



## ***Trichomonas vaginalis* and *Candida albicans* Infections among Women of Reproductive Age in Ihiala, Anambra State, Nigeria**

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### **Authors' contributions**

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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### **ABSTRACT**

**Background:** Co-infection of *T. vaginalis* and *C. albicans* continues to present major health, social and economic problems in Nigeria.

**Aim:** A study to determine the prevalence and co-infection of *T. vaginalis* and *C. albicans* infections among women of reproductive age in Ihiala, Anambra state.

**Study Design:** The study was a cross-sectional survey involving 734 women from four villages in Ihiala community.

**Duration:** The study was conducted between January and March 2015.

**Materials and Methods:** Wet mounts were used to examine the high vaginal swabs of the women. Jerky movement and flagella were used to confirm the presence of *T. vaginalis*. Germ-tube test was used to confirm *C. albicans* after culturing on sabouraud dextrose agar.

**Results:** Of the 734 women, 150(20.4%) were positive to *C. albicans*, 7(1.0%) were positive to *T. vaginalis* and 4(0.5%) had co-infection of *T. vaginalis* and *C. albicans*. Women in the age group 40-49 years had the highest infection 22(37.9) of *C. albicans* and the age group 20-29 years had the

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lowest 96(17.0). Women in age group, 30-39 years had the highest co-infection of *C. albicans* and *T. vaginalis* 2(1.8%), while age group 20-29 years had the lowest 2(0.4%). Co-infections were highest 2(6.7%) among traders and none among civil servants and students. Co-infection of the pathogens was observed in married women only, and highest 1(3.6%) among wives of polygamous men. Co-infections of *T. vaginalis* and *C. albicans* was highest 1(14.3%) among the women with non-formal education and none among the women with tertiary education. Co-infections of the pathogens were observed only among non-pregnant women.

**Conclusion:** Since most co-infections were observed among women with no formal education and those from polygamous families, mass education on reproductive health to the women will help reduce the scourge of these infections in the study area.

**Keywords:** *Trichomonas vaginalis*; *Candida albicans*; women; Ihiala; Anambra state.

## 1. INTRODUCTION

*Trichomonas vaginalis* is most likely the world's most common non-viral sexually transmitted infection. It is a serious public health issue affecting women of reproductive age [1]. Poor personal cleanliness, frequent sex partners, low socioeconomic level, and underdevelopment have all been linked to a higher risk of infection [2]. Other sexually transmitted infections, such as gonorrhoea, chlamydia, and sexually transmitted viruses, are also linked to *Trichomonas vaginalis* infection. Other viruses, such as herpes, human papillomavirus (HPV), and human immunodeficiency virus (HIV), become more susceptible as a result of the infection (HIV) [3].

The condition is caused by a pear-shaped protozoan called *T. vaginalis* [4], which causes frothy-greenish-foul-smelling vaginal discharge, vulvo-vaginal irritation, dysuria, lower stomach aches, discomfort, and psychosocial distress in female patients [5,6]. Other problems include cystitis, cervicitis, and urethritis [7]. These symptoms are generally increased during menses and pregnancy. Premature rupture of membranes, premature labour, slow labour, low birth weight, abortion, post-hysterectomy infection, infertility, and neoplastic change in cervical tissues are all described *Trichomonas vaginalis* complications in pregnant women. [8,9,10].

In Nigeria, spread of Sexually Transmitted Diseases (STDs) has been blamed on increase in poverty, unemployment and violence among women [11,12]. Sexual recklessness, lack of awareness, ignorance of the public health implications, poor sanitation and poor personal hygiene are other risk factors of trichomoniasis. Prevention of *T. vaginalis* infection has not been

a priority due to lack of understanding of its public health implications and lack of resources [13]. Trichomonal infections are most common in those between the ages of 16 and 35 [14], while they can also affect people in their later years [15].

Candidiasis on the other hand is associated with vaginal discharge and pruritis. The discharge appears like curded milk and deep erythema of vulva and vagina is often seen [16]. The incidence of the infection is almost doubled in pregnancy mostly in the third trimester. It has been estimated that up to 40% of pregnant women world-wide may have vaginal colonization by *Candida* species, a two-fold increase from the prevalence rate in non-pregnant women [17]. Among various pathogenic species of fungi, *Candida* is the most prominent cause of fungal infections [18]. The genus *Candida* is composed of heterogeneous group of organisms and more than 17 different species are implicated in human infections [19]. The most important pathogenic species is *Candida albicans*. The high frequency of STDs and reproductive tract infections is due to a combination of factors including low socioeconomic position, a lack of knowledge, a lack of diagnostic facilities, and a lack of appropriate treatment options [20,21].

The aim of this study was to investigate *T. vaginalis* and *C. albicans* infections among women of reproductive age in Ihiala. The specific objectives were to determine the prevalence of co-infection of *T. vaginalis* and *C. albicans* by age, marital status, gravidity, pregnancy status, educational status, occupational groups and contraceptive usage, and the treatment habits of *T. vaginalis* and *C. albicans* infections among women in the study area.

## 2. MATERIALS AND METHODS

### 2.1 The Study Area

The study was conducted in Ihiala community, Ihiala Local Government Area, Anambra State, Southeastern Nigeria. Ihiala is the local administrative capital of the area. The community is situated between Longitude 6<sup>o</sup>.86' East and Latitude 5<sup>o</sup>.85' North of the equator, within the tropical rainforest zone of Nigeria. It has about 7 months of wet season period (April to October) and 5 months of dry season (November to March) with a short period of harmattan season. The relative humidity of the area is about 70% in the dry season reaching 80% during wet season [22]. The annual rainfall is between 2,250 to 2,500mm. The average daily temperature range of the area during the dry season is 27-33<sup>o</sup>C and 21.1<sup>o</sup>C-30<sup>o</sup>C in wet season. The community is about 144m above the sea level. Ihiala has a population of 87,796 inhabitants [23]. The area is an Igbo speaking community living peacefully with people of other ethnic groups. The inhabitants are predominantly, farmers, traders, students and a few civil servants.

### 2.2 Study Design

The study was a cross-sectional survey of women of reproductive age in the community. Quantitative and qualitative methods were used to determine the prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates among the participants. The study was conducted from January to March 2015.

Ethical approval to conduct the study was obtained from the Ethics Committee of Our Lourdes Hospital, Ihiala, Anambra

### 2.3 Advocacy Visit and Mobilization of Participants

The participants were mobilized through town criers and public notices made in the churches within the villages. Health education on the health implications of infections with *T. vaginalis* and *C. albicans* was given to the participants. They were assured of confidentiality of individual test results. The study was conducted in line with the international guideline for experimentation [24].

### 2.4 Study Population and Sample Size Determination

A total of seven hundred and thirty-four (734) apparently healthy women without complains of

infection were enrolled. The sample size (n) was determined using Taro Yamane formula [25] for a finite population. The formula was given as:

$$n = \frac{N}{1 + N(e)^2}$$

Where:

- n = the minimum sample size
- N = the population size
- e = level of significance or limit of tolerable error (0.05)
- 1 = unity (a constant)

### 2.5 Sampling Technique

The women were assembled at the Akwa Health Centre, Ihiala and at Our Lady of Lourdes Hospital complex, Ihiala, for examination. Convenience sampling in which each woman was attended to as she arrived was used. Hospital examination rooms were used to provide each participant with maximum privacy. Trained nurses and laboratory scientists assisted in swab collection and examination.

### 2.6 Specimen Collection

#### 2.6.1 Collection of Bio-data of the participants

A total of 734 structured questionnaires were used to obtain the biodata of the participants such as age, village, pregnancy status, occupation, marriage structure, gravidity, educational status and marital status.

#### 2.6.2 Collection of high vaginal swabs

With the help of two medical laboratory scientists, samples of vaginal discharge were carefully obtained aseptically from the women's high vaginal area using well-labeled, sterile non-abrasive swab sticks. The vaginal swabs were tested in the medical laboratory shortly after they were collected. *T. vaginalis* was diagnosed using wet mounts, and *C. albicans* was cultured from swabs.

### 2.7 Parasitological Examination of the Specimens

#### 2.7.1 Diagnosis of *T. vaginalis* (Wet – Mount Preparation)

Each high vaginal swab from the women was mixed with a drop of sterile normal saline on a

clean grease-free slide, covered with cover slip and examined with x10 objectives lens of the microscope. The presence of yeast cell, *T. vaginalis* and other deposits were observed. *Trichomonas vaginalis* was identified by the jerky movement, the flagella, axostyle and its oval shape. Ten percent (10%) Potassium hydroxide (KOH) was added to the wet preparation to increase the detection sensitivity of yeast cells, making the recognition of mycelia (pseudohyphae) much easier [26].

### 2.7.2 Diagnosis of *C. albicans* (Culture Method)

Each high vaginal swab was inoculated on Sabouraud Dextrose Agar gel and incubated at the temperature of 37<sup>o</sup> C for 48hrs. Suspected yeast colonies and positive colonies were gram stained and further sub-cultured for germ tube test [26].

## 2.8 Data Analysis

The statistical differences in the prevalence and co-infection of *T. vaginalis* and *C. albicans* in the various groups were tested using chi-square test. Chi-square statistics was computed to accompany each cross tabulation. The statistical package used was SPSS version 21.0.

## 3. RESULTS

Out of the 734 women, 150(20.4%) were positive to *C. albicans*, 7(1.0%) were positive to *T. vaginalis* and 4(0.5%) had co- infection of *T. vaginalis* and *C. albicans* (Table 1).

The highest prevalence of *C. albicans* 22(37.9%) was observed in age group 40-49 years while age group 20-29 years had the least 2(9.1%). The age group 30-39 years had the highest prevalence of (Table 2). There was a significant difference in prevalence of *T. vaginalis* and *C. albicans* in age groups (P<0.05).

Ubahiekwem 3 (6.3%) had the highest prevalence of *T. vaginalis* while Ihudim village had the least 1 (3.3%). Akwa village had the highest prevalence of *C.albicans* 23 (41.1%), while the least was observed in Eziani, 102(17.0%). The co-infection was observed only among Ubahiekwem and Akwa village 3 (6.3%) and 1 (1.8%) respectively (Table 3). There was a significant difference in the prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates in the villages (P<0.05).

The traders had the highest prevalence of *T. vaginalis*, 4(13.3%) and co-infection of *T. vaginalis* and *C. albicans* 2(6.7%). There was a significant difference in the prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates among women from different occupational groups (P<0.05) (Table 4).

The married women had the highest prevalence of *C. albicans*, and the widows had the least 1(11.5%). Only the married women had co-infection of *T. vaginalis* and *C. albicans* 4(1.7%) among the marital groups (Table 5). The prevalence of *T. vaginalis* and co-infection rates had significant difference (P<0.05) while the prevalence in *C. albicans* had no significant difference (P>0.05).

The highest co-infection of 1(14.3%) was observed among the non-formal educational group and the least was among secondary educational group, 1(1.7%) (Table 6). The prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates among the educational groups showed a significant difference (P<0.05).

The women from polygamous families had higher co-infection rate 1(3.6%) of *T. vaginalis* and *C. albicans* than the women from monogamous families 3(1.5%). The prevalence of *T. vaginalis* and co-infection rates were significantly difference (P<0.05) while the prevalence in *C. albicans* had no significant difference in marital structure(P>0.05) (Table 7).

The pregnant women had higher infection of *T. vaginalis* 1(2.3%) than the non-pregnant women 6(0.9%). There was no co-infection of *T. vaginalis* and *C. albicans* among the pregnant women. The prevalence of *T. vaginalis* and co-infection rates showed no significant difference (P>0.05) while the prevalence in *C. albicans* showed significant difference (P<0.05) (Table 8).

*Trichomonas vaginalis* infection was highest among primigravida, 2(2.9%) and least 1(0.2%) among the nulligravida. *Candida albicans* infection was highest among the multigravida, 67(32.5%) and the least 9(12.9%) among the primigravida (Table 9). The multigravida had a higher rate of co-infection 3(1.5%) than the primigravida 1(1.4%). There was no significant difference in the prevalence of *T. vaginalis*, *C. albicans* and their co-infection rates among the study participants (P<0.05).

**Table 1. Prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates among the women participants from Ihiala**

No. Examined	No. infected with <i>C. albicans</i> (%)	No. infected with <i>T. Vaginalis</i> (%)	No. co-infected with <i>T. Vaginalis</i> and <i>C. albicans</i> (%)
734	150 (20.4)	7 (1.0)	4 (0.5)

**Table 2. Prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates by age groups**

Age Group (Years)	No. Examined	No. infected with <i>T. vaginalis</i> (%)	No. infected with <i>C. albicans</i> (%)	No. co-infected with <i>T. vaginalis</i> and <i>C.albicans</i> (%)
20-29	564	2 (0.4)	96 (17.0)	2 (0.4)
30-39	112	4 (3.6)	32 (28.6)	2 (1.8)
40-49	58	1 (1.7)	22 (37.9)	0
Total	734	7 (0.95)	150 (20.4)	4 (0.5)

$$\chi^2 = 3.876, df = 3, P < 0.05$$

**Table 3. Prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates by location**

Villages examined	No. Examined	No. infected with <i>T. vaginalis</i> (%)	No. infected with <i>C. albicans</i> (%)	No. co-infected with <i>T. vaginalis</i> and <i>C.albicans</i> (%)
Eziani	600	0	102 (17.0)	0
Ubahiekwem	48	3 (6.3)	18 (37.5)	3 (6.3)
Akwa	56	3 (5.4)	13 (41.1)	1 (1.8)
Ihudim	30	1 (3.3)	7 (23.3)	0
Total	734	7 (0.95)	150 (20.4)	4 (0.5)

$$\chi^2 = 3.876, df = 3, P < 0.05$$

**Table 4. Prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates by occupation**

Occupation	No. Examined	No. infected with <i>T. vaginalis</i> (%)	No. infected with <i>C. albicans</i> (%)	No. co-infected with <i>T. vaginalis</i> and <i>C.albicans</i> (%)
Students	600	0	102 (17.0)	0
Civil Servants	18	0	3 (16.7)	0
Trader	30	4 (13.3)	10 (33.3)	2 (6.7)
Farmers	86	3 (3.5)	35 (40.7)	2 (2.3)
Total	734	7 (0.95)	150 (20.4)	4 (0.5)

$$\chi^2 = 29,169, df = 3 P < 0.05$$

**Table 5. Prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates by marital status**

<b>Marital Status</b>	<b>No. Examined</b>	<b>No. infected with <i>T. vaginalis</i> (%)</b>	<b>No. infected with <i>C. albicans</i> (%)</b>	<b>No. co-infected with <i>T. vaginalis</i> and <i>C.albicans</i> (%)</b>
Singles	468	0	91 (19.4)	0
Married	234	6 (2.6)	55 (23.5)	4 (1.7)
Divorced	6	0	1 (16.7)	0
Widowed	26	1 (3.8)	3 (11.5)	0
<b>Total</b>	<b>734</b>	<b>7 (0.95)</b>	<b>150 (20.4)</b>	<b>4 (0.5)</b>

$$\chi^2 = 8.594, df = 3 P < 0.05$$

**Table 6. Prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates by educational status**

<b>Educational Status</b>	<b>No. Examined</b>	<b>No. infected with <i>T. vaginalis</i> (%)</b>	<b>No. infected with <i>C. albicans</i> (%)</b>	<b>No. co-infected with <i>T. vaginalis</i> and <i>C.albicans</i> (%)</b>
Non-formal	7	1 (14.3)	4 (57.1)	1 (14.3)
Primary	42	3 (7.1)	22 (52.4)	2 (4.8)
Secondary	60	3 (5.1)	25 (41.7)	1 (1.7)
Tertiary	625	0	99 (15.8)	0
<b>Total</b>	<b>734</b>	<b>7 (0.95)</b>	<b>150 (20.4)</b>	<b>4 (0.5)</b>

$$\chi^2 = 42.983, df = 3, P < 0.05$$

**Table 7. Prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates by marital structure**

<b>Marital Structure</b>	<b>No. Examined</b>	<b>No. infected with <i>T. vaginalis</i> (%)</b>	<b>No. infected with <i>C. albicans</i> (%)</b>	<b>No. co-infected with <i>T. vaginalis</i> and <i>C.albicans</i> (%)</b>
Monogamy	206	5 (2.4)	44 (21.4)	3 (1.5)
Polygamy	28	1 (3.6)	11 (39.3)	1 (3.6)

$$\chi^2 = 656, df = 1, P < 0.05$$

**Table 8. Prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates by pregnancy status**

<b>Pregnancy Status</b>	<b>No. Examined</b>	<b>No. infected with <i>T. vaginalis</i> (%)</b>	<b>No. infected with <i>C. albicans</i> (%)</b>	<b>No. co-infected with <i>T. vaginalis</i> and <i>C.albicans</i> (%)</b>
Pregnant	44	1 (2.3)	19 (43.2)	0
Non-Pregnant	690	6 (0.9)	131 (18.98)	4 (0.5)
<b>Total</b>	<b>734</b>	<b>7 (0.95)</b>	<b>150 (20.4)</b>	<b>4 (0.5)</b>

$$\chi^2 = 256, df = 3, P < 0.05$$

**Table 9. Prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates by gravidity**

<b>Gravidity No. examined</b>	<b>No. Examined</b>	<b>No. infected with <i>T. vaginalis</i> (%)</b>	<b>No. infected with <i>C. albicans</i> (%)</b>	<b>No. co-infected with <i>T. vaginalis</i> and <i>C. albicans</i> (%)</b>
Nulligravidae	458	1 (0.2)	4 (57.1)	0
Primigravidae	70	2 (2.9)	9 (12.9)	1 (1.4)
Multigravidae	206	4 (1.9)	67 (32.5)	3 (1.5)
Total	734	7 (0.95)	150 (20.4)	4 (0.5)

$\chi^2 = 6.675, df = 3, P < 0.05$

**Table 10. Prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates by clinical manifestation among the women participants**

<b>Clinical Manifestation</b>	<b>No. Examined</b>	<b>No. infected with <i>T. vaginalis</i> (%)</b>	<b>No. infected with <i>C. albicans</i> (%)</b>	<b>No. co-infected with <i>T. vaginalis</i> and <i>C. albicans</i> (%)</b>
Symptomatic	248 (33.8)	2 (0.8)	81 (32.7)	2 (0.8)
Asymptomatic	486 (66.2)	5 (1.0)	69 (14.2)	2 (0.4)
Total	734 (100)	7 (0.95)	150 (20.4)	4 (0.5)

$\chi^2 = 0.473, df = 2, P < 0.05$

**Table 11. Prevalence of *T. vaginalis* and *C. albicans* and their co-infection rates according to treatment habit of the women participants**

<b>Treatment Habit</b>	<b>No. Examined</b>	<b>No. infected with <i>T. vaginalis</i> (%)</b>	<b>No. infected with <i>C. albicans</i> (%)</b>	<b>No. co-infected with <i>T. vaginalis</i> and <i>C. albicans</i> (%)</b>
Use of Metronidazole	Yes 520	1 (0.2)	122 (23.5)	0
	No 214	6 (2.8)	28 (13.1)	4 (1.9)
Self-medication of antibiotics	Yes 534	6 (1.1)	138 (25.80)	3 (0.6)
	No 200	1 (0.5)	12 (6.0)	1 (0.5)
Use of Contraceptives	Yes 186	5 (2.7)	122 (65.6)	2 (1.1)
	No 548	2 (0.4)	28 (5.1)	2 (0.4)

$\chi^2 = 4.800, df = 3, P < 0.05$

Asymptomatic women 5 (1.0%) had higher prevalence of *T. vaginalis* than the symptomatic women (Table 10). The co-infection occurred more among the symptomatic women 2(0.8) than asymptomatic women 2(0.4). The prevalence of *C. albicans* showed significant difference ( $P>0.05$ ) while the prevalence in *T. vaginalis* and co-infection rates showed no significant difference ( $P<0.05$ ).

Women using Metronidazole always, had the highest prevalence of *C. albicans* infection, 122(23.5%) while those who do not take Metonidazole always, had the least, 28(13.1%). There was no statistical difference between them ( $P>0.05$ ).

#### 4. DISCUSSION

The high prevalence of *C. albicans* (20.4%) reported in this study could be due to the ability of the pathogen as a frequent colonizer and responsible for most cases of vulvovaginitis [27,28]. The result is higher than some studies [29,30,31,32,33] who reported 2.0%, 2.20%, 12.0%, 2.6%, and 17.8% respectively. However, this is lower than other studies [16,34,35] who reported the prevalence rate of 28.0%, 52.5% and 33.6% respectively. It has previously been observed that high incidence of *C. albicans* could be associated with severe immunosuppression or illness, and sexual activities of a woman [36]. Some studies noted that *Candida* species are part of the lower genital tract flora in 20-50% of healthy asymptomatic women and may spread due to sexual activities or immunosuppression [37,38].

High prevalence of *C. albicans* was recorded among age groups 20-29 years (17.0%), 30-39 years (28.6%) and 40-49 years (55.6%). This disagrees with the findings of a previous study [30] who observed that the infections were almost uniformly distributed in all age groups studied. But it agrees with [39] who reported the prevalence of *C. albicans* to be significantly higher in cohort of 30 years Old Dutch women and lower in the cohorts of 45-50, 55 and 60 years. The age range of between 20-49 years constitute the sexually active period of most women. Although *C. albicans* infection is not a conventional sexually transmitted disease, it is known to be spread in some cases through sexual intercourse, particularly among sexually active people and sex workers, according to prior studies [19]. It was also noted that the high prevalence rate discovered in the younger age

group was ascribed to the group being the most sexually active age range, bolstering the assumption that sexual activity may play a big role in the disease's transmission [40].

The low prevalence rate of trichomoniasis (1.0%) recorded in this study is in tandem with low prevalence of (2.3%) and (3.3%) reported in some studies [41,42]. Also, other investigations reported prevalence rate of 0.2% [43] and 1.2% [44]. However, some studies have reported higher prevalence rates of 24.1% [45] and 24% [46]. Low prevalence of *T. vaginalis* in this study could be attributed to multiple factors such as awareness of people towards unsafe sex. The use of protection during sex help in reducing transmission of pathogenic organisms especially *T. vaginalis* that affects the vaginal mucosa in women [47].

The co-infection rate of trichomoniasis and candidiasis in this study was 0.5%. This compares favorably with a previous observation [35] who reported co-infection rate of 0.6% of *T. vaginalis* and *C. albicans* in Lagos metropolis. Also, another study [48] reported co-infection rate of 3.2% which is slightly higher. However, higher rate of co-infection has been reported in other studies including 22% observed in Abakaliki, Ebonyi State, Nigeria [46]. Low co-infection rate of the parasites observed in this study could be attributed to improved life style, especially in personal hygiene and literacy in reproductive health, as well as the use of Metronidazole in treatment of other protozoa infections.

#### 5. CONCLUSION

Infections caused by *T. vaginalis* and *C. albicans*, as well as their co-infections, are found among women of reproductive age in Ihiala, according to the study. The information gathered will serve as a foundation for developing policies and control mechanisms to combat the illnesses. It will also be useful in training reproductive-aged women to be more aware of their reproductive health. As a result, it is advised that widespread identification and treatment, as well as health education, will aid in the eradication of illnesses in the study region.

#### CONSENT

Informed written consent was obtained from each woman before enrolment.



## COMPETING INTERESTS

Authors have declared that no competing interests exist.

## REFERENCES

1. Kissinger P. *Trichomonas vaginalis*: A review of epidemiologic, clinical and treatment issues. BMC Infectious Diseases. 2015;15:307
2. Crosby R, DiClemente RJ, Wingwood GM. Predictions of infection with *Trichomonas vaginalis*: A prospective study of low - income African-American adolescent females. Sexually Transmitted Infections. 2002;78:360-364.
3. Huppert JS. Trichomoniasis in teens: an update. Current Opinion in Obstetrics and Gynecology. 2009;21(5):371-378.
4. Schwebke JR. Update of trichomoniasis. Sexually Transmitted Infections. 2002;78: 378-379.
5. Faro S. Vaginitis. Differential diagnosis and management. New York: Parthenon Publishing Group; 2004.
6. Edrisian Q, Rezaiian M, Qorbani M, Keshavarz H, Mohebbali M. *Medical protozoology*. 1 edition Tehran: Tehran universal publication. 2006;85-95.
7. Alcamo IE. *Fundamentals of microbiology*. Jones and Bartlett Publishers, Boston. 2000;486-487.
8. Uneke CJ, Ugwuonu CDC, Ali E, Ali M. *Trichomonas vaginalis* infections among pregnant women in South Eastern Nigeria: The public health significance. The International Journal of Gynecology and Obstetrics. 2006;6(1):17-21.
9. Perazzi BE, Menghi CI, Coppolillo EF, Gatta C, Eliseth MC, De Torres RA. Prevalence and comparison of diagnostic methods for *Trichomonas vaginalis* infection in pregnant women in Argentina. Korean Journal of Parasitology. 2010;48(1):61-5.
10. Soper D. "Trichomoniasis: under control or under controlled?". American Journal of Obstetrics and Gynecology. 2004;190(1): 281–299.
11. Obiajuru IOC. The prevalence of *Trichomonas vaginalis* infection in Imo State Nigeria, Unpublished M. Sc Thesis Imo-State University Owerri Nigeria; 2004.
12. Ulogu IO, Obiajuru IOC, Ekejindu IM. Trichomoniasis among women in Nnewi Nigeria. Nigerian Journal of Parasitology. 2007;28(1):7-10.
13. Schwebke JR, Burgess D. "Trichomoniasis". Clinical Journal of Microbiology Reviews. 2004;17(4):794–803.
14. Naguils S. Epidemiology of trichomoniasis in normal females. Journal of Obstetrics Gynecology. 1966;27:607-620.
15. Onyido AE, Umeanaeto PU, Irikannu KC, Ekwunife CA, Ezeanya LC, Nwangwu UC, Ugha CN, Obiechina IO. Prevalence of *Trichomonas vaginalis* Among the Rural Women of Ekwulumili Community Anambra State, Southeastern Nigeria. Journal of Natural Sciences. 2014;12(5):129-134.
16. Khan AS, Amir F, Altaf S, Tanveer R. Evaluation of common organisms causing vaginal discharge. Journal of Ayub. Medicine Coll. Abbottabad. 2009;21(2): 90- 93.
17. Hay P, Czeizel AE. Asymptomatic *Trichomonas* and *Candida* colonization and pregnancy outcome. Basic Practice and Research Clinical Obstetrics Gynecology. 2007;21(3):403-409.
18. Sullivan D, Henman M, Moran G. Molecular genetic approaches to identification, epidemiology and taxonomy of non-*albicans* *Candida* species. Journal of Medical Microbiology. 1996;44:399-408.
19. Pfaller MA, Diekema DJ. Epidemiology of Invasive Candidiasis: A Persistent Public Health Problem. Virulence. 2007(2):119–128.
20. Tyadal SP, Zaaijamann JD, Kent HL. An unusual vaginal foreign body. African Medical Journal. 1992;61(33):108-110.
21. Burrow RC, Bueshing UJ. Bacterial vaginosis in virgins and sexually active females: Evidence against exclusive sexual transmission. Journal of Obstetrics Gynecology. 1999;8(5):97-99.
22. Microsoft Encarta. Microsoft Corporation; 2009.
23. National Population Commission. Population Census of Nigeria. Population distribution in Local Government Areas by Sex and Number of Households; 2006.
24. World Medical Association Declaration of Helsinki, WMADH. Ethical principles for medical research involving human subjects. Curable Sexually Transmitted Diseases. *Overview and Estimates*; 2000.
25. Uzoagulu AE. Practical guide to writing research project reports in tertiary institutions. Cheston Ltd. Enugu. 2011;57-58.

26. Cheesbrough M. *Microbiological test in: District laboratory practice for tropical countries part 2*, Cambridge Low Price Edition. Cambridge University Press. 2000; 69-71.
27. Donbraye E, Okonkwo IO, Okedeji IO. Detection and prevalence of *Trichomonas vaginalis* among pregnant women in Ibadan, South Western Nigeria. *World Applied Science Journal*. 2010;11(12):1512-1517.
28. Alli JAO, Okonko IO, Odu NN, Kolade AF, Nwanze JC. Detection and prevalence of *Candida* isolates among patients in Ibadan, Southwestern Nigeria. *Journal of Microbiology and Biotechnology Research*. 2011;1(3):176-184.
29. Choudhry S, Ramachandran VG, Das S, Bhattacharya SN, Mogha NS. Pattern of sexually transmitted infections and performance of syndromic management against etiological diagnosis in patients attending the sexually transmitted infection clinic of a tertiary care hospital. *Indian Journal of Sexual Transmitted Diseases*. 2010;31:104-8.
30. Konje JC, Otolorin EO, Ogunniyi JO, Obisesan KA, Ladipo OA. The prevalence of *Gardnerella vaginalis*, *Trichomonas vaginalis* and *Candida albicans* in the Cytology clinic at Ibadan Nigeria. *African Journal Medical Sciences*. 1991;20(1):29-34.
31. Nwokedi EE, Aniyam NN. A study of high vaginal swabs in Kano teaching hospital- A preliminary study. *Highland Medical Research Journal*. 2003;1:57-61.
32. Cronje HS, Joubert G, Muir A, Chapman RD, Divan P, Barn RH. Prevalence of vaginitis, syphilis and HIV infection in women in the Orange Free State. *South African Medical Journal*. 1994;84:602-605.
33. Di Bartolomeo S, Rodriguez Fermepin M, Sauka DH, Alberto de Torres R. Prevalence of associated micro-organisms in genital discharge, Argentina. *Review Saude Publication*. 2002;36(5):545-552.
34. Muvunyi CM, Hernandez CT. Prevalence of bacterial vaginosis in women with vaginal symptoms in south province, Rwanda. *African Journal Clinical Experimental Microbiology*. 2009;10(3):156-153.
35. Adeoye GO, Akande AH. Epidemiology of *Trichomonas vaginalis* among women in Lagos Metropolis, Nigeria. *Pakistan Journal of Biological Science*. 2007;10:2198-2201.
36. Horn DL, Neofytos D, Anaissie EJ, Fishman JA, Steinbach WJ, Olyaei AJ, Marr KA, Pfaller MA, Chang CH, Webster KM. Epidemiology and outcomes of candidemia in 2019 patients: data from the prospective antifungal therapy alliance registry. *Journal of Clinical Infectious Diseases*. 2009;48: 1695–1703.
37. McClelland RS, Richardson BA, Hassan WM, Graham SM, Kiarie J, Baeten JM, Mandaliya K, Jaoko W, Ndinya-Achola JO, Holmes KK. Prospective study of vaginal bacterial flora and other risk factors for vulvovaginal candidiasis. *Journal of Infectious Diseases*. 2009;15:199(12):1883-1890.
38. Akah PA, Nnamani CE, Nnamani PO. Prevalence and treatment outcome of vulvovaginal candidiasis in pregnancy in a rural community in Enugu State, Nigeria. *Journal of Medicine and Medical Sciences*. 2010;1(10):447-452.
39. Engberts MK, Vermeulen CF, Verbruggen BS, Van Haaften M, Boon ME, Heintz AP. *Candida* and squamous (pre) neoplasia of immigrants and Dutch women as established in population- based cervical screening. *Internet Journal of Gynecological Cancer*. 2006;16(4):1596-1600.
40. Ononge S, Wondabwa J, Kionto P, Busigye R. Clinical presentation and management of alleged sexually assaulted females at Muago Hospital, Kampala, Uganda. *African Journal of Health Science*. 2005;5(1):50-54.
41. Orji NM. Reproductive tract infections among females in Ihiala Local Government Area, Anambra State, Nigeria. *International Journal of Scientific Engineering and Applied Science*. 2015;1(3):22-28
42. Gundiri MA, Okwuosa VN. Prevalence of urinary and intestinal tracts parasites in Kwampe, Langtong, North Nigeria. *Nigerian Journal of Parasitology*. 2005;26:19-22.
43. Kalantari N, Ghaffari S, Bayani M. *Trichomonas*, *Candida*, and *Gardnerella* in cervical smears of Iranian women for cancer screening. *North America Journal of Medical Science*. 2014;6:25-90.
44. Nsagha DS, Zofou D, Nguedia Assob JC, Njunda AL, Nchang CD, MvoNgum N, Weledji EP, Ngowe NM. The epidemiology of *Trichomonas vaginalis*, *Gardnerella vaginalis* and *Candida albicans* co-Infections in women attending the Yaounde University Teaching Hospital. *American Journal of Epidemiology and Infectious Diseases*. 2015;3(2):28-31.
45. Jombo GT, Opayobi SO, Hobbs NM. Trichonomiasis. *The Internet Journal of parasitic diseases* 2007;8:67-88.

46. Alo MN, Anyim C, Onyebuchi AK, Okonkwo EC. Prevalence of asymptomatic Co-Infection of Candidiasis and Vaginal Trichomoniasis among Pregnant Women in Abakaliki, South-Eastern Nigeria. *Journal of Natural Sciences Research*. 2012;2(7): 2224-3186.
47. Evans BA, Bond PD, MacRae KD. Heterosexual relationship and Condom use in the spread of sexual transmitted diseases to woman. *Genito-urinary Medicine*. 1995; 71(5):291-294.
48. Olorode OA, Ogba ON, Ezenobi NO. Urogenital trichomoniasis in women in relation to candidiasis and gonorrhoea in University of Port-Harcourt Teaching Hospital. *African Journal of Microbiology Research*. 2014;8(2):2482-2485.

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