n (%) ^b	Uneducated	11 (1.82)	1 (2.88)	10 (0.39)	< 0.001
	Primary	200 (33.11)	66 (38.62)	134 (25.68)	
	secondary	305 (50.50)	136 (48.70)	169 (52.92)	
	University	88 (14.57)	54(9.80)	34 (21.01)	
Residence area, n (%) ^b					0.487
	Mbouda	424 (70.20)	188 (73.15)	236 (68.01)	
	Batcham	118 (19.54)	45 (17.51)	73 (21.04)	
	Galim	35 (5.79)	15 (5.84)	20 (5.76)	
	Babadjou	27 (4.47)	9 (3.50)	18 (5.19)	
Drinking, n (%) ^b					< 0.001
	Never drinker	119 (19.70)	40 (15.56)	79 (22.77)	
	Moderate alcohol drinker	299(49.50)	117 (45.53)	182 (52.45)	
	Excessive alcohol drinker	186 (30.79)	100 (38.91)	86 (24.78)	
Smoking, n (%) ^b					< 0.001
	Current smoker	21(3.48)	20 (7.78)	1(0.29)	
	Former smoker	69 (11.42)	47 (18.29)	22 (6.34)	
	Never smoker	514(85.10)	190 (73.93)	324 (93.37)	
BMI (kg/m ²) ^a ;		27.45±6.16	26.69 ± 5.22	28.02±6.72	0.006
BMI, n (%) ^b					0.004
	Underweight	16 (2.65)	4(1.56)	12 (3.46)	
	Normal weight	222 (36.75)	105(40.86)	117 (33.72)	
	Overweight	176 (29.14)	85 (33.07)	91 (26.22)	
	Obesity	190(31.46)	63 (24.51)	127 (36.60)	
Physical activity, n (%)					0.001
	Low	405 (67.05)	154 (59.92)	251 (72.33)	
	Moderate	147(24.34)	70 (27.24)	77 (22.19)	
	High	41 (6.79)	28 (10.89)	13 (3.75)	
	Very high	11 (1.82)	5 (1.95)	6 (1.73)	
Waist circumference (cm) ^a ;		83.68±16.64	85.27±15.15	82.51±17.59	0.039
Abdominal obesity, n (%) ^b					< 0.001
	No	360 (59.60)	202 (78.60)	158(45.53)	
	Yes	244(40.40)	55 (21.40)	189 (54.47)	
Diabetes, n (%) ^b		37 (6.13)	16 (6.23)	21 (6.05)	0.929
Hypertension, n (%) ^b		266 (44.04)	119 (46.30)	147 (42.36)	0.334
Hyper-LDL cholesterol, n (%) ^b		138 (22.85)	49 (19.07)	89 (25.65)	0.056
Hyper-total cholesterol, $n (\%)^b$		163 (26.99)	62 (24.12)	101 (29.11)	0.172
Hypo-total cholesterol, n (%) ^b		291 (48.18)	137 (53.31)	154 (44.38)	0.029
Hypertriglyceridemia, n (%) ^b		326 (53.97)	149 (57.98)	177 (51.01)	0.089
Hypo-HDL cholesterol, n (%) ^b		500 (82.78)	200 (77.82)	300 (86.46)	0.005
Diastolic blood pressure (mmHg) ^a ; [min-max]	g) ^a ; [min-max]	80.39±12.55 [40-120]	80.95±10.87 [54-112]	79.97±13.67 [40-120]	0.329
Systolic blood pressure (mmHg) ^a ; [min-max]	a; [min-max]	129.20±20.89 [80-195]	131.49±21.11 [100-195]	127.50±20.58 [80-188]	0.020
4	1				

TABLE 1: Features of persons who participated in the study according to sex in total participants.

4

BioMed Research International

Total $p(0)$		Normal participants 408 (67.55)	Participants with MetS	<i>p value</i> (between normal and MetS)
Total, n (%) Age (years) ^a		408 (07.55) 41.60±17.61	196 (32.45) 48.19±15.48	< 0.001
Educational level, n (%) ^b		41.00±17.01	40.17±13.40	< 0.001
Educational level, if (70)	Uneducated	6 (1.47)	5 (2.55)	< 0.001
	Primary	125 (30.64)	75 (38.27)	
	secondary	201 (49.26)	104 (53.06)	
	University	76 (18.63)	12 (6.12)	
Residence area, n (%) ^b			(**)	0.648
	Mbouda	292 (71.57)	132 (67.35)	
	Batcham	78 (19.12)	40 (20.41)	
	Galim	22 (5.39)	13 (6.63)	
	Babadjou	16 (3.92)	11 (5.61)	
Drinking, n (%) ^b	,			0.017
0	Never drinker	89 (21.81)	30 (15.31)	
Ν	Ioderate alcohol drinker	186 (45.59)	113 (57.65)	
E	xcessive alcohol drinker	133 (32.60)	53 (27.04)	
Smoking, n (%) ^b				0.080
	Current smoker	18 (4.41)	3 (1.53)	
	Former smoker	51 (12.50)	18 (9.18)	
	Never smoker	339 (83.09)	175 (89.29)	
BMI (kg/m ²) ^a		25.48 ± 4.91	31.56±6.46	< 0.001
BMI, n (%) ^b				< 0.001
	Underweight	16 (3.92)	0 (0.00)	
	Normal weight	205 (50.25)	17 (8.67)	
	Overweight	107 (26.23)	69 (35.21)	
	Obesity	80 (19.60)	110 (56.12)	
Physical activity, n (%)				0.007
	Low	257 (62.99)	148 (75.51)	
	Moderate	107 (26.23)	40 (20.41)	
	High	35 (8.58)	6 (3,06)	
	Very high	9 (2.21)	2 (1.02)	
Waist circumference (cm) ^a		77.50±13.89	96.56±14.40	< 0.001
Abdominal obesity, n (%) ^b				< 0.001
	No	351 (86.03)	9 (4.59)	
	Yes	57 (13.97)	187 (95.41)	
Diabetes, n (%) ^b		88 (21.57)	88 (44.90)	< 0.001
Hypertension, n (%) ^b		152 (37.25)	114 (58.16)	< 0.001
Diastolic blood pressure (mr	nHg)ª; [min-max]	79.11±12.17 [40-108]	83.04±12.94 [50-120]	< 0.001
Systolic blood pressure (mm	Hg) ^a ; [min-max]	127.32±21.10 [80-195]	133.12±19.92 [80-188]	0.001
Glycaemia (mg/dl) ^a ; [min-M	lax]	93.50±22.38 [55-283]	110.14±49.85 [54-448]	< 0.001
HDL cholesterol (mg/dl) ^a ; [r	nin-max]	34.11±14.70 [3-126]	31.00±16.24 [11-190]	0.023
LDL cholesterol (mg/dl) ^a ; [n	nin-max]	84.00±65.96 [10-357]	118.26±85.95 [20-175]	< 0.001
Total cholesterol (mg/dl) ^a ; [r	nin-max]	152.25±70.01 [37-187]	175.75±85.83 [50-618]	< 0.001
Triglyceride (mg/dl) ^a ; [min-	max]	163.70±71.72 [24-209]	178.43±71.14 [33-479]	0.018
hs-CRP (mg/l)		2.12±3.94 [0.005-27]	13.72±26.96 [0.005-192]	< 0.001

TABLE 2: Features of participants according to metabolic syndrome status in total participants.

^aAge, BMI, diastolic blood pressure; systolic blood pressure; glycaemia; HDL cholesterol; LDL cholesterol; total cholesterol; and triglyceride were expressed as mean ± SD. BMI, body mass index.

^aWelch*t*-test; ^bChi-square.

MetS, metabolic syndrome; HDL, high-density lipoprotein; LDL, low-density lipoprotein; n, size; min, minimum; max, maximum.

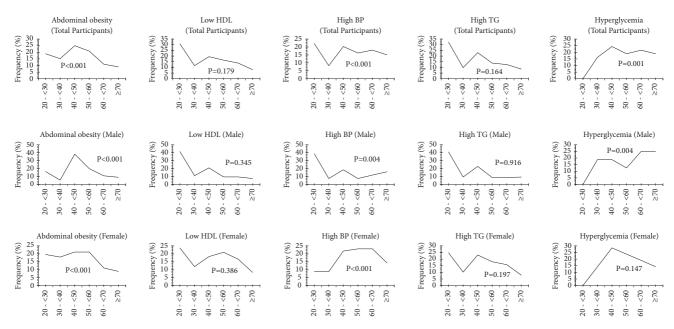


FIGURE 3: Frequency (%) of MetS components by sex and age groups in the study population. P value (between age groups); MetS, metabolic syndrome; HDL, high-density lipoprotein; n, size; BP, blood pressure.

low-HDL (82.78%) and hypertriglyceridemia (53.97%). The most affected groups were 20-<30 years (HDL, 30.80%, and hypertriglyceridemia, 32.21%) and 40-<50 years (HDL, 19.40%, and hypertriglyceridemia, 23.01%). In the male and female participants, low-HDL was the common abnormality affecting participants aged 20-<30 years. Figure 3 also shows that abdominal obesity significantly decreases with age, with a maximum among participants aged 40-<50 years in the total, male, and female participants. The majority of abnormalities associated with MetS were expressed in their highest frequencies in participants aged 20-<30 years for some parameters and 40-<50 years for others [also see Table S2].

In this study, we examined the frequency of the number of MetS components by sex (Figure 4) and age group (Figure 5). In our study population, 34.27% (31.13% in males and 36.60% in females) of the participants had at least three MetS components, which is the definition of MetS (Figure 4). Those with two risk factors represent 29.97% of the population and are at risk of developing MetS. The participants aged 20-<30 years and 40-<50 years accumulated more risk factors compared to the other age groups among both males and females. In total participants, the variation of severe MetS (five associated anomalies) was significant in all age groups (p<0.001) (Figure 5(a)) [see Table S3 for more details].

The association between sociodemographic parameters, hs-CRP levels, and MetS in total participants was studied and presented in Table 3. Participants with obesity (OR: 16.34; 95% CI: 9.21-28.96) and overweight (OR: 7.45; 95% CI: 4.17-13.30) had a higher risk of developing MetS. Participants aged 50-<60 years (OR: 5.66; 95% CI: 3.17-10.12) had a higher MetS risk (Table 3). This study has shown that participants with the highest hs-CRP (hs-CRP >11 mg/l) had a 4.37-fold increased risk of MetS compared to those with the lowest hs-CRP (hs-CRP [0-11] mg/l).

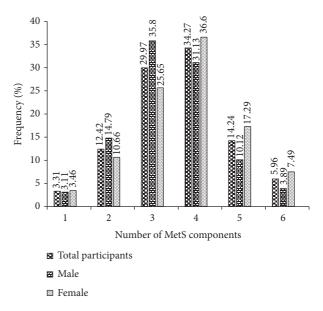


FIGURE 4: Frequency (%) of the number of metabolic syndrome components by sex in the study population. MetS, metabolic syndrome; 0: no metabolic syndrome components; 1: one metabolic syndrome component; 2: two metabolic syndrome components; 3: three metabolic syndrome components; 4: four metabolic syndrome components; 5: five metabolic syndrome components.

Logistic regression analysis was used to study the most common MetS definition parameters among the participants (Table 4). According to the results, in the entire population, abdominal obesity (OR: 353.13; 95% CI: 136.16-915.81), low-HDL (OR: 9.28; 95% CI: 3.98-21.62), high-TG (OR: 5.62; 95% CI: 2.69-11.74), high blood pressure (OR:

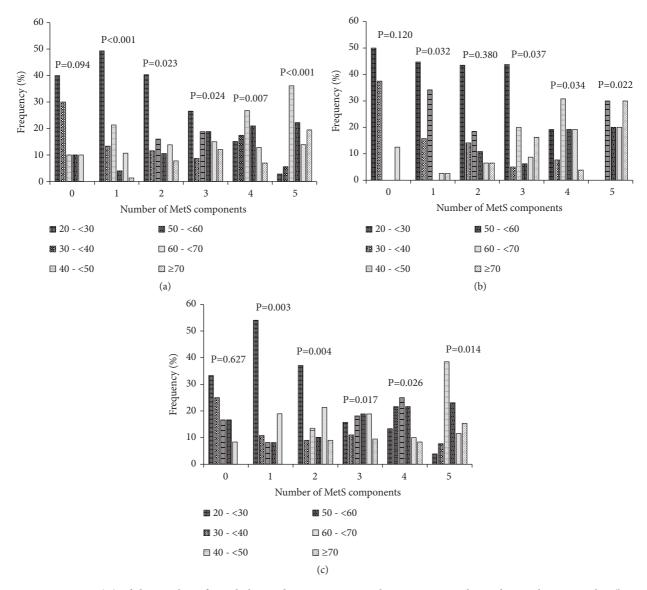


FIGURE 5: Frequency (%) of the number of metabolic syndrome components by age group in the study population. P value (between age groups); (a) overall population; (b) female participants; (c) male participants; MetS, metabolic syndrome; 0: no metabolic syndrome components; 1: one metabolic syndrome component; 2: two metabolic syndrome components; 3: three metabolic syndrome components; 4: four metabolic syndrome components; 5: five metabolic syndrome components.

4.43; 95% CI: 2.27-8.63), and hyperglycemia (OR: 4.24; 95% CI: 2.11-8.52) are the most common abnormalities affecting participants. The results suggest that abdominal obesity is the strongest predictor of MetS in our study participants (Table 4).

Table S4 shows the average frequency of food intake of the participants according to metabolic syndrome. MetS participants consumed more fishes (p<0.001), fufu corn (p=0.010), and cabbage (p=0.045) than normal participants who in turn consumed more pasta (p<0.001) and sugar products (p=0.027) than MetS participants.

Associations between the frequency of foods intake and metabolic syndrome prevalence in total participants are shown in Table S5. A high frequency of fish intake was associated with lower odds of having MetS after adjustment for confounding factors.

4. Discussion

In the present study, we evaluated the prevalence of MetS, its individual components, and associated risk factors among Bamboutos Division's adult population using a Joint Interim Statement of the International Diabetes Federation Task Force on Epidemiology and Prevention definitions parameters [13]. Results on features of persons who participated in the study according to sex showed a significant difference between the two groups in terms of educational level, drinking and smoking status, physical activities, abdominal obesity, and serum hs-CRP. Female participants (45.87±16.87 years) were more likely than males participants (40.87±17.29 years) to be older, with lower levels of SBP. Sociodemographic characteristics of population vary geographically and depend on the socioeconomic and sociopolitical development of each

			Metabolic syndrome	
	Categories/groups	OR	95% CI	p value
Age (years)				
	20 - <30	(ref)		
	30 - <40	4.13	2.22-7.69	< 0.001
	40 - <50	4.37	2.53-7.55	< 0.001
	50 - <60	5.66	3.17-10.12	< 0.001
	60 - <70	2.80	1.50-5.20	0.001
	≥70	3.04	1.51-6.13	0.001
Educational level, n (%) ^b				
	Uneducated	(ref)		
	Primary	0.72	0.21-2.44	0.597
	Secondary	0.62	0.18-2.08	0.440
	University	0.19	0.05-0.72	0.014
Residence area, n (%) ^b				
	Mbouda	(ref)		
	Batcham	1.13	0.73-1.74	0.568
	Galim	1.30	0.63-2.67	0.463
	Babadjou	1.52	0.68-3.37	0.301
Drinking, n (%) ^b				
	Never drinker	(ref)		
	Moderate alcohol drinker	1.80	1.12-2.89	0.015
	Excessive alcohol drinker	1.18	0.70-1.99	0.530
Smoking, n (%) ^b				
	Never smoker	(ref)		
	Former smoker	2.11	0.55-8.01	0.270
	Current smoker	3.09	0.09-10.65	0.073
BMI, n (%) ^b				
	Normal weight	(ref)		
	Underweight	0.00	0.00-0.00	0.971
	Overweight	7.45	4.17-13.30	< 0.001
	Obesity	16.34	9.21-28.96	< 0.001
Physical activity, n (%)				
	Low	(ref)		
	Moderate	0.65	0.42-0.98	0.041
	High	0.30	0.12-0.72	0.007
	Very high	0.38	0.08-1.81	0.227
hs-CRP (mg/l)	[0-11[(ref)		
-	>11	4.37	2.45-7.77	< 0.001

TABLE 3: Association between sociodemographic parameters, hs-CRP levels, and metabolic syndrome in total participants.

BMI, body mass index; hs-CRP, high sensitivity C-reactive protein; n, size; OR, odds ratio; CI, confidence interval; ref, reference=1.

TABLE 4: Association between the variables related to metabolic syndrome in the total participants.

	Metabolic syndrome		
	OR	95% CI	p value
Abdominal obesity	353.13	136.16-915.81	< 0.001
Low HDL	9.28	3.98-21.62	< 0.001
High TG	5.62	2.69-11.74	< 0.001
High blood pressure	4.43	2.27-8.63	< 0.001
Hyperglycemia	4.24	2.11-8.52	< 0.001
High total-cholesterol	2.13	1.08-4.21	0.028

HDL, high-density lipoprotein; TG, triglyceride; n, size; OR, odds ratio; CI, confidence interval.

region. The sex differences on serum hs-CRP level observed in this study could be explained by endogenous synthesis of hormone like oestrogen that might play a role in the inflammatory process in women. This could also be explained by the fact that women compared to men have a large amount of adipose tissue source of proinflammatory cytokine [18].

According to the MetS status, participants with MetS were more likely older than the normal participants. These results are in agreement with those obtained by Li and collaborators among adults in China [19]. Participants with metabolic syndrome had higher levels of total cholesterol, glycaemia, DBP, waist circumference, BMI, and serum hs-CRP and were more likely to have abdominal obesity (95.41%), obesity (56.12%), diabetes (44.90%), and hypertension (58.16%) compared to normal participants. Previous studies revealed that metabolic dysfunctions such as high blood pressure, hyperglycemia, and obesity are the factors related to the MetS [20–22].

The overall prevalence of MetS in this study was 32.45 % with highly significant female predominance (46.11% for females and 14.01 % for males). The overall metabolic prevalence obtained in this study is similar to the prevalence rate of Dandji et al. (38.98%) among adult men of Dschang Health District [12], Lee et al. (30.52%) in south Korea [4], Brini et al. (35.73%) in Morocco [10], and Sabir et al. (35.1%) in northwestern Nigeria [23]. However, it is high compared to the prevalence in the Eastern Cape, South Africa (21.8%) by Owolabi et al. [9]. High prevalence of the MetS in Mbouda adults may be caused by high prevalence of obesity, hypertension, and diabetes in this population. This may also be due to the lifestyle and some genetics factors. High prevalence of the MetS in females compared to males corroborates with the prevalence rate of Brini et al. (40.12% among women and 18.56% among men) in Morocco [10]. This finding is consistent with that of Belfki et al. in Tunisia [24] and differs with the finding of Santos et al. among South European population [25]. These results might be due to different cutoff points set as criteria for metabolic syndrome like abdominal obesity, low-HDL cholesterol, and hypertriglyceridemia. Women in the menopausal state had a decline in circulating oestrogen levels. This decrease in oestrogen concentration may increase cardiovascular diseases in women through effects on adiposity, lipid metabolism, and prothrombotic state [26]. This is consistent with other results of this study, which have shown that female participants were more likely to be older than male participants.

The study of the variation in the frequency of the MetS components by sex and age group has shown that low HDL (82.78%) and hypertriglyceridemia (53.97%) were the most common abnormalities in the entire participants. It has also revealed that participants aged 20-<30 years and 40-<50 years were the most affected and that abdominal obesity significantly decreases with age with a maximum among participants. Cameroon's population is currently very young, and the above results are mainly driven by the constant rise in diabetes, obesity, and hypertension in Cameroonian population [27–29]. Diabetes and obesity increase the risk of adipose tissue insulin resistance, which plays important role in the pathophysiology of the MetS [30]. High prevalence

of MetS in females throughout the age groups could be explained by the high prevalence of obesity among female participants in this study. It may be also attributable to the steep increase in blood pressure in women after menopause, which initiates a more rapid decrease in endothelial function [31].

Regarding the trend curves of abdominal obesity and hyperglycemia as a function of age groups in males in Figure 3, abdominal obesity would be due to waist circumference values. Waist circumference tended to be higher in younger adults than in older men [32, 33]. A hormonal process involving endogenous oestrogen, which would provoke hyperglycemia in men, could explain the trend curves of hyperglycemia as a function of age groups in males [34].

This study reveals that 29.97% of the population are at risk of developing MetS. Other studies showed that MetS is emerging alarmingly in low-income countries [35, 36]. This may be due to increasing urbanization, westernization of lifestyle including unhealthy diet, physical inactivity, and lack of awareness about metabolic syndrome.

The results on the study of associations between MetS and sociodemographic parameters showed that obesity (OR: 16.34; 95% CI: 9.21-28.96), overweight (OR: 7.45; 95% CI: 4.17-13.30), and participants aged 50-<60 years (OR: 5.66; 95% CI: 3.17-10.12) had a higher MetS risk. They also show that participants with the highest hs-CRP (hs-CRP >11 mg/l) had a 4.37-fold increased risk of MetS compared to those with the lowest hs-CRP (hs-CRP [0-11], mg/l). Considering the everincreasing body of evidence regarding MetS, chronic lowgrade inflammation may have an important role in the pathogenesis of metabolic disorders [37]. Concerning parameters used for MetS definition, abdominal obesity (OR: 353.13; 95% CI: 136.16-915.81), low HDL (OR: 9.28; 95% CI: 3.98-21.62), high-TG (OR: 5.62; 95% CI: 2.69-11.74), high blood pressure (OR: 4.43; 95% CI: 2.27-8.63), and hyperglycemia (OR: 4.24; 95% CI: 2.11-8.52) are the most common abnormalities affecting participants. Obesity is known as a risk factor of the MetS and our study results are consistent with those of Brini et al. [10], Moreira et al. [38], and Carnethon et al. [39]. The results suggest that abdominal obesity is the strongest predictor of MetS in our study participants.

The present study also suggested that the frequent consumption of fishes was associated with lower odds of having MetS. The possible explanation for this association is that fish proteins are easily digestible, rich in essential amino acids, and have been seen to slow absorption and synthesis of lipids and promote the lipid excretion [40].

The main strength of the current study is that it is one of the large-sample-size studies regarding MetS in Cameroon. The results of this study would certainly contribute to the sensitization and the prevention of the MetS in Bamboutos Division. However, several limitations should be considered. First, the cross-sectional design limits the ability to address causal relationships between risk factors and metabolic syndrome. Second, the prevalence of metabolic syndrome was based on a single assessment of blood samples, which may lead to minor inaccuracies. Third, because the sociodemographic characteristics and dietary information were obtained through a questionnaire, this may lead to recall bias. Fourth, the species of fish, preparation methods, seasonal variation, and possible contaminants of fish consumed could not be examined in the current study and assessment of these factors will provide additional information regarding fish-MetS associations.

5. Conclusion

The present study discloses high prevalence of metabolic syndrome among our study population and significant females predominance compared to males. In addition, onethird of the study population were at risk of metabolic syndrome. Low-HDL cholesterol, hypertriglyceridemia, high blood pressure, and abdominal obesity were the common abnormalities among participants. Our results suggest specific association between risk factors including sociodemographic features abdominal obesity, low-HDL cholesterol, high blood pressure, and hyperglycemia with metabolic syndrome. The results also suggest that abdominal obesity is the strongest predictor of metabolic syndrome in our study participants. The findings highlight the need for evidencebased prevention, diagnosis, and management of metabolic syndrome and its associated factors among Bamboutos Division adults in Cameroun.

Abbreviations

ATP III:	Adult Treatment Panel III
BMI:	Body mass index
BP:	Blood pressure
CDC:	Centers for Disease Control
CNEC:	Cameroon National Ethics Committee
DBP:	Diastolic blood pressure
FPG:	Fasting plasma glucose
HDL:	High-density lipoprotein
hs-CRP:	High sensitivity C-reactive protein
HIV:	The human immunodeficiency virus
IDF:	International Diabetes Federation
LDL:	Low-density lipoprotein
MetS:	Metabolic syndrome
SBP:	Systolic blood pressure
SD:	Standard Deviation
TG:	Triglyceride.

Data Availability

All data generated or analyzed during this study are included in this published article and the supporting file.

Conflicts of Interest

The authors declare that there are no conflicts of interest.

Authors' Contributions

Wiliane J. T. Marbou performed the sampling and data collection. Both Wiliane J. T. Marbou and Victor Kuete designed the study, participated in the analysis of data, drafted the manuscript, and read the manuscript and approved the final version prior to submission.

Acknowledgments

The authors would like to thank Dr. NJIFOU NJIMAH Amadou (Director of the Mbouda District Hospital), Dr. AKANA Cyriaque (Director of the Mbouda ADLUCEM Hospital), all the staffs, and participants for their support towards the successful completion of the study.

Supplementary Materials

Supporting file (DOCX): Table S1: variation in the prevalence of metabolic syndrome by sex and age group in the entire population; Table S2: frequency (%) of metabolic syndrome components by sex and age group in the study population; Table S3: frequency (%) of the number of metabolic syndrome components by sex and age group in the study population; Table S4: average frequency of foods intake of participants according to metabolic syndrome; Table S5: associations between the frequency of foods intake and metabolic syndrome in total participants (n = 604). (Supplementary Materials)

References

- [1] B. Naby and M. John, "IDF Afria," https://www.idf.org/ournetwork/regions-members/africa/welcome.
- [2] A. D. Kaze, A. E. Schutte, S. Erqou, A. P. Kengne, and J. B. Echouffo-Tcheugui, "Prevalence of hypertension in older people in Africa: A systematic review and meta-analysis," *Journal of Hypertension*, vol. 35, no. 7, pp. 1345–1352, 2017.
- [3] S. Biadgilign, T. Mgutshini, D. Haile, B. Gebremichael, Y. Moges, and K. Tilahun, "Epidemiology of obesity and overweight in sub-Saharan Africa: A protocol for a systematic review and meta-analysis," *BMJ Open*, vol. 7, no. 11, Article ID e017666, 2017.
- [4] S. E. Lee, K. Han, Y. M. Kang et al., "Trends in the prevalence of metabolic syndrome and its components in South Korea: Findings from the Korean National Health Insurance Service Database (2009–2013)," *PLoS ONE*, vol. 13, no. 3, Article ID e0194490, 2018.
- [5] E. S. Ford, W. H. Giles, and W. H. Dietz, "Prevalence of the metabolic syndrome among US adults: findings from the third national health and nutrition examination survey," *Journal of the American Medical Association*, vol. 287, no. 3, pp. 356–359, 2002.
- [6] C. Pitsavos, D. Panagiotakos, M. Weinem, and C. Stefanadis, "Diet, exercise and the metabolic syndrome," *The Review of Diabetic Studies*, vol. 3, no. 3, pp. 118–126, 2006.
- [7] L. Fezeu, B. Balkau, A.-P. Kengne, E. Sobngwi, and J.-C. Mbanya, "Metabolic syndrome in a sub-Saharan African setting: Central obesity may be the key determinant," *Atherosclerosis*, vol. 193, no. 1, pp. 70–76, 2007.
- [8] A. A. Sabir, A. Jimoh, S. O. Iwuala et al., "Metabolic syndrome in urban city of North-Western Nigeria: Prevalence and determinants," *Pan African Medical Journal*, vol. 23, 2016.
- [9] E. O. Owolabi, D. T. Goon, O. V. Adeniyi, A. O. Adedokun, and E. Seekoe, "Prevalence and correlates of metabolic syndrome among adults attending healthcare facilities in eastern cape,

South Africa," *The Open Public Health Journal*, vol. 10, pp. 148–159, 2017.

- [10] O. El Brini, O. Akhouayri, A. Gamal, A. Mesfioui, and B. Benazzouz, "Prevalence of metabolic syndrome and its components based on a harmonious definition among adults in Morocco," *Diabetes, Metabolic Syndrome and Obesity: Targets and Therapy*, vol. 7, pp. 341–346, 2014.
- [11] G. Omuse, D. Maina, M. Hoffman et al., "Metabolic syndrome and its predictors in an urban population in Kenya: A cross sectional study," *BMC Endocrine Disorders*, vol. 17, no. 1, p. 37, 2017.
- [12] M. Dandji, F. Zambou, D. Dangang, F. Nana, and F. Tchouanguep, "Prevalence of metabolic syndrome in adult men of the dschang health district in Western-Cameroon," *World Journal of Nutrition and Health*, vol. 6, no. 1, pp. 1–10, 2018.
- [13] K. G. Alberti, R. H. Eckel, S. M. Grundy et al., "Harmonizing the metabolic syndrome: a joint interim statement of the international diabetes federation task force on epidemiology and prevention; National heart, lung, and blood institute; American heart association; World heart federation; International atherosclerosis society; and international association for the study of obesity," *Circulation*, vol. 120, no. 16, pp. 1640–1645, 2009.
- [14] R. Tsobou, P. M. Mapongmetsem, and P. Van Damme, "Medicinal plants used for treating reproductive health care problems in cameroon, central africal," *Economic Botany*, vol. 70, no. 2, pp. 145–159, 2016.
- [15] CDC, "About Adult BMI," https://www.cdc.gov/healthyweight/ assessing/bmi/adult_bmi/index.html.
- [16] W. T. Friedewald, R. I. Levy, and D. S. Fredrickson, "Estimation of concentration of low density lipoprotein cholesterol in plasma without use of ultracentrifuge," *Clinical Chemistry*, vol. 18, pp. 499–502, 1972.
- [17] P. S. Jellinger, Y. Handelsman, P. D. Rosenblit et al., "Special advisor michael davidson: 1 CPG for managing dyslidemia and prevention of CVD," *Endocrine Practice*, vol. 23, no. 2, 2017.
- [18] T. S. Han, N. Sattar, K. Williams, C. Gonzalez-Villalpando, M. E. J. Lean, and S. M. Haffner, "Prospective study of C-reactive protein in relation to the development of diabetes and metabolic syndrome in the Mexico City diabetes study," *Diabetes Care*, vol. 25, no. 11, pp. 2016–2021, 2002.
- [19] Y. Li, L. Zhao, D. Yu, Z. Wang, G. Ding, and Y. Li, "Metabolic syndrome prevalence and its risk factors among adults in China: A nationally representative cross-sectional study," *PLoS ONE*, vol. 13, no. 6, Article ID e0199293, 2018.
- [20] J. Kaur, "A comprehensive review on metabolic syndrome," *Cardiology Research and Practice*, vol. 2014, Article ID 943162, 21 pages, 2014.
- [21] N.-K. Ki, H.-K. Lee, J.-H. Cho, S.-C. Kim, and N.-S. Kim, "Factors affecting metabolic syndrome by lifestyle," *Journal of Physical Therapy Science*, vol. 28, no. 1, pp. 38–45, 2016.
- [22] M. A. Cornier, D. Dabelea, T. L. Hernandez et al., "The metabolic syndrome," *Endocrine Reviews*, vol. 29, no. 7, pp. 777– 822, 2008.
- [23] A. A. Sabir, A. Jimoh, S. O. Iwuala et al., "Metabolic syndrome in urban city of North-Western Nigeria: Prevalence and determinants," *Pan African Medical Journal*, vol. 23, no. 19, 2016.
- [24] H. Belfki, S. B. Ali, H. Aounallah-Skhiri et al., "Prevalence and determinants of the metabolic syndrome among Tunisian adults: Results of the Transition and Health Impact in North Africa (TAHINA) project," *Public Health Nutrition*, vol. 16, no. 4, pp. 582–590, 2013.

- [25] A. C. Santos, M. Severo, and H. Barros, "Incidence and risk factors for the metabolic syndrome in an urban South European population," *Preventive Medicine*, vol. 50, no. 3, pp. 99–105, 2010.
- [26] J. G. Schneider, C. Tompkins, R. S. Blumenthal, and S. Mora, "The metabolic syndrome in women," *Cardiology in Review*, vol. 14, no. 6, pp. 286–291, 2006.
- [27] R. M. Kamadjeu, R. Edwards, J. S. Atanga, N. Unwin, E. C. Kiawi, and J.-C. Mbanya, "Prevalence, awareness and management of hypertension in Cameroon: findings of the 2003 Cameroon Burden of Diabetes Baseline Survey," *Journal of Human Hypertension*, vol. 20, no. 1, pp. 91-92, 2006.
- [28] World Diabetes Foundation, "Cameroon National Diabetes and Hypertension Programme, WDF16-1429 — World diabetes foundation," https://www.worlddiabetesfoundation.org/ projects/cameroon-wdf16-1429.
- [29] S.-P. Choukem, A.-P. Kengne, M.-L. Nguefack et al., "Fouryear trends in adiposity and its association with hypertension in serial groups of young adult university students in urban Cameroon: a time-series study," *BMC Public Health*, vol. 17, no. 1, 2017.
- [30] J. Denis McGarry, "Dysregulation of fatty acid metabolism in the etiology of type 2 diabetes," *Diabetes*, vol. 51, no. 1, pp. 7–18, 2002.
- [31] A. Stefanska, K. Bergmann, and G. Sypniewska, "Metabolic syndrome and menopause: pathophysiology, clinical and diagnostic significance," *Advances in Clinical Chemistry*, vol. 72, pp. 1–75, 2015.
- [32] E. Oda and R. Kawa, "Age- and gender-related differences in correlations between abdominal obesity and obesity-related metabolic risk factors in Japanese," *Internal Medicine*, vol. 48, no. 7, pp. 497–502, 2009.
- [33] I. Wakabayashi and T. Daimon, "Age-dependent decline of association between obesity and hyperglycemia in men and women," *Diabetes Care*, vol. 35, no. 1, pp. 175–177, 2012.
- [34] M. R. Meyer, D. J. Clegg, E. R. Prossnitz, and M. Barton, "Obesity, insulin resistance and diabetes: sex differences and role of oestrogen receptors," *Acta Physiologica*, vol. 203, no. 1, pp. 259–269, 2011.
- [35] R. BeLue, T. A. Okoror, J. Iwelunmor et al., "An overview of cardiovascular risk factor burden in sub-Saharan African countries: a socio-cultural perspective," *Globalization and Health*, vol. 5, no. 1, p. 10, 2009.
- [36] M. G. Saklayen, "The global epidemic of the metabolic syndrome," *Current Hypertension Reports*, vol. 20, no. 2, p. 12, 2018.
- [37] M. N. Chowta, P. M. Adhikari, R. Sinha, S. D. Acharya, H. N. Gopalakrishna, and J. T. Ramapuram, "Highly sensitive C reactive protein in patients with metabolic syndrome and cardiovascular disease," *Annals of Tropical Medicine and Public Health*, vol. 5, no. 2, pp. 98–102, 2012.
- [38] G. C. Moreira, J. P. Cipullo, L. A. S. Ciorlia, C. B. Cesarino, and J. F. Vilela-Martin, "Prevalence of metabolic syndrome: Association with risk factors and cardiovascular complications in an urban population," *PLoS ONE*, vol. 9, no. 9, Article ID e105056, 2014.
- [39] M. R. Carnethon, C. M. Loria, J. O. Hill, S. Sidney, P. J. Savage, and K. Liu, "Risk factors for metabolic syndrome: the coronary artery risk development in young adults (CARDIA) study, 1985-2001," *Diabetes Care*, vol. 27, no. 11, pp. 2707–2715, 2004.
- [40] D. El Khoury and G. H. Anderson, "Recent advances in dietary proteins and lipid metabolism," *Current Opinion in Lipidology*, vol. 24, no. 3, pp. 207–213, 2013.