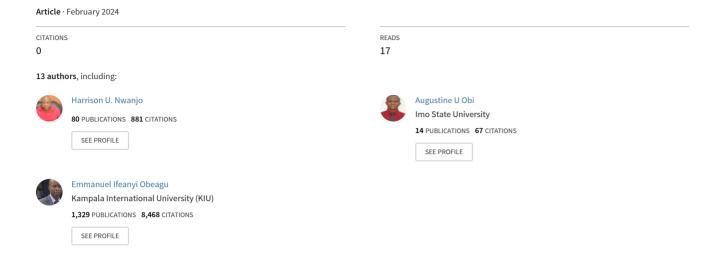
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EVALUATION OF LIPID PROFILE OF PATIENTS WITH DIABETES ATTENDING ABIA STATE UNIVERSITY TEACHING HOSPITAL ABA, NIGERIA

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ABSTRACT

The evaluation of lipid profile in diabetic patients attending Abia State Teaching Hospital Aba, was carried out. Venous blood samples were collected from 150 participants who gave consent. This comprises 100 diabetic patients as test and 50 healthy subjects as control. Cholesterol, Triglyceride, High density lipoprotein and Lowdensity lipoprotein were determined using semi-auto analyzer (Rato 9200). Data from this study were analyzed using statistical package for the social sciences (SPSS). Result shows that, serum cholesterol (4.96±0.91mmol/l), triglyceride (1.84±0.42mmol/l), low density lipoprotein(2.40±0.66mmol/l) levels were higher in study population compared with control (4.48±0.41mmol/l, 1.63±0.09mmol/l, 1.67±0.22mmol/l, 176.18±25.26x109/l, 11.85±0.63secs, respectively) (p<0.05 in each case). From the findings, management of conditions related to cardiovascular disease, artherosclerotic disease, anemia and stress in diabetics may benefit patients if lipid profile, and some hematological parameters are included as part of their routine laboratory investigations.

KEYWORDS: Lipid Profile, Diabetes mellitus, cardiovascular disease, artherosclerotic disease.

INTRODUCTION

Diabetes mellitus (DM), is a group of metabolic disorders in which there is high blood sugar level over a prolonged period and it is commonly referred to as diabetes. [1-6] Frequent urination, increased thirst, and increased hunger are symptoms of high blood sugar. [7-12] Many complications are resulted as a cause of untreated diabetes. [13-18] Diabetic ketoacidosis, hyperosmolar hyperglycemic state, or death are as a result of acute complications. However, cardiovascular disease, stroke, chronic kidney disease, foot ulcers, and damage to the eyes are included as long-term complication. [19-24]

Diabetes mellitus results from either the pancreas is not generating enough insulin or the cells of the body do not respond appropriately to the insulin produced. [19]

Reduced HDL cholesterol, a predominance of small dense LDL particles, and elevated triglyceride levels are abnormalities associated with a clustering of interrelated plasma lipid and lipoprotein in Insulin resistance and form 2 diabetes.^[1] Every single of these dyslipidemic

structures is related with an amplified risk of cardiovascular disease. Increased hepatic secretion of large triglyceride-rich VLDL and impaired clearance of VLDL appears to be of central importance in the pathophysiology of this dyslipidemia. Small dense LDL particles arise from the intravascular processing of specific larger VLDL precursors. Typically, reduced plasma HDL levels in type 2 diabetes are manifest as reductions in the HDL2b subspecies and relative or absolute increases in smaller denser HDL3b and HDL3c. Although behavioral interventions such as diet and exercise can improve diabetic dyslipidemia, for most patients, pharmacological therapy is needed to reach treatment goals.

All the abnormalities occur in many patients despite normal LDL cholesterol levels. These changes are also a feature of the insulin resistance syndrome (also known as the metabolic syndrome), which underlies many cases of type 2 diabetes. In fact, pre-diabetic individuals often exhibit an atherogenic pattern of risk factors that includes higher levels of total cholesterol, LDL

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cholesterol, and triglycerides and lower levels of HDL cholesterol than individuals who do not develop diabetes. Insulin resistance has striking effects on lipoprotein size and subclass particle concentrations for VLDL, LDL, and HDL.

MATERIALS AND METHODS STUDY AREA

The study was carried out at Abia State University Teaching Hospital (ABSUTH), Aba city in Abia state, South East of Nigeria..

STUDY POPULATION

The size of population was calculated using the method of Aroye 2004 with the formula $n=(z^2pq)/d^2$), and one hundred and fifty(150) subjects were recruited into the study. This comprises fifty(50)non-diabetic subjects as control and one hundred(100) diabetic subjects as test subjects.

SELECTION CRITERIA

INCLUSION: Those selected are:

- (i) Male and Female subjects of age 18 years to 74 years,
- (ii) Diabetic patients with blood sugar 10mmol/l and above.
- (iii) The subjects that gave their consent.

EXCLUSION;The excluded subjects are;

- (i) Male and Female subjects below the age of 18 years
- (ii) Subjects of blood sugar below 10mmol/l
- (iii)The subjects that did not give consent.

LABORATORY PROCEDURES

The reagents were commercially purchased and the manufacturers' standard operating procedures(S.O.P) were strictly adhered to.

DETERMINATION OF CHOLESTEROL: (enzymatic endpoint method) of Turfitt 1994.

Assay procedure; 1000µl of cholesterol liquor and 10µl of sample, standard and control were added into test tubes respectively, and incubated for 10 minutes at 37°c. The absorbance of standard and test were read(measured)

against reagent blank within 60 minutes at 500nm wavelength.

Conc. of cholesterol=<u>Abs. of test</u> x Conc. of Std. Abs. of Std.

Abs=absorbance Conc.=concentration Std=standard.

DETERMINATION OF TRIGLYCERIDE (enzymatic colorimetric method) of Trinder [25]. Assay procedure:

 $1000\mu l$ of triglyceride liquor and $10\mu l$ of sample, standard and control were added into test tubes respectively, and incubated for 10 minutes at $37^{\circ}c$. The absorbance of standard and test were read(measured) against reagent blank within 60 minutes at 500 nm wavelength.

Conc. of triglyceride = Abs. of test x Conc. of Std.

Abs. of Std.

Abs=absorbance Conc.=concentration Std=standard.

DETERMINATION OF HIGH-DENSITY LIPOPROTEIN(HDL)

ASSAY PROCEDURE; 100µl of precipitating reagent and 100µl of sample were added into dry test-tube and incubated at room temperature for 5mins. The solution was centrifuged at 3000 rpm (revolution per minute). 50µl of the sample supernatant, standard solution and distilled water (as blank solution) were added into 1000µl of cholesterol reagent respectively. The solutions were incubated for 15mins at 25°c. The absorbance was read at 505nm against the blank within 60mins.

Concentration of HDL cholesterol(mg/dl)=<u>Abs.T</u> x Conc. Std x2 Abs.Std

CALCULATION OF LDL (Freidewald's Formula)

Total cholesterol-(<u>triglyceride</u>) – HDL Cholesterol

RESULTS

Table 1: Mean \pm SD values of lipid profile of the studied population.

Parameters	Diabetic patients (n=100)	Controls (n=50)	t-value	p-value
Cholesterol(mmol/l)	4.96±0.91	4.48±0.41	3.584	0.001
Lowe r95% C.I	0.21	0.27		
Upper 95% C.I	0.75	0.69		
Triglyceride(mmol/l)	1.84±0.42	1.63±0.09	3.420	0.001
Lowe r95% C.I	0.08	0.12		
Upper 95% C.I High density	0.33	0.29		
Lipoprotein(mmol/l)	1.63±0.43	1.99±0.32	-5.204	0.001
Lowe r95% C.I	- 0.49	- 0.48		
Upper 95% C.I	- 0.22	- 0.23		
Low density	2.40±0.66	1.67±0.22	7.574	0.001

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Lipoprotein(mmol/l)			
Lowe r95% C.I	0.54	0.58	
Upper 95% C.I	0.92	0.88	
Atherogenic index	0.053±0.01	-0.087±0.72	0.001

Values of serum lipid profile levels of the studied population

The serum Cholesterol, Triglyceride and low-density lipoprotein were statistically higher (p=<0.001, p=0.001, p=<0.001 respectively) in the studied population compared with controls. The serum High density lipoprotein was statistically lower(p=<0.001) in the studied population compared with control. Athrogenic index was statistically higher(p=0.001) in the studied population compared with control.

DISCUSSION

In this present work, the serum Cholesterol, Triglyceride, Low density lipoprotein, blood platelet and prothrombin time levels were higher in studied population compared with controls. There was higher level of hemoglobin in male studied population compared with female studied population. The higher levels of Cholesterol, Triglyceride and Low-density lipoprotein, could possibly be due to altered metabolism of triglyceride-rich lipoproteins which is crucial in the pathophysiology of the atherogenic dyslipidemia of diabetes. Alterations include both increased hepatic secretion of Very Low-Density Lipoprotein (VLDL) and impaired clearance of VLDL and intestinally derived chylomicrons. An important consequence of retarded clearance is prolonged plasma retention of both VLDL and postprandial chylomicrons as partially lipolyzed remnant particles.

CONCLUSION

From the study since cholesterol, triglyceride, low density lipoprotein was higher in diabetic patients and are associated with risk factors as coronary artery disease, cardiovascular disease, artherosclerotic diseases, atherothrombotic complications and deficiency of vitamin k, care has to be taken in managing the disease.

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