

Current Status of Microsporidium Among Hospitalised Human Immunodeficiency Virus (HIV/AIDS) Infected Patients, Federal Medical Centre, Keffi, Nigeria

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Abstract: Microsporidia are obligate intracellular parasitic fungi causing chronic diarrhea, particularly among immunocompromised patients. Microsporidial infections have been recognized as an increasingly important infection, particularly among those hospitalised with HIV/AIDS infected patients at the federal medical centre (FMC), Keffi, Nasarawa State. One health facility was used to engage HIV/AIDS infected patient accessing antiretroviral therapy (ART) to identify the presence of microsporidium and to determine the associated risk factors. 252 stool samples were examined for microsporidial spores by modified giemsa staining technique. The overall prevalence rate of 15.08% was recorded. Based on age group, microsporidia infection was common among patients aged between 61-70years 15.0% while sex-related, the male had 30.43% rate of infection. Though, marital status, occupation, and widow/widower had 15.79% and artisans (33.33%) respectively. The vulnerability was determined by their fate and status. However, chi-square result showed no significant relationship observed ($P>0.05$) between the age, and sex- distribution of microsporidium. The proportion among occupation and marital related distribution of microsporidium among HIV/AIDS infected patients were diarrhoic. In relation to viral load, all positive HIV/AIDS infected patients with microsporidia spore had viral load above normal. Twenty-five (25) HIV/AIDS infected patients had 10,000ml/viral load in replicates. Microsporidium is therefore, identified and recognized as an invasive opportunistic infection among HIV/AIDS infected patients which should be considered in a routine checks among HIV/AIDS infected patients.

Keywords: Current Microsporidium Status, Among Hospitalised, Hiv/Aids, Infected Patients, Medical Centre

1. Introduction

Microsporidia, classified as highly specialized fungi, are unicellular and obligate intracellular opportunistic pathogens which can infect a wide range of vertebrate and invertebrate hosts such as fish, insects, farm and companion pets [9, 11]. The phylum microsporidia is consisting of more than 170 genera and 1300 species. Among these genera, eight of them have been responsible for human infections, including *Enterocytozoon*, *Pleistophora*, *Encephalitozoon*, *Vittaforma*, *Trachipleistophora*, *Brachiola*, *Nosema* and *Microsporidium* [8, 5, 1]. *Enterocytozoon bienewsi* (*E. bienewsi*) and the *Encephalitozoon* species (*E. cuniculi*, *E. intestinalis* and *E. hellem*) are the four major species infecting humans. *E.*

bieneusi which is responsible for more than 90% of cases with microsporidiosis in humans is most commonly diagnosed [4, 14].

Microsporidial infections have been reported to occur in severely immunocompromised individuals, mainly HIV/AIDS patients, but cases in HIV-negative people, including travellers and elderly people, are continually increasingly recognised worldwide as opportunistic infection agent. [5]. Of the several species that infect man, *Enterocytozoon bienewsi* were the first documented case and the most commonly recognised microsporidia that caused gastrointestinal disease in immunocompromised patients particularly in HIV/AIDS patients. [6].

This parasite is commonly observed in HIV-infected patients with high viral load of 10,000ml and CD₄

lymphocytes count of less than 50 cells/mm³ who complain of chronic diarrhoea, nausea, mal-absorption and severe weight loss. Whereas *Encephalitozoon intestinalis* causes both a disseminated and intestinal infections associated with nephritis, sinusitis or bronchitis [6, 3]. As a zoonotic pathogen, the main transmission way of *E. bieneusi* is faecal-oral route or oral-oral route because its spores are shed into environment via faeces. Therefore, the way of consumption of contaminated food and water is the main route of *E. bieneusi* infection [5] this will help determine current status of microsporidium in Human Immunodeficiency Virus infected patients attending Federal medical centre, Keffi.

2. Materials and Method

2.1. Study Area and Population

The study area this research was conducted among 252 consenting HIV/AIDS positive patients cover the period in 2021 at the antiretroviral therapy (ART) clinic of the federal medical centre (FMC) Keffi, Nasarawa state, Nigeria. The general topography of Nasarawa State is that of hills/dissected terrain, undulating plains and lowlands (Nasarawa State Government 2022). The State has a climate typical of the tropical zone with a maximum and minimum temperature of 81.7° F and 16.7° F respectively with moderate rainfall which varies from 131.73cm in some places to 145cm in others (Nasarawa State Government 2022). The State has a total land area of 27,117 km² (10,470 sq mi) and a population of 1,869,377 according to 2006 census and the main economic activity is agriculture.

2.2. Study Design

This study was a Cross-sectional study which includes both male and female HIV/AIDS hospitalised infected patients at the federal medical centre, Keffi, Nasarawa state Nigeria. Consent was sought among infected individuals and stool samples were collected and examined. The viral load of HIV/AIDS patients were taken between the periods of study 2021.

2.3. Sample Collection

About 10g of fresh stool samples were collected from each individual HIV/AIDS patients and each sample were well labelled with the encryption of the patient's identity; sex, age, occupation and date of collection within 14days. A clean dry, mouthed sample container with 10% formalin saline was given to each person's for collection of stool sample and stored at room temperature until it was processed.

3. Method

3.1. Giemsa Stain Method

The slides were prepared from homogenized stool samples and one drop of methanol onto the slide was fixed and was allowed to dry for 1-2 minutes. The slides were stained in 10% giemsa for 1hr and was washed with tap water and allowed to air dry. Slides were examined using x100magnification (oil immersion). Giemsa stained spores were oval, with the cytoplasm staining light grey-blue with a dark stained nucleus spores.

The versant HIV-1RNA 3.0 bDNA system tests was used for direct quantification of type-1 human immunodeficiency virus in plasma from individuals with HIV by amplification of the signal emitted by the nucleic acid, using a buyer system 34 bDNA analyser was used for viral load test [7].

3.2. Data Analysis

The Statistical Package (SPSS version 22.0) was used for data analysis. Prevalence of infection was given in percentages in line with the variables. Chi-square and 95% confidence interval analysis were the statistical tools used to determine significance at a cut-off value of 95% ($p = 0.05$).

4. Result

Age and the sex-related distribution of Microsporidium among HIV/AIDS infected patients.

Table 1. Age and Sex-related Distribution of Microsporidium among HIV/AIDS infected Patients.

Age group	Sex				Total
	Male		Female		
	No examined (%)	No positive (%)	No examined (%)	No positive (%)	
11 – 20	4	2 (50.00)	12	0 (0.00)	2 (12.50)
21 – 30	8	4 (50.00)	18	4 (22.22)	8 (30.77)
31 – 40	11	0 (0.00)	73	10 (13.70)	10 (11.90)
41 – 50	20	6 (30.00)	60	10 (16.67)	16 (20.00)
51 – 60	0	0 (0.00)	42	0 (0.00)	0 (0.00)
61 – 70	3	2 (66.67)	1	0 (0.00)	2 (50.00)
Total	46	14 (30.43)	206	24 (11.65)	38 (15.08)

The two hundred and fifty two (252) HIV/AIDS examined, were patients hospitalized and their stools sampled for examinations thus, 38 HIV/AIDS persons among the large number examined were identified/established to be affected with Microsporidium Spp. between the ages of 11-20 years in both sexes, 2 (12.50%) were positive with the high number

10 (11.90%) of HIV/AIDS individuals suffering from the prolific diarrhoic microsporidium particularly among females aged between 31-40 years and 41-50 years 16 (20.00%), availing the state of vulnerability in the age category, and between aged 61-70 years, 2 (66.67%) were infiltrated with microsporidium among the least most infected in the age

group considered vulnerable with no significant relationship in the infection of microsporidium among the ages and sexes of the infected AIDS patient's. ($X^2=6.10<11.07$, $df = 5$, $P<0.05$), table 1.

Marital status and sex-related distribution of microsporidium among HIV/AIDS infected persons were very much determined with the status of their viral load as discrepancies can be seen among the infected individuals so much that widows/widowers of both sexes 6 (15.79%) had

viral load between 520ml-56,600ml in contrast to the value/ml volume of the viral load among the married with 56,300ml-3,900ml, as well among the singles (56,600ml-92,300ml) this, clearly showed the status and the potency of the immune system among group of individuals, the higher the viral load, the lower the CD4 counts which means the state of vulnerability among the infected HIV/AIDS patients was simply established.

Table 2. Marital status and sex-related Distribution of Microsporidium among HIV/AIDS infected Patients.

Marital Status	Sex		Female		Total
	Male		No examined (%)		
	No examined (%)	No positive (%)	No examined (%)	No positive (%)	No positive for both sexes
Widow/Widowers	12	4 (33.33)	8	2 (25.00)	6 (15.79)
Married	22	4 (18.18)	119	14 (11.76)	18 (12.77)
Singles	12	6 (50.00)	79	8 (10.13)	14 (15.38)
Total	46	14 (30.43)	206	24 (11.65)	38 (15.08)

The prevalence of microsporidium spp. among the occupation and sex-related HIV/AIDS infected person was obvious, considering the occupation of the individuals that mostly exposed them to such infection, particularly farmers who by the nature of tiling the land, are always accessible and most vulnerable to the soil in their dried muddy/loamy state. Farmers in both sexes 6 (25.00%) were positive with microsporidium with their viral load 513ml-92,300ml as it differs among individuals for student category, 14 (15.38%)

were stigmatized with the viral load between 20ml-56,600ml meaning that, the low their CD4 counts in contrast to the civil servants 16 (12.31%) with viral load 15ml-1.710ml which differ with the higher viral load in the unemployed who most are left with no choice to life and among the unemployed 1 (25.00%) had over 1,890ml-58,400ml with the artisans 2,590ml-319,000ml given a wide margin among the categories in the structure though, the proportion of the infection was relatively significant to the status of the viral load (table 3).

Table 3. Occupation and sex-related Distribution of Microsporidium among HIV/AIDS infected Patients.

Occupation	SEX		Female		Total
	Male		No examined (%)		
	No examined (%)	No positive (%)	No examined (%)	No positive (%)	No positive for both sexes
Farmers	10	3 (30.00)	14	3 (21.43)	6 (25.00)
Students	9	3 (33.33)	82	11 (13.41)	14 (15.38)
Public/Civil Servants	23	7 (30.43)	107	9 (8.41)	16 (12.31)
Unemployed	2	0 (0.00)	2	1 (50.00)	1 (25.00)
Artisans	2	1 (50.00)	1	0 (0.00)	1 (33.33)
Total	46	14 (30.43)	206	24 (11.65)	38 (15.08)

However microsporidium related viral load among HIV/AIDS showed that, more of the suppressive tendencies of the virus can be expressed by simply silencing the ability of the immune capacity of the individual whose condition seems deteriorating, meaning that the CD4 counts showed no capacity to replicate hence, stagnation of the immune system which would have assume to boost the status of the patients health was due to negligence and lack of adherence strictly to antiretroviral drugs treatment. Of the 234 HIV/AIDS patients,

8.55% had less viral/ml volume in contrast to 10 persons with HIV/AIDS who had 100% viral load between 10,000ml-30,000 and 8 HIV/AIDS with 30,000 and 319,000ml higher per value/ml volume and very low was their CD4 counts exhibiting more clinical symptoms HIV/AIDS full blown vulnerability, at this time most patients were passing out putrid diarrhoea, pale with unsteady movement, blisters of the mouth-lips and other undesirable symptoms.

Table 4. Microsporidium and viral load among HIV/AIDS infected Patients.

Viral load range	No. of HIV/AIDS Patients	No Positive with microsporidium spp (%)
20ml-10,000ml	234	20 (8.55)
10,000ml-30,000ml	10	10 (100.00)
30,000ml-319000ml	8	8 (100.00)
Total	252	38 (15.08)

Though, in the sex related distribution of microsporidium, among the 46 HIV/AIDS infected individuals, 30.43% males had viral load value/ml between 513ml-92,300ml and 11.65%

females (926ml-319,000ml) most suppressed with the illness possibly that, females are most vulnerable due to their physiology and co-relate the ages between 51-60 without

microsporidium had their viral load within the range of 20ml-23ml, this can be seen clearly that the infected patients had less viral load unlike those with microsporidium accelerate the illness (table 6). Notwithstanding, artisans show more to have harbour more of the virus of 2590-319000 value/ml

(table 7). Though, microsporidia are important disease organisms that are considered opportunistic and this has led in recent times the improvement on the diagnostic tools to combat the organism as it show's further momentum among HIV/AIDS infected patients (table 5).

Table 5. Sex related distribution of *Microsporidium* and viral load among HIV/AIDS infected Patients.

Gender	No of HIV/AIDS Patients	No Positive with microsporidium spp (%)	Viral load range
Male	46	14 (30.43)	513ml-92,300ml
Female	206	24 (11.65)	926ml-319000ml
Total	252	38 (15.08)	

Table 6. Age related distribution of *Microsporidium* and viral load among HIV/AIDS infected Patients.

Age group	No of HIV/AIDS Patients examined	No Positive with microsporidium spp (%)	Viral load range
11 – 20	16	2 (12.50)	520ml -56,600ml
21 – 30	26	8 (30.77)	513ml-92,300ml
31 – 40	84	10 (11.90)	520ml -56,600ml
41 – 50	80	16 (20.00)	56,300ml -58,400ml
51 – 60	48	0 (0.00)	20ml-23ml
61 – 70	4	2 (50.00)	926ml-319000ml
Total	252	38 (15.08)	

Table 7. Occupational distribution of *Microsporidium* and viral load among HIV/AIDS infected Patients.

Occupation	No of HIV/AIDS Patients	No Positive with microsporidium spp (%)	Viral load range
Farmers	24	6 (25.00)	513ml-92,300ml
Students	91	14 (15.38)	20ml-56,600ml
Public /Civil Servants	130	16 (12.31)	15ml-1710ml
Unemployed	4	1 (25.00)	1890ml-58,4000ml
Artisans	3	1 (33.33)	2590ml-319000ml
Total	252	38 (15.08)	

5. Discussion

This present study showed the overall prevalence of microsporidia among HIV/AIDS positive patients attending Federal Medical Centre, Keffi (FMC), and the general prevalence rate was 15.0% and agree with the previous case study carried out among indigenous groups in Malaysia where the prevalence rate of microsporidium ranged from 15.0% to 21.2% [15, 12, 2]. The present study also showed lower prevalence rate (15.0%) compare to the study conducted in Ilorin, Nigeria with 42.4% [17]. This low prevalence could be related to location in which the study was conducted.

This study has demonstrated the effectiveness of anti-retroviral therapy which is associated with the restoration of immune response accompanying the resolution of opportunistic infection including microsporidia [13].

5.1. Age and Sex Related Distribution of *Microsporidium* Among HIV/AIDS Patients

The intensity of spores in this present study was high amongst the old age group 61-70years with 50.0% and those with very high viral load of 10,000ml and above, suggesting a relationship with the level of viral load in HIV/AIDS infected patients been similar with the study carried out in Ilorin, Nigeria [17] who reported high prevalence rate among the old aged group 52-61years with 77.3%. Samie A *et al.* [18] reported high prevalence rate of microsporidium among

aged 1-10years with 52.6% in contrast with the present study.

5.2. Marital Status and Sex Related Distribution of *Microsporidia* Among HIV/AIDS Infected Patients

Microsporidia was common among the males with 30.43% compared to the female with 11.65%. Those widower and singles had prevalence rate of microsporidia to about 50.0% and 33.33% compared to 18.18% among the married, there was however no significant difference ($X^2_{cal} = 6.10 < X^2_{tab} = 11.07$; $df = 5$; $p > 0.05$).

The present study is in agreement with the study carried out [16] with male had 35.50% and female 24.40%

The present study disagreed with the earlier studies carried out [18] who reported high prevalence rate (34.1) in the females compared to 26.5% among HIV/AIDS patients that indicated as male.

5.3. Occupation and *Microsporidium* Related HIV/AIDS Infection

The occupation of the HIV/AIDS infected patients significantly affected the prevalence of microsporidial infection with Artisans having infected with microsporidium (33.33%). Artisans were more likely to eat food and drink water from questionable sources as they carry out their work activities. This may be the reason for the high rate of prevalence in this particular group. This agrees with the findings of Speich B *et al.* [19] who observed similar risk

work factor among HIV/AIDS individuals. However, contrary to the findings in North-central, Nigeria [10].

6. Conclusion

In conclusion, this study shows that microsporidia are important pathogens capable of causing opportunistic infections in severely immunodeficiency HIV-infected patients. The prevalence of microsporidia was high and was associated with high viral load in the present study.

The improvement in diagnostic methods and greater awareness has resulted in microsporidia infections being increasingly recognized in humans. The presence of microsporidia in water sources and in pets and food producing animals, along with epidemiological risk factors that have associated exposure to water and eating undercooked meat with microsporidiosis in HIV-infected individuals have further raised the concern that microsporidia infections may be food and water borne parasitic zoonosis. Additional epidemiological studies focusing on risk factors associated with microsporidiosis will define more clearly the sources of microsporidia in the environment that pose a risk for transmission so that better preventive strategies can be implemented.

7. Recommendation

1. I recommend that routine laboratory screening need to be performed for microsporidia in the hospital setting and those positive results be made notifiable.
2. Continued studies also are needed to identify with better accuracy the presence of viable and infectious microsporidia that may pose a risk for transmission from various environmental sources.
3. Methods to remove or inactivate microsporidia in water sources and food still need to be developed, and more effective, less toxic drugs are needed for effectively treating microsporidiosis in human and animals.

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