

Prevalence of Hepatitis C Virus (HCV) seropositivity in a cohort of people living with HIV and AIDS in Abeokuta, Ogun State, Southwestern Nigeria

¹Ojo DA, ^{1&2}Akingbade OA, ³Okerentugba PO, and ³Okonko IO

¹Department of Microbiology, Federal University of Agriculture, Abeokuta, Ogun State, Nigeria

E-mail: daojo3@yahoo.com; Tel: +234803 392 8703

²Department of Microbiology, Federal Medical Centre Idi Aba, Abeokuta, Ogun State, Nigeria

³Medical Microbiology Unit, Department of Microbiology, University of Port Harcourt, P.M.B. 5323, Choba, East-West Road, Port Harcourt, Rivers State, Nigeria;

iheanyi.okonko@uniport.edu.ng; Tel.: +234 803 538 0891

Abstract: Viral hepatitis is also a major public health concern as hepatitis B virus (HBV) afflicts an estimated 350 million people, and hepatitis C virus (HCV) affects 150 million people worldwide. Therefore, this study was carried out to estimate the prevalence of HCV seropositivity in a cohort of people living with HIV and AIDS in Abeokuta, Ogun State, Southwestern Nigeria. In this cohort study, we collected blood samples from 183 HIV-positive patients between January 2012 and January 2013. Standard enzyme immunoassays were used to determine the serological prevalence of hepatitis C virus among HIV-positive individuals. Among the HIV positive individuals, we found that 43 patients were infected with hepatitis C virus. The overall HCV-HIV prevalence is 23.5%. The result showed a significance difference ($P < 0.05$) between HIV-infected subjects and subjects positive to HCV. A higher percentage of males (28.9%) were infected with HCV than females (19.0%). The highest HIV prevalence (53.3%) was observed in age groups 55 years and above. The findings of this present study have demonstrated that co-infection of HIV and HCV is on the increase in Nigeria. Thus, the phenomenon of HIV and HCV co-infection is a cause for concern.

[Ojo DA, Akingbade OA, Okerentugba PO, and Okonko IO. **Prevalence of Hepatitis C Virus (HCV) seropositivity in a cohort of people living with HIV and AIDS in Abeokuta, Ogun State, Southwestern Nigeria.** *N Y Sci J* 2013;6(7):19-23]. (ISSN: 1554-0200). <http://www.sciencepub.net/newyork>. 4

Keywords: AIDS, HIV, HCV, Cohort study, Prevalence, Nigeria

1. INTRODUCTION

In Nigeria, it is estimated that 3.6% of the population were living with the virus in 2009, and the country had the world's second highest number of HIV and Acquired Immune Deficiency Syndrome (AIDS) related deaths (220,000) after South Africa (CIA, 2009; Tremeau-Bravard et al., 2012). Viral hepatitis is also a major public health concern as hepatitis B virus (HBV) afflicts an estimated 350 million people, and hepatitis C virus (HCV) affects 150 million people worldwide (WHO, 2011; Tremeau-Bravard et al., 2012). Both viruses are endemic in sub-Saharan Africa where an estimated 75 million people (over 35 million in Nigeria) live with hepatitis C virus (Madhava et al., 2002; Modi and Feld, 2007; Tremeau-Bravard et al., 2012).

Transmission of HCV predominantly occurs parenterally as a result of blood transfusion and exposure to blood derivatives, and the disease was first recognized in recipients of blood and blood products such as factor VIII and immunoglobulins. Transplanted organs and needles stick injuries have also been implicated in transmission. Mc Lean *et al.* (1997) reported transmission in drug misusers and patients in dialysis and surgical units. Sexual contact has also been incriminated in the transmission of HCV

(Alter *et al.*, 1982). There is also a growing evidence of vertical transmission (mother to baby).

HCV has been shown to have a worldwide distribution, occurring among persons of all ages, genders, races and regions of the world (WHO, 1996). Recent report by the World Health Organization (WHO) estimated that 170 million persons, or about 3% of the world's population, are infected with HCV who are at risk of developing liver cirrhosis, cancer or both (WHO, 1996). Slightly different prevalence was reported from different regions of the world.

Co-infection with HCV increases the risk for hepatotoxicity of HAART and likelihood of onset of an AIDS-defining illness, compared with infection with HIV-1 alone (Greub 2000, Feld et al. 2005; Forbi et al., 2007). Although the HIV co-infection with HCV has been recognized worldwide in individuals exposed to blood-borne diseases, limited data are available on the extent of co-infection and effect of these viruses on the immune system in developing countries (Forbi et al., 2007). Nigeria belongs to the group of countries highly endemic for viral hepatitis (Odemuyiwa et al. 2001). Few studies have been done on HIV and HCV separately in Nigeria but the knowledge about the interrelationship between these viruses and their effect on the immune system remains unclear (Forbi et al., 2007).

A clearer picture of HIV and HCV prevalence in Africa is important in order to better educate the population, and control these epidemics. Studies are crucial and necessary to give us a better understanding of the epidemiology of the diseases in developing countries like Nigeria (Tremeau-Bravard et al., 2012). Although screening for HBV markers in patients was introduced over two decades ago, transfusion-associated hepatitis due to HCV has continued to occur (Egah *et al.*, 2004). Therefore, this study was carried out to estimate the prevalence of HCV seropositivity in a cohort of people living with HIV and AIDS in Abeokuta, Ogun State, Southwestern Nigeria.

2. MATERIALS AND METHODS

2.1. Study Area

This study was conducted at the Federal Medical Centre, Idi-Aba, in Abeokuta, the capital city of Ogun State, Southwest Nigeria between January 2012 and January 2013.

2.2. Study Population

A total of one hundred and eighty three HIV-infected individuals were enrolled in this study. Of which, 83 were males and 100 were females. Subject were confirmed HIV-infected, ages 15 years and above. The ethical approval was granted by the Ethical Review Committee of the hospital. Informed consent was obtained from each patients and relevant confidentiality was maintained throughout the study.

2.3. Sample Collection

One hundred and eighty-three (183) blood samples were collected for this study. Venous blood was obtained into non-anticoagulated tubes. The samples were centrifuged at 2000 resolution per minutes (rpm) for 5 minutes to obtain sera. The sera were stored at -20°C for serologic assay of HIV and HCV.

2.4. Serologic Assay

SD HIV-1 and HIV-2 screening kit (manufactured by Bioline Standard Diagnostic Inc, Korea) was used in this study. This is an immunochromatography (rapid) method for quantitative detection of antibodies of all isotopes (IgG, IgM, IgA) specific to HIV-1 and HIV-2 simultaneously in serum. Red colour in the control and patient windows indicated a positive result while presence of the red colour in the control and its absence in the patient window indicates a negative result.

Each serum sample was screened for antibodies to HCV using the membrane based immunoassay technique (ACON Laboratories, Inc. San Diego, USA). The test line region of the strip had been

pre-coated with recombinant HCV antigen. Both are based on chromatographics capillary migration to form colour line. The presence of the colour line indicated a positive result while the absence indicated a negative result.

2.5. Data Analysis

The study was carried out and the proportion of subjects with HIV and HCV status were calculated. The prevalence level of HIV and HCV were cross tabulated with age and sex. Relevant chi-square statistics were computed using SPSS 20.0 window packages to accompany each cross tabulation.

3. RESULTS ANALYSIS

One hundred and eighty-three (183) HIV-infected subjects were examined in this study (Table 1 and 2). Of which, 83 (45.4%) were males while 100 (54.6%) were females. Table 1 shows the prevalence of HCV amongst HIV-infected subjects in relation to sex. The sex-specific prevalence showed that males had higher prevalence of HCV (28.9%) than females with 19.0% prevalence for HCV. However, there was significant difference ($P < 0.05$) between sex and HCV seropositivity among HIV-infected subjects.

Table 1: Prevalence of HCV amongst HIV-Infected Subjects in relation to Sex

Sex	No. Tested (%)	No. Positive for HCV (%)
Males	83(45.4)	24 (28.9)
Females	100(54.6)	19 (19.0)
Total	183 (100.0)	43 (23.5)

The age-specific prevalence of HCV amongst HIV-infected subjects is shown in Table 2. It showed that HCV were higher in age group 55 years and above (53.3%) than other age groups. However, there was no significant difference ($P > 0.05$) between age and HCV seropositivity amongst HIV-infected subjects.

Table 2: Prevalence of HCV amongst HIV-Infected Subjects in relation to Age

Age (year)	Groups	No. Tested (%)	No. Positive for HCV (%)
15-24		33(18.0)	8 (24.2)
25-34		42(23.0)	6 (14.3)
35-44		64(35.0)	17 (26.6)
45-54		29(15.8)	4 (13.8)
55 and above		15(8.2)	8 (53.3)
Total		183 (100.0)	43 (23.5)

4. DISCUSSION

This study showed a high prevalence of HCV (23.5%) among HIV-positive subjects. The study also showed a higher prevalence of co-infections of HCV with HIV in males than their female counterparts.

There was also higher infection rates among the age group 55 years and above compared to other age groups. This is in agreement with previous reports in Northern Nigeria (Nwokoedi *et al.*, 2006) and elsewhere (Lincoln *et al.*, 2003; Michael *et al.*, 2012). Ruan *et al.* (2004) reported 71.0% of intravenous drug abusers (IDAs) in China had antibodies to HCV. HCV—HIV co-infection among IDAs was 11.3% in a study by Ruan *et al.* (2004). Kaur and Marshalla (1998) screened 233 serum samples for HCV and found that 0.8% for HCV. Garg *et al.* (2001) evaluated blood donors for HCV and the incidence of HCV was 0.29%. Nanu *et al.* (1997) screened blood donors and reported that HCV rates to be 1.49% among donors, and those with multiple infections were uncommon. Patel (2004) screened blood donors in Mumbai over a 6-year period, from 1994 to 1999, and found that 0.78% had antibodies to HCV. Gupta *et al.* (2004) screened blood units in Ludhiana, during the period 2001—2003 and reported that 1.09% were HCV positive.

The finding of 23.5% prevalence rate of HCV in this study further confirm the presence of hepatitis C infection in Nigeria (Mutimer *et al.*, 1994; Mwangi, 1999; Egah *et al.*, 2004). Forbi *et al.* (2007) found an HCV seroprevalence rate of 11.1% in the group of HIV-1 infected individuals sampled. Previous studies in Nigeria had reported an overall HCV prevalence of 2.9% among blood donors in Rivers state of Nigeria (Kaote *et al.* 2005). Agwale *et al.* (2004) had recorded an HCV seroprevalence rate of 8.2% among HIV infected Nigerians. There is a clear indication of increased HCV infection in HIV infected individual in Nigeria. It is known that HIV/HCV co-infected individuals accelerate rapidly to end-stage liver disease, AIDS defining clinical event and death (Greub 2000, Monga *et al.*, 2001). Unfortunately at this time, no effective vaccine has been developed against HCV infection. Forbi *et al.* (2007) reported a case of a 35 years old female client co-infected with HIV and HCV who virologically failed therapy (CD4+ decline from 199 to 66) after four months on HAART. Forbi *et al.* (2007) also recorded that, the rate of increase in CD4+ cells post-HAART does not change in HIV and hepatitis coinfection but HCV appears to hinder virological response to therapy. Although there have been case reports of clearance of HCV viraemia after initiation of HAART (Ranieri *et al.* 2003), majority of available data indicates that HAART results in net increase in HCV viraemia (Chung *et al.* 2002). Forbi *et al.* (2007) have recorded that 7.2% of individual do have triple co-infection with HIV/HBV/HCV.

Gender-specific prevalence showed that males (28.9%) had higher seropositivity for HIV-HBV co-infections than their female counterparts with 19.0% prevalence. The difference was however,

significant ($P < 0.05$); the reason for this difference might be due to lesser number of males in this study. In this study, more females than males were infected with HIV only. This gender disparity is consistent with the sex distribution seen in other studies in Nigeria and reflects the national distribution where a little over 55% of people infected with HIV are women (Otegbayo *et al.*, 2008; Adewole *et al.*, 2009; Ogbuji and Oke, 2010; Aliyu *et al.*, 2010; Tremeau-Bravard *et al.*, 2012). This is comparable to what was reported in a study by Tremeau-Bravard *et al.* (2012) that more women than men were infected with HIV only. However, in this present study, more males (28.9%) are co-infected with HIV and hepatitis than females (19.0%). This distribution has been also found in other parts of Nigeria (Uneke *et al.*, 2005; Forbi *et al.*, 2007; Tremeau-Bravard *et al.*, 2012), and could be explained by the fact that the male population is more prone to have multiple sexual partners than females (NCCA, 2001; Tremeau-Bravard *et al.*, 2012).

The age distribution in our study does not vary between the HIV and HIV/hepatitis population. Almost half of the studied population is in the 35–44 years age group, which mirrors the overall demographic in Nigeria (CIA, 2009; Tremeau-Bravard *et al.*, 2012). Analysis of the age related prevalence of HCV in this study showed that age group 50 years and above had the highest prevalence of 53.3% compared to other age groups. This agrees favourably with the findings of Okonko *et al.* (2012) who reported a higher prevalence of HCV among older age group 40 years and above. This pattern indicates that most HCV transmission occurred in the recent past (i.e. 20–40 years ago), primarily among young adults. This slightly agrees with the pattern observed in the United States where highest prevalence was observed among persons 30–49 years old (Alter *et al.*, 1999; Okonko *et al.*, 2012). Another pattern that emerges is observed in Egypt, where the prevalence of HCV infection increases steadily with age and high rates of infection are observed among persons in all age groups (Mohammed *et al.*, 1996; Okonko *et al.*, 2012). The reason for these observed differences in the prevalence pattern of HCV infection in different parts of the world is not immediately known to this study.

5. CONCLUSION

The present study have demonstrated that co-infection of HIV and HCV is on the increase in Nigeria. Thus, the phenomenon of HIV and HCV co-infection is a cause for concern. The medical community in Nigeria therefore needs to be alert to this phenomenon as smart treatment options would need to be instituted in such individuals if treatment is to be meaningful.

ACKNOWLEDGEMENTS

We sincerely thank the management and staff of the HIV Clinic of Federal Medical Center, Abeokuta.

Correspondence to:**Iheanyi O. Okonko**

Medical Microbiology Unit,
Department of Microbiology,
University of Port Harcourt,
PMB 5323 Choba, East-West Road,
Port Harcourt, Rivers State, Nigeria;
E-mail: mac2finney@yahoo.com,
iheanyi.okonko@uniport.edu.ng;
Tel.: +234 803 538 0891

REFERENCES

1. Adewole OO, Anteyi E, Ajuwon Z, Wada I, Elegba F, Ahmed P, et al. Hepatitis B and C virus co-infection in Nigerian patients with HIV infection. *J Infect Dev Ctries*. 2009;3(5):369-375.
2. Agwale SM, Tanimoto L, Womack C, Odama L, Leung K, Duey D, Negedu-Momoh R, Audu I, Mohammed SB, Inyang U, Graham B, Ziermann R. 2004. Prevalence of HCV coinfection in HIV-infected individuals in Nigeria and characterization of HCV genotypes. *J Clin Virol* 31(Suppl. 1): S3-6.
3. Aliyu MH, Varkey P, Salihu HM, Iliyasu Z, Abubakar IS. The HIV/AIDS epidemic in Nigeria: progress, problems and prospects. *Afr J Med Med Sci*. 2010 Sep;39(3):233-239.
4. Alter MJ, Gerety R.J., Smallwood L. 1982. Sporadic non-A non-B hepatitis: frequency & epidemiology in an urban United States population. *J. infect Dis*; **145**:886-93.
5. Alter MJ, Kruszon – Moran D, Nainan O.V. 1999. Prevalence of hepatitis C infection in the United States. *N Eng J Med*. **341**: 556 – 62.
6. Chung RT, Evans SR, Yang Y, Theodore D, Valdez H, Clark R, Shikuma C, Nevin T, Sherman KE, AIDS Clinical Trials Group 383 Study Team. 2002. Immune recovery is associated with persistent rise in hepatitis C virus RNA, infrequent liver test flares, and is not impaired by hepatitis C virus in co-infected subjects *AIDS* **16**: 1915-1923.
7. CIA-The World Factbook [database on theInternet]. Washington, DC: Central Intelligence Agency. 2009 [cited 2011 Jul 23]. Available from:<https://www.cia.gov/library/publications/theworld-factbook/>.
8. Egah DZ, Mandong BM, Iya D, Gomwalk NE, Audu ES, Banwat EB, Onile BA. 2004. Hepatitis C Virus Antibodies among Blood Donors in Jos, Nigeria. *Annals of African Medicine* **3(1)**: 35-37
9. Feld JJ, Ocama P, Ronald A. 2005. The liver in HIV in Africa. *Antivir Ther* **10**: 953-965.
10. Forbi JC, Gabadi S, Alabi R, Iperepolu HO, Pam CR, Entonu PE, et al. The role of triple infection with hepatitis B virus, hepatitis C virus, and human immunodeficiency virus (HIV) type-1 on CD4+ lymphocyte levels in the highly HIV infected population of North-Central Nigeria. *Mem Inst Oswaldo Cruz*. 2007 Jun;102(4):535-537.
11. Garg S, Mathur DR, Garg DK. Comparison of seropositivity of HIV, HBV, HCV and syphilis in replacement and voluntary blood donors in Western India. *Ind J Pathol Microbiol* 2001;44:409—412.
12. Greub, G., Ledergerder, B. and Bathegay, M. (2002). Clinical progression, survival and immune recovery during antiretroviral therapy in patients with HIV I and hepatitis C virus coinfection: the Swiss HIV control study. *Lancet*. 356: 1800-1805.
13. Gupta N, Kumar V, Kaur A. Seroepidemiology of HIV, HBV, HCV and syphilis in voluntary blood donors. *Ind J Med Sci* 2004;58:306—307.
14. Kaur H, Marshalla R. Seroepidemiology of HIV, HBV, HCV and treponemal infections. *J Commun Dis* 1998;30:29—31.
15. Koate BB, Buseri FI, Jeremiah ZA. Seroprevalence of hepatitis C virus among blood donors in Rivers State, Nigeria. *Transfus Med*. 2005 Oct;15(5):449-451.
16. Lincoln, D., Petoumenos, K. and Dore, G.J. (2003). HIV/HBV and HIV/HCV coinfection and outcomes following high active antiretroviral therapy. *Human Immunodeficiency Virus Medicine*. 4: 241-249.
17. Madhava V, Burgess C, Drucker E. Epidemiology of chronic hepatitis C virus infection in sub-Saharan Africa. *Lancet Infect Dis*. 2002 May;2(5):293-302.
18. Mc Lean FM, Morgan – Copner P, Peutherer JF. 1997. Arboviruses. In D. Greenwood, R.C.B. Slack, J.F., Peutherer (eds). *Medical microbiology* 15th Edith church 1 Livingstone pp. 485 – 505.
19. Michael, D., Dieterich, D. and Touly, M. (2012). Uses of CD4 Cell Count in Human immunodeficiency virus. *Haematology*. 45: 2-15.
20. Modi AA, Feld JJ. Viral hepatitis and HIV in Africa. *AIDS Rev*. 2007 Jan-Mar;9(1):25-39.
21. Mohammed MK., Hussein MH., Massoud AA. 1996. Study of the risk factors for viral hepatitis C infection among Egyptians applying for work abroad. *J Egypt Public Health Assoc*. 71:113-42
22. Monga HK, Rodriguez-Barradas MC, Breaux K, Khattak K, Troisi CL, Velez M, et al. Hepatitis C virus infection-related morbidity and mortality among patients with human immunodeficiency virus infection. *Clin Infect Dis*. 2001 Jul 15;33(2):240-247.
23. Mutimer D. J, Olomu A, Skidmore S. et al. Viral hepatitis in Nigeria: sickle cell disease and

- commercial blood donors. *Quarterly J Med* 1994; 87:407- 411.
24. Mwangi JW. 1999. Viral markers in a blood donor population. *East Afr Med J* 79: 35-37.
25. Nanu A, Sharma SP, Chatterjee K, Jyoti P. Markers for transfusion transmissible infections in north Indian voluntary and replacement blood donors: prevalence and trends 1989—1996. *Vox Sang* 1997;73:70—73
26. Nigeria Common Country Assessment (NCCA)[database on the Internet]. World Health Organization. 2001 [cited 2012 Feb]. Available from: www.ng.undp.org/documents/CCA_2001.pdf.
27. Nwokedi, E.E., Ilyasu, Z., Emokpae, M.A., Dutse, A.I. and Taura, A.A. (2006). Hepatitis C virus infection among Teaching Hospital patients in Kano, Nigeria: A retrospective study. *Annals African Medicine*. 5(4): 185-87.
28. Odemuyiwa SO, Mulders MN, Oyedele OI, Ola SO, Odaibo GN, Olaleye DO, Muller CP 2001. Phylogenetic analysis of new hepatitis B virus isolates from Nigeria supports endemicity of genotype E in West Africa. *J Med Virol* 65: 463-469.
29. Ogbuji QC, Oke AE. Quality of life among persons living with HIV infection in Ibadan, Nigeria. *Afr J Med Med Sci*. 2010 Jun;39(2):127-135.
30. Okonko IO, Oyediji TO, Anugweje KC, Adeniji FO, Alli JA, Abraham OA. 2012. Detection of HCV antibody among intending blood donors. *Nature and Science* ; 10(1):53-58
31. Otegbayo JA, Taiwo BO, Akingbola TS, Odaibo GN, Adedapo KS, Penugonda S, et al. Prevalence of hepatitis B and C seropositivity in a Nigerian cohort of HIV-infected patients. *Ann Hepatol*. 2008 Apr-Jun;7(2):152-156.
32. Patel Y. Seroprevalence of HIV, HBV, HCV and syphilis in blood donors. *Indian Journal of Medical Sciences* 2004;58:255—257.
33. Ranieri R, Santambrogio C, Veronelli A, Pontiroli AE 2003. Hepatitis C viremia persistently suppressed by HAART. *Clin Infect Dis* 36: 1086-1087.
34. Ruan YH, Hong KX, Liu SZ, He YX, Qin GM, Chen KL, et al. Community based survey of HCV and HIV co-infection in injection drug abusers in Sichuan Province of China. *World J. Gastroenterology* 2004;10:1589—1593.
35. Tremeau-Bravard A, Ogbukagu IC, Ticao CJ, Abubakar JJ. 2012. Seroprevalence of hepatitis B and C infection among the HIV-positive population in Abuja, Nigeria. *African Health Sciences*; 12(3): 312 - 317
36. Uneke CJ, Ogbu O, Inyama PU, Anyanwu GI, Njoku MO, Idoko JH. Prevalence of hepatitis- B surface antigen among blood donors and human immunodeficiency virus-infected patients in Jos, Nigeria. *Mem Inst Oswaldo Cruz*. 2005 Feb;100(1):13-16.
37. WHO-Hepatitis C [database on the Internet]. World Health Organization. 2011 Jun [cited 2011 July]. Available from: <http://www.who.int/mediacentre/factsheets/fs164/en/index.html>.
38. World Health Organization (WHO). 1996. Fighting disease, fostering development, World health Organization Report, Geneva; 1996.

5/15/2013