

Assessment of Cardiovascular Disease Risk Factors of Adult Male and Female Diabetic Patients Attending Nnamdi Azikiwe University Teaching Hospital, Nnewi, Anambra State, Nigeria.

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Abstract: Diabetes mellitus is a prime risk factor for cardiovascular disease and is a major cause of death in both developing and developed countries.

Objective: The study was carried out to assess the cardiovascular risk factors and to predict the gender that may likely die of cardiovascular disease in the next ten (10) years in Nnewi, Anambra State, Nigeria.

Method: A total of 50 adult diabetic subjects comprised of 31 males and 19 females, who attend Nnamdi Azikiwe University Teaching Hospital (NAUTH) Nnewi, took part in this study. The study design was a case study. Ethical approval was obtained from the Ethics Review Committee and written informed consent was obtained from each participant. Also, questionnaire was used to obtain their biodata and the risk levels of both modifiable and non-modifiable factors. Fasting blood samples were drawn from the subjects for biochemical assays. A Framingham cardiovascular risk score was used to determine the risk of the individuals to cardiovascular disease (CVD). Student t-test was used for data analysis.

Results: The age (years) of males was the same with that of the females at $p > 0.05$. The mean Systolic blood pressure (SBP) and diastolic blood pressure (DBP) values were significantly higher in male subjects compared to the female subjects ($p < 0.05$). The mean FBS was not significantly different between both genders ($p > 0.05$). The serum levels of high density lipoprotein (HDL) and total cholesterol (TC) were not significantly different both genders ($p > 0.05$). The Framingham cardiovascular risk score was significantly higher in male subjects than in female subjects ($p < 0.05$).

Conclusion: The findings observed a greater risk of death from cardiovascular events within ten years in men than in women of the same age bracket.

Keywords: Cardiovascular disease, Framingham risk calculator, gender.

1. INTRODUCTION

Heart disease is a general term for a variety of heart conditions; it includes coronary diseases such as angina and myocardial infarction also known as heart attack.¹ There are other examples of CVD such as stroke, hypertensive heart disease, rheumatic heart disease, cardiomyopathy, heart arrhythmia and venous thrombosis^{1,2}, with each disease having

its underlying mechanism. Coronary artery disease, stroke and peripheral artery disease are linked to atherosclerosis, which may be induced by high blood pressure, smoking, diabetes, physical inactivity, obesity, high blood cholesterol, poor diet and excessive alcohol intake.¹

CVD is the leading cause of mortality globally.¹ Coronary artery disease and stroke account for 80 % of CVD death in males and 75 % of CVD deaths in females.¹ Most CVD affects older adults. Studies have it that 11 % of people between 20 and 40 years have CVD 37 % of people between 40 and 60 years have CVD, 71 % of people between 60 and 80 have CVD and 85 % of people over 80 years have CVD in developed world.³ But the average age of death in developing world is around 68 %.⁴ Report has it that Africa has witnessed tremendous lifestyle changes and urbanization; which increased the incidence of CVD.⁵

Atherosclerosis and hypertension are the most common cause of about eighty percent of the coronary heart disease and the cardiovascular disease cases are as a result of behavioral risk factors.⁶ The most important behavioral risk factors are unhealthy diet, physical inactivity and smoking which lead to effects such as increased blood pressure, increased blood sugar levels, increased blood lipid levels, overweight and obesity.⁶ Other risk factors not preventable are increasing age, male gender, family history and genetic disposition.^{7,8} Deaths from cardiovascular events have been reported to progressed with age in both genders.⁹ The disease onset is 7 to 10 years in men than in women.¹⁰

It has been reported that 90 % of CVD is preventable.¹¹ A diet high in fruits and vegetables intake and low intake of sweets, red meat and trans- fatty acid^{12, 13} have been shown to reduce blood pressure¹⁴ and improve metabolic syndrome.¹⁵ Moderate evidence has it that high salt intake increases cardiovascular mortality.¹⁶

The Framingham risk score is a gender specific algorithm used to estimate the ten years cardiovascular risk of an individual.^{17, 18} The Framingham risk score gives the result of the patient's risk score and predicts the risk of death due to cardiovascular disease by comparing the values to an average value for a person of the same sex and age.¹⁹ Therefore, as a result of the increase number of people suffering from hypertension and diabetes, it is pertinent to urgently assess the risk of death from cardiovascular disease in individual living in Nnewi, hence the aim of this study.

2. MATERIALS AND METHODS

A total of 50 adults diabetic subjects (31 males and 19 females) aged 35 years and above, who attended NAUTH, Nnewi Anambra State were randomly recruited for this study. Questionnaire was used to collect individual Biodata and lifestyle status which were used in the Framingham risk calculator.

5ml of venous blood sample was collected from each subject. FBS was determined by glucose oxidase system using the one-touch glucose monitoring system (Life Scan Inc, Johnson- Johnson Company, USA)²⁰ and diabetes defined using WHO recommendations.²¹

The remaining blood sample (4ml) was dispensed into a plain tube, allowed to clot, centrifuged and the serum dispensed into plain tube for lipid profile assay. Total cholesterol was estimated by enzymatic hydrolysis and oxidation of cholesterol by the method of Allain *et al*,²² Serum, HDL was estimated by precipitation method described by Assmann *et al*,²³ Serum Triglyceride was estimated by enzymatic hydrolysis by the method of Buccolo and David²⁴, HDL was estimated by precipitation method described by Assmann *et al*,²⁵ and LDL was estimated and was calculated as described by Kaplan *et al*,²⁷ The reagents used were prepared by Randox Laboratory Ltd., UK.

High blood pressure was defined according to the WHO guideline: a SBP \geq 140mm/Hg or DBP \geq 90mm/Hg or being on treatment.²⁶ The SBP and DBP of the subjects were measured and recorded accordingly using sphyngnomanometer.

The data collected from this study were keyed into the risk score Framingham calculator, which provides coronary heart disease risk at 10 years in percent.²⁷ The statistical analysis was done using SPSS version 16. Data collected were expressed as mean \pm SD and subjected to student unpaired t- test. The acceptable level of significance was $p < 0.05$.

3. RESULTS

Table-1 shows the mean risk factors studied. The mean age (years) of males was not significantly different from that of the females at $p > 0.05$. The mean SBP, DBP and the serum levels of triglycerides and LDL were significantly higher in

male subjects than in female subjects ($p < 0.05$) respectively. There were no significant different ($p > 0.05$) in the mean FBS, total cholesterol and HDL between the genders, although the male subjects had higher serum values. The Framingham cardiovascular risk score was significantly higher in male subjects than in female subjects ($p < 0.05$).

Table-1: Means of Framingham Cardiovascular Risk Factors

Risk Factors	Males (n=19)	Females (n=31)	p- value
Framingham Score (%)	23.94 ± 8.72	15.45 ± 6.95	<0.05
Age (Years)	56.68 ± 7.42	54.84 ± 8.34	>0.05
Total chol (mmol/l)	4.51 ± 1.34	4.45 ± 1.45	>0.05
HDL (mmol/l)	0.94 ± 0.44	0.87 ± 0.46	>0.05
FBS (mmol/l)	6.27 ± 2.39	6.92 ± 3.44	>0.05
SBP (mmHg)	139.05 ± 6.12	135.51 ± 5.80	<0.05
DBP (mmHg)	81.74 ± 13.30	74.87 ± 13.42	>0.05
TG (mmol/l)	1.36 ± 0.45	1.29 ± 0.05	<0.05
LDL (mmol/l)	2.50 ± 1.08	2.30 ± 0.98	<0.05

4. DISCUSSION

The study gives clue on the risk of death from cardiovascular causes in the next ten years of individuals with some cardiovascular risk factors. The Framingham cardiovascular risk score used in this study were based on individual age, gender, total cholesterol, HDL- cholesterol, smoking status, systolic blood pressure and state of any medication to treat high blood pressure. The information was aided by questionnaire. The DBP was not used in the calculation of cardiovascular risk score. This is because it does not independently predict cardiovascular risk.²⁶

The findings from this study revealed that SBP value was significantly higher in men than in women. This confirmed the work done by some researchers^{9, 28, 29} Reports have it that men had a higher risk of dying from cardiovascular causes than women of similar age range.⁹ The mechanism underlying the differential age effect is not well understood.³⁰ The age of both genders was the same in this study. Age is among the important factor in developing cardiovascular disease as one aged.³¹ It is associated with loss of arterial elasticity and reduced arterial compliance which may lead to coronary artery disease.³

In this study, the Framingham cardiovascular risk score was significantly higher in male subjects than in female subjects after the computation of risk factors obtained in the study. Ezeanyika *et al*,⁹ reported that men are likely to die of cardiovascular event compared to females of similar age range with high blood pressure in the next five years using a risk score calculator.

5. CONCLUSION

The study revealed that male diabetics may be more prone to cardiovascular events than none diabetics due to higher increase in cardiovascular risk factors such as increase in percentage Framingham risk score, age, SBP, LDL and triglycerides. Therefore, it is needful for individuals as they are ageing gracefully should modify lifestyles and habits that may predispose them to cardiovascular disease and improve longevity with quality of life.

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REFERENCES

- [1] World Health Organization (WHO). *Global Atlas on Cardiovascular Disease Prevention and Control* (PDF). World Health Organization in collaboration with the World Heart Federation and the World Stroke Organization) 2011; pp. 3–18. ISBN 978924156437-3.
- [2] Global Burden of Disease (GBD). Mortality and Causes of Death, Collaborators (17 December 2014). Global, regional, and national age-sex specific all-cause and cause-specific mortality for 240 causes of death, 1990-2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet* 2013; 385 (9963): 117–171.
- [3] Go A.S, Mozaffarian D, Roger V.L, Benjamin E.J, Berry J.D, Borden W.B, Bravata D.M. Heart disease and stroke statistics 2013 update: a report from the American Heart Association. *Circulation* 2013; 127 (1): e6- e245.
- [4] Fuster, Board on Global Health; Valentin; Academies, Bridget B. Kelly. Institute of Medicine of the National, eds. *Promoting cardiovascular health in the developing world: a critical challenge to achieve global health*. Washington, D.C.: National Academies Press. pp. Chapter 2, 2010 ; ISBN 9780309147743.
- [5] Kabiri N, Asgary S, Madani H, Mahzouni P. Effects of *Amaranthus caudatus* extract And lovastatin an atherosclerosis in hypercholesterolemic rabbits. *Journal of Medicinal plants Respiration* 2010; 4 (5): 355 -381.
- [6] Guldiken S, Demir M, Anka E, Tugrul, A. The level of serum high sensitive C - reactive protein in women with hyperthyroidism. *Journal of Clinical Endocrinology and Metabolism* 2005;3: 85-88.
- [7] Micha R, Michas G, Mozaffarian D. Unprocessed and processed meats and risk of coronary artery disease and type 2 Diabetes- an updated review of the evidence. *Current atherosclerosis reports* 2012; 14 (6): 515-524.
- [8] Finks SW, Airee A, Chow SL, Macaulay TE, Moranville MP, Rogers KC, Trujillo TC. Key articles of dietary interventions that influence cardiovascular mortality. *Pharmacotherap* 2012; 32 (4): e54-87.
- [9] Ezeanyika L.U.S, Ugwu C.E, Nwanguma B.C, Onah L.E, Ojobo C, Abba V, Okpanachi G. Assessment of cardiovascular disease risk factors of an urban Nigerian hypertensive population using a risk score calculator. *Pakistan journal of Medical Sciences* 2008; 24 (3) 390 – 394.
- [10] Mendis, Shanthi, Puska, Pekka; Norrving, Bo. *Global atlas on cardiovascular disease prevention and control* (1 ed.). Geneva: World Health Organization in collaboration with the World Heart Federation and the World Stroke Organization 2011; p. 48. ISBN 9789241564373.
- [11] McGill HC, McMahan CA, Gidding SS. Preventing heart disease in the 21st century: implications of the Pathobiological Determinants of Atherosclerosis in Youth (PDAY) study. *Circulation* 2008; 117 (9): 1216–1227.
- [12] Willett, WC. Dietary fats and coronary heart disease. *Journal of internal medicine* 2012; 272 (1): 13–24.
- [13] Chowdhury, Rajiv; Warnakula, Samantha; Kunutsor, Setor; Crowe, Francesca; Ward, Heather A.; Johnson, Laura; Franco, Oscar H.; Butterworth, Adam S.; Forouhi, Nita G.; Thompson, Simon G.; Khaw, Kay-Tee; Mozaffarian, Dariush; Danesh, John; Di Angelantonio, Emanuele. Association of Dietary, Circulating, and Supplement Fatty Acids With Coronary Risk. *Annals of Internal Medicine*, 2013; 160 (6): 398–406.
- [14] Sacks FM, Svetkey LP, Vollmer WM, et al. Effects on blood pressure of reduced dietary sodium and the Dietary Approaches to Stop Hypertension (DASH) diet. DASH-Sodium Collaborative Research Group. *New England Journal of Medicine* 2001; 344 (1): 3–10.
- [15] Azadbakht L, Mirmiran P, Esmailzadeh A, Azizi T, Azizi F. Beneficial effects of a Dietary Approaches to Stop Hypertension eating plan on features of the metabolic syndrome. *Diabetes Care* 2005; 28 (12): 2823–31.
- [16] Bochud, M; Marques-Vidal, P; Burnier, M; Paccaud, F. *Dietary Salt Intake and Cardiovascular Disease: Summarizing the Evidence*. *Public Health Reviews* 2012; 33: 530–552.
- [17] Wilson P.W, D’Agostino R.B, Levey D, Belanger A.M, Silbershatz H, Kannel W.B. Predictions of coronary heart disease using risk factors categories. *Circulation* 1998; 97 (18): 1837 – 1847.

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- [18] D'Agostino R.B, Vasan R.S, Pencina M.J, Wolf P.A, Cabain M, Massaro J.M, Kannel W.B. General cardiovascular risk profile for use in primary care: the Framingham Heart study. *Circulation* 2008; 117 (6): 743 – 753.
- [19] Pocock S.J, McCormack V, Gueyffier F, Boutitie F, Fagard R, Boisel. A score for predicting risk of death from cardiovascular disease in adults with raised blood pressure based on individual patients data from randomized controlled trials. *British medical journal* 2001; 323: 75 – 81.
- [20] Mark S.V, Dawson A. Rapid stick methods for determining of blood glucose concentration. *British Medical Journal* 1965;1: 293.
- [21] World Health Organization (WHO). Part 1: Diagnosis and Classification of Diabetes mellitus, Report of WHO consultation. Geneva: WHO 1999.
- [22] Alain B. Leucocyte typing: Human leucocyte differentiation antigens detected by monoclonal antibodies: Specification, classification, nomenclature. Report on the first international references workshop sponsored by INSERM, WHO and IUIS, Berlin. *Springer* 1984; 45-48.
- [23] Assmann G, Jabs H U, Nolte W, Schriewer H. LDL- cholesterol determination in blood serum following precipitation of LDL with polyvinyl sulphate. *Clinica Chimica Acta* 1984; 140: 77 -88.
- [24] Buccolo G and David H. Quantitative determination of serum triglycerides by the use of Enzymes. *Journal of Clinical Chemistry* 1973; 19: 476-482.
- [25] Kaplan A, Szabo L, Opheim K.. Clinical chemistry interpretation and techniques. 3rd ed. Published by Lea and Febiger at 600 Washington Square Philadelphia, USA 1983; pp 307 – 316.
- [26] World Health Organization (WHO). International society of Hypertension (ISH). Statement on management of Hypertension. *Journal of Hypertension* 2003; 21: 1983 – 1992.
- [27] Aberg JA. Lipid Management in Patients Who Have HIV and Are Receiving HIV Therapy. *Endocrinology and metabolism clinics of North America* 2009; 38 (1): 207 – 222.
- [28] Kannel W.B. Historic perspectives on the relative contributions of diastolic/systolic blood pressure elevation to cardiovascular risk profile. *American Heart Journal* 1999; 138: 205 – 210.
- [29] Shapo L, Pomerleau J, M-ckee M. Epidemiology of hypertension and associated cardiovascular risk factors in a country in transition; A population based survey in Tirana city, Albania. *Epidemiology of Community Health* 2003; 57: 734 – 739.
- [30] Mufunda J, Body mass index and blood pressure: where are we?. *Journal of Human Hypertension* 2007; 21: 5 – 7.
- [31] Mosca L, Manson J.E, Suterland S.E, Langer R.D, Menolio T, Barreitt-connor E. Cardiovascular disease in women: a statement of health care professional from the American Heart Association *Circulation* 1997; 96: 2468- 2482.
- [32] Finegold J.A and Ascaria P, F. Mortality from ischaemic heart disease by country region and age statistics from world Health Organisation and United Nation. *International Journal of Cardiology* 2002; 168 (2): 934 - 945.