## **East African Scholars Journal of Medical Sciences**

Abbreviated Key Title: East African Scholars J Med Sci ISSN: 2617-4421 (Print) & ISSN: 2617-7188 (Online) Published By East African Scholars Publisher, Kenya

Volume-6 | Issue-3 | Mar-2023 |

#### **Original Research Article**

DOI: 10.36349/easms.2023.v06i03.005

OPEN ACCESS

# **Evaluation of Metabolic Syndrome Using Lipid Accumulation Products,** Visceral Adiposity Index and Body Mass Index in Apparently Healthy Students of University of Maiduguri

Alhaji Haruna Musa<sup>1</sup><sup>(1)</sup>\*, Ijagila, I. N<sup>1</sup>, Dungus, M. M<sup>2</sup>

<sup>1</sup>Department of Medical Laboratory Science, Faculty of Allied Health Sciences, College of Medical Sciences, University of Maiduguri, Maiduguri, Borno State, Nigeria

<sup>2</sup>Department of Chemical Pathology, University of Maiduguri Teaching Hospital, 600104, Maiduguri, Borno, Nigeria

Article History Received: 13.02.2023 Accepted: 18.03.2023 Published: 26.03.2023

Journal homepage: https://www.easpublisher.com



Abstract: Introduction: Metabolic Syndrome (MetS) is a group of conditions that together raise your risk of cardiovascular disease, diabetes mellitus, dyslipidemia and hypertension. The diagnosis of MetS was established according to the revised criteria of the NCEP/ATP III (MS-NCEP/ ATP III). Although BMI is the most common screening measures to identify people who are at relatively high risk of MetS, this tool is not particularly the most effective tool. Thus, we seek to find single parameter with the strongest diagnostic accuracy for MetS in a sample of healthy, unrelated adults. Materials & Methods: The study was conducted on 200 apparently healthy male and female students of university of Maiduguri. Random sampling techniques were used to recruit the subjects aged 18-41 years with mean age of 25.65±5.56 years for males and 24.11±4.60 years for females. Examination of subjects consisted of physical examination with measurement of anthropometric and clinical parameters, filling out a questionnaire, and evaluation of serum lipid levels. Result: The result shows the mean±SD for the components of MetS includes BP (SBP=105.67±9.82; DBP=70.43±6.16); TG=1.69±0.72; HDL=1.10±0.37; FBG=3.88±0.72; WC=87.76±8.41 for males and BP (SBP=104.69±9.93; DBP=69.93±6.44), TG=1.82±0.69; HDL=1.13±0.39; FBG=3.64±0.79; WC=86.93±9.89 for females. The result shows prevalence of MetS in males was 8.1% and 21.5% in females. The result also shows that LAP has the highest diagnostic accuracy among the study population (AUC=0.856) than VAI (AUC=0.820) and BMI (AUC=0.523). LAP was also found to have highest diagnostic accuracy in males (AUC=0.908) than VAI (AUC=0.850) and BMI (AUC=0.498) while in females VAI was found to have highest diagnostic accuracy with AUC (0.865) than LAP (AUC=0.721) and BMI (AUC=0.436). Conclusion: This study therefore shows LAP has the highest diagnostic accuracy and particularly the most effective tool in identifying males who are at the risk of MetS. While VAI has the highest diagnostic accuracy and particularly the most effective tool in identifying females who are at the risk of MetS.

Keywords: Metabolic syndrome, Lipid, Visceral, Healthy, Apparently.

Copyright © 2023 The Author(s): This is an open-access article distributed under the terms of the Creative Commons Attribution 4.0 International License (CC BY-NC 4.0) which permits unrestricted use, distribution, and reproduction in any medium for non-commercial use provided the original author and source are credited.

#### **1. INTRODUCTION**

Metabolic Syndrome (MetS) is a group of conditions that together raise your risk of cardiovascular disease, diabetes mellitus, dyslipidemia and hypertension [1]. MetS has been one of the major public health as well as clinical challenges both in developed and developing countries, [2, 3]. The global prevalence of adults with MetS is estimated to be 20– 25% and is rising [4, 5]. The etiology of MetS is not fully understood, but aging, inflammation, obesity, sedentary lifestyle, and genetics are implicated as predisposing factors [6]. MetS has been diagnosed or defined using the revised diagnostic criteria of National Cholesterol Education Program/Adult Treatment Panel III (NCEP/ATP III)[6].According to the program MetS is defined when at least three of the following five risk

<sup>\*</sup>Corresponding Author: Alhaji Haruna Musa

Department of Medical Laboratory Science, Faculty of Allied Health Sciences, College of Medical Sciences, University of Maiduguri, Maiduguri, Borno State, Nigeria, ORCID ID: 0000-0003-1087-1056

determinants are present: waist circumference  $\geq 90$  cm in men and  $\geq 80$  cm in women; blood pressure > 130/85mmHg or patient is taking antihypertensive medications; high-density lipoprotein cholesterol (HDL-C) < 0.9 mmol/L in men and < 1.03 mmol/L in women; fasting plasma glucose  $\geq 5.6$  mmol/L or patient is undergoing regular treatment for diabetes mellitus; and triglyceride level  $\geq 1.70$  mmol/L. Nevertheless, it would be useful if a simpler index is available for easy diagnosis of individuals at risk of MetS in clinical settings [8].

Accumulation of body fat as visceral adipose tissue (VAT) rather than subcutaneous adipose tissue (SAT) was more closely correlated with CVD [9]. Thus, evaluation of obesity may play an important role in management of CVD. Computed tomography (CT) and magnetic resonance imaging (MRI) are the gold standard for measuring visceral fat, but they are inconvenient and expensive [9-11].

Hypertension is the most common cause of cardiovascular disease. The major causes of hypertension are Age (more common after 50 years of age), family history, genetic composition, Environment (unhealthy diets, stress), excessive salt intake, the consumption of tobacco and alcohol, physical inactivity and obesity [12]. Studies have indicated that hypertension is associated with metabolic disturbances and may be considered a metabolic disorder [10].

Anthropometric measures of central adiposity, such as BMI or abdominal circumference have been used routinely to evaluate MetS but neither could distinguish accumulation of VAT from SAT [9, 12]. biochemical and anthropometric Thus. both measurements have been used to calculate lipid accumulation product (LAP), a novel index of central lipid accumulation based on a combination of waist circumference (WC) and serum triglycerides (TG). Several studies have shown that LAP is a simple, accurate and inexpensive index and is thus widely used as a marker of metabolic disorders and a predictor of MetS [8, 13, 14].

Furthermore, in order to predict VATassociated cardio metabolic risk, a mathematical model, Visceral Adiposity Index (VAI) was developed that uses both anthropometric measurements like Body Mass Index (BMI) expressed in kg/m2 and Waist Circumference (WC) expressed in cm and functional (triglycerides [TG] and high-density lipoprotein [HDL] cholesterol [17, 18]. The use of VAI as a routine indicator for the evaluation of visceral adipose function has been reported to be useful for cardiometabolic risk assessment because of its higher sensitivity and specificity than classical parameters (WC, BMI, and lipids) [17]. Body Mass Index (BMI) has been the most commonly used anthropometric index to define and classify adults into obese, overweight, or normal weight in clinical practices. It is the weight in kilograms divided by the square of height in meters. It is also used in interpreting the individual's fatness and risk factors for the development or the prevalence of many noncommunicable diseases [18]. Thus, early and accurate identification of high-risk individuals for MetS could be important to predict and prevent CVD and type 2 diabetes. However, there are paucity of epidemiological data on the clinical significance of LAP and VAI in the evaluation of MetS among apparently healthy individuals <40 years in Maiduguri, Borno state, Nigeria.

## 2. MATERIALS AND METHODS

The study was conducted on 200 apparently healthy male and female students of university of Maiduguri. Random sampling techniques were used to recruit the subjects aged 18-41 years. Examination of subjects consisted of physical examination with measurement of anthropometric and clinical parameters, filling out a questionnaire, and evaluation of serum lipid levels. Individuals presenting the following conditions were excluded from the study: severe obesity (BMI>40 kg/m2), previous liver failure diagnosis, chronic kidney disease requiring renal replacement therapy, corticosteroid and immunosuppressant use. hypertension and cognitive impairments.

### 2.1 Measurements

Anthropometric measurements were made by well-trained examiners using standardized instruments. Body weight was measured on a digital scale with participants wearing light clothing and no shoes. Height was determined without shoes using a standard stadiometer. Body mass index was calculated as weight (kg) divided by height squared (m2). WC (cm) was measured using non-elastic tape at the midway point between the last rib and the iliac crest after normal expiration. Blood pressure was measured on the right arm using an automated device (DINAMAP-R (Criticon, Tampa, Florida)) after the participant rested for 10 min in a seated position. Blood pressure was measured 3 times with a 3-min interval between each measurement, and the average of the three measurements was recorded for Systolic (SBP) and diastolic blood pressures (DBP).

#### 2.2 Biochemical Analyses

Blood samples were collected from venous vessels after 12-h fasting. Plasma glucose was determined in duplicate by a glucose oxidase method, TG and high-density lipoprotein cholesterol (HDL-C) levels were measured by an enzymatic method using Hitachi chemistry autoanalyzer Cobas C311.

#### 2.3 Assessment of Visceral Adiposity Index

VAI score was calculated using the following sex-specific equations, when Triglycerides (TG) levels expressed in mmol/l and HDL is HDL-Cholesterol levels expressed in mmol/l:

- Males: VAI = (WC /39.68 + (1.88 × BMI)) × (TG /1.03) × (1.31 /HDL),
- Females: VAI = (WC/36.58 + (1.89 × BMI)) × (TG/ 0.81) × (1.52 /HDL) [16].

The diagnosis of MetS was established according to the revised criteria of the NCEP/ATP III (MS-NCEP/ ATP III): any three or more of the following criteria:

- i. WC>102 cm (men) or >88 cm (women),
- ii. Fasting TG≥150 mg/dl,
- iii. SBP≥130 and/or DBP≥85 mmHg,
- iv. Fasting HDL-C<40 mg/dl (men) or <50 mg/dl (women), and
- v. Fasting plasma glucose (FPG)>5.6 mmol/L [7, 19]. WC was measured at the midpoint between the lower rib and the iliac crest.

#### 2.4 Assessment of Lipid Accumulation Product

The total-body lipid accumulation is described as:

- LAP for men=(waist circumference [cm] -65)x (triglyceride concentration [mmol/l]);
- LAP for women=(waist circumference [cm] -58)x(triglyceride concentration [mmol/l]) [13, 15, 19].

### **3. STATISTICAL ANALYSIS**

Data were analyzed using spss vs 20.0 and the results were presented as mean  $\pm$  S.D. Prevalence rates were expressed as percentages. The areas under the curves (AUCs) for ROC curves were determined for each continuous variable to identify the predictors of MSNCEP/ATP III and MS-IDF. AUCs are provided with S.E.M. and 95% confidence intervals (95% CI). ROC curves, a plot of the sensitivity (SEN) (true positive) versus 1-specificity (SP) (false positive) for each potential predictor tested, determine the ability of a screening measure for correctly identifying

individuals based on their classification by a reference test.

#### **4. RESULTS**

In the present study, 200 participants of both sexes who fulfilled the inclusion criteria were enrolled into the study. There were 135 males and 65 female participants with mean age of 25.65±5.56 years for males and 24.11±4.60 years for females. The anthropometric and clinical characteristics of the participants are shown in table 1. The values for the components of MetS includes BP (SBP=105.67±9.82; DBP=70.43±6.16); TG=1.69±0.72; HDL=1.10±0.37; FBG=3.88±0.72; WC=87.76±8.41 for males and BP (SBP=104.69±9.93; DBP=69.93±6.44), TG=1.82±0.69; HDL=1.13±0.39; FBG=3.64±0.79; WC=86.93±9.89 for females (table1). Prevalence of MetS according to the IDF definition was 8.1% in males and 21.5% in females as shown in table2. Performance of the three markers for identifying persons with MetS according to the IDF criteria was assessed as shown in table3. In males, the performance of LAP to detect MetS-IDF was better in comparison with VAI and BMI. The LAP cut-off value in male was 50.00 with sensitivity (0.909), specificity (0.200) and AUC (0.908) exhibiting high performance as compared with VAI (sensitivity (0.906), specificity (0.240) and AUC (0.850)) and BMI (sensitivity (0.545), specificity (0.444) and AUC (0.498)) as shown in table3 and figure 1. In females, the performance of LAP to detect MetS-IDF was better compared with VAI and BMI. LAP cut-off value was 41.50 with sensitivity (0.929), specificity (0.400) and AUC (0.721) as compared with VAI (sensitivity (0.786), specificity (0.373) and AUC (0.865)) and BMI (sensitivity (0.429), specificity (0.490) and AUC (0.436)) as shown in table3 and figure 2. Figure 3 shows the performance of LAP, VAI and BMI of the study subjects.

Variables	Whole subjects	Males	Females	
	Mean ± SD	Mean ± SD	Mean ± SD	
Age	25.15±5.31	$25.65 \pm 5.56$	24.11±4.60	
SBP	$105.4 \pm 9.84$	105.67±9.82	104.69±9.93	
DBP	70.17±6.25	70.43±6.16	69.93±6.44	
WC	$87.49 \pm 8.90$	87.76±8.41	86.93±9.89	
BMI	22.52±3.36	22.59±2.86	22.36±4.23	
VAI	2.29±1.11	2.21±1.11	$2.49 \pm 1.12$	
LAP	43.68±25.63	39.06±23.69	53.26±26.99	
TC	3.62±0.74	3.60±0.71	3.66±0.79	
TG	1.73±0.71	1.69±0.72	$1.82 \pm 0.69$	
HDL	1.11±0.37	1.10±0.37	1.13±0.39	
LDL	$1.78\pm0.44$	$1.80\pm0.42$	$1.76\pm0.46$	
FBG	3.80±0.75	3.88±0.72	3.64±0.79	

 Table 1: Clinical characteristics of the study population

	Metabolic Syndrome				
Sex	Yes (%)	No (%)	Total		
Male	11(8.1)	124(91.9%)	135		
Female	14(21.5%)	51(78.5%)	65		
Total	25(12.5%)	175(87.5%)	200		

 Table 2: Prevalence of Metabolic syndrome among the study subjects (n=200)

## Table 3: Performance of Markers for identifying persons with Metabolic Syndrome according to sex

Makers	Population	Cut-off	Sensitivity	Specificity	AUC(95%CI)	p-values
LAP	Male	50.00	0.909	0.200	0.908(0.857-0.960)	0.000
	Female	41.50	0.929	0.100	0.721(0.600-0.842)	0.012
VAI	Male	2.54	0.906	0.240	0.850(0.724-0.976)	0.000
	Female	2.52	0.786	0.373	0.865(0.765-0.842)	0.000
BMI	Male	22.75	0.545	0.444	0.498(0.307-0.689)	0.981
	Female	21.60	0.429	0.490	0.436(0.276-0.605)	0.468

 Table 4: Performance of Markers for identifying persons with Metabolic Syndrome among the study subjects

Markers	AUC	<b>P-Values</b>
LAP	0.856	0.000
VAI	0.820	0.000
BMI	0.523	0.795







#### **5. DISCUSSION**

This study is the first to identify the most efficient MetS risk assessment systems LAP, VAI or BMI, in healthy individuals in Maiduguri. Although BMI is the most common screening measures used to identify people who are at relatively high risk of MetS, it is not particularly the most effective tool [20] as it does not give a clear idea about central obesity [18] and it is also not gender-specific entity. Consequent to this limitation, we conducted a cross-sectional populationbased survey on metabolic syndrome in Maiduguri, Borno state on healthy individuals in order to identify single parameter/ index as surrogates with high efficiency in predicting metabolic syndrome. Obesity, notably abdominal obesity (Visceral adiposity), is the most important cardiovascular and MetS risks factor. Visceral adipose tissues have been found to be correlated with plasma triglyceride (TG). TG is also a reliable predictor for these cardiometabolic syndromes [21]. WC has also been reported as a robust predictor for cardiometabolic risk and a simple measure of truncal fat. It reflects both abdominal subcutaneous adipose tissue and visceral adipose tissue, and therefore represents the main component of MetS [22]. However, waist circumference is unable to distinguish between visceral adipose tissue and subcutaneous adipose tissue. Therefore, it is important to identify a routinely applicable indicator for evaluation of visceral adiposity. Furthermore, the use of triglyceride levels in with waist circumference, termed combination hypertriglyceridemic waist, has been shown to be able to identify individuals with the greatest amount of visceral fat and is found to be associated with increased risk of MetS [8]. The prevalence of MetS within our study subjects was found to be 12.5% and gender wise it is 21.5% in females and 8.1% in males. This prevalence is low compared to the one reported in Turkey where the prevalence was reported to be 33.9% within the adult population and gender wise was 54.5% in women and 45.1% in men [2, 28]. This difference may be due to inclusion of overweight and obese subjects among the previous study. This difference may also be due to influence of ethnic food and cultural ties which are associated with feeding lifestyle of the people. Our finding was also not in agreement with report of Roomi where overall prevalence was 21.7%, male 21.9% and females 14.1% [29]. Similarly Aisha reported higher prevalence of MetS it overall prevalence of 25.78%, 22.41% in males and 32.26% in females in northwestern Nigeria [30]. The differences observed compared to our population of study could also be due to individual lifestyle such as smoking and alcohol use which are potential risk factors of obesity and MetS.

Receiver operating characteristic (ROC) curves were plotted to assess the performance of MetS predictors by gender and among study subjects. The power of MetS prediction was quantified by the area under the curve (AUC) with 95% confidence intervals, i.e. a larger AUC reflecting better predictive accuracy.

In this study, we found that LAP has the strongest diagnostic accuracy (AUC = 0.856) for MetS among the healthy, unrelated adults than VAI (AUC=0.820) and BMI (AUC=0.523). This report is similar to findings of [8] and also to reports of [23]. Our finding is also in agreement with report of [24]. Our AUC results for the MetS predictors also shows that LAP has the highest diagnostic accuracy for MetS (AUC=908) than VAI and BMI in males than females. Our results were similar to those previously reported by [23] in cross-sectional study of healthy Argentinian men and report of [25] who found that LAP had the highest diagnostic accuracy for metabolic syndrome, with an AUC of 0.91 and 0.90 in males and females. Our result was also similar to those reported by [26] and [27].

### **6. CONCLUSIONS**

This study therefore shows LAP has the highest diagnostic accuracy and particularly the most effective tool in identifying males who are at the risk of MetS. While VAI has the highest diagnostic accuracy and particularly the most effective tool in identifying females who are at the risk of MetS. These simple clinical tools may help, in a primary care setting, to identify subjects at risk of metabolic syndrome and who require further biochemical evaluation.

### ACKNOWLEDGEMENTS

We thank the participants. We also want to acknowledge the contribution of the department of Chemical Pathology Laboratory University of Maiduguri Teaching Hospital for allowing us to use their facilities for the analysis.

Funding: This is a self-sponsored work.

Conflict of Interests: None declared.

**Ethics Approval:** Institutional research board and human ethics committee of University of Maiduguri, Borno state of Nigeria.

## REFERENCES

- Ray, L., Ravichandran, K., & Nanda, S. K. (2018). Comparison of lipid accumulation product index with body mass index and waist circumference as a predictor of metabolic syndrome in Indian population. *Metabolic syndrome and related disorders*, *16*(5), 240-245. DOI: 10.1089/met.2017.0119
- Pekgor, S., Duran, C., Berberoglu, U., & Eryilmaz, M. A. (2019). The role of visceral adiposity index levels in predicting the presence of metabolic syndrome and insulin resistance in overweight and obese patients. *Metabolic syndrome and related disorders*, 17(5), 296-302. DOI: 10.1089/met.2019.0005.
- Tabassum, M., Mozaffor, M., Rahman, M. M., & Huda, R. M. (2022). Lipid Accumulation Product: An Effective Obesity Index to Predict Metabolic Syndrome. *Journal of Bangladesh College of Physicians and Surgeons*, 40(1), 5-9. DOI: https://doi.org/10.3329/jbcps.v40i1.57053
- Bijari, M., Jangjoo, S., Emami, N., Raji, S., Mottaghi, M., Moallem, R., ... & Saberi, A. (2021). The accuracy of visceral adiposity index for the screening of metabolic syndrome: a systematic review and meta-analysis. *International Journal of Endocrinology*, 2021. Article ID 6684627, 14. https://doi.org/10.1155/2021/6684627
- Fatahi, A., Doosti-Irani, A., & Cheraghi, Z. (2020). Prevalence and incidence of metabolic syndrome in Iran: a systematic review and metaanalysis. *International journal of preventive medicine*, 11, 64.
- Taverna, M. J., Martínez-Larrad, M. T., Frechtel, G. D., & Serrano-Ríos, M. (2011). Lipid accumulation product: a powerful marker of

metabolic syndrome in healthy population. *European Journal of Endocrinology*, 164(4), 559-567.

- Grundy, S. M., Brewer, H. B., Cleeman, J. I., Smith, S. C., & Lenfant, C. (2004). Institute/American Heart Association Conference on Scientific Issues Related to Definition of Metabolic Syndrome: Report of the National Heart. *Lung, and Blood. Circulation, 109*, 433-438. (doi:10.1161/01.CIR.0000111245.75752.C6)
- Chiang, J. K., & Koo, M. (2012). Lipid accumulation product: a simple and accurate index for predicting metabolic syndrome in Taiwanese people aged 50 and over. *BMC cardiovascular disorders*, *12*, 1-6. http://www.biomedcentral.com/1471-2261/12/78
- Zhao, S., Ren, Z., Yu, S., Chi, C., Tang, J., Maimaitiaili, R., ... & Zhang, Y. (2021). Association between lipid accumulation product and target organ damage in elderly population: The Northern Shanghai Study. *Clinical Interventions in Aging*, 1769-1776.
- Shah, R. V., Murthy, V. L., Abbasi, S. A., Blankstein, R., Kwong, R. Y., Goldfine, A. B., ... & Allison, M. A. (2014). Visceral adiposity and the risk of metabolic syndrome across body mass index: the MESA Study. *JACC: Cardiovascular Imaging*, 7(12), 1221-1235. doi:10.1016/j.jcmg.2014.07.017
- Neeland, I. J., Turer, A. T., Ayers, C. R., Powell-Wiley, T. M., Vega, G. L., Farzaneh-Far, R., ... & de Lemos, J. A. (2012). Dysfunctional adiposity and the risk of prediabetes and type 2 diabetes in obese adults. *Jama*, 308(11), 1150-1159. doi:10.1001/2012.jama.11132
- Camhi, S. M., Bray, G. A., Bouchard, C., Greenway, F. L., Johnson, W. D., Newton, R. L., ... & Katzmarzyk, P. T. (2011). The relationship of waist circumference and BMI to visceral, subcutaneous, and total body fat: sex and race differences. *Obesity*, 19(2), 402-408. doi:10.1038/oby.2010.248
- 13. Kahn, H. S., & Valdez, R. (2003). Metabolic risks identified by the combination of enlarged waist and elevated triacylglycerol concentration. *The American journal of clinical nutrition*, 78(5), 928-934.
- 14. Kaneva, A. M., & Bojko, E. R. (2021). Ageadjusted cut-off values of lipid accumulation product (LAP) for predicting hypertension. *Scientific Reports*, 11(1), 1-6. https://doi.org/10.1038/s41598-021-90648-y
- 15. Kahn, H. S. (2006). The lipid accumulation product is better than BMI for identifying diabetes: a population-based comparison. *Diabetes care*, 29(1), 151-153.
- Amato, M. C., Giordano, C., Galia, M., Criscimanna, A., Vitabile, S., Midiri, M., ... & AlkaMeSy Study Group. (2010). Visceral Adiposity Index: a reliable indicator of visceral fat

function associated with cardiometabolic risk. *Diabetes care*, *33*(4), 920-922.

- 17. Amato, M. C., Giordano, C., Pitrone, M., & Galluzzo, A. (2011). Cut-off points of the visceral adiposity index (VAI) identifying a visceral adipose dysfunction associated with cardiometabolic risk in a Caucasian Sicilian population. *Lipids in health and disease*, *10*, 1-8. http://www.lipidworld.com/content/10/1/183.p1-8
- Thamilovia, S. A., & Mageshwari, S. U. (2020). Visceral adiposity index - predictive index of cardiovascular diseases. *Int J Health Sci Res.*, 10(12), 252-256.
- Taverna, M. J., Martínez-Larrad, M. T., Frechtel, G. D., & Serrano-Ríos, M. (2011). Lipid accumulation product: a powerful marker of metabolic syndrome in healthy population. *European Journal of Endocrinology*, 164(4), 559-567.
- Vieira, J. N., Braz, M. A. D., Gomes, F. O., Silva, P. R. D., Santos, O. T. D. M., Rocha, I. M. G. D., ... & Fayh, A. P. T. (2019). Cardiovascular risk assessment using the lipid accumulation product index among primary healthcare users: a crosssectional study. *Sao Paulo Medical Journal*, 137, 126-131.
- Gallagher, E. J., LeRoith, D., & Karnieli, E. (2008). The metabolic syndrome from insulin resistance to obesity and diabetes. *Endocrinology and Metabolism Clinics of North America*, 37, 559–579. (doi:10.1016/j.ecl.2008.05.002)
- 22. Klein, S., Allison, D. B., Heymsfield, S. B., Kelley, D. E., Leibel, R. L., Nonas, C., Kahn, R., & Association for Weight Management and Obesity Prevention; NAASO; Obesity Society; American Society for Nutrition; American Diabetes Association. (2007). Waist circumference and cardiometabolic risk: a consensus statement from shaping America's health: Association for Weight Management and Obesity Prevention; NAASO, the Obesity Society; the American Society for Nutrition; and the American Diabetes Association. The American journal of clinical nutrition, 85(5), 1197-1202. (doi:10.2337/dc07-9921)
- Tellechea, M. L., Aranguren, F., Martínez-Larrad, M. T., Serrano-Ríos, M., Taverna, M. J., & Frechtel, G. D. (2009). Ability of lipid accumulation product to identify metabolic syndrome in healthy men from Buenos Aires. *Diabetes Care*, 32(7), e85-e85.
- Nascimento, J. X. P. T., Chein, M. B. D. C., de Sousa, R. M. L., Ferreira, A. D. S., Navarro, P. A., & Brito, L. M. O. (2015). Importance of lipid accumulation product index as a marker of CVD risk in PCOS women. *Lipids in health and disease*, 14, 1-8. DOI 10.1186/s12944-015-0061-y
- Taverna, M. J., Martínez-Larrad, M. T., Frechtel, G. D., & Serrano-Ríos, M. (2011). Lipid accumulation product: a powerful marker of

metabolic syndrome in healthy population. *European Journal of Endocrinology*, 164(4), 559-567.

- 26. Xiang, S., Hua, F., Chen, L., Tang, Y., Jiang, X., & Liu, Z. (2013). Lipid accumulation product is related to metabolic syndrome in women with polycystic ovary syndrome. *Experimental and Clinical Endocrinology & Diabetes*, 121(02), 115-118.
- 27. Chan, L., Xue, H., Xiaoya, Z., Jiajia, X., Wei, R., Linman, L., ... & Lan, L. (2016). Lipid accumulation product: a simple and accurate index for predicting metabolic syndrome in patients with adult growth hormone deficiency. *Experimental and clinical endocrinology & diabetes*, *124*(04), 220-224.
- Onat, A., Yüksel, M., Köroğlu, B., Gümrükçüoğlu, H. A., Aydın, M., Çakmak, H. A., ... & Can, G. (2013). Turkish Adult Risk Factor Study survey 2012: overall and coronary mortality and trends in the prevalence of metabolic syndrome. *Turk Kardiyoloji Dernegi Arsivi: Turk Kardiyoloji Derneginin Yayin Organidir*, 41(5), 373-378.
- 29. Roomi, M. A., & Mohammadnezhad, M. (2019). Prevalence of Metabolic Syndrome Among Apparently Healthy Workforce. *J Ayub Med Coll Abbottabad*, 31(2), 252-254. PMID: 31094127.
- Nalado, A. M., Musa, B. M., Gezawa, I. D., Muhammad, H., Ibrahim, D. A., & Uloko, A. E. (2015). Prevalence of Metabolic Syndrome among Apparently Healthy Adults in a Rural Community, in North-western Nigeria. *Niger J Med*, 24(4), 323-30. PMID: 27487609.

**Cite This Article:** Alhaji Haruna Musa, Ijagila, I. N, Dungus, M. M (2023). Evaluation of Metabolic Syndrome Using Lipid Accumulation Products, Visceral Adiposity Index and Body Mass Index in Apparently Healthy Students of University of Maiduguri. *East African Scholars J Med Sci*, 6(3), 100-107.