

Evaluation of dyslipidemia prevalence among undergraduate university students.

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Abstract:

Dyslipidemia has grown to be a significant global public health issue particularly in developing nations like ours, as a result of rising trends in dietary habits, physical inactivity, and obesity, which each play essential roles in the development of cardiovascular diseases. The purpose of this cross-sectional study was to determine the prevalence of dyslipidemia among undergraduate students at Nnamdi Azikiwe University in Nnewi. Two hundred (200) students between the ages of 18 and 30 who appeared to be in good health were selected at random. After that, each participant provided a fasting blood sample of five milliliters (5ml) for the evaluation of serum total cholesterol (TC), triglycerides (TG), low density lipoprotein cholesterol (LDL-C), and high-density lipoprotein cholesterol (HDL-C) using standard laboratory techniques. Additionally, the body mass index (BMI) and the systolic and diastolic blood pressure readings were calculated. 44 percent of participants were males and 56 percent were females, with an average age of 22.41 ± 1.68 years. While 3% of the participants were underweight, 27% of them were overweight, and 6% were pre-obese, the majority of them (64%) had a normal BMI. Three percent (3%) of the participants had pre-hypertension, whereas 97% of the participants had normal blood pressure. The prevalence of hypercholesterolemia, hypertriglyceridemia, low HDL-C, and high LDL-C among the participants, as defined by the World Health Organization, was 39.5%, 26.5%, 4%, 12.5%, and 16.5%, respectively. SBP Vs DBP, DBP Vs Weight, DBP Vs BMI, Weight Vs BMI, TC Vs TG, TC Vs HDL, and TC Vs LDL all had significant positive correlations ($p < 0.05$) while SBP Vs TG, SBP Vs HDL, and TC Vs LDL all had significant negative correlations ($r = -0.148$, $p = 0.036$, and $r = -0.203$, $p = 0.004$). The female sex was strongly related with dyslipidemia (48.2%), whereas the male sex was not (28.4%). As a result, there is a pressing need to educate the public about the essential changes in food habits and overall lifestyle to maintain cardiovascular health.

Key Words: dyslipidemia; BMI; blood pressure; age; lipid profile

Introduction

The abnormal amount of lipids (e.g. triglycerides, cholesterol, and/or fats phospholipids) in the blood is known as dyslipidemia. Elevated plasma levels of total cholesterol (TC), low-density lipoprotein cholesterol (LDL-C), and triglycerides (TG), as well as reduced plasma levels of HDL cholesterol (HDL-C), are all characteristics of dyslipidemia (Manjunath *et al.*, 2013; Bibbins-Domingo *et al.*, 2016). Type 2 diabetes mellitus, hypothyroidism, excessive alcohol consumption, cholestatic liver disease,

renal disease, smoking, and obesity are all conditions that can result in dyslipidemia (Okaka and Eiya, 2013). Despite being a modifiable risk factor for cardiovascular disease (CVD), dyslipidemia is becoming more common. Cardiovascular disease is the leading cause of morbidity and mortality worldwide (GBD 2017 Causes of Death Collaborators, 2018), with a high prevalence in most countries but a higher burden on the populations of developing countries like Nigeria as a result of dietary changes, physical inactivity, and increased alcohol consumption, among other factors (Munsinguz *et al.*, 2020). Prevalence varies by region in

Sub-Saharan Africa, with rates of more than 50% in Nigeria, Ghana, and Senegal, and 39.30 percent in Nigeria (Fatou *et al.*, 2016). Dyslipidemia is growing more frequent, although most people are unaware of it because they are not identified and thus not treated. This increases the risk to an already vulnerable group, exacerbating the morbidity, death, and medical expenses associated with cardiovascular disease.

The burden of cardiovascular disease is increasing in emerging nations like Nigeria due to the increasing prevalence of cardiovascular risk factors (Nnamudi *et al.*, 2020). The heart and blood arteries make up the cardiovascular system (Farley *et al.*, 2012). On the other hand, cardiovascular diseases are a group of conditions that affect the heart and blood arteries. Atherosclerosis is the main underlying pathology of cardiovascular diseases, which continue to be a major cause of morbidity and premature death in the world. CVDs accounted for 11% of all fatalities in Nigeria in 2018 (WHO, 2018). Men appear to have a higher mortality rate than women of the same age (Ezeugwunne *et al.*, 2017), with coronary artery disease and stroke accounting for 80% of CVD fatalities in men and 75% of CVD fatalities in women (WHO, 2011). Atherosclerosis is linked to dyslipidemia, which raises the risk of cardiovascular disease (CVD). Numerous risk factors for CVDs can be categorized as modifiable or non-modifiable. Risk factors that can be changed include unhealthy eating habits, inactivity, drinking alcohol improperly, and smoking, which can lead to dyslipidemia, hypertension, high blood sugar, and other conditions, as well as overweight and obesity (WHO, 2018). These factors can be avoided with a balanced diet, regular exercise, and a well-balanced lifestyle. Several research (Oguoma *et al.*, 2015; Ezeugwunne *et al.*, 2017; González-Rivas *et al.*, 2018; Vizentin *et al.*, 2019; Nnamudi *et al.*, 2020) have found varied levels of abnormalities in lipid levels (dyslipidemia) in apparently healthy persons. In a study of a Nigerian adult population, Oguoma *et al.* discovered a prevalence of low HDL (17.8%), hypertriglyceridemia (23.2%), hypercholesterolemia (38.1%), and central obesity (52.2%) (Oguoma *et al.*, 2015). Also, a study examining the frequency and pattern of dyslipidemia in Nigeria discovered that dyslipidemia was common in all of Nigeria's geographic zones. Overall 60 percent of the apparently healthy Nigerians had dyslipidemia, while 89 percent of diabetic Nigerians had dyslipidemia (Oguejiofor *et al.*, 2012). Furthermore, Olamoyegun and colleagues discovered elevated TC, high LDL-C, elevated TG, and low HDL-C in 5.3 percent, 19.3 percent, 4.4 percent, and 76.3 percent of their semi-urban inhabitants in Nigeria (Olamoyegun *et al.*, 2016). Furthermore, varying prevalence of prediabetes and prehypertension has also been recorded in the current area among the study population (Obiora *et al.*, 2017; Oponi *et al.*, 2017). In light of the foregoing, it is critical to address the issue of dyslipidemia and, by extension, cardiovascular disease by involving the population at risk in a well-designed study that will raise awareness of the risk factors for dyslipidemia and its associated diseases and provide guidance to health policymakers, laboratory scientists, and clinicians.

Materials And Methods

Study design

This cross-sectional study determined the prevalence of dyslipidemia among Undergraduate University students at Nnamdi Azikiwe University, Nnewi Campus.

Sample size

Using the procedure outlined by Charan and Biswas, the sample size was estimated (2013);

$$N = \frac{(Z^2 pq)}{d^2}$$

Where:

N= Desired number of sample when population of the facility is limited

Z = The standard normal variance where confidence level is 1.96 at 95%

p = prevalence rate of dyslipidemia in Nigeria is 85.9% (Fatou *et al.*, 2016)

q = 1 – p

d = 5% i.e degree of precision as desired by the researcher.

Applying the method,

$$N = \frac{Z^2 \times P \times (1-P)}{D^2}$$

$$N = \frac{1.96^2 \times 0.859 \times (1-0.859)}{0.05^2}$$

$$N = 186 \pm 10\%$$

Participants' Recruitment

200 college students, both male and female, between the ages of 18 and 30, who appeared to be in good health, were enrolled in the study.

Inclusion criteria

The study included undergraduates between the ages of 18 and 30 from the Faculty of Basic Medical Sciences, Faculty of Health Sciences and Technology, and Faculty of Medicine at Nnamdi Azikiwe University, Nnewi campus.

Exclusion criteria

This study excluded smokers, alcoholics, and individuals with pre-existing diseases (diabetes, hypertension, and heart disease).

Ethical consideration

The faculty of Health Science and Technology's ethics committee approved the study before it began (NAU/FHST/2021/MLS77). Also, prior to the start of the study, the participants were asked to give their written informed consent.

Collection Of Samples

Five milliliters (5ml) of venous fasting blood was aseptically taken from each person after a 10- to 12-hour fast through the antecubital vein using a plastic syringe that had little to no stasis and placed in a plain container. Before being centrifuged for five minutes at a speed of 1000 rpm, the blood sample was given time to clot and retracted. The sera were then separated into plain tubes for lipid profile estimation and the samples that wouldn't be evaluated immediately were frozen at -8°C.

Laboratory Methods

According to Allain *et al.* (1974), Fossati *et al.* (1982), and Burstein *et al.* (1980), TC, TG, and HDL-C levels were measured using enzymatic techniques, while LDL-C was estimated using the formula provided by Assmann *et al.* (1984). Fasting total cholesterol >5.2 mmol/l, fasting TG >1.7 mmol/l, LDL \geq 3.5 mmol/L, and HDL <0.9 mmol/L were all considered to be dyslipidemia. **Anthropometrics indices and Blood pressure reading**

Body mass index (BMI) was computed using the following formula: BMI= weight (Kg) / height (m²). Weight was measured using an automatic weighing scale, and height was measured using a measuring tape fastened to a piece of wood. Body mass index (BMI) values \geq 25 and above 30 kg/m² were used to characterize overweight and generalized obesity, respectively.

After a 10-minute rest, the subject's systemic blood pressure was assessed on the left arm while they were seated using an OMRON automatic digital blood pressure monitor and an appropriate-sized cuff. Blood pressure measurements were done, including systolic and diastolic rates.

Hypertension was defined as systolic and/or diastolic blood pressures of ≥ 140 mmHg and ≥ 90 mmHg respectively.

Statistical Analysis

The Statistical Package for the Social Sciences (SPSS) version 23.0 was used to analyze the results. The paired student t-test and Pearson correlation were used to statistically analyze the data, which was presented as mean \pm SD, with level of significance assumed at $p < 0.05$.

Results

The anthropometric indices of the study subjects are shown in TABLE 1.

VARIABLES	LEVELS	n=200	PERCENTAGE (100%)
SEX	Male	88	44%
	Female	112	56%
AGE	<25	178	89%
	>25	22	11%
BMI	Normal	128	64%
	Underweight	6	3%
	Overweight	54	27%
	Pre-obesity	12	6%
BLOOD PRESSURE	Normal	194	97%
	Pre-hypertension	6	3%

Table 1: Anthropometric Indices of the Students Studied

The study involved 200 students (88 men and 112 females) between the ages of 18 and 30. Female students accounted for 56 percent of the total, while male students accounted for 44 percent. 89 percent of the students who took part were under the age of 25, while 11 percent were beyond the age of 25. Students with a normal BMI, underweight, overweight, pre-obese, normal blood pressure, and pre-hypertensive blood pressure account for 64 percent, 3 percent, 27 percent, 6 percent, 97 percent, and 3 percent of the population, respectively.

Male systolic blood pressure (119.05 ± 9.73), female systolic blood pressure (116.82 ± 9.09), male diastolic blood pressure (70.59 ± 8.03), female diastolic blood pressure (69.33 ± 9.99), male body mass index (22.86 ± 3.96) and female body mass index (23.48 ± 3.84) had no sex differences ($P > 0.05$). Male height (1.77 ± 0.08) and male weight (72.64 ± 11.07) had significantly higher mean \pm SDs than female height (1.68 ± 0.08) and female weight (66.40 ± 10.32), respectively ($p < 0.05$). See table 2.

Parameters	Total Participants (n=200)	Males n=88	Females n=112	t-value	P-value
SBP (mmHg)	117.80 + 9.42	119.05 + 9.73	116.82 + 9.09	1.665	0.097
DBP (mmHg)	69.89 + 9.18	70.59 + 8.03	69.33 + 9.99	0.964	0.336
HEIGHT (m)	1.72 + 0.090	1.77 + 0.08	1.68 + 0.08	7.529	0.000*
WEIGHT (Kg)	69.15 + 11.07	72.64 + 11.07	66.40 + 10.32	4.106	0.000*
BMI (Kg/m ²)	23.21 + 3.89	22.86 + 3.96	23.48 + 3.84	-1.102	0.272

*Statistically significant at $p < 0.05$.

Table 2: Blood Pressure and Anthropometric Parameters in participants studied Grouped by Sex (Mean \pm SD)

There were no sex differences in the mean and standard deviation of male high density lipoprotein cholesterol (1.21 ± 0.27) compared to female high density lipoprotein cholesterol (1.22 ± 0.23) ($P > 0.05$). Male total cholesterol (4.56 ± 0.66) and male low density lipoprotein cholesterol (2.75 ± 0.62) mean \pm SD were significantly lower than female total

cholesterol (4.86 ± 0.84) and female low density lipoprotein cholesterol (3.08 ± 0.83), respectively ($p < 0.05$). Furthermore, the male triglycerides (1.34 ± 0.34) had a significantly higher mean \pm SD (1.24 ± 0.20) than the female triglycerides ($p < 0.05$). See table 3.

Parameters	Total participants (n=200)	Males N=88	Females N=112	t-value	P-value
TC (mmol/l)	4.73 + 0.78	4.56 + 0.66	4.86 + 0.84	-2.759	0.006*
TG (mmol/l)	1.29 + 0.27	1.34 + 0.34	1.24 + 0.20	2.523	0.012*
HDLC (mmol/l)	1.21 + 0.25	1.21 + 0.27	1.22 + 0.23	-0.359	0.720
LDLC (mmol/l)	2.93 + 0.76	2.75 + 0.62	3.08 + 0.83	-3.146	0.002*

*Statistically significant at $p < 0.05$.

Table 3: Lipid profile levels in participants studied based on Sex (mean \pm SD)

Parameters	r-value	p-value
SBP Vs DBP	0.323	0.000*
SBP Vs TG	-0.148	0.036*

SBP Vs HDL	-0.203	0.004*
DBP Vs Weight	0.206	0.003*
DBP Vs BMI	0.156	0.028*
Height Vs Weight	0.416	0.000*
Height Vs BMI	-0.235	0.001*
Weight Vs BMI	0.645	0.000*
TC Vs TG	0.152	0.032*
TC Vs HDL	0.207	0.003*
TC Vs LDL	0.934	0.000*

*Statistically significant at p<0.05.

Table 4: Association between Parameters of the Studied Participants

Significant positive correlations were found between SBP and DBP (r= 0.323; p= 0.000), DBP and weight (r= 0.206; p=0.003), DBP and BMI (r= 0.156; p=0.028), height and weight (r=0.416; p=0.000), weight and BMI (r= 0.645; p=0.000), TC and TG (r=0.152; p=0.032), TC and HDL (r=0.207; p=0.003), TC and LDL (r=0.934, p=0.000) while SBP Vs TG (r=-0.148; p=0.036), SBP Vs HDL (r=-0.203; p=0.004), and Height Vs BMI (r=-0.235; p=0.001) all had statistically significant negative relationships.

Parameters	r-value	p-value
SBP Vs DBP	0.277	0.009*
SBP Vs BMI	0.212	0.048*
SBP Vs TG	-0.276	0.009*
Height Vs Weight	0.464	0.000*
Weight Vs BMI	0.537	0.000*
TC Vs TG	0.280	0.008*
TC Vs LDL	0.881	0.000*

*Statistically significant at p<0.05.

Table 5: Association between Parameters Studied in the Male Participants

The study parameters showed significant positive correlations between SBP and DBP (r=0.277; p=0.009), SBP and BMI (r=0.212; p=0.048), Height and Weight (r=0.464; p=0.000), Weight and BMI (r=0.537; p=0.000), TC and TG (r=0.280; p=0.008), TC and HDL (r=0.266; p=0.012), TC and LDL (r=0.881; p=0.000), while SBP and TG had a strong negative correlation (r=-0.276; p=0.009).

Parameters	r-value	p-value
SBP Vs DBP	0.351	0.000*
SBP Vs HDL	-0.292	0.002*
DBP Vs Weight	0.234	0.013*
DBP Vs BMI	0.264	0.005*
Height Vs Weight	0.219	0.020*
Height Vs BMI	-0.347	0.000*
Height Vs TG	-0.227	0.016*
Weight Vs BMI	0.836	0.000*
TC Vs LDL	0.955	0.000*

*Statistically significant at p<0.05.

Table 6: Association between Parameters Studied in the Female Participants.

Significant positive associations were found in the study parameters between SBP and DBP (0.351; p=0.000), DBP and BMI (0.264; p=0.005), Height and Weight (0.219; p=0.020), Weight and BMI (0.836; p=0.000), TC and LDL (0.955; p=0.000), and DBP and Weight (0.234; 0.013). Also, significant negative correlations were seen between SBP and HDL (r= -0.292; p=0.002), height and BMI (r= -0.347; p=0.000), and height and TG (r= -0.227; p=0.016), but not with the other parameters (p>0.05).

PREVALANCE OF	N=200	PERCENTAGE (100%)
Hypercholesterolemia	53	26.5%
Hypertriglyceridemia	8	4%
Low HDL-C	25	12.5%
High LDL-C	33	16.5%
Dyslipidemia (at least one factor affected)	79	39.5%

Table 7: Prevalence of Dyslipidemia Among Studied Group

Hypercholesterolemia, hypertriglyceridemia, low HDL-C, and high LDL-C were found to be prevalent in 26.5 percent, 4 percent, 12.5 percent, and 16.5 percent of the population, respectively.

Dyslipidemia is a global epidemic that is a major cause of cardiovascular disease.

Dyslipidemia was quite prevalent among the individuals in this study (39.5 percent). This increase in dyslipidemia prevalence could be due to

Discussions:

changes in eating habits, decreased physical activity, and work intensity. The current result is higher than previous findings in Eastern Ethiopia (34.8%; Sufa *et al.*, 2019), Africa (25.5%; Noubiap *et al.*, 2018), China (32.2%; Liu *et al.*, 2018); and Iran (30.0%; Najafipour *et al.*, 2016). However, the prevalence recorded in the present study is lower compared to previous studies reported in Lithuania (89.7%; Rinkūnienė *et al.*, 2015), South Africa (85.0%) (Dave *et al.*, 2016), India (78.4%) (Banerjee *et al.*, 2014), Poland (77.2%) (Pająk *et al.*, 2016), India (50.7%) (Wankhade *et al.*, 2018), Uganda (63.3%) (Bakesiima *et al.*, 2018), Palestine (66.4%) (Ali *et al.*, 2019) and South Africa (67.3%) (Reiger *et al.*, 2017). This disparity could be attributed to differences in cutoffs applied in the different studies.

Elevated LDL-C (16.5%) and high total cholesterol (26.5%) were the two most common components of dyslipidemia, which is consistent with earlier findings in Nigeria reported by Okaka and Eiya (Okaka and Eiya, 2013) and Oguoma *et al.* (Oguoma *et al.*, 2015). (Okaka and Eiya, 2013). The increasing consumption of simple carbohydrates and saturated fat in modern diets may be the cause of this phenomenon.

The prevalence of high total cholesterol (26.5%) in this study is lower than that in the prior study published in Ethiopia (33.7%) and Iran (29.6%) (Sufa *et al.*, 2019), but is higher than the study results in Ethiopia (5.2%) (Gebreyes *et al.*, 2018), Northern Ghana (4.02%) (Agongo *et al.*, 2018), South Western Uganda (6%) (Asiki *et al.*, 2015), South East Nigeria (8%; 11.4%) (Okwara *et al.*, 2021; Anyabolu, 2017) and South East coastal region in China (8.4%) (Lin *et al.*, 2019).

The prevalence of high LDL-C (16.5%) in this study is higher than the previous finding reported in Ethiopia (14.1%) (Gebreyes *et al.*, 2018), Northern Ghana (5.55%) (Agongo *et al.*, 2018) and in South East coastal region in China (13.9%) (Lin *et al.*, 2019). But lower than the findings reported in Thailand (56.5%) (Narindrarangkura *et al.*, 2019), Uganda (60.9%) (Lumu *et al.*, 2017), Ghana (61.0%) (Micah and Nkum, 2012), Senegal (66.3%) (Doupa *et al.*, 2014), Jordan (75.9%) (Abujbara *et al.*, 2018), and India (47.8%) (Banerjee *et al.*, 2014)

The current study's prevalence of low HDL-C (12.5%) is somewhat comparable to earlier studies conducted in various African nations, notably Malawi (15.9%) (Amberbir *et al.*, 2018), Ghana (17.0%) (Micah and Nkum, 2012), and Africa (18.5%) (Micah and Nkum, 2012), unlike many prior studies in the South East coastal region of China (23.1%) (Lin *et al.*, 2019), South East Nigeria (34.4%) (Anyabolu, 2017), Northern Ghana (60.30%) (Agongo *et al.*, 2018), Ethiopia (68.7%) (Gebreyes *et al.*, 2018), and South Western Uganda (71.3%) (Asiki *et al.*, 2015).

The prevalence of high triglycerides (4%) in this study is higher than the results from Northern Ghana reported before (2.12%) (Agongo *et al.*, 2018), but lower than those reported in Thailand (49.9%) (Narindrarangkura *et al.*, 2019), South Africa (59.3%) (Reiger *et al.*, 2017), Brazil (65.3%) (Feitosa *et al.*, 2017), Senegal (7.1%) (Doupa *et al.*, 2014), Ethiopia (21.0%) (Gebreyes *et al.*, 2018), Malawi (28.7%) (Amberbir *et al.*, 2018), Venezuela (39.7%) (González-Rivas *et al.*, 2018), Jordan (41.9%) (Abujbara *et al.*, 2018), and Uganda (42.1%) (Lumu *et al.*, 2017).

Males (28.4%) had a lower prevalence of dyslipidemia in this study than females (48.2%), which was 39.5 percent. There were no sex differences in the mean and standard deviation of male high density lipoprotein cholesterol (1.21±0.27) compared to female high density lipoprotein cholesterol (1.22±0.23). Male total cholesterol (4.56 0.66) and male low density lipoprotein cholesterol (2.75±0.62) had mean and standard deviations that were significantly lower than those of female total cholesterol (4.86±0.84) and female low density lipoprotein cholesterol (3.08±0.83), respectively (P <0.05). Additionally, the mean and standard deviation of the male triglycerides (1.34±0.34 Vs 1.24 ±0.20) were significantly higher (P <0.05).

Significant positive correlations were found between SBP and DBP, DBP and weight, DBP and BMI, height and weight, weight and BMI, TC and TG, TC and HDL, TC and LDL while SBP and TG, SBP and HDL as well as Height and BMI all had statistically significant negative relationships.

Conclusion

Although the prevalence of dyslipidemia among the undergraduate students studied was not as high as that reported in Lithuania, South Africa, India, Poland, Uganda, Palestine, and South Africa, it was higher than that reported in Ethiopia, Africa, China, and Iran. The most common kind of dyslipidemia was high total cholesterol, followed by raised LDL-C. Lifestyle and diet appear to be driving a rise in prevalence, particularly among women. Among other interventions, population-wide awareness, specialized education on increased cholesterols and related dangers should be encouraged.

References

1. Manjunath, C.N., Rawal, J.R., Irani, P.M., Madhu K. (2013). Atherogenic dyslipidemia. *Indian Journal of Endocrinology Metabolism*; 17(6): 969-976.
2. Bibbins-Domingo, K., Grossman, D.C., Curry, S.J., Davidson, K.W., Epling, J.W., Garcia, F.A., et al. (2016). US Preventive Services Task Force. Screening for lipid disorders in children and adolescents: *US Preventive Services Task Force recommendation statement. J Am Med Assoc*; 316(6): 625-633.
3. Okaka, E.I., Eiya, B.O. (2013). Prevalence and pattern of dyslipidemia in a rural community in Southern Nigeria. *African Journal of Medicine and Health Science*; 12:80-84.
4. GBD (2017) Causes of Death Collaborators Global, regional, and national age-sex-specific mortality for 282 causes of death in 195 countries and territories, 1980-2017: a systematic analysis for the Global Burden of Disease Study 2017. *Lancet*; 392 1736-1788.
5. Munsinguz, G., Ndejjo, R., Ssinabulya, I., Bastiaens, H., Van Marwijk, H., Wanyenze, R.K. (2020). Cardiovascular risk factors mapping and distribution among adults in Munkono and Buikwe districts in Uganda: small area analysis. *BMC Cardiovascular disorders*; 20(1):1
6. Fatou, C., Fatou, D.A., Alassane, D., Abdou, S.M., Arame, N., Abdourahmane, S., Souleymane, T., Dominique, D., Gaston, N.S., Niama, D.S., Méissa, T. (2016). Prévalence des dyslipidémies au laboratoire de biochimie du CHU Aristide le Dantec de Dakar, Sénégal. *The Pan African Medical Journal*; 25: 67.
7. Nnamudi, A.C., Orhue, J.E.N., Ijeh, I.I. (2020). Assessment of the Levels of Cardiovascular Risk Markers in Hyperglycemic Young Nigerian Adults. *European Journal of Biology and Biotechnology*; 1(3): 1-7.
8. Farley, A., Mchafferty, E., Hendry, C. (2012). The Cardiovascular system. *Nursing Standard*; 27(9): 35-39.
9. WHO (2018). Non-communicable diseases (NCD) country profile.
10. World Health Organization in collaboration with the World Heart Federation and the World Stroke Organization) 2011; pages. 3–18.
11. Ezeugwunne, I.P., Ezeora, J.C., Ogbodo, E.C., Analike, R.A., Okwara, J.E., Oguaka, V.N., Amah, U.K., Meludu, S.C. (2017). Assessment of Cardiovascular Disease Risk Factors of Adult Male and Female Diabetic Patients Attending Nnamdi Azikiwe University Teaching Hospital, Nnewi, Anambra State, Nigeria. *International Journal of Novel Research in Healthcare and Nursing*; 4(2): 68-72.
12. Ezeugwunne, I.P., Ogbodo, E.C., Nwankwo, E.C., Analike, R.A., Onah, C.E., Okwara, J.E., Amah, U.K., Oguaka, V.N.,

- Asebioyo, S.J., Meludu, S.C. (2017). Evaluation of Cardiovascular Status of Apparently Healthy Sedentary Subjects in Nnewi, Anambra State Using Framingham Risk Score Calculator. *International Journal of Innovative Studies in Sciences and Engineering Technology*; **3**(8): 1-6.
13. Oguoma, V.M., Nwose, E.U., Skinner, T.C., Digban, K.A., Onyia, I.C., Richards, R.S. (2015). Prevalence of cardiovascular disease risk factors among a Nigerian adult population: relationship with income level and accessibility to CVD risks screening. *BioMed Central Public Health*; **15**:397.
 14. González-Rivas, J.P., Nieto-Martínez, R., Brajkovich, I., Ugel, E., Rísquez6, A. (2018). Prevalence of Dyslipidemias in Three Regions in Venezuela: The VEMSOLS Study Results. *Arquivos Brasileiros de Cardiologia*; **110**(1):30-35.
 15. Vizontin, P.N., Cardoso, S.M.P., Maia, G.A.C., Alves, P.I., Aranha, L.G., Giannini, T.D. (2019). *Arquivos Brasileiros de Cardiologia*; **112**(2):147-151.
 16. Ogueji, O.C., Onwukwe, C.H., Odenigbo, C.U. (2012). Dyslipidemia in Nigeria: Prevalence and pattern. *Annals of African Medicine*; **11**:197-202.
 17. Olamoyegun, M. A., Oluyombo, R., Asaolu, S. O. (2016). Evaluation of dyslipidemia, lipid ratios, and atherogenic index as cardiovascular risk factors among semi-urban dwellers in Nigeria. *Annals of African Medicine*; **15**(4):194–199.
 18. Obiorah, M.O., Ogbodo, E.C., Amah, U.K., Ezeugwunne, I.P., Analike, R.A., Onah, C.E., Okwara, J.E., Egbe, J.U., Oha, P.C., Ajulu, C.A., Ugwu, M.C., Meludu, S.C. (2017). Prevalence of Prehypertension and Assessment of Cardiovascular Function among Prehypertensive Undergraduate Students in Nnewi, Anambra State, Nigeria. *International Journal of Community Research*; **6**(2): 16 – 25.
 19. Oponi, B.B., Ogbodo, E.C., Amah, U.K., Ezeugwunne, I.P., Ihim, A.C., Onah, C.E., Analike, R.A., Okwara, J.E., Madukwe, D.U.P., Obiorah, M.O., Ugwu, M.C., Meludu, S.C. (2017). Prevalence Of Prediabetes And Assessment Of Associated Risks Of Cardiovascular Disease In Nnewi, Anambra State, Nigeria. *International Journal of Community Research*; **6**(2): 36 – 43.
 20. Charan, J., Biswas, T. (2013). How to calculate sample size for different study designs in medical research?. *Indian Journal of Psychological Medicine*; **35**(2):121–126.
 21. Allain, C.C., Poon, L.S., Chan, C.S.G., Richmond, W., Fu, W. (1974). Enzymatic determination of total serum cholesterol. *Clinical chemistry*; **20**: 470-475.
 22. Fossati, P., Prencipe, L. (1982). Serum triglycerides determined colorimetrically with an enzyme that produces hydrogen peroxide. *Clinical chemistry*; **28**: 2077-2080.
 23. Burstein, M., Scholnick, H.R., Morfin, R. (1980). Rapid method for the isolation of lipoproteins from human serum by precipitation with polyanions. *Scandinavian Journal Clinical Laboratory Investigation*; **40**: 583-595.
 24. Assman, G., Jabs H.U., Kohnert, U., Nolte, W., Schriewer, h. (1984). LDL-Cholesterol determination in blood serum following precipitation of LDL with polyvinylsulfate. *Clinical Chemistry Acta*; **140**: 77-83.
 25. Kesteloot, H., Oviasu, V. O., Obasohan, A. O., Olomu, A., Cobbaert, C., Lissens, W. (1989). Serum lipid and apoprotein levels in a Nigerian population sample. *Atherosclerosis*; **78**: 33-38.
 26. Sufa, B., Abebe, G., Cheneke, W. (2019). Dyslipidemia and associated factors among women using hormonal contraceptives in Harar town, Eastern Ethiopia. *BMC Research Notes*. **12**:120
 27. Liu, X., Yu, S., Mao, Z., Li, Y., Zhang, H., Yang, K., (2018). Dyslipidemia prevalence, awareness, treatment, control, and risk factors in Chinese rural population: the Henan rural cohort study. *Lipids in Health and Disease*; **17**: 119
 28. Najafipour, H., Shokoohi, M., Yousefzadeh, G., Azimzadeh, B.S., Kashanian, G.M., Bagheri, M.M. (2016). Prevalence of dyslipidemia and its association with other coronary artery disease risk factors among urban population in Southeast of Iran: results of the Kerman coronary artery disease risk factors study (KERCADRS). *Journal of Diabetes & Metabolic Disorders*; **15**: 49
 29. Rinkūnienė, E., Laucevičius, A., Petrulionienė, Z., Dženkevičiūtė, V., Kutkienė, S., Skujaitė, A., (2015). The prevalence of dyslipidemia and its relation to other risk factors: a nationwide survey of Lithuania. *Clinical Lipidology*; **10**(3): 219–225.
 30. Dave, J.A., Levitt, N.S., Ross, I.L., Lacerda, M., Maartens, G., Blom, D. (2016). Anti-Retroviral Therapy Increases the Prevalence of Dyslipidemia in South African HIV-Infected Patients. *PLoS ONE*; **11**(3): e0151911.
 31. Banerjee, R., Bhattacharjee, S., Ray, K., Roy, J.K., Datta, S., Banerjee, I. (2014). Dyslipidemia and its relationship with cardiovascular risk factors in a selected population of Siliguri city, west bengal, India. *American Journal of the Medical Sciences*; **5** (1): 1–8.
 32. Pająk, A., Szafraniec, K., Polak, M., Polakowska, M., Kozela, M., Piotrowski, W., (2016). Changes in the prevalence, treatment, and control of hypercholesterolemia and other dyslipidemias over 10 years in Poland: the WOBASZ study. *Polskie Archiwum Medycyny Wewnętrznej*; **126** (9) :642-652.
 33. Wankhade, P.S., Pedhambkar, R.B., Pagare, R.S., Pedhambkar, B.S. (2018). Prevalence and risk factors of dyslipidemia among male industrial workers in India. *International Journal of Community Medicine Public Health*; **5**(4):1458–1465.
 34. Bakesiima, R., Byakika-Kibwika, P., Tumwine, J.K., Kalyango, N., Nabaasa, G., Najjingo, I (2018). Dyslipidaemias in women using hormonal contraceptives: a cross sectional study in Mulago Hospital Family Planning Clinic, Kampala, Uganda. *British Medical Journal Open*; **8**: e022338
 35. Ali, I., Kharm, A., Samara, M., Odeh, S., Jaradat, N., Zaid, A.N., Ahmad, M.A.S. (2019). Prevalence of Dyslipidemia in Undiagnosed Palestinian Men: A Cross-Sectional Study. *Journal of Lipid*; **2019**: 3473042.
 36. Reiger, S., Jardim, T.V., Abrahams-Gessel, S., Crowther, N.J., Wade, A., Gomez-Olive, F.X. (2017). Awareness, treatment, and control of dyslipidemia in rural South Africa: The HAALSI (Health and Aging in Africa: A Longitudinal Study of an INDEPTH Community in South Africa) study. *PLoS ONE*; **12**(10): e0187347
 37. Gebreyes, Y. F., Goshu, D. Y., Geletew, T. K., Argefa, T. G., Zemedu, T. G., Lemu, K. A., Waka, F. C., Mengesha, A. B., Degefu, F. S., Deghebo, A. D., Wubie, H. T., Negeri, M. G., Tesema, T. T., Tessema, Y. G., Regassa, M. G., Eba, G. G., Beyene, M. G., Yesu, K. M., Zeleke, G. T., Mengistu, Y. T., Belayneh, A. B. (2018). Prevalence of high blood pressure, hyperglycemia, dyslipidemia, metabolic syndrome and their determinants in Ethiopia: Evidences from the National NCDs STEPS Survey, *PLoS one*; **13**(5): e0194819.
 38. Agongo, G., Nonterah, E. A., Debpuur, C., Amenga-Etego, L., Ali, S., Oduro, A., Crowther, N. J., Ramsay, M., as members of AWI-Gen and the H3Africa Consortium (2018). The burden of dyslipidaemia and factors associated with lipid levels among

- adults in rural northern Ghana: An AWI-Gen sub-study. *PloSone*; **13**(11), e0206326.
39. Asiki, G., Murphy, G. A., Baisley, K., Nsubuga, R. N., Karabarinde, A., Newton, R., Seeley, J., Young, E. H., Kamali, A., Sandhu, M. S. (2015). Prevalence of dyslipidaemia and associated risk factors in a rural population in South-Western Uganda: a community based survey. *PloS one*; **10**(5): e0126166.
 40. Anyabolu, E. N. (2017). Dyslipidemia in people living with HIV-AIDS in a tertiary hospital in South-East Nigeria. *The Pan African medical journal*; **28**:204.
 41. Lin, H. Q., Wu, J. Y., Chen, M. L., Chen, F. Q., Liao, Y. J., Wu, Y. T., Guo, Z. J. (2019). Prevalence of dyslipidemia and prediction of 10-year CVD risk among older adults living in southeast coastal regions in China: a cross-sectional study. *Clinical interventions in aging*; **14**: 1119–1129.
 42. Okwara J.E., Ike N.A., Ogbodo E.C., Okwara, N.A., Igwebuobi, C.F., Odumodu, I.O., Amah, A.K., Okezie, O.A., Onyenekwe, C.C. (2021). Assessment of cardiovascular disease awareness and risk factors in a market population in Nnewi, Nigeria. *Advances in Bioresearch*; **12** (4):236-243.
 43. Narindrangkura, P., Bosl, W., Rangsinsin, R., Hatthachote, P. (2019). Prevalence of dyslipidemia associated with complications in diabetic patients: a nationwide study in Thailand. *Lipids in Health and Disease*; **18**(90).
 44. Lumu, W., Kampiire, L., Akabwai, G. P., Ssekitoleko, R., Kiggundu, D. S., & Kibirige, D. (2017). Dyslipidaemia in a Black African diabetic population: burden, pattern and predictors. *BMC research notes*; **10**(1): 587.
 45. Micha, R., Michas, G., Mozaffarian, D. (2012). "Unprocessed red and processed meats and risk of coronary artery disease and type 2 diabetes--an updated review of the evidence". *Current Atherosclerosis Reports*; **14** (6): 515–524.
 46. Doupa, D., Seck, S.M., Dia, C.A., Diallo, F.A., Kane, M.O., Kane, A. (2014). Dyslipidemia, obesity and other cardiovascular risk factors in the adult population in Senegal. *Pan African Medical Journal*; **19**:181.
 47. Abujbara, M., Batieha, A., Khader, Y., Jaddou, H., El-Khateeb, M., Ajlouni, K. (2018). The Prevalence of Dyslipidemia among Jordanians. *Journal of lipids*, 2018, 6298739.
 48. Amberbir, A., Singano, V., Matengeni, A., Ismail, Z., Kawalazira, G., Chan, A.K. (2018). Dyslipidemia among rural and urban HIV patients in south-east Malawi. *PLoS ONE*; **13**(5): e0197728
 49. Feitosa, A.C.R., Barreto, L.T., Silva, I.M., Silva, F.F., Filho, G.S.F. (2017). Impact of the Use of Different Diagnostic Criteria in the Prevalence of Dyslipidemia in Pregnant Women. *Arq Bras Cardiol*; **109**(1):30–38.



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