

Dietary patterns and metabolic morbidities correlate among adolescents and young adults with HIV in Lagos, Nigeria: A cross-sectional study design Journal of Multimorbidity and Comorbidity Volume 15: 1–13 © The Author(s) 2025 Article reuse guidelines: sagepub.com/journals-permissions DOI: 10.1177/26335565251341388 journals.sagepub.com/home/cob



Mobolaji Olagunju^{1,*}, Abideen Olurotimi Salako^{2,3,*}, Titilola Abike Gbaja-biamila^{2,4}, Tomilola Musari-Martins², Priscilla Ngozi Ezemelue², Babasola Opaneye^{2,5}, Abubakar AhmadRufai Abubakar², Ibukun Oluwatomisin Odusote², Feyikemi Fasina², Adeniyi Adeyinka², Oreoluwa Alabi², Agatha Nkiru David², Lilian Ezechi⁶ and Oluwatosin O. Odubela²

Abstract

Background: The relationship between dietary patterns (DP) and health outcomes (elevated blood pressure, dyslipidaemia, hyperglycaemia, and body mass index) among adolescents and young adults (AYA) with HIV is not well understood. We aimed to identify dietary patterns and determinants associated with metabolic syndrome components among adolescents and young adults living with HIV in Lagos, Nigeria.

Methods: We conducted a cross-sectional study among 180 participants at an ART clinic in Lagos. Information on sociodemographic and clinical characteristics, as well as 72-hour dietary recall were collected. Anthropometric measurements (BMI (kg/m²)) and blood pressure readings were collected. Blood samples were assayed for fasting blood sugar and lipid profiles. Statistical analysis was done using SPSS version 27, WHO Anthro Plus software.

Results: The male-to-female ratio was I:I. Four major DPs identified were DP I (higher consumption of beverages, vitamins, and vegetables); DP 2 (high intake of minerals, and fruits); DP3 (higher intake of carbohydrates, fat and oil); and DP4 (higher intake alcohol and vegetables). DPI was associated with dyslipidaemia, underweight BMI and hyperglycaemia were associated with DP4, while DP2 had lower odds of overweight and elevated blood pressure.

Conclusion: This outcome provides valuable insights into DPs and their association with metabolic co-morbidity among AYA with HIV. This will inform nutritional counselling and interventions to promote quality of life and health.

Corresponding author:

Abideen Olurotimi Salako, Nigerian Institute of Medical Research, 6 Edmund Crescent Yaba, Lagos 101245, Nigeria. Email: salako.abideennaheem@gmail.com



Department of Epidemiology and Health Statistics, School of Public Health, Nanjing Medical University, Nanjing, China

²Clinical Sciences Department, Nigerian Institute of Medical Research Yaba, Lagos, Nigeria

³Global Paediatric Medicine (Global Scholar), Graduate School of Biomedical Sciences, St. Jude Children's Research Hospital, Memphis, TN, USA

⁴Department of Infectious Diseases, Washington State University, Pullman, WA, USA

⁵Wirral University Teaching Hospital, NHS Foundation Trust, Birkenhead Wirral London, UK

⁶Federal College of Education (Technical), Lagos, Nigeria

^{*}Joint First Author.

Keywords

adolescents, young adults, dietary pattern, metabolic disorders, HIV, Lagos, Nigeria

Received: 5 February 2025; revised: 6 April 2025; accepted: 23 April 2025

Key messages

What is already known this topic

Adolescents and Young Adults living with HIV (AYA with HIV) are at an increased risk of developing comorbid metabolic disorders, majorly originating from Antiretroviral therapy.

What does this study add(s)

Our study explored the contribution of diets to the burden of these comorbid metabolic disorders among AYA with HIV. The study participants on diets rich in minerals, fruits, and vegetables were found to be protective against hypertension and overweight.

How affect research, practice or policy

This will guide the development of nutritional guidelines for AYA with HIV to curtail the adversity of metabolic disorders.

Introduction

Adolescence and youth are critical in the growth and development phase towards the transition to adulthood. ^{1,2} This age group (adolescents and young adults) faces various challenges that could undermine achieving maximal human capabilities and capacities, particularly nutrition. ^{2,3} The nutritional status among this age group is influenced by risk factors, such as peer influences, parental modelling, food availability, food preferences, cost, convenience, personal and cultural beliefs, mass media, and body image that modulates their overall well-being. ^{4,5} Special populations such as adolescents and young adults (AYA) living with chronic diseases are of particular interest due to the interplay between immune status and nutrition. ³

Research has shown varied dietary patterns among AYA with HIV, ranging from diets rich in fruits, vegetables, lean protein, and whole grains to those high in processed and fast foods with low nutritional value. The dietary patterns among AYA with HIV can have significant long-term health implications, such as elevated blood pressure,

hyperglycaemia, dyslipidaemia and unhealthy BMI (underweight/overweight/obesity), which can undermine the gains of antiretroviral therapy (ART) and their overall wellbeing. ^{10,11} Balanced diets are essential for enhancing immune response, improving ART efficacy, and maintaining this cohort's optimal health. ^{12,13}

Adolescents and young adults, especially those with HIV, often receive insufficient attention regarding their diet, resulting in a poor understanding of their dietary behaviours.^{3,14,15} Emerging adulthood, a key stage for developing lifelong healthy habits, requires clear nutritional guidelines and education to promote better dietary choices. Targeted strategies, including nutrition-focused interventions, are essential to enhancing the well-being of AYA with HIV(3,14,15).

Therefore, considering the importance of diet to health in this cohort, our study aims to investigate the relationship between specific dietary patterns and metabolic comorbidities among AYA with HIV. The study will also characterize the common dietary patterns, as well as determine the association between dietary patterns and metabolic disorders components (hyperglycaemia, hypertension, dyslipidaemia and nutritional disorders (underweight/overweight/obesity) among AYA with HIV. We hypothesize that certain dietary patterns are associated with elevated blood pressure, dyslipidaemia, hyperglycaemia, and unhealthy (underweight/overweight) BMI) among AYA with HIV. This will provide valuable information that could inform nutritional guidance/counselling and the development of interventions to promote healthy dietary habits and lifestyles to enhance response to ART, immune function, and ultimately, improve the health and overall quality of life among AYA with HIV.

Methods

This was a prospective quantitative cross-sectional study among adolescents and young adults (AYA) with HIV (aged 10 to 24 years) who attended the ART Clinic at the Clinical Sciences Department of the Nigerian Institute of Medical Research (NIMR), Yaba, Lagos, Nigeria, between July and October 2019. The ART clinic is in the commercial capital of Nigeria and offers comprehensive HIV care and treatment to adults, adolescents, children and pregnant women. The ART clinic operates an adolescent-friendly clinic on the second Saturday of every month with about four hundred AYA with HIV currently in care. A cross-sectional design was chosen to provide a snapshot of the dietary patterns and

their associations with metabolic comorbidities in AYA with HIV. This study design was done to ensure efficient data collection and cost-effectiveness. Eligible participants were AYA with HIV who had been on antiretroviral drugs for at least six months and consented (consent \pm assent) to participate in the study. Excluded study participants were AYA with HIV with significant co-morbidities (seizure disorders, sickle cell anaemia, chronic hepatitis B infection, and those with acute illness) and those who were pregnant.

A sample size of 189 was determined using the Raosoft© online calculator. The calculation was performed with a 5% margin of error, 95% confidence level, a response distribution of 50% and a population of 300 AYAs with HIV attending the facility based on the ART clinic database at the conception of the study. Furthermore, an attrition rate of 10% was considered. A convenient sampling technique was used to select the required number of study participants. A case report form was used to obtain information on sociodemographic and clinical characteristics, clinical examination, laboratory parameters, and dietary patterns.

Socio-demographic and clinical characteristics

Age, gender, education, occupation, clinical parameters (age at HIV diagnosis, duration and type of ART, and family history of comorbidities [hypertension and diabetes mellitus]), physical examination (anthropometric measurements, blood pressure readings) and recent laboratory parameters (HIV RNA viral load and CD4 count within the 6 months of the study period).

Dietary information was obtained from respondents using the 72-hour dietary recall tool, which was administered three times to the study respondents (twice on a weekday and once on a weekend) to provide a detailed view of dietary intake among study participants. This information was analyzed and categorized into five classes of food (carbohydrates, fat and oil, minerals, vitamins, and proteins), with alcohol, beverages, fruits, and vegetables being extra classes.

Clinical examination

The anthropometric measurements (weight in kilograms, height in centimetres) were obtained with the patient in light clothing and barefooted using the Seca® combination weighing scale and stadiometer. The body mass index (BMI), expressed in kilograms per square meter, was calculated from the weight and height measurements. The BMI measurements were converted into z-scores using the WHO AnthroPlus software, a tool specifically designed for assessing the growth and development of children and adolescents. The z-scores provide a standardized way to compare the adolescents' growth to a reference population.

Two blood pressure readings were obtained one minute apart, using an electronic sphygmomanometer (Omron M3 Intellisense TM, Model: M3 (HEM-7131-E [Omron

Healthcare Co. Ltd]) with the participant comfortably seated and the right arm placed at the level of the participant's heart. The mean of the two readings was recorded as the participant's blood pressure. A third measurement was conducted after another 20 minutes if a high reading was obtained. The mean of the two closest readings is recorded as the participant's blood pressure. Point Hypertension was defined as a blood pressure reading \geq 95th percentile for age, sex, and height or readings above 130/80mmHg irrespective of participant's age or gender. ¹⁶

Laboratory parameters

After a 12-hour fast, approximately 6 ml of venous blood was collected from the antecubital fossa of each participant to determine their fasting lipid profiles [total cholesterol, triglycerides, high-density lipoprotein-cholesterol (HDL-c), and low-density lipoprotein-cholesterol (LDL-c)] using Roche C311 Clinical Chemistry autoanalyzer. Hypertriglyceridaemia (TG \geq 110mg/dl) and/or low HDL-c (\leq 40mg/dl) were used to define dyslipidemia as these are the two major predictive parameters for the development of cardiovascular disease. Age and gender-specific lipid profile cutoff points were used (normal and abnormal categories).

Dietary pattern assessment

Dietary patterns were determined using exploratory factor analysis (EFA). Kaiser–Meyer–Olkin (KMO; value of p > 0.05) and Bartlett's test of sphericity (value of p < 0.05) were used to check the adequacy of the sample-to-factors ratio. The existence of a correlation between food items was determined for each step of factor analysis. Then, EFA was done under orthogonal rotation with the varimax method to enhance the interpretability and independent dietary patterns. In determining the final dietary patterns, we considered factors with a communality above 0.2 and an eigenvalue greater than 1. These factors account for more variance than a single variable and are deemed significant in principal components analysis that reflect distinct consumption behaviours within the population.

Nutritional status assessment

The nutritional status of the participant was assessed through anthropometric measurements which include height, and weight, informing the calculation of the Body Mass Index (BMI). The World Health Organization (WHO) classification for adolescents was adopted, using age- and sex-specific growth charts. BMI is categorized as follows: underweight is below the 5th percentile, normal weight is between the 5th and 85th percentiles, overweight is from the 85th to less than the 97th percentile, and obesity is at or above the 97th percentile. While for

the young adults aged 20–24, the WHO standard adult BMI classification was applied. The categories were underweight (BMI < 18.5 kg/m^2), normal weight (BMI 18.5– 24.9 kg/m^2), overweight (BMI 25.0– 29.9 kg/m^2), and obesity (BMI $\geq 30.0 \text{ kg/m}^2$).

In this study, we identified distinct dietary patterns among AYA with HIV and determined how these eating habits influenced the study participants' nutrition and health. To achieve this, we employed regression models to evaluate the relationship between dietary patterns (exposure variables) and nutritional status as well as metabolic disorders (outcome variables). This approach provided a robust framework to understand the potential impact of dietary patterns on the health of study participants.

Statistical analysis

A descriptive analysis was performed to characterize the dataset, employing frequencies, percentages, means, and

standard deviations. The distribution of each variable was assessed to determine the normality, with outliers identified as values outside of the range (mean \pm 3SD). The chi-square test was used to assess the association between variables and gender, while binary logistic regression analysis was used to determine the association between dietary patterns (DPs) with sociodemographic characteristics and metabolic disorders. Multivariate logistic regression analysis was done to assess further correlation between the DPs and metabolic disorders while controlling for covariates in the models (Model 1 was adjusted for age and sex; Model 2 adjusted for age, sex, education, ethnicity and religion). Missing values and outliers for interested variables below a threshold of 5% of the total population were replaced with the mean (normally distributed data) or median (non-normally distributed data). Statistical analysis was done using SPSS version 27, WHO Anthro Plus software, and R studio, with statistical significance set at a p-value ≤ 0.05 .

Table 1. Gender-specific classification of sociodemographic and HIV characteristics.

Characteristics	Male N (%)	Female N (%)	Total N (%)	p-value
Age group (years)				0.106
10 – 14	42 (46.7)	40 (44.4)	82 (45.6)	
15 – 19	38 (42.2)	44 (48.9)	82 (45.6)	
20 – 24	12 (13.3)	4 (4.0)	16 (8.9)	
Education level				0.120
None	1 (1.1)	2 (2.2)	3 (1.7)	
Primary	12 (13.3)	12 (13.3)	24 (13.3)	
Secondary	73 (81.1)	63 (70.0=	136 (75.6)	
Tertiary	4 (4.4)	13 (13.3)	17 (9.4)	
Religion	. ,	, ,	, ,	0.269
Christianity	74 (82.2)	78 (86.7)	152 (84.4)	
Islam	16 (17.8)	12 (13.3)	28 (15.6)	
Ethnicity	` ,	,	,	0.524
Hausa	3 (3.3)	2 (2.2)	5 (2.8)	
Igbo	36 (40.0)	33 (36.7)	69 (38.3)	
Yoruba	39 (43.3)	34 (37.8)	73 (40.6)	
Edo	4 (4.4)	6 (6.7)	10 (5.6)	
Others	8 (8.9)	15 (16. 7)	23 (12.8)	
Family history of DM				0.500
Yes	83 (92.2)	8 (91.1)	15 (8.3)	
No	22 (24.4)	82 (91.1)	165 (91.7)	
CD4 counts	` ,	,	,	0.367
< 500	22 (24.4)	25 (27.8)	47 (26.1)	
≥ 500	68 (75.6)	65 (72.2)	133 (73.9)	
Viral load	, ,	, ,	, ,	0.171
≤1000	76 (84.4)	70 (77.8)	146 (81.1)	
>1000	14 (15.6)	20 (22.2)	34 (18.9)	
ART duration	, ,	` ,	, ,	0.046*
>5 years	77 (85.6)	67 (74.4)	144 (80.0)	
≤5 years	13 (14.4)	23 (25.6)	39 (20.0)	

Ethical issues

Ethical approval was obtained from the Institutional Review Board (IRB) of the Nigerian Institute of Medical Research, Lagos before the commencement of the study. All enrolled participants and/or caregivers were provided informed written consent at enrollment into the study.

Results

One hundred and eight nine (189) AYA with HIV were enrolled into the study. Nine⁹ individuals were excluded due to incomplete data. One hundred and eighty (180) AYA with HIV had complete data with a male-to-female ratio of 1:1. A

significant proportion (164, 91.2%) of the study participants were adolescents. Most of the participants had a secondary level of education (75.9%), were Christians (84.4%), were on ART for more than 5 years (80%), achieved virological suppression (81.1%) and had CD4 counts above 500 cells/ml (73.9%) Table 1.

Overall, underweight (43.3%) and normal (38.9%) BMI categories constituted the dominant BMI categories, with males having a higher proportion of underweight BMI categories compared to female counterparts (55.6% vs 31.1%). Females were predisposed to having normal (41.1% vs 36.7%) or overweight (27.8% vs 7.8%) BMI compared to their male counterparts and the gender differences were found to be statistically significant (Table 2).

Table 2. Gender-specific metabolic disorders.

Characteristics	Male N (%)	Female N (%)	Total N (%)	p-value
Body mass index (BMI)				<0.001*
Underweight	50 (55.6)	28 (31.1)	78 (43.3)	
Normal	33 (36.7)	37 (41.1)	70 (38.9)	
Overweight	7 (7.8)	25 (27.8)	32 (17.8)	
Elevated blood pressure				0.397
Yes	9 (10.7)	7 (43.8)	16 (8.9)	
No	75 (89.3)	77 (50.7)	152 (84.4)	
Lipid profile				
High density lipoprote	ein-cholesterol (HDL-c)			0.049*
Normal	44 (48.9)	56 (62.2)	100 (55.6)	
Abnormal	46 (51.1)	34 (37.8)	80 (44.4)	
Low density lipoprotein-	-cholesterol (LDL-c)			0.360
Normal	87 (96.7)	85 (94.4)	172 (95.6)	
Abnormal	3 (3.3)	5 (5.6)	8 (4.4)	
Triglycerides (TG)				0.235
Normal	68 (75.6)	73 (81.1)	141 (78.3)	
Abnormal	22 (24.4)	17 (18.9)	39 (21.7)	
Total cholesterol (TC)				0.235
Normal	68 (75.6)	73 (81.1)	141 (78.3)	
Abnormal	22 (24.4)	17 (18.9)	39 (21.7)	
Non high density lipo	protein-cholesterol (non-HDI	c)		0.391
Normal	84 (93.3)	82 (91.1)	166 (92.2)	
Abnormal	6 (6.7)	8 (8.9)	14 (7.8)	
TC/HDL-c				0.369
Normal	64 (71.1)	63 (71.1)	131 (72.8)	
Abnormal	26 (28.9)	26 (28.9)	49 (27.2)	
TG/HDL-c				0.218
Normal	55 (61.1)	61 (67.8)	116 (64.4)	
Abnormal	35 (38.9)	29 (32.2)	64 (35.6)	
LDL-c/HDL-c				0.363
Normal	67 (74.4)	70 (76.1)	137 (76.1)	
Abnormal	23 (25.6)	20 (22.2)	43 (23.9)	
Hyperglycemia				0.614
Normal	53 (94.6)	42 (95.5)	95 (52.8)	
Abnormal	3 (5.4)	2 (4.5)	5 (2.8)	

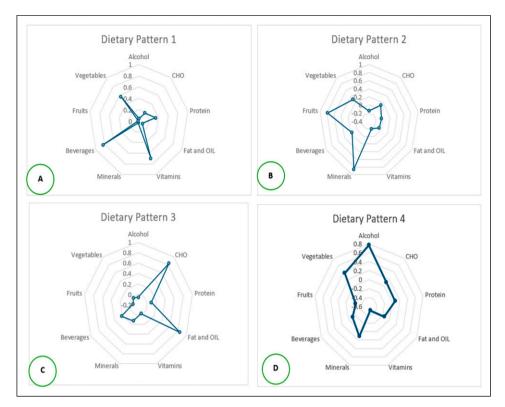


Figure 1. Dietary patterns among study participants following meal 72 hour recall. A: Dietary Pattern 1; B: Dietary Pattern 2; C: Dietary Pattern 3; D: Dietary Pattern 4

The prevalence of hypertension in the study was 9.5%, with the lipid profile revealing abnormal HDL-c (44.4%), LDL-c (4.4%), Triglycerides (21.7%) and Total Cholesterol (21.7%) levels among study participants. Only 2.8% of the study population had abnormal fasting hyperglycaemic levels (Table 2).

Figure 1 depicts the four dietary patterns identified among AYA with HIV enrolled in the study. These dietary patterns account for approximately 65% of the study population (Supplemental data Table S1, S2 and S3). Dietary pattern 1 (23.5%) is characterized by higher consumption of beverages, vitamins, and vegetables while dietary pattern 2 (16.4%), contains predominantly minerals and fruits but is low in vegetables, carbohydrates, protein, and other essential nutrients. Dietary pattern 3 (13.2%) consists of high consumption of carbohydrates, fats and oil, along with low micronutrient content, while dietary pattern 4 (11.6%) had a high composition of alcohol, vegetables, and minerals, but low protein and carbohydrate content. (Figure 1).

Dietary patterns and BMI (underweight and overweight)

Underweight BMI was associated with dietary patterns 1 (OR:1.176, 95% CI: 1.001 – 1.303) and 4 (OR: 1.109, 95%

CI: 1.082 – 1.986). Dietary pattern 4 retained a positive association with being underweight using both models 1 and 2. None of the dietary patterns displayed any association with overweight except for dietary pattern 2 (using Model 2), which showed reduced odds of being overweight (OR: 0.376, 95% CI: 0.234 – 0.852). Table 3

Dietary patterns and blood pressure

Dietary pattern 2 was shown to protect study participants from being hypertensive irrespective of the equations utilized (p <0.05). All other dietary patterns did not reveal any association with blood pressure. Table 4

Dietary patterns and hyperglycaemia

Only dietary pattern 4 was associated with hyperglycaemia irrespective of the model utilized (p < 0.05). Table 4

Dietary patterns and dyslipidaemia

Dietary pattern 1 was exclusively associated with lipid profile derangement with TC/HDL and LDL-c/HDLc ratios showing significant associations (Table 5).

 Table 3. Association between sociodemographic status and dietary patterns.

		Dietary pattern I							
Variables		Mean (95% CI)	ф	Dietary pattern 2	ф	Dietary pattern 3	ф	Dietary pattern 4	ф
Age	10-14 15-19 20-24	0.148 (-0.110 - 0.406) -0.34 (-0.318 - 0.050) -0.732 (-0.444 - 0.298)	0.189	0.048 (-0.171 - 0.267 0.020 (-0.218 - 0.258) -0.347 (-0.5340.160)	0.344	-0.039 (-0.249 - 0.169) 0.097 (-0.141 - 0.335) -0.293 (-0.678 - 0.093)	0.323	-0.097 (-0.263 - 0.069) -0.009 (-0.243 - 0.223) 0.547 (-0.281 - 1.374)	0.061
Sex	Male Female		0.456	$0.049 \; (-0.189 - 0.287) \\ -0.049 \; (-0.225 - 0.128)$	0.515	-0.037 (-0.253 - 0.179) 0.037 (-0.166 - 0.241)	0.619	0.061 (-0.172 - 0.295) -0.061 (-0.244 - 0.121)	0.412
Education level	Primary Secondary Tertiary	0.179 (-0.356 - 0.714) -0.013 (-0.178 - 0.152) 0.103 (-0.401 - 0.422)	0.686	-0.179 (-0.445 - 0.086) 0.074 (-0.109 - 0.259) -0.334 (-0.6340.034)	0.184	-0.291 (-0.693 - 0.111) 0.098 (-0.071 - 0.266) -0.233 (-0.784 - 0.318)	0.123	-0.066 (-0.343 - 0.211) 0.028 (-0.146 - 0.201) -0.249 (-0.793 -0.295)	0.530
Ethnicity	Yoruba Igbo Hausa Edo	-0.108 (-0.335 - 0.132) 0.007 (-0.231 - 0.244) -0.033 (-1.856 - 1.789) 0.391 (-0.118 - 0.899)	909.0	-0.122 (-0.328 - 0.085) 0.111 (-0.158 - 0.379) 0.732 (-1.345 - 2.807) 0.025 (-0.386 - 0.436)	0.297	-0.049 (-0.282 - 0.182) 0.013 (-0.248 - 0.278) 0.088 (-0.392 - 0.567) 0.182 (-0.474 - 0.839)	7967	-0.209 (-0.425 - 0.007) 0.112 (-0.123 - 0.347) -0.048 (-0.508 - 0.413) 0.294 (-0.611 - 1.199)	0.205
Religion	Christian Islam	-0.028 (-0.181 - 0.125) 0.153 (-0.321 - 0.627)	0.380	0.084 (-0.083 - 0.251) -0.458 (-0.6740.242)	0.008*		0.046*	0.027 (-0.134 - 0.188) 0.027 (-0.134 - 0.188) -0.147 (-0.531 - 0.237)	0.398

Discussion

Our study explores the association between dietary patterns and metabolic disorders (BMI, elevated blood pressure, dyslipidaemia, hyperglycemia) among AYA with HIV at a large ART clinic in Lagos, Nigeria. We identified four distinct dietary patterns with varying associations with metabolic disorders in the study population. Dietary pattern 1 was positively associated with higher likelihood of underweight and dyslipidaemia. Dietary pattern 2 was protective against elevated blood pressure and overweight, while dietary pattern 4 was associated with study participants being underweight and hyperglycaemic. Dietary pattern 3 did not show any association with any component of metabolic disorders.

The prevalence of underweight at 43.3% among AYA with HIV is within the prevalence reported (2 - 63%)among adolescents with HIV in LMICs in a recent systematic review. 17 However, our prevalence was higher compared with the reported prevalence of 27% in Nigeria, 18 20 - 27.2% in different studies across Ethiopia^{8,19,20} and 18% in Uganda.²¹ The wide disparity in the prevalence could be attributed to the variation in study design, study settings, socioeconomic status and cultural values/disposition. Our findings buttress the burden of nutritional challenges among the HIV cohort, even in the era of test and treat protocol. The high demand for nutrients (required growth and development) in this age group irrespective of HIV status, is inundated with food insecurity and unbalanced energy expenditures. The prolonged span of undernutrition in the foetal and under-5 age group, along with the catabolic effects of HIV could explain the high prevalence of underweight. The disproportionate underweight burden among males is in concordance with prior studies in LMICs irrespective of HIV status. ^{17,22–24} This could be attributed to the increased energy requirements and physical exertions among males amid poor food choices and socioeconomic status.²⁵

Dietary pattern 4 was characterized by low content of carbohydrates and proteins which are essential for growth and development but remain difficult to access due to prevailing food insecurity, food inflation and poor food choices in LMICs. 4,5,25 In addition, underweight BMI status was also associated with alcohol consumption among AYA in our study similar to findings in previous studies. 26-29 Although our findings differ from previous studies that have associated overweight/obese BMI status among adolescents with alcohol consumption, most of these studies did not include AYA with HIV population. Given the background HIV infection, poor appetite, opportunistic infections (in the gastrointestinal tract), and malabsorption of nutrients may result in the underweight status of our study populace. Future studies are required to delineate the impact

of alcohol and dietary patterns on weight changes among AYA with HIV.

The prevalence of 17.8% for overweight among the AYA with HIV, with the female gender accounting for a higher proportion of the individuals who were overweight. This is higher than the reported overweight prevalence (6.1 - 15.6%) in a systematic review and meta-analysis by Gebrie et al. ¹⁷ The increased prevalence of overweight could be attributed to the positive effects of ART, creating a significant reversal of the catabolic effects of HIV infection and a corresponding reduction in inflammatory markers with resultant improved appetite and nutrient uptake.

The protective effect of dietary pattern 2 (rich in minerals and fruits) in preventing overweight affirms previous studies corroborating the importance of micronutrient intake which occurs by modulating the body's metabolism, immune system, systemic inflammation, and oxidative stress. ^{34,35} Although it is noteworthy that the current study focuses on dietary intake among the study population with chronic illness (HIV), other variables beyond the scope of our study could predispose to overweight/ obesity. These factors include genetics, other chronic diseases, drugs, and environmental factors. ^{17,36–38}

The burden of hypertension (9.5%) is within the pooled prevalence (0.2-24.8%) reported in a systematic review and meta-analysis in sub-Saharan Africa.³⁹ Our prevalence differed from the reported prevalence by (6.9%) by Odubela et al., 40 (10.5%) by David et al., 41 and 11% by Nsanya et al.. 42 although there were variations in the study populations. In addition, our finding is lower than the prevalence estimates (10-40%) in Africa's general population of adolescents and young adults, 29% by Lubega et al.⁴³ and 49% reported by Migisha et al.⁴⁴ in Uganda. The variability in the prevalence of hypertension could be due to genetic, racial, lifestyle and environmental factors. Dietary pattern 2 was protective against elevated blood pressure. This dietary pattern displays the interplay between diet composition and hypertension. 45–47 Our finding affirms the effectiveness of dietary and lifestyle modifications in the prevention and management of elevated blood pressure and its complications.

The prevalence of hyperglycaemia among the participants was 2.8%. This finding is within the range of 2 - 28% in sub-Saharan Africa, 41,48-51 0.4-7% in America, 52-55 Europe, 56,57 and 2.6 - 43% from the Asian continent. 58,59 Significant contributors to hyperglycaemia include chronic inflammatory response (arising from HIV infection) and effects of ART (mitochondrial toxicity, altered endoplasmic reticulum function, alteration in adipocytokines & lipid metabolism, impaired insulin signaling pathways) in addition to other traditional risk factors. 49,60-64 Excessive alcohol consumption can induce insulin resistance as alcohol can interfere with the liver's ability to regulate blood

Table 4. Regression model to assess the association between dietary patterns and metabolic disorders.

		Elevated - BP		Hyperglycemia		Underweight -BMI		Overweight -BMI	
Variable		OR (95% CI)	p-value	OR (95% CI)	ρ-value	OR (95% CI)	p-value	OR (95% CI)	ρ-value
Unadjusted model	Dietary pattern 1	1.129 (0.664-1.829)	0.634	1.012 (0.359-2.384)	0.980	1.176 (1.001-1.303)	0.021*	1.003 (0.998-1.243)	0.064
	Dietary pattern 3	0.738 (0.419-1.249)	0.275	1.111 (0.413-2.907)	0.835	2.019 (0.983-3.572)	0.911	1.318 (0.768-2.019)	0.452
	Dietary pattern 4	1.018 (0.567-1.621)	0.947	2.259 (1.256-4.232)	*900.0	1.109 (1.082-1.986)	*I 00.0	1.845 (0.320-2.682)	0.649
		AOR (95% CI)	p-value	AOR (95% CI)	p-value	AOR (95% CI)	p-value	AOR (95% CI)	p-value
Model I	Dietary pattern I	1.177 (0.684-1.948)	0.537	1.152 (0.342-3.449)	0.803	1.018 (0.748-3.651)	0.714	0.998 (0.557-2.956)	0.828
	Dietary pattern 2	0.399 (0.127-0.913)	0.028*	1.772 (0.846-3.757)	0.108	0.976 (0.612-1.254)	0.524	1.123 (0.769-2.598)	0.821
	Dietary pattern 3	0.744 (0.421-1.256)	0.288	1.047 (0.325-2.873)	0.934	1.934 (0.678-2.287)	0.109	1.210 (0.854-1.714)	0.270
	Dietary pattern 4	0.984 (0.551-1.575)	0.951	2.115 (1.093-4.422)	0.027*	1.654 (1.432-1.989)	0.029*	0.109 (1.082-1.986)	0.451
		AOR (95% CI)	p-value	AOR (95% CI)	p-value	AOR (95% CI)	p-value	AOR (95% CI)	p-value
Model 2	Dietary pattern I	1.236 (0.692-2.152)	0.457	4.581 (0.592-244.308)	0.212	0.812 (0.567-1.163)	0.220	0.987 (0.743-1.312)	0.513
	Dietary pattern 2	0.468 (0.079-0.776)	0.040*	1.038 (0.115-3.338)	0.955	0.823 (0.678-1.257)	0.340	0.376 (0.234-0.852)	0.015*
	Dietary pattern 3	0.654 (0.343-1.157)	0.168	0.627 (0.055-6.329)	0.688	1.789 (0.543-2.146)	0.210	2.123 (0.567-2.874)	0.347
	Dietary pattern 4	1.098 (0.539-2.078)	0.782	3.012 (1.359-4.584)	0.035*	1.567 (1.102-2.229)	0.045*	1.345 (0.987-1.832)	0.065

Model 1: Adjusted for age and sex. Model 2: Adjusted for age, sex, education level, ethnicity and religion. p-value < 0.05 – statistically significant.

	Dietary pattern I		Dietary pattern 2		Dietary pattern 3		Dietary pattern 4	
	OR (95% CI)	p-value						
HDL-c (< 40)	1.049 (0.780-1.411)	0.749	1.037 (0.769-1.395)	0.807	0.785 (0.573-1.059)	0.121	1.227 (0.912-1.677)	0.183
LDL-c (≥ 130)	1.081 (0.507-2.077)	0.826	1.223 (0.594-2.086)	0.511	1.194 (0.583-2.279)	0.609	0.649 (0.240-1.419)	0.361
TG (≥ 200)	0.847 (0.317-1.866)	0.707	0.237 (0.028-1.043)	0.131	1.131 (0.489-2.370)	0.760	0.723 (0.238-1.645)	0.531
TC (≥ 200)	0.936 (0.512-1.590)	0.817	1.057 (0.574-1.687)	0.835	0.835 (0.463-1.442)	0.534	0.671 (0.314-1.243)	0.266
Non-HDL-c	0.902 (0.490-1.541)	0.722	1.323 (0.803-2.06)	0.220	0.819 (0.453-1.417)	0.493	0.790 (0.387-1.385)	0.473
TC/HDL-c (≥ 5.0)	1.506 (1.089-2.104)	0.014*	0.909 (0.627-1.263)	0.590	0.896 (0.635-1.246)	0.522	1.289 (0.939-1.774)	0.112
TG/HDL-c (>3.0)	1.323 (0.976-1.807)	0.073	0.81(0.569-1.119)	0.227	0.951 (0.695-1.292)	0.751	1.182 (0.873-1.606)	0.276
LDL-c/HDL-c (>3.5)	1.499 (1.072-2.115)	0.019*	0.979 (0.674-1.364)	0.908	0.787 (0.541-1.119)	0.194	1.187 (0.848-1.639)	0.300

Table 5. Regression association between abnormal lipid indices and dietary patterns.

glucose, predisposing to diabetes mellitus.⁶⁵ This aligns with previous studies on the relationship between blood sugar, metabolic disorders and alcohol intake in the general population.^{66–68}

The pattern of dyslipidaemia with abnormal HDL-c ratios found in our study aligns with reports from across Africa. 69-73 However, a reverse pattern of higher prevalence of hypertriglyceridaemia followed by low HDL-c was reported in other studies in Africa, ^{74,75} Asia ^{58,76} and Europe. 57 This variation could be due to genetic differences, dietary status and the class of ART. In consonance with prior studies, beverage consumption (DP 1) was associated with dyslipidaemia. 77-79 Our finding buttresses the adverse implications of high beverage intake and dyslipidaemia. Thus, the evidence supports the need to improve the reach and quality of health information available to AYA to promote healthy eating habits and curtail adverse health conditions of metabolic disorders such as dyslipidaemia, obesity, diabetes mellitus and cardiovascular diseases.80,81

Strength and limitations

This study is one of the first to describe dietary patterns and their association with common metabolic disorders (elevated blood pressure, hyperglycaemia, abnormal body weight and dyslipidaemia) among AYA with HIV in Africa. Food recall, the nature of the study design (cross-sectional), and the number of food servings limit the generalization of our findings. The study did not account for environmental or sociocultural factors that may impact food intake, quality and quantity.

Conclusion

This study provides valuable insights into the dietary patterns and its association with metabolic disorders. It also provides evidence of the association between dietary patterns and metabolic disorders among AYA with HIV. The study findings affirm the need for comprehensive care to ensure the gains of HIV care are consolidated and promote wholesome health among AYA with HIV. This will entail continuous nutritional counselling and prompt monitoring of modifiable and non-modifiable risk factors to curtail the burden of non-communicable diseases. This finding will help inform nutritional guidance/counselling and the development of interventions to promote healthy dietary habits, prevent the onset of NCDs and, ultimately, improve the quality of life of AYA with HIV.

Acknowledgements

Our sincere appreciation to the adolescents and young adults with HIV (AYA with HIV) and their caregivers/parents who participated in the study. Also, we appreciate the unflinching trust in the staff at the Clinical Sciences Department (CSD) at the Nigerian Institute of Medical Research (NIMR) by the AYA, parents/caregivers for over two decades. Furthermore, our depth of gratitude goes to the nurses, counsellors/social workers, members of the data unit, and the laboratory team at the Centre for Human Virology and Genomics, who have continued to work tirelessly towards improving the health and well-being of people living with HIV, especially AYA with HIV.

ORCID iD

Abideen Olurotimi Salako https://orcid.org/0000-0002-3018-3712

Ethical statement Ethical approval

All procedures performed in studies involving human participants followed the institutional and/or national research committee's ethical standards, the 1964 Helsinki Declaration, and its later amendments or comparable ethical standards. This article does not contain any studies with animals performed by any of the authors.

Ethical approval was obtained from the Institutional Review Board (IRB), NIMR, Lagos, IRB/19/071.

Consent to participate

The purpose, processes, and expected outcome of the study were explained to participants and their caregivers (for those younger than 18 years), and their assent and informed consent were obtained before the commencement of the study. Confidentiality was maintained, and the freedom to withdraw at any time from the study without negative consequences was emphasized.

Authors contributions

Conceptualization: AOS, MO, OOO, OO, TM, LE, NAD, TAG. Methodology and Data Collection: AOS, OOO, TM, PNE. AAA, TAG. Data Analysis-, MO, AOS, OOO, OO, AAA, TAG. Writing (Original Draft): AOS, MO, OOO, FF, AA, Writing – review & editing: AOS, OOO, OO, PNE, AAA, TM, FF, AA, LE, NAD, TAG. Supervision: AOS, OOO, LE, NAD, TAG.

Funding

No financial support or grant.

Declaration of conflicting interests

The authors declare no competing interest.

Data Availability Statement

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

Supplemental Material

Supplemental material for this article is available online.

References

- Patton GC, Sawyer SM, Santelli JS, et al. Our future: a Lancet commission on adolescent health and wellbeing. *The Lancet* 2016; 387(10036): 2423–2478.
- 2. Kleinert S and Horton R. Adolescent health and wellbeing: a key to a sustainable future. *The Lancet* 2016; 387(10036): 2355–2356.
- 3. Lassi Z, Moin A and Bhutta Z. Nutrition in Middle Childhood and Adolescence. In: Bundy DAP, Nd Silva, Horton S, et al., editors. *Child and Adolescent Health and Development*. Washington D.C; 2017. Available from: https://www.ncbi.nlm.nih.gov/books/NBK525242/
- Vahdat M, Hosseini SA, Khalatbari Mohseni G, et al. Effects of resistant starch interventions on circulating inflammatory biomarkers: a systematic review and meta-analysis of randomized controlled trials. *Nutr J* 2020; 19(1): 33.
- Bakhtiary M, Morvaridzadeh M, Agah S, et al. Effect of Probiotic, Prebiotic, and Synbiotic Supplementation on Cardiometabolic and Oxidative Stress Parameters in Patients With Chronic Kidney Disease: A Systematic Review and Meta-analysis. Clinical Therapeutics 2021; 43(3): e71–e96.

- 6. Neta A da CP de A, Steluti J, Ferreira FEL, et al. Dietary patterns among adolescents and associated factors: longitudinal study on sedentary behavior, physical activity, diet and adolescent health [Internet]. SciELO journals, 2022. Available from: https://scielo.figshare.com/articles/dataset/Dietary_patterns_among_adolescents_and_associated_factors_longitudinal_study_on_sedentary_behavior_physical_activity_diet_and_adolescent_health/19904714
- Malindisa E, Dika H, Rehman AM, et al. Dietary patterns and diabetes mellitus among people living with and without HIV: a cross-sectional study in Tanzania. Front Nutr 2023; 10: 1105254.
- Shiferaw H and Gebremedhin S. Undernutrition Among HIV-Positive Adolescents on Antiretroviral Therapy in Southern Ethiopia. AHMT 2020; 11: 101–111.
- Martín-Cañavate R, Sonego M, Sagrado MJ, et al. Dietary patterns and nutritional status of HIV-infected children and adolescents in El Salvador: A cross-sectional study. In: Van Wouwe JP (ed). *PLoS One* 2018; 13(5): e0196380.
- Al-Jawaldeh A and Abbass MMS. Unhealthy Dietary Habits and Obesity: The Major Risk Factors Beyond Non-Communicable Diseases in the Eastern Mediterranean Region. Front Nutr 2022; 9: 817808.
- Obeagu EI, Obeagu GU, Odo EO, et al. Nutritional Approaches for Enhancing Immune Competence in HIV-Positive Individuals: A Comprehensive Review. *IDOSR-JAS* 2024; 9(1): 40–50.
- Basta D, Latinovic OS and Silvestri G. Potential Advantages of a Well-balanced Nutrition Regimen for People Living with Human Immunodeficiency Virus Type -1. *J AIDS HIV Treat* 2024; 6(1): 11–27.
- Fathima A, Madhu M, Kumar UdayaV, et al. Nutritional Aspects of People Living with HIV (PLHIV) Amidst COVID-19 Pandemic: an Insight. *Curr Pharmacol Rep* 2022; 8(5): 350–364.
- 14. UNICEF. Adolescents Living with HIV: Developing and Strengthening Care and Support Services, Geneva: UNICEF Regional Office for Central and Eastern Europe and the Commonwealth of Independent States (CEECIS). UNICEF; 2016. Available from: https://www.unicef.org/ceecis
- Rezazadeh L, Ostadrahimi A, Tutunchi H, et al. Nutrition interventions to address nutritional problems in HIV-positive patients: translating knowledge into practice. *J Health Popul Nutr* 2023; 42(1): 94.
- Flynn JT, Kaelber DC, Baker-Smith CM, et al. Clinical Practice Guideline for Screening and Management of High Blood Pressure in Children and Adolescents. *Pediatrics* 2017; 140(3): e20171904.
- 17. Gebrie M, Perry L, Xu X, et al. Nutritional status and its determinants among adolescents with HIV on anti-retroviral treatment in low- and middle-income countries: a systematic review and meta-analysis. *BMC Nutr* 2023; 9(1): 60.
- Fagbamigbe AF, Adebowale AS and Ajayi I. An assessment of the nutritional status of ART receiving HIV-orphaned and

- vulnerable children in South-West Nigeria. *Heliyon* 2019; 5(12): e02925.
- Teshome MS, Gissa SB, Tefera BZ, et al. Undernutrition and its predictors among people living with HIV/AIDS attending antiretroviral therapy clinic in Jimma University Specialized HosPITAL. *Int J Nutr Metab*; 9(8): 67–74.
- Demilew YM and Emiru AA. Under nutrition and associated factors among school adolescents in Dangila Town, Northwest Ethiopia: a cross sectional study. *Afr H Sci* 2018; 18(3): 756.
- Lwanga F, Wanyenze KR, Matovu KBJ, et al. Nutritional Status of HIV-infected Adolescents Enrolled into an HIV-care Program in Urban and Rural Uganda: A Cross-sectional Study. World Journal of Nutrition and Health: 3(2): 35–40.
- Manyanga T, El-Sayed H, Doku DT, et al. The prevalence of underweight, overweight, obesity and associated risk factors among school-going adolescents in seven African countries. BMC Public Health 2014; 14(1): 887.
- Darling AM, Sunguya B, Ismail A, et al. Gender differences in nutritional status, diet and physical activity among adolescents in eight countries in sub-Saharan Africa. *Tropical Med Int Health* 2020; 25(1): 33–43.
- Bener A. Prevalence of Obesity, Overweight, and Underweight in Oatari Adolescents. Food Nutr Bull 2006; 27(1): 39–45.
- Pawlińska-Chmara R, Wronka I, Suliga E, et al. Socioeconomic factors and prevalence of underweight and overweight among female students in Poland. HOMO 2007; 58(4): 309–318.
- Tolstrup JS, Heitmann BL, Tjønneland AM, et al. The relation between drinking pattern and body mass index and waist and hip circumference. *Int J Obes* 2005; 29(5): 490–497.
- 27. Dumesnil C, Dauchet L, Ruidavets JB, et al. Alcohol Consumption Patterns and Body Weight. *Ann Nutr Metab* 2013; 62(2): 91–97.
- MacInnis RJ, Hodge AM, Dixon HG, et al. Predictors of increased body weight and waist circumference for middleaged adults. *Public Health Nutr* 2014; 17(5): 1087–1097.
- Lean MEJ, Vlachou P, Govan L, et al. Different associations between body composition and alcohol when assessed by exposure frequency or by quantitative estimates of consumption. J Human Nutrition Diet 2018; 31(6): 747–757.
- 30. Wannamethee SG, Shaper AG and Whincup PH. Alcohol and adiposity: effects of quantity and type of drink and time relation with meals. *Int J Obes* 2005; 29(12): 1436–1444.
- Breslow RA. Drinking Patterns and Body Mass Index in Never Smokers: National Health Interview Survey, 1997-2001. American Journal of Epidemiology 2005; 161(4): 368–376.
- Brumby S, Kennedy A and Chandrasekara A. Alcohol Consumption, Obesity, and Psychological Distress in Farming Communities—An Australian Study. *The Journal of Rural Health* 2013; 29(3): 311–319.
- 33. O'Donovan G, Stamatakis E and Hamer M. Associations between alcohol and obesity in more than 100 000 adults in England and Scotland. *Br J Nutr* 2018; 119(2): 222–227.

- Bradley M, Melchor J, Carr R, et al. Obesity and malnutrition in children and adults: A clinical review. *Obesity Pillars* 2023; 8: 100087.
- Pineda E, Stockton J, Scholes S, et al. Food environment and obesity: a systematic review and meta-analysis. *BMJ Nutri*tion 2024: e000663.
- Endy EJ, Yi SY, Steffen BT, et al. Added sugar intake is associated with weight gain and risk of developing obesity over 30 years: The CARDIA study. *Nutrition, Metabolism* and Cardiovascular Diseases 2024; 34(2): 466–474.
- Narciso J, Silva AJ, Rodrigues V, et al. Behavioral, contextual and biological factors associated with obesity during adolescence: A systematic review. In: Carraça E (ed). *PLoS One* 2019; 14(4): e0214941.
- Endalifer ML and Diress G. Epidemiology, Predisposing Factors, Biomarkers, and Prevention Mechanism of Obesity: A Systematic Review. *Journal of Obesity* 2020; 2020: 1–8.
- Chen A, Waite L, Mocumbi AO, et al. Elevated blood pressure among adolescents in sub-Saharan Africa: a systematic review and meta-analysis. *The Lancet Global Health* 2023; 11(8): e1238–e1248.
- 40. Odubela O, Odunukwe N, Peer N, et al. Prevalence of hypertension among antiretroviral therapy naïve patients in Lagos, Nigeria. *Clin Hypertens* 2023; 29(1): 29.
- David AN, Gbaja-Biamilla TA, Odubela OO, et al. Prevalence of hypertension among adolescents living with human immunodeficiency virus in Lagos, Nigeria. Edorium J Pediatr 2; 5: 1–8.
- 42. Nsanya MK, Kavishe BB, Katende D, et al. Prevalence of high blood pressure and associated factors among adolescents and young people in Tanzania and Uganda. *J of Clinical Hypertension* 2019; 21(4): 470–478.
- 43. Lubega G, Mayanja B, Lutaakome J, et al. Prevalence and factors associated with hypertension among people living with HIV/AIDS on antiretroviral therapy in Uganda. *Pan Afr Med J* 2021; 15: 38. Available from: https://www.panafricanmed-journal.com/content/article/38/216/full
- 44. Migisha R, Ario AR, Kadobera D, et al. High blood pressure and associated factors among HIV-infected young persons aged 13 to 25 years at selected health facilities in Rwenzori region, western Uganda, September–October 2021. Clin Hypertens 2023; 29(1): 6.
- 45. Reddy KS and Katan MB. Diet, nutrition and the prevention of hypertension and cardiovascular diseases. *Public Health Nutr* 2004; 7(1a): 167–186.
- Bazzano LA, Green T, Harrison TN, et al. Dietary Approaches to Prevent Hypertension. *Curr Hypertens Rep* 2013; 15(6): 694–702.
- Aljuraiban GS, Gibson R, Chan DSM, et al. The Role of Diet in the Prevention of Hypertension and Management of Blood Pressure: An Umbrella Review of Meta-Analyses of Interventional and Observational Studies. *Advances in Nutrition* 2024; 15(1): 100123.
- 48. Frigati LJ, Jao J, Mahtab S, et al. Insulin Resistance in South African Youth Living with Perinatally Acquired HIV

- Receiving Antiretroviral Therapy. AIDS Research and Human Retroviruses 2019; 35(1): 56-62.
- Nkinda L, Buberwa E, Memiah P, et al. Impaired fasting glucose levels among perinatally HIV-infected adolescents and youths in Dar es Salaam, Tanzania. Front Endocrinol 2022; 13: 1045628.
- Micondo HK, Oyenusi EE, Dainguy ME, et al. Fasting Blood Glucose Profile of Children Living with HIV taking First-Line Antiretroviral Treatment in Abidjan, Cote D'Ivoire: A Cross-Sectional Study. Rom J Diabetes Nutr Metab Ds 2020; 27(2): 90–98.
- Ohuche IO, Chikani UN, Oyenusi EE, et al. Correlates of fasting blood glucose among children living with hiv in a Nigerian tertiary hospital: a cross-sectional study. BMC Pediatr 2020: 20(1): 458.
- 52. Geffner ME, Patel K, Jacobson DL, et al. Changes in insulin sensitivity over time and associated factors in HIV-infected adolescents. *AIDS* 2018; 32(5): 613–622.
- Hazra R, Hance LF, Monteiro JP, et al. Insulin Resistance and Glucose and Lipid Concentrations in a Cohort of Perinatally HIV-infected Latin American Children. *Pediatric Infectious Disease Journal* 2013; 32(7): 757–759.
- Paganella MP, Cohen RA, Harris DR, et al. Association of Dyslipidemia and Glucose Abnormalities With Antiretroviral Treatment in a Cohort of HIV-Infected Latin American Children. *JAIDS Journal of Acquired Immune Deficiency Syndromes* 2017; 74(1): e1–8.
- 55. Borkowska T, Chkhartishvili N, Karkashadze E, et al. The prevalence of hyperglycemia and its impact on mortality among people living with HIV in Georgia. Apetrei C, editor. *PLoS ONE* 2022; 17(10): e0276749.
- Espiau M, Yeste D, Noguera-Julian A, et al. Metabolic Syndrome in Children and Adolescents Living with HIV. *The Pediatric Infectious Disease Journal* 2016; 35(6): e171–e176.
- Blázquez D, Ramos-Amador JT, Saínz T, et al. Lipid and glucose alterations in perinatally-acquired HIV-infected adolescents and young adults. BMC Infect Dis 2015; 15(1): 119.
- Santiprabhob J, Tanchaweng S, Maturapat S, et al. Metabolic Disorders in HIV-Infected Adolescents Receiving Protease Inhibitors. *BioMed Research International* 2017; 2017: 1–14.
- 59. Seang K, Javanbakht M, Lee SJ, et al. Differences in prevalence and risk factors of non-communicable diseases between young people living with HIV (YLWH) and young general population in Cambodia. In: Page K (ed), *PLoS One* 2022; 17(6): e0269989.
- Hu FB. Globalization of Diabetes. *Diabetes Care* 2011; 34(6): 1249–1257.
- 61. Hivert MF, Sun Q, Shrader P, et al. Circulating IL-18 and the risk of type 2 diabetes in women. *Diabetologia* 2009; 52(10): 2101–2108.
- Limone P, Biglino A, Valle M, et al. Insulin resistance in HIVinfected patients: relationship with pro-inflammatory cytokines released by peripheral leukocytes. *Journal of Infection* 2003; 47(1): 52–58.

- 63. Pham TM, Carpenter JR, Morris TP, et al. Ethnic Differences in the Prevalence of Type 2 Diabetes Diagnoses in the UK: Cross-Sectional Analysis of the Health Improvement Network Primary Care Database. CLEP 2019; 11: 1081–1088.
- 64. Gutierrez AD and Balasubramanyam A. Dysregulation of glucose metabolism in HIV patients: epidemiology, mechanisms, and management. *Endocrine* 2012; 41(1): 1–10.
- 65. Salama MS, Isunju JB, David SK, et al. Prevalence and factors associated with alcohol consumption among persons with diabetes in Kampala, Uganda: a cross sectional study. BMC Public Health 2021; 21(1): 719.
- Liu L, Wang Y, Lam K, et al. Moderate Wine Consumption in the Prevention of Metabolic Syndrome and its Related Medical Complications. *EMIDDT* 2008; 8(2): 89–98.
- Koppes LLJ, Dekker JM, Hendriks HFJ, et al. Moderate Alcohol Consumption Lowers the Risk of Type 2 Diabetes. *Diabetes Care* 2005; 28(3): 719–725.
- 68. Athyros VG, Liberopoulos EN, Mikhailidis DP, et al. Association of Drinking Pattern and Alcohol Beverage Type With the Prevalence of Metabolic Syndrome, Diabetes, Coronary Heart Disease, Stroke, and Peripheral Arterial Disease in a Mediterranean Cohort. *Angiology* 2007; 58(6): 689–697.
- Thapsuwan S, Phulkerd S, Chamratrithirong A, et al. Relationship between consumption of high fat, sugar or sodium (HFSS) food and obesity and non-communicable diseases.
 BMJNPH 2024; 7(1): 78–87.
- Gbaja-Biamila TA, Odubela OO, Salako AO, et al. Dyslipidaemia among antiretroviral therapy-experienced adolescents and young people living with HIV in Lagos, Nigeria. *IRJMBS*; 6(3): 20–27.
- 71. Ige OO, Yilgwan CS, Ebonyi AO, et al. Serum lipid and glucose profiles in HIV-positive Nigerian children. *J Virus Erad* 2017; 3(3): 157–162.
- Innes S, Abdullah KL, Haubrich R, et al. High Prevalence of Dyslipidemia and Insulin Resistance in HIV-infected Prepubertal African Children on Antiretroviral Therapy. *Pediatric Infectious Disease Journal* 2016; 35(1): e1-e7.
- 73. Nampijja D. Dyslipidemia and its Correlates among HIV Infected Children on HAART Attending Mbarara Regional Referral Hospital. *ICPJL* 2017; 4(3). Available from: https://medcraveonline.com/ICPJL/dyslipidemia-and-its-correlates-among-hiv-infected-children-on-haart-attending-mbarara-regional-referral-hospital.html
- Tadesse BT, Foster BA, Chala A, et al. HIV and cART-Associated Dyslipidemia Among HIV-Infected Children. *JCM* 2019; 8(4): 430.
- Viljoen E, MacDougall C, Mathibe M, et al. Dyslipidaemia among HIV-infected children on antiretroviral therapy in Garankuwa, Pretoria. South African Journal of Clinical Nutrition 2020; 33(3): 86–93.

 Mandal A, Mukherjee A, Lakshmy R, et al. Dyslipidemia in HIV Infected Children Receiving Highly Active Antiretroviral Therapy. *Indian J Pediatr* 2016; 83(3): 226–231.

- Welsh JA, Sharma A, Cunningham SA, et al. Consumption of Added Sugars and Indicators of Cardiovascular Disease Risk Among US Adolescents. *Circulation* 2011; 123(3): 249–257.
- Hert KA, Fisk PS, Rhee YS, et al. Decreased consumption of sugar-sweetened beverages improved selected biomarkers of chronic disease risk among US adults: 1999 to 2010. *Nutrition Research* 2014; 34(1): 58–65.
- 79. Haslam DE, Peloso GM, Herman MA, et al. Beverage Consumption and Longitudinal Changes in Lipoprotein

- Concentrations and Incident Dyslipidemia in US Adults: The Framingham Heart Study. *JAHA* 2020; 9(5): e014083.
- 80. Malik VS and Hu FB. Sugar-Sweetened Beverages and Cardiometabolic Health: An Update of the Evidence. *Nutrients* 2019; 11(8): 1840.
- 81. Van Rompay MI, McKeown NM, Goodman E, et al. Sugar-Sweetened Beverage Intake Is Positively Associated with Baseline Triglyceride Concentrations, and Changes in Intake Are Inversely Associated with Changes in HDL Cholesterol over 12 Months in a Multi-Ethnic Sample of Children. *The Journal of Nutrition* 2015; 145(10): 2389–2395.