

MATERNAL AND FETAL OUTCOMES AMONG HIV POSITIVE PREGNANT MOTHERS ATTENDING ANTENATAL CLINIC AT UNIVERSITY OF MAIDUGURI TEACHING HOSPITAL MAIDUGURI NIGERIA

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ABSTRACT

Objective: To determine the pregnancy outcome among HIV positive women attending ANC at the University of Maiduguri Teaching Hospital, Maiduguri. **Method:** A prospective cohort study of 500 women attending the ANC between 12th August 2009 – 21st December 2010 was carried out. Initial HIV screening was done using single rapid tests, while diagnosis for those found positive was done with an additional second rapid test, while confirmation is done with 'Western Blot' technique. Socio demographic and obstetrics variables were obtained and analyzed. **Results:** Out of the 500 pregnant mothers that were counseled and tested for HIV, 52 were found to be HIV positive giving a sero prevalence rate of 10.4%. The HIV positive women are used as cases while out of the remaining cohort 162 patients that are matched for age and parity were used as control. The perinatal transmission rate was 11.5%. The HIV positive women were more educated and younger than the control. More than 90% of the HIV positive women booked for ANC between 2nd and 3rd trimester with average gestational age at booking of 29.3 weeks. Elective caesarean section (ELCS) was offered to 11(21.2%) and all the babies delivered through ELCS were negative for HIV. Prematurity, low birth weight and birth asphyxia were more in HIV positive women than control. Advanced maternal age ($X^2=33.53 P=0.000$), Low CD4 count ($X^2=15.58 P=0.016$), high maternal viral load ($X^2=21.85 P=0.005$), prematurity ($X^2= 9.872 P= 0.007$), low birth weight ($X^2 = 63.80 P= 0.000$) and birth asphyxia($X^2 = 24.149 P= 0.000$) were the major determinants of perinatal transmission of HIV infection in this study. **Conclusion:** The Seroprevalence and MTCT of HIV infection in pregnancy is high in our environment. HIV infection was also found to be associated with increases risk of elective caesarean section and neonatal asphyxia. Efforts should be intensified to improve the situation.

Keywords: HIV, Maternal Outcome, Fetal Outcome, Maiduguri

INTRODUCTION

Since the report of the first HIV/AIDS case in Nigeria, in a 13 year old girl in 1986, the rate of the infection increases rapidly,¹ with the national Sero-prevalence among pregnant women attending ANC rising from less than

1% in 1990 to 5.8% in 2001.² A slight downward trend has been observed in the last four surveys with a sero prevalence rate of 5.0% 4.4%, 4.6% and 3.4% in 2003, 2005, 2008 and 2013 respectively.^{3,4} The burden of HIV is higher in sub-Saharan

Africa than the rest of the world because of the high level of heterosexual transmission, high female to male ratio, and high total fertility rate (TFR), and high prevalence of breast feeding.⁵

HIV infection has been reported to have little effect on pregnancy outcome in the developed world.⁵⁻⁸ However, adverse pregnancy outcome have been reported in a number of African studies.⁹⁻¹² These complication rates vary across studies and may reflect the extent of the epidemic and the nature of HIV - related diseases in different communities. Untreated maternal HIV infection was observed to be associated with adverse pregnancy outcomes in form of increased maternal and fetal morbidities. Most common complications reported were, maternal anaemia, preterm labour, premature rupture of membranes, abortions, IUGR, IUFD, low birth weight, still birth and increase risk of perinatal transmission of HIV.¹³⁻¹⁷

The objective of this study therefore is to determine the maternal and fetal outcome of pregnancies complicated by HIV.

METHOD

This was a prospective hospital based cohort study that was carried out at the department of Obstetrics and Gynaecology of the University of Maiduguri Teaching Hospital from 12th August 2009 - 21st December 2010 after approval was granted by the research and ethical committee of the hospital.

Five hundred (500) pregnant women were recruited using convenient sampling technique as the study population. After counseling, the HIV testing (VCT) was offered to the study participants using the opt-out method. All study participants were tested after obtaining informed consent and emphasis was placed on confidentiality, benefit of the test and were informed that they are free to opt out or withdraw from the study at any point if they so

wish. Those women found to be positive were used as cases and 162 of those negative were randomly selected and used as control. These women were followed up through delivery and up to the first 12 weeks post partum to obtain their pregnancy outcome including the status of their babies.

A pretested questionnaire was used to obtain the socio demographic and Obstetrics data of the study population. The information obtained were coded, entered into a computer and analysed using SPSS version 11 statistical package (SPSS, Chicago, Ill, USA). Test of significance was done using Chi square test. A p-value of <0.05 was considered to be significant.

The diagnosis of HIV infection was made in two stages i.e. the initial screening using Rapid tests" determine" (revised serial method) which detects antibodies to HIV using latex agglutination. For those found positive with determine a second rapid test "stat pak" was used for the diagnosis ,where these two test were found to be discordant a 'tie breaker (third rapid test) using 'Unigold' was done. The rapid tests were all carried out in the ANC clinic and the results are available within 30 minutes.

Those found positive were confirmed with 'Western Blot' technique. This was done in the main immunology laboratory of the Hospital. Patients who tested positive by western blot were offered post test counseling and enrolled in to the PMTCT programme to continue their ANC follow up according to the national protocol,⁴ while negative mothers were offered post test counseling on how to remain negative and continue with their routine ANC follow up. According to the Nigerian national guideline 2007³ in addition to the normal criteria for initiation of ART in adults infected with HIV, pregnancy constitutes an indication for ART prophylaxis. Treatment is offered as per the WHO clinical staging and eligibility

criteria.³ Mothers who are not eligible for their own disease are offered prophylaxis with Highly Active Antiretroviral therapy (HAART) and exposed infants received single dose nevirapine suspension 2mg/kg within 72 hours of birth and Zidovudine 4mg/Kg twice daily for six weeks.

The infants of the HIV positive mothers were followed up to first 12 weeks of age to determine their HIV sero status using polymerase chain reaction (PCR) after separation amplification and reading using the PCR machine (Amplicor HIV -1 DNA Assay version 1.5 ,Roche molecular system USA). An infant is considered HIV positive if the PCR is reactive for viral DNA at 0, 6 and/or 12 weeks of age.

RESULTS

Out of the 500 pregnant mothers that were counseled and tested for HIV; 52 were found to be HIV positive giving a Sero prevalence rate of 10.4%. The mean age was 28.6±6.6 (16 - 41 years) and the mean parity was 3.2±2.0.

Table 1 illustrates the socio demographic characteristics of the two groups. The peak age specific incidence (38.5%) of HIV was in the 25-29 year age group (p=0.01). However there was no statistically significant incidence differences between the two groups in the other age groups. There were significantly fewer primigravidas 4(7.7%) among the HIV positive pregnant mothers than HIV negative 36 (22.2%) (p = 0.04). Overall more of HIV negative mothers were less educated compared to HIV positive mothers (p = 0.001).

Sixteen (30.8%) of HIV positive pregnant women were known HIV patients receiving treatment and care at the adult ARV clinic before transfer to the ANC for antenatal care, while the remaining 36(69.2%) were new HIV positive diagnosed at booking in the antenatal

clinic. Forty eight (92.3%) of the HIV women booked for ANC between 2nd and 3rd trimester with average booking gestational age of 29.3±2.4 weeks.

The Obstetrics performance of the HIV infected and uninfected pregnant mothers were compared as shown on table 2. There was no statistically significant difference in terms of ANC complications, gestational age at delivery and postpartum complications among the two groups. Anaemia and preterm labour were each observed among 3 (5.8%) of the infected mothers. Spontaneous onset of labour was observed among 39 (95.1%) of those HIV infected and 147 (90.7%) of those uninfected, while induction of labour was done to 2 (4.9%) of the infected and 15 (9.3%) of the uninfected mothers. Only 2 HIV infected mothers were offered episiotomy compared to 45 of those uninfected (P = 0.000). Elective caesarean section was offered to 11(21.2%) compared to none among the uninfected with p < 0.001, while spontaneous vaginal delivery occurred among 39(76.9%) and 153(94.4%) with p = 0.005.

Table 3 detailed the fetal outcome among the two groups. There was no statistically significant difference between the two groups in respect to fetal survival at birth, fetal sex, and fetal weight. Fetal APGAR at 5th minutes of birth revealed mild asphyxia among the HIV infected mothers of 17 (32.7%) compared to 6(3.7%) of uninfected pregnant mothers p < 0.001.

Of the 52 babies delivered by the HIV positive mothers, 6 were found to be HIV positive both by PCR 1 & 2 at 0, 6 and/or 12th week of birth giving a transmission rate of 11.5%.

Table 1: Socio demographic characteristics of the 52 HIV infected and 162 uninfected control among the study population

Parameters	HIV positive %		HIV negative %		P value
Age					
<20	2	3.8	12	7.4	0.42
20-24	10	19.2	48	29.6	0.21
25-29	20	38.5	27	16.7	0.01
30-34	11	21.2	48	29.6	0.31
35-40	8	15.4	18	11.1	0.52
>40	1	1.9	95	58.6	0.33
Total	52	100	162	100	
Education					
Nil	9	17.3	78	48.1	0.001
Primary	13	25.0	21	13.0	0.11
Secondary	18	34.6	33	20.4	0.10
Post secondary	6	11.5	15	9.3	0.70
Tertiary	6	11.5	15	9.3	0.70
Total	52	100	162	100	
Gravidity					
Primigravida	4	7.7	36	22.2	0.04
2-4	27	51.9	84	51.9	0.99
5-7	20	38.5	39	24.1	0.11
8	1	1.9	3	1.9	0.98
Total	52	100	162	100	
Gest.age at booking					
1 st trimester	4	7.7	6	3.7	0.37
2 nd trimester	29	55.8	105	64.8	0.34
3 rd trimester	19	36.5	51	31.5	0.58
Total	52	100	162	100	

Table 2: Obstetrics performance of the 52 HIV infected and 162 uninfected Control

ANC complications	HIV infected	%	Uninfected	%	P value
Nil	45	86.5	150	90.5	0.31
Anaemia	3	5.8	3	1.9	0.29
Preterm labour	3	5.8	0	0	0.07
IUGR	1	1.9	0	0	0.31
Malaria	0	0	103	63.8	0.17
UTI	0	0	3	1.9	0.37
Total	52	100	162	100	
G/age at delivery					
Preterm	6	11.6	6	3.7	0.13
Term	44	84.6	147	90.7	0.34
Post term	2	3.8	9	5.6	0.68
Total	52	100	162	100	
Onset of labour					
Spontaneous	39	95.1	147	90.7	<0.001
Induced	2	4.9	15	9.3	<0.001
Total	41*	100	162	100	
Duration of labour					
Normal	40	97.6	153	94.4	<0.001
Prolong	1	2.4	9	5.6	<0.001
Total	41*	100	162	100	
Intrapartum intervention					
Episiotomy: Yes	2	3.8	45	27.8	<0.001
No	50	96.2	117	72.2	<0.001
Total	52	100	162	100	
Mode of delivery					
SVD	40	76.9	153	94.4	0.005
Elective C/S	11	21.2	0	0	0.000
Emergency C/S	1	1.9	3	1.9	0.98
Ass.vag. delivery	0	0	6	3.7	0.16
Total	52	100	162	100	
Postpartum complications					
Nil	48	92.3	156	96.3	0.38
Peuerperal pyrexia	3	5.8	31.9	0.29	
Anaemia	1	1.9	31.9	0.10	
Total	52	100	162	100	

* 11 of the patients had elective c/s hence N= 41

Table 3: fetal outcome among the 52 HIV infected and 162 uninfected control

Fetal survival	HIV infected	%	HIV uninfected	%	P value
Alive	51	98.1	162	100	0.31
Still birth	1	1.9	0	0	0.31
Total	52	100	162	100	
Fetal APGAR					
Normal apgar	31	59.6	156	96.3	<0.001
Mild asphyxia	17	32.7	6	3.7	<0.001
Moderate asphyxia	3	5.8	0	0	0.07
Total	*51	100	162	100	
Fetal sex					
Male	27	51.9	84	51.9	0.99
Female	25	48.1	78	48.1	0.99
Total	52	100	162	100	
Fetal birth weight					
Normal weight	45	86.5	156	96.3	0.07
Low birth weight	7	13.5	6	3.7	0.07
Total	52	100	162	100	

* N= 51 HIV infected 1 was a still birth

DISCUSSION

The HIV Seroprevalence rate found among the study population is 10.4%, which is similar to 11.8% and 10.5% reported from the same institution in 2003¹⁷ and 2004¹⁸ respectively. Our findings though within the range of 1.2% - 12% reported in most states of Nigeria³, is however higher than the national Seroprevalence rate of 3.2% reported in 2013⁴. Other Hospital based studies from Nigeria have equally reported much lower prevalence rates^{13,19} compared to our finding.

Previous studies have documented that HIV is common among young and those of low parity.^{20,22, 23} This study also showed that HIV positive mothers were significantly younger and of lower parity. This is likely so because of the ethnicity and cultural peculiarities of the study population in which early marriage is common.⁴ Our study revealed that HIV positive mothers were less likely not to have western education than their negative

counterparts. This is in contrast to a report from Kigali, Rwanda among Antenatal clinic attendants.²¹ With low literacy level our women tend to marry early compared to their educated counterparts who are of liberal mind increasing their risk opportunity for infection. Also our women are not courageous enough to negotiate for sex because of our cultural sensitivities.

Although there isn't much significant difference in antenatal complications among the two study group, anemia was observed to be commoner among HIV positive mothers than HIV negative ones. Several investigators^{19, 21, 24 - 28} have reported higher prevalence of anemia among HIV positive pregnant women than HIV negative and can be because of the effect of the virus or the drugs used to treat the virus.. The low episiotomy rate of 3.8% found in this study is in conformity with the Nigerian national guideline to minimize intrapartum interventions that increase the risk of perinatal transmission.³

Elective caesarean section though not a popular method of preventing perinatal transmission of HIV in developing countries²⁹, was offered to 21.2% of our patients, this is similar to reported in Makurdi³⁰ (21.7%), but much higher than 10.3% reported from Kano¹³, Nigeria. The justification for the high elective caesarean section rate in his study was informed by the recommendation that this option of delivery should be offered to some HIV positive mothers in our environment to prevent mother to child transmission of HIV.²³ All the babies delivered through elective caesarean section were found to be HIV negative, hence the choice of this mode of delivery by their mothers was a wise decision aim at preventing transmission of HIV to their neonates. A much larger trial have also advocated elective caesarean section as an effective way to minimize perinatal transmission of HIV particularly for those with detectable viral load.³¹⁻³³

Several studies have previously established that HIV-1 infection in pregnancy is associated with prematurity and low birth weight.^{19, 26, 34-}

³⁶Though the difference in birth weight was not statistically significant between the two study groups, babies born to HIV positive mothers were found to weigh less than those of HIV negative mothers with mean of 2.87kg and 3.43kg respectively.

The high prevalence of mild birth asphyxia found among the cases in this study is in agreement with earlier reports in some parts of Nigeria.^{19, 20, 26} This may be attributed to general reservation to avoid unnecessary intrapartum interventions and fetal manipulations that are presume to increase the risks of perinatal transmission of HIV.

CONCLUSION

The Seroprevalence and Mother to Child Transmission (MTCT) of HIV are high in our environment. HIV infection was also found to be associated with increases risk of elective caesarean section and neonatal asphyxia. Efforts should be intensified to improve the situation.

REFERENCES

1. Manual on the management of HIV/AIDS in pregnancy. Society of gynaecology and Obstetrics of Nigeria (SOGON) 2005.
2. Akani CI, Ojule AC, Oporum HC, John CT. Seroprevalence of HIV antibodies in pregnant women in Port Harcourt, Nigeria. *Nig J Med* 2006; 15(1): 44- 48.
3. National guidelines on the prevention of mother to child transmission of HIV in Nigeria, Federal ministry of health 2007.
4. National HIV/AIDS and Reproductive Health Survey - plus (NARHS - plus) 2013
5. Nigeria Demographic and Health Survey. Abuja, Nigeria: National Population Commission, Federal Republic of Nigeria and MEASURE DHS+ ORC Macro; 2009.
6. Johnstone FD. Pregnancy outcome and pregnancy management in HIV- infected women. In: Johnson MA Johnson FD (Eds). HIV infection in women. Edinburg, Churchill Livingstone, 1993; 187-198.
7. Bakas C, Zarou DM, de Caprariis PJ. First trimester spontaneous abortions and incidence of HIV seropositivity. *J Reprod Med*, 1996; 41(1): 15- 18.
8. Johnstone FD. HIV and pregnancy. *Br J Obstet Gynecol* 1996; 103: 1184- 1190.
9. Brocklehurst P, French R. The association between maternal HIV infection and perinatal outcome: review of the literature and meta analysis. *Brit J Obstet*

- Gynaecol 1998; 105: 839-848.
10. McIntyre JA. Pregnancy and HIV infection at Baragwanath Hospital, 1987-1993. Eight International Conference on AIDS and STD in Africa, Marrakesh, 1993; Abstract
 11. Temmerman M, Plummer FA, Mirza NB et al. Infection with HIV as a risk factor for adverse pregnancy outcome. *AIDS* 2007; 162: 30-34.
 12. Musana JW, Oswang SB, Khisa W, Kiarie JW. Pregnancy outcome in mothers with Advanced Human Immunodeficiency Virus disease. *East Afr Med J* 2009; 86(10): 480-5
 13. Tukur J, Galadanchi H, Adeleke SI, Mukhtar-Yola M. Outcome of delivery among HIV positive mothers at Aminu Kano Teaching Hospital, Kano. *Nig J Med* 2007; 16(1):34-7.
 14. Bergstrom S, sonnetborg A, Osman NB, Libombo A. HIV infection and maternal outcome of pregnancy in Mozambican women: a case control study. *Genito Urin Med*, 2009; 71: 323- 324.
 15. Langston C, Lewis DE, Hammill HA et al. Excess intrauterine fetal demise associated with maternal HIV infection. *J Infect Dis* 1995; 172: 1451-1460.
 16. Shearer WT , Langston C, Lewis DE et al. Early spontaneous abortion and fetal thymic abnormalities in maternal to fetal HIV infection. *Acta Paediatr* 1997; Suppl 421: 60-64.
 17. Chama CM, Audu BM, Kyari O. Prevention of MTCT of HIV at Maiduguri, Nigeria. In *J Obstet Gynaecol* 2004; 266- 269.
 18. Chama C, Gashau W, Oguche S. The value of highly active antiretroviral therapy in the prevention of mother to child transmission of HIV . *J Obstet Gynaecol* 2007; 27(2): 134-137
 19. Samuel NO. Pregnancy outcome in HIV seropositive women in Abakiliki, Nigeria. *Orient J Med* 2005; 17(3&4):25-30
 20. Onah HE, Obi SN, Agbata TA, Oguano TC. Pregnancy outcome in HIV-positive women in Enugu, *Nig J Obstet Gynaecol* 2007; 27:271-4
 21. Leroy V, Ladner J, Nyiraziraje M, Declercq A, Bazubagira A, Van de Perre P. Karita study group. Effect of HIV-1 infection on pregnancy outcome in women in Kigali, Rwanda. *AIDS* 1998; 12:643-650
 22. Gomutbutra V. Characteristics of pregnancy with human immunodeficiency virus (HIV) and perinatal transmission in Nakornping Hospital. *J Med Assoc Thai* 2008; 91(2):142-5
 23. Chama CM, Moruppa JY. The safety of elective caesarean section for the prevention of Mother-to-Child transmission of HIV-1. *J Obstet Gynaecol* 2008; 28(2):194-7.
 24. Candice B, Hester K, Gayle L. A comparison of HIV positive and negative pregnant women at a public sector hospital in South Africa. *J Clin Nurs* 2006; 15(6):735-741
 25. Nanche D, Bardaji A, Lahuerta M, Berenquera A, Mandomando I, Sanz, S et al. Impact of maternal human immunodeficiency virus infection on birth outcome & infants survival in rural Mozambique. *Am J Trop Hyg* 2009; 80(5): 870-6.
 26. Olagbusi BN, Ezeanochie MC, Ande AB, Oboro VO. Obstetric and Perinatal outcome in HIV positive Women receiving HAART in Urban Nigeria. *Arch Gynecol Obstet* 2009; 3(2) 37 - 39 (Epub ahead of print)
 27. Jessie L, Aleyamma TK, Lilly V, Jessica B, Geetha G, Sreekanth C et al. HIV and Obstetrics complications and fetal outcome in Vellore. *India Trop Doct* 2008; 38:144-146
 28. Bardequez A, Palambo P Wesley Y, Holland B, Denny T, Connor E. Pregnancy outcome of the HIV infected

- gravida: 9th Int Conf AIDS 1993; 9: 458(6-11) PO-B23-1936 Abstract.
29. Newell ML. Current issues in the prevention of Mother-to-child Transmission of HIV-1 infection. *Tans R Soc Trop Med Hy* 2006; 100(1):1-5.
 30. Swende TZ, Agida ET, Jogo AA. Elective caesarean section at The Federal Medical Centre Makurdi, North central Nigeria. *Nig J Med* 2007; 16(4):372-4
 31. Abarzua F, Nunez F, Hubinont C, Bernard P, Yombi JC, Vandercan B. Human immunodeficiency virus (HIV) infection In pregnancy: antiretroviral treatment (ART) and mode of delivery. *Rev Chilena Infectol* 2005; 22(4):327-37.
 32. Suy A, Hernandez S, Thorne C, Lonca M, Lopez M, Coll O. Current guidelines on management of HIV infected pregnant Women: impact on mode of delivery. *Eur J Obstet Gynaecol Reprod Biol* 2008; 13(2):127-32.
 33. Read JS, Newell MK. Efficacy and safety of caesarean delivery for prevention of mother to child transmission of HIV-1 *Cochrane Database Syst Rev*. 2005;(4):CD005479.
 34. Taha TET, Sufia SD, Hafizur MR, Jin S, Johnstone K, Newton K. Trends in Birth Weight and Gestational Age for Infants Born to HIV-infected, Antiretroviral Treatment-Naïve Women in Malawi. *Paediatr Infect Dis J* 2012; 31(5): 481 - 486
 35. Sukwa TY, Bakketeig L, Kanyama I, Samdal HH. Maternal HIV infection and pregnancy outcome. *Centr Afr J Med*, 1996. 42(8): 233- 235.
 36. Iroha EO, Ezeaka VC, Akinsulie AO, Temiye FO, Adetifa IM. Maternal HIV infection and Intra-uterine growth: A prospective Study in Lagos Nigeria. *West Afr J Med*. 2007; 26(2):121-5.