

Original Research Article

Evaluation of serum sex hormones and CD₄⁺ count among HIV patients on HAART, HAART naïve patients and apparently healthy subjects in Sokoto, Nigeria

Armiya'u Ahmed Yelwa^{1*}, Abdallah Suleiman Mainasara¹, Shehu Abubakar Akuyam²,
Balarabe Isah Adamu³, Zayyanu Usman Umar⁴, Sharafudeen Dahiru Abubakar²,
Yahaya Muhammad⁵, Solomon Matthias Gamde⁶, Aisha Abdulazziz⁶

¹Department of Chemical Pathology, Usmanu Danfodiyo University Sokoto, Nigeria

²Department of Medical Laboratory Science, Ahmadu Bello University, Zaria, Kaduna State, Nigeria

³Department of Community Medicine, College of Health Sciences Usmanu Danfodiyo University Sokoto, Nigeria

⁴Department of Physiology, College of Health Sciences Usmanu Danfodiyo University Sokoto, Nigeria

⁵Department of Chemical Pathology, Rasheed Shekoni Teaching Hospital Dutse, Jigawa, Nigeria

⁶Department of Histopathology, College of Health Sciences, Usmanu Danfodiyo University, Sokoto State, Nigeria

Received: 01 January 2020

Accepted: 03 February 2020

*Correspondence:

Dr. Armiya'u Ahmed Yelwa,

E-mail: ahmedarmiyau@yahoo.com

Copyright: © the author(s), publisher and licensee Medip Academy. This is an open-access article distributed under the terms of the Creative Commons Attribution Non-Commercial License, which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

ABSTRACT

Background: Acquired Immunodeficiency Syndrome (AIDS) is a chronic disease associated with Human Immunodeficiency Virus (HIV) which progressively induces depletion of CD₄⁺ T cells, and increased vulnerability to opportunistic infections. Previous reported studies associated HIV-infected men with sexual dysfunction; hypogonadism is the most common endocrinological disorders. Its prevalence remains poorly defined and widely ranging from different studies.

Methods: This study evaluated a total of 135 serum sex hormones (testosterone, estrogen, follicle stimulating hormone and luteinizing hormone) and its correlation with CD₄⁺ counts among HIV patients on HAART, HAART naïve patients and negative control subjects (n=45). CD₄⁺ cell counts were estimated using standard flow cytometry method and serum sex hormones by competitive enzyme immunoassay technique.

Results: There were significantly lower testosterone and CD₄⁺ levels (p<0.05) among HIV positive HAART naïve men compared to negative control. LH and FSH indicated significant increased (p<0.05) among HIV positive men on HAART.

Conclusions: Antiretroviral therapy improves sexual functions in males infected with human immunodeficiency virus. Hence, further study to evaluate its effects on other sexual behaviors.

Keywords: Acquired Immunodeficiency syndrome, Antiretroviral therapy, Antiretroviral naïve patients, Sex hormones

INTRODUCTION

Nigeria has the second largest Human Immunodeficiency Virus (HIV) epidemic in the world.¹ In accordance with its population size, 1.9 million Nigerians were living with

HIV in 2018.² HIV belongs to the family of retrovirus that causes Acquired immunodeficiency syndrome (AIDS). A chronic viral disease associated with the progressively induced depletion of CD₄⁺ T cells and increased vulnerability to opportunistic infections.³ HIV

infection is associated with functional derangement of virtually every endocrine organ system of human body.⁴ Functional derangement of sex hormones may result either from primary testicular failure or inadequate signaling from the pituitary or hypothalamus.⁵ High prevalence of hypo-gonadism was reported in several studies conducted in HIV infected males during pre-antiretroviral therapy.^{3,6}

Highly active antiretroviral therapy (HAART) has been extremely successful in suppressing HIV infection restoring immune function.⁷ However, HAART has also been implicated to cause sexual dysfunction indirectly affecting people living with HIV/AIDS (PLWHAs) and their inability to adhere to antiretroviral therapy.⁸ Sexuality is an intrinsic part of person's wellbeing, knowledge gaps exist on our understanding of issues of sexuality outside the risky behaviors paradigm among PLWHA on HAART in Sokoto and Nigeria at large. Very few studies have been conducted in Nigeria on the prevalence of hypogonadism in HIV infected males. This research would explore sex profile hormones among HIV-infected men in Nigeria is poorly defined and evaluate it correlation to CD₄ count.

METHODS

This study was conducted from 3rd July, 2017 to March, 2018 at the Specialist Hospital Sokoto, Nigeria. Sokoto State is located at the extreme part of North-Western Nigeria between longitude 3 and 7o East and latitude 10 and 14o North of the equator. Sokoto shares borders with Niger Republic to the North, Kebbi State to the South-West and Zamfara State to the east.⁹ The state covers a total land area of about 32,000 square kilometers and a population of 4,602,298 million based on 2013 projection.¹⁰ Sokoto state has semi-arid climate and vegetation is largely Sudan Savannah with an annual rainfall of between 500 and 1300 mm and temperature range between 15 and over 40°C during warm days.¹⁰

Inclusion criteria

- HIV seropositive male aged 15-60 years presented with no clinical conditions likely to affect serum concentrations of sex hormones
- Apparently healthy male subjects as negative controls.

Exclusion criteria

- HIV-positive patients with history of concomitant comorbidities such as diabetes mellitus, chronic kidney disease (serum creatinine >1.5 mg%), chronic liver disease, past history of meningitis, stroke, cryptococcal infection and other related conditions.
- HIV-positive patients with established cases of sexual dysfunction and/or infertility before commencement of HAART therapy.

- HIV-positive patients with substance abuse opiates (including heroin and methadone) or marijuana
- HIV-positive patients who declined to give consent for inclusion.

A total of 135 male subjects aged 15-60 years were recruited for this study. These consisted of 45 HIV patients on HAART, 45 HAART-naïve HIV patients attending the ART Clinic in Specialist Hospital, Sokoto, Nigeria and 45 apparently healthy individuals (negative controls).

Ethical approval with registration number: SHS/SUB133/VOL.1 was obtained from the Ethical and Research Committee of Specialist Hospital Sokoto, Nigeria.

The sample size for the study was calculated using the formula below¹¹

$$n = \frac{z^2 p \times q}{d^2}$$

Where,

n = the desired sample when the population is greater than 10,000.

Z = the desired normal deviate, usually set at 1.96 which corresponds to the 95% confidence level.

P = the current prevalence rate of HIV in Sokoto which is 5.6%.¹²

q = 1-p

d = degree of accuracy desired, usually set at 0.05.

The calculated sample size was 81. However, 10% (≈9 patients) were added as attrition rate. Therefore, the final calculated sample size was 90.

Experimental design

A simple random sampling technique was used to recruit 45 HIV infected patients on HAART and 45 HIV infected patients attending Antiretroviral Therapy (ART) Clinic of Specialist Hospital Sokoto that were yet to commence HAART and 45 Negative Control Subjects were also selected from staff of Specialist Hospital Sokoto (Table 1).

Sample collection

About 5ml of venous blood sample was collected at the clinic using a sterile disposable syringe and needle. 4ml of the blood was transferred into plain tubes and allowed to clot at room temperature and then centrifuged at 4000 rpm for 5 minutes. The sera were harvested and placed into other plain tubes, stored at -20°C until the time of

analysis. The remaining 1 ml was transferred into a sterile EDTA specimen bottle and used for the estimation of CD4+ count within 3 hours of the blood collection.

Table 1: A cross sectional descriptive design of experimental groups and descriptions (n = 45).

Groups	Description of experimental groups
Group I	HIV infected patients on HAART
Group II	HAART-naïve HIV patients
Group III	Apparently healthy subjects (negative control)

Laboratory analysis

Partec, Germany flow cytometer was used to obtain CD4 T cell count.¹³ Free testosterone, estrogen, luteinizing hormone and follicle stimulating hormone were estimated using method of competitive enzyme immunoassay technique as described by.¹⁴

Principles of flow cytometer

Flow cytometer was used to obtain CD4 T cell count. In flow cytometry, cells are separated in aqueous suspension and stained with fluorescent dyes. Cells in flow cuvette are individually illuminated by excitation light source of the laser (488nm). This excitation causes dye molecules to fluorescence at characteristic color of emission. The fluorescent signals are then displayed and analyzed in histograms.¹³

Principles of enzyme immunoassay (determination of sex hormone profile)

Serum sex hormone was carried out using standard method of estimation of testosterone, estrogen, serum luteinizing hormone, and follicle stimulating hormone.¹⁵⁻¹⁸

Statistical analysis

The data generated were analyzed using Statistical Package for Social Sciences (SPSS) version 22.0. Serum sex hormones were analyzed and expressed as Mean±SEM. The results obtained were compared between different groups using ANOVA. A p-value of p < 0.05 was considered significant.

Table 4: Comparison of serum sex hormones and CD4+ count among male subjects.

Parameters	Group I	Group II	Group III
Testosterone (ng/ml)	3.40±0.70*	3.43±0.59*	10.34±0.56*
Oestrogen (pg/ml)	83.84±7.82	104.15±10.32	102.32±6.25
LH (MIU/ml)	10.79±2.93*	7.03±0.88*	3.94±0.54
FSH (MIU/ml)	12.05±2.86*	5.51±1.00	4.05±0.78
CD4+ (cell/mm3)	280.63±42.41*	245.79±45.48*	790.32±36.50*

Values are mean ± SEM, n= number of subjects, the values bearing asterisk differ significantly with the respective control at p <0.05 (*), using ANOVA. Group I= HIV-positive on HAART, Group II = HIV-positive HAART-naïve and Group III = HIV-negative control.

RESULTS

Majority of the HIV infected men (66.7%) were married while 25.2% were unmarried (Table 2). Among the tribes (Hausa, Fulani, Igbo and Yoruba), Hausa (73.3%) constitute the largest population of HIV infected men and Fulani (3.7%) the least in Sokoto metropolis (Table 2).

Table 2: Demographic characteristics of the study population.

Characteristics	n	Percentage (%)
Marital status		
Married	90	66.7
Single	34	25.2
Widowed	9	6.6
Divorce	2	1.5
Total	135	100
Ethnicity		
Hausa	99	73.3
Fulani	5	3.7
Yoruba	11	8.1
Igbo	14	10.5
Others	6	4.4
Total	135	100

Table 3: Age (years) and gender distribution of HIV-positive on HAART, HIV-positive, HAART-naïve and HIV-negative controls.

Group	Group I	Group II	Group III	Total number of groups
Age group(years)	Male	Male	Male	
15-24	12	5	2	39
25-34	5	4	7	41
35-44	5	5	8	32
45-54	1	6	5	17
55-64	0	3	1	6
Total	23	23	23	135

Group I= HIV on HAART, GROUP II= HIV-positive HAART-naïve, Group III= controls, HAART= highly active antiretroviral therapy, HIV= Human immune virus.

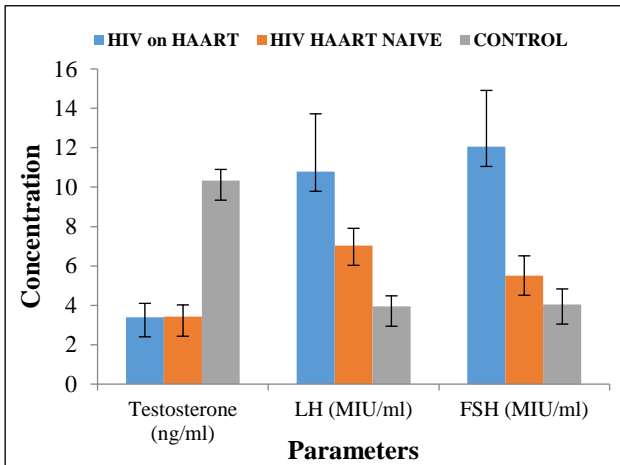


Figure 1: Comparison of serum sex hormone among male HIV on HAART, HIV HAART-Naive and controls.

Table 5: Correlation of CD4+ count with sex hormones among HIV positive men on HAART, HAART-naïve and HIV negative controls.

Parameters	r-value	p-value	Remark
Testosterone (ng/ml)	0.682**	<0.001	SS
Oestrogen (pg/ml)	0.025	0.140	NSS
LH (MIU/ml)	-0.181	0.025	SS
FSH (MIU/ml)	-0.271*	0.027	SS

**=correlation is significant at 0.01 level (2-tailed), LH= Luteinizing hormone, FSH= Follicular stimulating hormone.

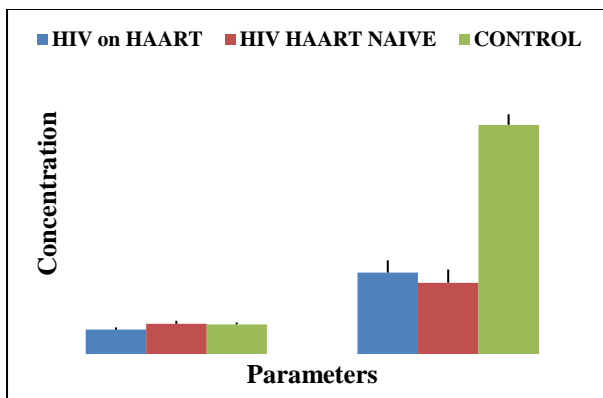


Figure 2: Comparison of serum sex hormone and CD4+ among male HIV on HAART, HIV HAART-naïve and controls.

HIV-infection in men is more prevalent among the age 45-54 years HAART naïve men and 15-24 years HIV patients on HAART (Table 3). There was significant decrease in testosterone among HIV infected men on HAART (3.40±0.70) and HAART naïve patients (3.43±0.59) compared with negative control (10.34±0.56) (p < 0.05, Table 4). There was significant increase in estrogen, follicle stimulating hormone and luteinizing hormones compared with the negative control (p < 0.05,

Table 4). However, these profiles were more elevated in patients on HAART compared to HAART naïve (Figure 1). The CD4+ count among HIV infected men on HAART (280.63±42.41) and HAART-naïve (245.79±45.48) were significantly decreased compared with negative control controls (790.32±36.50) (Figure 2). In addition, there was significant correlation between CD4 count and sex hormone profiles except estrogen (Table 5).

DISCUSSION

This study indicated a significant decrease in serum levels testosterone in HIV infected male on HAART and HAART naïve male compared with negative control (p<0.05). This finding is in agreement with previous reports.¹⁹ Hypogonadism is common among HIV infected male with incidence between 29-50% without HAART and 20-30% on HAART.²⁰

There were significant increase in sex hormone profile levels of FSH, LH and estrogen among HIV positive and HAART naïve males compared with negative control. These significant differences might be due to the relationship between LH, FSH, estrogen and the testosterone. LH and FSH regulate the secretion and release of testosterone.

The finding also shows significant decreased levels of CD4+ count among HIV on HAART and HAART naïve in male (p < 0.05).This is in agreement with previous report.²¹ HIV infection and its stages of progression decreased CD4+ count and have been reported to be associated with changes in serum sex hormones level.⁶

CONCLUSION

HIV is associated with changes in sex hormones which may lead to sexual dysfunction in infected individuals and probably antiretroviral therapy may improve sexual functions. Further study is needed to evaluate its effect on other sexual behavior.

ACKNOWLEDGEMENTS

Authors acknowledge the efforts of the management and Staff of Medical Laboratory Services, Specialist Hospital Sokoto, for the provision of ethical clearance, facilities and equipment.

Funding: No funding sources

Conflict of interest: None declared

Ethical approval: The study was approved by the Institutional Ethics Committee

REFERENCES

1. United Nation intervention for AIDS/HIV. AIDS info. Access 18 December 2019.

2. PEPFAR. Annual Report to Congress, 2019;1-106.
3. Aggarwal J, Taneja RS, Gupta PK, Wali M, Chitkara A, Jamal A. Sex hormone Profile in Human Immunodeficiency Virus-infected men and its correlation with CD4 cell counts. *Indian J Endocrinol Metab.* 2018;22:328-34.
4. Sinha U, Sengupta N, Mukhopadhyay P, Roy KS. Human immunodeficiency virus endocrinopathy. *Indian J. Endocrinol Metab.* 2011;15:251-60.
5. Mylonakis E, Koutkia P, Grinspoon S. Diagnosis and treatment of androgen deficiency in human immunodeficiency virus-infected men and women. *Clin Infect Dis.* 2001;33:857-64.
6. Dobs AS, Dempsey MA, Ladenson PW, Polk BF. Endocrine disorders in men infected with human immunodeficiency virus. *Am J Med.* 1988;84:611-6.
7. World Health Organization. Antiretroviral drugs for treating pregnant women and preventing HIV infections in infants: recommendations for a public health approach 2010. Accessed on 10 June 2015.
8. Panos G, Samonis G, Alexiou VG, Kavarnou GA, Charatsis G, Falagas ME. Mortality and morbidity of HIV infected patients receiving HAART: a cohort study. *Current HIV Research.* 2008;6(3):257-60.
9. Sokoto State Business Directory. A Publication of the Commerce Department, Ministry of Commerce, Industry and Tourism, Sokoto; 2007:14-18.
10. 10. United Nations Population Fund. Annual report. 605 Third Avenue New York, NY, 10158; 2013:1-56.
11. Cochran WG. Sampling techniques. 3rd edition. New York, John Wiley and Sons; 1977:122.
12. National Agency for the Control of Aids. National health reproductive survey. Available at <http://www.naca.gov.ng>. Accessed on 23rd March 2017.
13. Cassens U, Golde W, Kuling G, Schlenke P, Lehman LG, Traore Y, et al. Simplified volumetric flow cytometry allows feasible and accurate 55 determination of cd4 t-lymphocytes in immunodeficient patients worldwide. *Antiviral Therapy.* 2004;9:395-405.
14. Tietz NW. Clinical guide to laboratory tests. Testosterone kits book. 3rd ed. Philadelphia, PA: WB Saunders Co.; 1995:186-188.
15. Horton R, Tait JF. Androstenedione production and Interconversion rates measured in peripheral blood and studies on the possible site of conversion to testosterone. *J Clinic Investi.* 1996;45:310-03.
16. Abraham GE. The application of natural steroid radioimmunoassay to gynecologic endocrinology. *Clinic Endocrinol.* 1981;33:475-529.
17. Kosasa TS. Measurement of human luteinizing hormone. *J Reprod Med.* 1981;26:201-6.
18. Wennink JM, Delemarre VW, Schoemaker HA, Schoemaker R. Luteinizing hormone and follicle stimulating hormone secretion patterns in girls throughout puberty measured using highly sensitive immune radiometric assay. *Clinic Endocrinol.* 1990;33:333-44.
19. Goldmeier LH, Mackie D, Scullard NE. Antiretroviral therapy is associated with sexual dysfunction and with increased serum estradiol levels in men. *Int J STD AIDS.* 2004;15:234-7.
20. Crum NF, Furtek KJ, Olson PE, Amling CL, Wallace MR. A review of hypogonadism and erectile dysfunction among HIV-infected men during the pre- and post-HAART eras: diagnosis, pathogenesis, and management. *AIDS Patient Care STDS.* 2005;19:655-71.
21. Tindall B, Forde S, Goldstein D, Ross MW, Cooper DA. Sexual dysfunction in advanced HIV disease. *AIDS Care.* 2004;6:105-7.

Cite this article as: Yelwa AA, Mainasara AS, Akuyam SA, Adamu BI, Umar ZU, Abubakar SD, et al. Evaluation of serum sex hormones and CD4⁺ count among HIV patients on HAART, HAART naive patients and apparently healthy subjects in Sokoto, Nigeria. *Int J Res Med Sci* 2020;8:891-5.