



Prevalence of Vaginitis among Women Attending Antenatal Care Centre of Wudil General Hospital, Kano State Nigeria

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Abstract

Vaginitis is usually caused by a change in the normal balance of vaginal microbes, an infection or reduced oestrogen levels after menopause leading to variety of infections by bacteria, fungi and viruses. The study aimed to determine the prevalence of vaginitis and its associated risk factors among women attending antenatal care center of Wudil general hospital, Kano Nigeria. A total of 352 High Vaginal Swab and serum samples were collected from women attending antenatal care centre of Wudil general hospital. The samples were processed and screened for the presence of *Candida albicans*, *Trichomonas vaginalis* and *Treponema pallidum* through culture and microscopy and serum venereal disease research laboratory (VDRL) test. The risk factors were assessed using a designed questionnaire and analysed using Multivariate logistic modelling. The prevalence of *C. albicans*, *T. vaginalis* and *Treponema pallidum* were found to be 32.38%, 18.46 %, and 4.83%, respectively ($p \leq 0.05$). The result showed that the risk factors that are significantly associated with the prevalence of vaginitis were active sex age (26 - 30) years, educational status and number of participants husband's wives. The study identifies that *C. albicans* is the major cause of vaginitis, and is insignificantly associated with active sex age educational status and number of participants husband's wives.

Keywords: *Candida albicans*, *Trichomonas vaginalis*, *Treponema pallidum*, Pregnant Women

INTRODUCTION

Vaginitis is a common medical problem in women that is associated with substantial discomfort and frequent medical visits, roughly 5 to 10 million hospital visits per year are attributed to vaginitis (Novakowska *et al.*, 2004). Moreover, the availability of over-the-counter medications for fungal vaginitis has increased the likelihood that women will come to medical attention with partially or inadequately treated infections (Adetunde *et al.*, 2011).

The vaginal ecosystem undergoes major compositional changes throughout a woman's life. From childhood until puberty, the limited presence of estrogen implies a low bacterial content in the vagina, which occurs during reproductive years (Mohanty *et al.*, 2007). On the other hand, during menopause, as estrogen levels drop, not only the epithelial fragility of vaginal mucosa is affected, due to the decrease of thickness of the different cellular layers, it also becomes susceptible to infectious diseases (Deorukhkar and Saini, 2013). Therefore the vagina loses much of its self-cleaning ability

and natural defenses hence the privilege of vaginitis.

Vaginitis may be due to infectious agents including bacteria such *Treponema pallidum*; protozoans like *Trichomonas vaginalis* and fungi such as *Candida albicans* (Abdulsadah *et al.* 2014). Accurate diagnosis of the etiological agent and determination of the susceptibility pattern of the agent to certain antibiotic pose a significant effect while dealing with antimicrobial drugs resistance (Dharma *et al.*, 2013).

Candida albicans, *Treponema pallidum* and *Trichomonas vaginalis* are the most common cause of vaginitis causing vaginal discharge, vaginal and vulval pruritis, dysuria and dyspareunia (Bradshaw *et al.*, 2012). *Candida* yeast is the most common cause of opportunistic mycoses worldwide (Ahmad *et al.*, 2016).

Culture is considered as the "gold standard" in diagnosing *Candida vaginitis*. It can also be diagnosed by microscopic detection of yeast cells on a vaginal smear along with the presence of a white, mucous-like vaginal discharge on physical examination.

Saline or 10% KOH wet mount may show yeast cells and pseudohyphae (Bradshaw *et al.*, 2012). *Candida* culture is usually done on Sabouraud's dextrose agar and Gram stain of the colony shows gram positive yeast cells. Germ tube test can be used for differentiation of *Candida albicans* species from nonalbicans species. *Candida* spp. can also be identified serologically, using specific antisera. DNA probes are also available for *Candida* spp detection (Dariane *et al.*, 2012).

Trichomonas vaginitis is more often difficult to diagnose because of its heterogeneous presentation. Vaginal pH is often more than 4.5. Many cases give a positive amine (KOH) test ("whiff" test). *T. vaginalis* was traditionally diagnosed via a wet mount, in which polymorphonuclear response and a "corkscrew" motility can be observed (Alli *et al.*, 2011). The most common method of diagnosis is by culture. Despite the low cost, culture of trichomonas is not routinely practised in many laboratories. Culture requires Diamond's medium. The In Pouch TV culture system is now found to be a good alternative to traditional culture techniques. Other methods like rapid antigen testing and transcription-mediated amplification are not in widespread use (Kissinger and Adamski, 2013). The presence of *T. vaginalis* can also be diagnosed by direct fluorescent antibody test, enzyme immune assay and PCR.

Treponema pallidum is most commonly spread bacteria through sexual activity. It may also be transmitted from mother to baby during pregnancy or at birth, resulting in congenital syphilis (Akinbiyi *et al.*, 2008). Diagnosis is usually made by using blood tests; the bacteria can also be detected using dark field microscopy.

While interventions aiming at promoting safer sex, such as condom use, and vaginitis etiological diagnosis and treatment, in high risk populations are widely accepted and advocated for (Abdulsadah *et al.*, 2014). It's also possible that sexual practices influence *C. albicans* infection, as well as other reproductive tract infections (Gill *et al.*, 2011; Ballini *et al.*, 2012). While the requirements for *T. vaginalis* to develop itself in the genital tract vary from those for *C. albicans* and *T. pallidum*, the coexistence of these two or three microorganisms in the genital tract has been discovered in several women (Levi *et al.*, 2011; Lopez-Monteon *et al.*, 2013). There has been no record of this incident in the general hospital in Wudil, as far as we know.

As a result, the aim of this analysis was to evaluate the etiological agent of vaginitis, the rate of incidence, and potential risk factors among Wudil general hospital's female patients.

MATERIAL AND METHODS

Ethical clearance for the study was obtained from Kano state ministry of health. High Vaginal Swab and serum samples were aseptically collected through the assistance of the medical personnel from 352 randomly selected women attending antenatal care centre of Wudil general hospital, Kano.

The samples were aseptically collected using sterile cotton swab stick and venipuncture and labelled appropriately. Patient's information regarding their ages, pregnancy age, educational level and number of participants' husband's wives were documented using questionnaire. Samples collected were immediately transported to the laboratory unit of Wudil general hospital for immediate analysis (Gill *et al.*, 2011; López-Monteon *et al.*, 2013).

Identification of *Trichomonas vaginalis*

The presence of *Trichomonas vaginalis* in the sample was detected using wet mount preparations according to the methods described by Akinbo *et al.* (2017). A small portion of the swab was suspended in one drop of 0.85% physiological saline on a slide and covered with a cover slip. The wet mount preparation was then examined for the characteristic morphology and darting motility under X10 and X40 objectives.

Isolation and Identification of *Candida albicans*

The swab samples were inoculated each on the surface of previously dried Sabouraud dextrose agar plate as described by Uzoh *et al.* (2016). The plates were incubated at room temperature for 48 hours and were examined for white cream colonies characteristic of *Candida* spp. Small inoculum of suspected *Candida* cultures was inoculated into 0.5 ml of human serum in a test tube and was incubated at 37° C for 3 hours. After incubation, a loop-full of culture was placed on a glass slide, overlaid with a cover-slip and was then examined microscopically for the presence or absence of germ-tubes. Formation of germ tubes was seen as long tube like projections extending from the yeast cells with no constriction or septa at the point of attachment to the yeast cells. This is a confirmatory test for the identification of *Candida albicans* (Saha *et al.*, 2018).

Venereal Diseases Research Laboratory Test for Antibodies/Antigens of *Treponema pallidum*

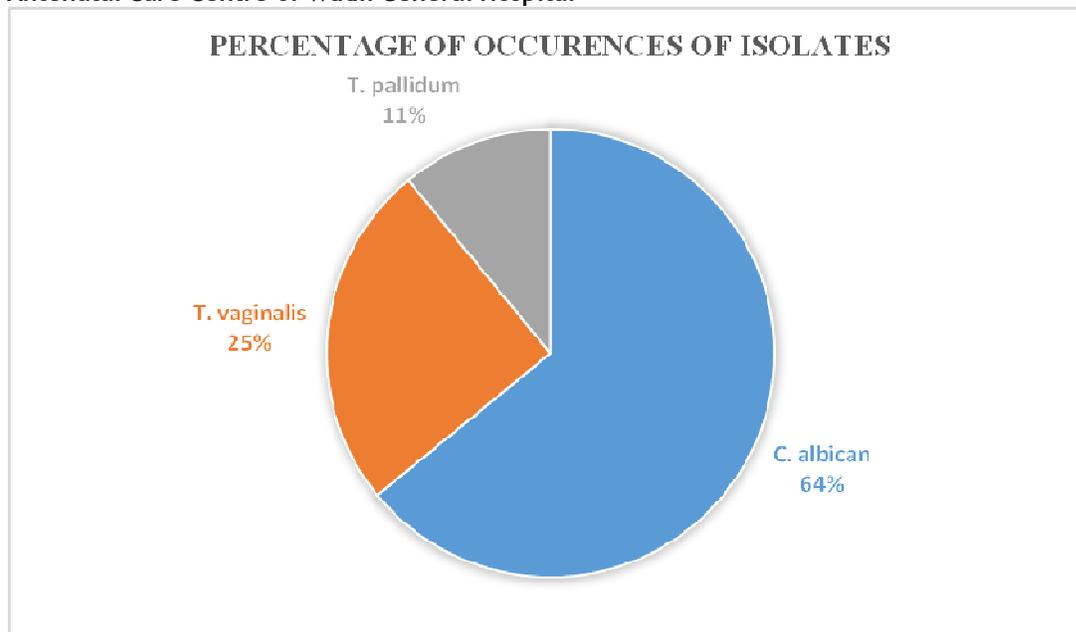
The VDRL test was carried out to detect antibodies/antigens of *Treponema pallidum* according to the methods described by Rapikit Neclife, (2019). The blood sample was centrifuge to obtain the serum. Two drops of serum was placed into the sample window using provided disposable dropper. Appearance of

two colour line in both control and test region indicated a positive results.

RESULTS

The result of the study revealed that out of the 352 studied samples, 114 (32.38%) were identified as *C. albicans*, 65 (18.46%) as *T. vaginalis* and 17 (04.83%) were *Treponema pallidum* positive (VDRL+) (Table 1).

TABLE 1: Prevalence of *C. albicans*, *T. vaginalis* and *T. pallidum* among Women Attending Antenatal Care Centre of Wudil General Hospital



Findings also shows that age range 26-30 had the highest prevalence rate of 71 (20.17%) of which 41 (11.6%) were identified as *C. albicans*, 23 (6.53) as *T. vaginalis* and 7 (1.99%) *T. pallidum* while least positive prevalence 18 (5.11%) was at age range of 31-35 with 6 (1.70%) for *C. albicans*, 7 (1.99%) *T. vaginalis* and 5 (1.42%) *T. pallidum* (Table 2)

TABLE 2: Prevalence *C. albicans*, *T. vaginalis* and *T. pallidum* in Relation to Age

Age group (years)	No. sampled	No. Positive (%)	<i>C. albicans</i> (%)	<i>T. vaginalis</i> (%)	<i>T. pallidum</i> (%)
16-20	81	29 (8.23)	21 (5.97)	8 (2.27)	0 (0)
21-25	93	44 (12.5)	28 (7.95)	13 (3.69)	3 (0.85)
26-30	103	71 (20.17)	41 (11.65)	23 (6.53)	7 (1.99)
31-35	43	18 (5.11)	6 (1.70)	7 (1.99)	5 (1.42)
>35	32	34 (9.66)	18 (5.11)	14 (3.98)	2 (0.57)
Total	352	196 (55.68)	114 (32.38)	65 (18.46)	17 (4.83)

Moreover, highest positive prevalence of 83 (23.58%) was among women with none number of co-wives, 54 (15.35%) were identified as *C. albicans*, 23 (6.33%) as *T. vaginalis* and 6 (1.70%) *T. pallidum* whereas category women

with three co-wives revealed the least positive prevalence of 10 (2.84%), 6 (1.70%) were identified as *C. albicans* and 4 (1.14%) as *T. vaginalis* (Table 3).

TABLE 3: Prevalence of *C.albicans*, *T.vaginalis* and *T. Pallidum* in Relation to Number of Co-wives

No. of Co-Wives	No. sampled (%)	No. Positive (%)	<i>C. albicans</i> (%)	<i>T. vaginalis</i> (%)	<i>T. pallidum</i> (%)
None	147 (41.76)	83 (23.58)	54 (15.35)	23 (6.53)	6 (1.70)
One	128 (36.36)	71 (20.17)	35 (9.94)	27 (7.67)	9 (2.56)
Two	59 (16.76)	32 (9.09)	19 (5.39)	11 (3.12)	2 (0.57)
Three	18 (5.11)	10 (2.84)	6 (1.70)	4 (1.14)	0 (0)
Total	352 (100)	196 (55.68%)	114 (32.38)	65 (18.46)	17 (4.83)

Furthermore, highest positive prevalence of 84 (23.85%) was found among women with secondary education, 47 (13.35%) were identified as *C. albicans*, 26 (7.38%) as *T. vaginalis* and 11 (3.12%) as *T. pallidum*. While women with Islamiyya education has the least

positive prevalence of 20 (5.69%), 11 (3.13%) and 9 (2.56%) (Table 4).

Findings also shows the co-prevalence of 53 (15.06%) for *C. albicans* and *T. vaginalis*, 23 (6.53%) for *C. albicans* and *T. pallidum*, 16 (4.55%) for *T. vaginalis* and *T. pallidum* (Table 5).

TABLE 4: Prevalence of vaginitis in Relation to Educational Status of the Participants

Status	No. sampled (%)	No. Positive (%)	<i>C. albicans</i> (%)	<i>T. vaginalis</i> (%)	<i>T. pallidum</i> (%)
Islamiyya	37 (10.51)	20 (5.69%)	11 (3.13)	9 (2.56)	0 (0)
Primary	118 (33.52)	61 (17.32%)	41 (11.64)	18 (5.11)	2 (0.57)
Secondary	151 (42.90)	84 (23.85%)	47 (13.35)	26 (7.38)	11 (3.12)
Tertiary	46 (13.07)	31 (8.81%)	15 (4.26)	12 (3.41)	4 (1.14)
Total	352 (100)	196 (55.68%)	114 (32.38)	65 (18.46)	17 (4.83)

TABLE 5: Co infection OF *C. albicans*, *T. vaginalis* and *T. pallidum* among the Participants.

Agents	Co-infection(%)
<i>C. albicans</i>	53 (15.06)
<i>T. vaginalis</i>	
<i>C. albicans</i>	23 (6.53)
<i>T. pallidum</i>	
<i>T. vaginalis</i>	16 (4.55)
<i>T. pallidum</i>	
<i>C. albicans</i>	0(0)
<i>T. vaginalis</i>	
<i>T. pallidum</i>	

DISCUSSION

Result of this study revealed a significant difference between the prevalence of *C. albicans* compared to *T. vaginalis* and *T. Pallidum* at (P<.05) base on the frequency of occurrence. This is line with the study of Alo *et al.*, (2012) in Abakaliki and Aniebue *et al.* (2018) in Enugu with (43.00%) and (49.53%) all in Nigeria, this is attributed to the nature of tight under wares use in the vaginal area leading to formation of moisture in combination with organic releases that favour the growth and multiplication of *C. albicans* in the area. Less prevalence of *T. pallidum* among the participants was in agreement with the finding

of Bakare *et al.*, (2002) in Ibadan, Dada *et al.*, (1998) in Lagos, Aboyeji and Nwabuisi, (2003) and that of Taiwo *et al.*, (2006) all in Nigeria who’s reported a prevalence range of 0.125% to 4.1%.

The higher percentage prevalence of all the three agent of vaginitis found within the age range of 26 - 30 years agree with the finding of Garba *et al.* (2014) in Nigeria who reported vaginitis to be most prevalent among 26-30 age group (35.8%) and least in >40 years age group (10.5%). This prevalence might be attributed to most active reproductive age group and high sexual exposure.

Throughout this period, the ovary produces excessive amount of estrogen, which favours the growth of microbes by maintaining the acidic pH in the vagina and enhancing the yeast adherence to vaginal epithelial cells (Adetunde *et al.* 2011). This also explains why there is lower prevalence at age 45years and above.

The higher prevalence of *C. albicans* among those without a partner could be due to their lifestyle rather than cross infection, as one would predict given the higher prevalence of *T. vaginalis* and *T. pallidum* among those with one partner.

Correspondingly, women with secondary educational level had the highest vaginitis prevalence of 23.85% which differs from Ibrahim *et al.* (2014), who recorded the highest prevalence of 54% in those with primary education in Maiduguri, Nigeria. The low economic status, lack of education, lack of a female consultant at the health service center, hesitance to approach medical service, and sociocultural structure might be the cause of higher prevalence of vaginitis among less educated women while life style is the major cause of higher prevalence in educated women. Correlation of *C. albicans* and *T. vaginalis* co-infection shows a prevalence of co-infection of 53(15.06%). This high rate of co-infection

recorded in this work is lower than that observed by Alo *et al.*, (2012) and (2016) who recorded a high prevalence co-infection rate of (43.00%) and (32.14%) in Abakaliki and Ebonyi respectively. Also correlation of *C. albicans* and Syphilis co-infection shows a prevalence of co-infection of 23(6.06%)

CONCLUSION

C. albicans was the major etiological agent of vaginitis isolated in this study followed by *T. vaginalis* and to a lesser effect *T. pallidum*. Co-infections of *C. albicans* with *T. Vaginalis* were high compared to other possible co-infection and no case of co-infection with both the three agents recorded. Since high prevalence were noticed within the age (26 - 30) when one is very sexually active, which concurs with the utmost of one's reproductive life, there is need for a quick and rapid cognizance campaign for infection prevention and control as this will lead to barrenness, fatal wastage and even divorce if left untreated. The low economic status, lack of hygiene education, lack of a female consultant at the health service center, hesitance to approach medical service, and sociocultural structure were the predisposing factors for vaginitis in the study area.

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