

# Evaluating the Antifungal, Antibacterial, Serum Electrolytes, Anti-inflammatory Markers of Zingiber Officinale Crude Extract on Candida Albicans Infected Wistar Rats

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## ABSTRACT

**Background:** Vaginal candidiasis is a fungal infection mainly caused by *Candida albicans*, an opportunistic pathogen responsible for candidiasis in humans. The proliferation of *C. albicans* is favored by environmental conditions such as low pH, high humidity, and moisture, which facilitate its colonization and overgrowth. Disruption of the normal vaginal microflora or local immune defenses can trigger infection, resulting in the clinical manifestations associated with vaginal candidiasis.

**Aim:** The study was carried out to aimed to assess the antifungal and antibacterial potential, serum electrolytes and anti-inflammatory markers of *Zingiber officinale* (ginger) using standard methods.

**Method:** The study was carried out in two phases, in vivo and in vitro. In the in vitro experiment, *Candida albicans* was cultured in SDA at 25 °C and exposed to the drug fluconazole and serial concentrations of the extract ranging from 300 mg/ml to 9.37 mg/ml and readings taken. In the in vivo experimental, 30 albino wistar rats were divided into 6 groups which includes; normal control (uninfected untreated), negative control (infected untreated), infected + standard drug (fluconazole [1000 mg/kg body weight {BW}]), and infected + extract at 250 mg/kg BW, 500 mg/kg BW and 1000 mg/kg BW; and its effect on some biochemical markers were assessed.

**Result:** The extract demonstrated strong antifungal efficacy, with a minimum inhibitory concentration (MIC) of 18.75 µg/mL and a fungicidal effect confirmed by the minimum bactericidal concentration (MBC), indicating complete elimination of *C. albicans*. The assessment of electrolyte parameters revealed significant alterations associated with *Candida albicans* infection and subsequent treatment with *Zingiber officinale*. Sodium (Na<sup>+</sup>) levels ranged from 131.50 ± 2.10 mmol/L in the negative control group to 144.00 ± 1.00 mmol/L in the 500 mg/kg ginger-treated group. The assessment of inflammatory and anti-inflammatory cytokines (TNF-α, IL-6, and IL-4) revealed the negative control group exhibited elevated pro-inflammatory cytokines TNF-α (34.80 ± 2.40 pg/ml) and IL-6 (165.40 ± 8.70 pg/ml) with a comparatively low anti-inflammatory cytokine IL-4 (28.60 ± 3.90 pg/ml).

**Conclusion:** The reduced sodium concentration observed in the infected untreated group suggests that *Candida albicans* infection may disrupt electrolyte balance, possibly through impaired renal regulation, dehydration, or inflammatory-mediated sodium loss. Elevated TNF-α and IL-6 are well established indicators of systemic inflammation and immune activation. Administration of plant extract at 250, 500, and 1000 mg/kg resulted in consistent reductions in TNF-α levels and IL-6 levels, while IL-4 levels increased. The standard drug group showed comparable effects.

**Keywords:** *Candida albicans*; Antifungal; Antibacterial; Anti-inflammatory; *Zingiber officinale*.

## INTRODUCTION

Nowadays infectious diseases are day by day increasing in prevalence (Rodriguez-Morales et al., 2023). Fungal infections were less common than bacterial and viral infections in the past but have been responsible for a dramatic increase of diseases in recent decades (Banerjee et. al., 2021). Yeasts especially *Candida* species are the most

common fungi that have been isolated from human infections, in recent years (Katsipoulaki et al., 2024). *Candida albicans* is known as the most common human pathogen between 150 species of *Candida* yeast and cause of 60 percent of candidiasis infection due to most virulence. However, incidence of infection by other species such as *C. tropicalis*, *C. glabrata* and *C. krusei* has grown increasing (Gómez-Gaviria & Mora-Montes, 2020). *Candida* is an opportunistic fungus that is capable of inducing acute and chronic infections of the mouth, vagina, lung and gastrointestinal tract with different body reactions such as severe infections, purulent and granulomatous inflammation (Ostrosky-Zeichner & Sobel, 2023). It can be found in the mouth, vaginal and intestine microflora and cause fatal infections and mortality in people with weakened immune systems such as AIDS, cancer, bone marrow or other organs transplant.

Pregnancy, oral contraceptive drugs, antibiotics, diabetes, long-term skin contact with water, treatment with topical corticosteroids, some diseases of the endocrine system are involved in causing illness by yeast (Zapata et al., 2023). Studies in the last decades have clearly indicated the unpleasant side effect of synthetic drugs. Recent studies considered to the use of medicinal plants due to increasing drug resistance to chemical drugs (Moiketsi et al., 2023).

Using detergents, frequent shower or internal disorders can disrupt the vaginal flora, consisting of lactic acid bacteria such as lactobacilli and resulting an overgrowth of *Candida* cells and vaginal candidiasis (Chen et al., 2017). Medical benefits of medicinal plants in various drugs and as pharmaceutical products are important (Nath et al., 2023). Medicinal plants are applied in various fields of medicine, industry, agriculture, food and numerous applications (Hatami et al., 2024). The main causes of this trend are low side effects of natural products and high side effect of synthetic drugs and environmental pollution (Abdallah et al., 2023).

*Zingiber officinale* (known as Chitta in Hausa language and Ginger in English) belongs to the family Zingiberaceae. They are tuberous or non-tuberous rhizomes, which have a particular aroma and possess different medicinal properties. It is a flowering plant whose rhizome (ginger root) is widely used as a spice and for its medicinal properties. Ginger has been used in traditional medicine systems, including Ayurveda, Chinese medicine, and Western herbalism, for thousands of years. In the traditional medicine, the gnarly, thick underground stem (rhizome) of ginger is commonly used (Yadufashije et al., 2020).

## MATERIALS AND METHODS

### Reagents and Plant Materials

Solvents and chemicals used in the experiment were analytical grade and purchased from Jos, Nigeria. Distilled water was used throughout the experiment. Fresh matured samples of *Zingiber officinale* (Ginger) plant was harvested from a farm at Bethany Christian Academy Barkin Ladi, Plateau State. The plant material was identified and authenticated and the voucher number was obtained at the Federal college of Forestry Jos, Plateau State, Nigeria.

### Antifungal and Antibacterial Activity

The inhibition zone diameter (IZD), minimum inhibitory concentration (MIC) and minimum bacteriocidal concentration (MBC) were assessed using disk diffusion method as described by Scorzoni et al., (2007).

2.3 Determination of in vivo serum electrolytes (Potassium, sodium, chloride, and  $\text{HCO}_3^-$ ) and anti-inflammatory markers (IL-4, IL-6 and  $\text{TNF-}\alpha$ ) were assessed using Enzyme-Linked Immunosorbent Assay (ELISA).

## RESULTS

### Phytochemical Constituents of the Plant Extract

Table 1 shows some phytochemicals that were found to be present in the aqueous and extract of *Zingiber officinale* (Ginger). Alkaloids, flavonoids, tannins, saponins, terpenoids, cardiac glycosidase, balsam, carbohydrates, phenols, resins and anthraquinone were found to be present while anthraquinone and balsam were

absent in the aqueous extract while some phytochemicals that were found to be present in the metanol extract of *Zingiber officinale* (Ginger) were alkaloids, flavonoids, tannins, saponins, terpenoids, cardiac glycosidase, balsam, carbohydrates, phenols, resins, anthraquinone and balsam.

Table 2 shows the quantity of some phytochemicals that were present in the aqueous extract of *Zingiber officinale*. Alkaloids had the highest amount with a percentage of  $6.11 \pm 0.03\%$  and steroids was found to be the least in quantity with  $0.06 \pm 0.01\%$  while some of the phytochemicals that were present in the methanol extract of *Zingiber officinale* as shown on the table. Flavonoids had the highest amount with a percentage of  $8.60 \pm 0.23\%$  and anthraquinone was found to be the least in quantity with  $2.17 \pm 0.01\%$ .

Table 1. Phytochemical Content of Aqueous and Methanol Extract of Ginger (*Zingiber officinale*)

Chemical Constituents	Aqueous <i>Z.officinale</i>	Methanol <i>Z.officinale</i>
Alkaloids	+	+
Flavonoids	+	+
Tanins	+	+
Saponin	+	+
Terpenoids	+	+
Cardiac glycosidase	+	+
Balsam	-	+
Carbohydrates	+	+
Phenols	+	+
Resins	+	+
Anthraquinone	-	+

Key: +=Present -=Absent

Table 2. Quantitative Phytochemical Constituents of Aqueous and Methanol Extract of *Zingiber officinale* in Percentage (%)

S/N	Phytochemical	Aqueous <i>Z. officianale</i>	Methanol <i>Z. officinale</i>
1	Alkaloids	$6.11 \pm 0.03$	$5.12 \pm 0.03$
2	Saponins	$4.01 \pm 0.05$	$3.12 \pm 0.11$
3	Phenols	$5.61 \pm 0.01$	$7.11 \pm 0.10$
4	Tannins	$5.10 \pm 0.02$	$6.11 \pm 0.12$
5	Flavonoids	$2.30 \pm 0.03$	$8.60 \pm 0.23$
6	Anthraquinone	$0.42 \pm 0.03$	$2.17 \pm 0.01$

7	Reducing sugars	0.13 ± 0.02	3.27 ± 0.11
8	Terpenoids	0.51 ± 0.03	4.83± 0.12
9	Steroids	0.06 ± 0.01	2.19 ± 0.02

All values are mean ± SEM

### MIC Inhibition Zone Diameter (IZD) of Zingiber officinale against C.albicans

Table 3 shows the result of the inhibition zone diameter (IZD) of Zingiber officinale against C. albicans by well diffusion method. The result showed that the rate of inhibitory zones fluctuated between 25-30 mm,  $p > 0.05$  in the preparation of C.albicans compared to the drug control (fluconazole).

Table 4 shows the result of the minimal inhibitory concentration (MIC) and the minimal bactericidal concentration (MBC). The MIC of Zingiber officinale was recorded at 18.75 mg/ml which indicates significant positive effect on the microorganism (C.albicans) when compared with the drug control group. The result of the MBC indicates a significant positive bactericidal effect recorded at concentration group (9.37, 18.75, 37.5, 75, 150, 300) mg/ ml.

Table 3. MIC Inhibition Zone Diameter (IZD) of Zingiber officinale against C.albicans

Concentration (µg/mL)	IZD (mm)	Relative efficacy Vs Control (%)
300	30	107.14
200	27	96.43
100	25	89.29
Drug control (fluconazole)	28	100.00

Concentration dependent inhibitory effect; 300 µg/mL dose more effective than standard drug control with significance  $P > 0.05$ ,  $n=3$ .

Table 4. In vitro Minimal inhibitory concentration (MIC) and Minimal bactericidal concentration (MBC) of Zingiber officinale against C.albicans

Concentration (µg/ml)	MIC	MBC
300	-	-
150	-	-
75	-	-
37.5	-	-
18.75	-	-
9.37	+	-

KEY: += Slight growth, -= No growth

### In vivo Serum Electrolytes Analysis

Table 5 shows the result of serum electrolytes which includes; sodium (Na), potassium (K), chloride (Cl) and bicarbonate (HCO<sub>3</sub>).

Table 5. Serum Electrolytes

GROUP	Na (mmol/L)	K (mmol/L)	Cl (mmol/L)	HCO <sub>3</sub> (mmol/L)
NEG. CONT.(mg/kg)	131.50 ± 2.10	5.45 ± 0.42	94.00 ± 1.90	20.30 ± 1.70
NORM. CONT. (mg/kg)	137.00±0.58	4.33±0.28	99.00±0.58	25.33±0.88
STD (mg/kg)	140.33±1.67	4.83±0.18	99.66±0.33	26.33±1.45
250 (mg/kg)	139.33±0.33	4.73±0.38	101.00±0.58	24.00±1.16
500 (mg/kg)	144.00±1.00	4.66±0.23	102.00±0.58 <sup>a</sup>	24.00±1.53
1000 (mg/kg)	141.67±0.67	4.66±0.37	103.33±0.881 <sup>a</sup>	25.66±1.33
p-value	0.0051	0.0180	0.0336	0.0418

Values are expressed as mean ± SEM, n = 3.

<sup>a</sup>Values are statistically significant when compared to Normal Control.

### In vivo Anti-inflammatory Markers

Table 6 shows the result of anti-inflammatory markers which includes; Tumor Necrosis Factor alpha (TNF-α), Interleukin (IL-4) and Interleukin (IL-6).

Table 6. Anti-Inflammatory Markers

GROUP	TNF-a (pg/ml)	IL-4 (pg/ml)	IL-6 (pg/ml)
NEG. CONT. (mg/kg)	34.80 ± 2.40	28.60 ± 3.90	165.40 ± 8.70
NORM. CONT. (mg/kg)	24.86±0.69	45.95±5.59	114.51±3.42
STD (mg/kg)	21.47±0.89	49.90±3.39	101.94±1.99
250 (mg/kg)	20.85±0.97	39.86±0.72	103.31±3.19
500 (mg/kg)	20.50±1.35	41.35±0.53	108.89±1.15
1000 (mg/kg)	21.58±0.75	41.25±1.18	113.55±2.14
p-value	0.0574	0.1795	0.0159

Values are expressed as mean ± SEM, n = 3

<sup>a</sup>Values are statistically significant when compared to Normal Control

## DISCUSSION

A medicinal plant is any plant which contains substances that can be used for therapeutic purpose, also serve as precursors for the synthesis of useful drugs (Erb & Kliebenstein, 2020). Preliminary phytochemical screening of the extract of *Z. officinale* revealed the presence of various bioactive components of which alkaloids, flavonoids, tannins, saponins, terpenoids, cardiac glycosidase, balsam, carbohydrates, phenols and resins were the most prominent. Phytochemicals such as alkaloids, terpenoids, saponins and phenolics are considered strong antimicrobial and antifungal agents that can aid in solving the problem of antibiotic resistance (Anwer et al., 2017). Alkaloids, phenols, tannins and flavonoids were found in high amounts. Tannins have been found to possess antifungal activity as reported by Agi and Azike, (2019). Alkaloids which are one of the largest groups of phytochemicals in plants have led to the development of powerful pain killer medications as well as antifungal as reported by (Doughari, 2016). It has been recognized that flavonoids show antioxidant activity and their effects on human nutrition and health are considerable. The mechanisms of action of flavonoids are through scavenging or chelating process. Flavonoids are important in plant biochemistry and play an important role in plant physiology, acting as antioxidants, enzyme inhibitors, pigments and light screens (Ghasemzadeh et al., 2010). Flavonoids have been shown to have antioxidant, antibacterial anti-inflammatory, antiallergic activity etc (Riyazi et al., 2007).

The assessment of electrolyte parameters revealed significant alterations associated with *Candida albicans* infection and subsequent treatment with *Zingiber officinale*. Sodium ( $\text{Na}^+$ ) levels ranged from  $131.50 \pm 2.10$  mmol/L in the negative control group to  $144.00 \pm 1.00$  mmol/L in the 500 mg/kg ginger-treated group. The reduced sodium concentration observed in the infected untreated group suggests that *Candida albicans* infection may disrupt electrolyte balance, possibly through impaired renal regulation, dehydration, or inflammatory-mediated sodium loss (Liamis et al., 2010). The normal control group exhibited higher sodium levels, reflecting physiological electrolyte balance. Ginger-treated groups showed marked increases in sodium levels, indicating a restorative effect on electrolyte homeostasis. This improvement may be attributed to the ability of ginger to enhance renal function and reduce inflammation-induced electrolyte disturbances (Rahmani et al., 2014; Nile & Park, 2015). Notably, the p-value (0.0051) indicates a statistically significant difference among groups ( $p < 0.05$ ), confirming that ginger treatment significantly influenced sodium regulation. Potassium ( $\text{K}^+$ ) levels were highest in the negative control group, suggesting impaired renal excretion or cellular ion imbalance during infection. Elevated potassium is often associated with renal dysfunction and tissue breakdown (Palmer & Clegg, 2016). Ginger-treated groups demonstrated potassium levels closer to normal control values, suggesting a normalization of potassium homeostasis. Therefore, with statistical significance ( $p = 0.0180$ ) indicates that these changes can be conclusively attributed to treatment. Chloride ( $\text{Cl}^-$ ) levels were lowest in the infected untreated group, indicating electrolyte imbalance and possible metabolic disturbances. Ginger-treated groups showed progressive increases in chloride levels, suggesting improved electrolyte regulation and fluid balance. This may reflect the role of ginger in modulating renal ion transport and maintaining osmotic balance (Nile & Park, 2015). The p-value (0.0336) indicates a statistically significant difference among groups ( $p < 0.05$ ), demonstrating that ginger treatment significantly affected chloride levels. Bicarbonate ( $\text{HCO}_3^-$ ) levels were reduced in the negative control group, suggesting the presence of metabolic acidosis or acid–base imbalance, which is commonly observed in systemic infections (Kraut & Madias, 2012). Ginger-treated groups showed improved bicarbonate levels, indicating a potential role in restoring acid base equilibrium. This effect may be linked to improved renal function and reduced systemic inflammation. These differences were statistically significant ( $p = 0.0418$ ). The overall electrolyte profile indicates that *Candida albicans* infection disrupts electrolyte and acid base balance, as evidenced by reduced sodium and chloride levels, elevated potassium levels, and decreased bicarbonate concentrations. These alterations may result from renal dysfunction, dehydration, or systemic inflammatory responses associated with infection (Liamis et al., 2010; Kraut & Madias, 2012). Treatment with *Zingiber officinale* extract (250 mg/kg–1000 mg/kg) however generally improved electrolyte balance, including significant increases in sodium and chloride levels, normalization of potassium levels, and improved bicarbonate concentrations. Among the treatment groups, the 1000 mg/kg dose showed the most pronounced improvement, with the highest sodium and chloride levels, suggesting that higher doses may be more effective in restoring electrolyte homeostasis. Importantly, statistically significant differences were observed across groups, indicating that ginger treatment had a measurable effect on these electrolytes. These findings suggest that ginger may help correct electrolyte imbalances induced by *Candida albicans* infection, potentially through improved kidney

function, enhanced fluid regulation, and its anti-inflammatory and antioxidant properties (Rahmani et al., 2014; Nile & Park, 2015).

The effects of ginger treatment on inflammatory and anti-inflammatory cytokines (TNF- $\alpha$ , IL-6, and IL-4) are summarized in Table 6. The negative control group (infected untreated) exhibited elevated pro-inflammatory cytokines TNF- $\alpha$  ( $34.80 \pm 2.40$  pg/ml) and IL-6 ( $165.40 \pm 8.70$  pg/ml) with a comparatively low anti-inflammatory cytokine IL-4 ( $28.60 \pm 3.90$  pg/ml). Elevated TNF- $\alpha$  and IL-6 are well-established indicators of systemic inflammation and immune activation (Cerami, 1989; Kishimoto, 2005). Administration of ginger at 250, 500, and 1000 mg/kg resulted in consistent reductions in TNF- $\alpha$  levels ( $20.85 \pm 0.97$ ,  $20.50 \pm 1.35$ , and  $21.58 \pm 0.75$  pg/ml, respectively) and IL-6 levels ( $103.31 \pm 3.19$ ,  $108.89 \pm 1.15$ , and  $113.55 \pm 2.14$  pg/ml), while IL-4 levels increased ( $39.86 \pm 0.72$ ,  $41.35 \pm 0.53$ , and  $41.25 \pm 1.18$  pg/ml). The standard drug group showed comparable effects (TNF- $\alpha$ :  $21.47 \pm 0.89$  pg/ml; IL-6:  $101.94 \pm 1.99$  pg/ml; IL-4:  $49.90 \pm 3.39$  pg/ml). Modulation of these cytokines is a recognized mechanism for evaluating anti-inflammatory potential (Dinarello, 2000). Statistical analysis revealed a significant reduction in IL-6 ( $p = 0.0159$ ), whereas TNF- $\alpha$  ( $p = 0.0574$ ) and IL-4 ( $p = 0.1795$ ) did not reach significance. Nevertheless, the observed biological trends remain meaningful, reflecting relevant anti-inflammatory activity (Altman, 1995). The suppression of TNF- $\alpha$  aligns with prior studies highlighting its central role in mediating inflammation and tissue damage during infection (McInnes & Schett, 2007). IL-6 inhibition, particularly at 250 mg/kg, underscores ginger's anti-inflammatory potential, consistent with its established capacity to regulate pro-inflammatory signaling pathways (Kishimoto, 2005). Elevated IL-4 levels further suggest an immune-regulatory effect, promoting an anti-inflammatory environment and suppression of pro-inflammatory cytokine production (Paul, 1991; O'Garra, 2007).

## CONCLUSION

In conclusion, anti-Candida albicans activity of the methanol extract of *Zingiber officinale* (Ginger) was evaluated and the results revealed that the plant act significantly against *Candida albicans*. This might be as a result of the phytochemicals found to be present that makes it act against microorganisms. The findings of the present study suggested that *Z. officinale* could be a potential source of natural antioxidant that could have great importance as therapeutic agents in preventing or slowing the progress of aging and age associated oxidative stress related degenerative diseases. Systemic infection with *Candida albicans* induces widespread physiological derangements due to a combination of inflammatory immune activation, oxidative stress, and organ dysfunction. In this study, untreated infected rats (negative control) exhibited multiple pathological changes. Treatment with graded doses of ginger extract (250 mg/kg –1000 mg/kg) resulted in coordinated improvements across all domains, indicative of multi-system modulation of host responses by the bioactive constituents of *Zingiber officinale*. Sodium and chloride levels were significantly restored in ginger-treated rats, indicating improved renal electrolyte regulation and fluid balance. Electrolyte disturbances during systemic infection are linked to inflammation-mediated renal tubular dysfunction and dysregulated ion transport. Ginger likely improved electrolyte homeostasis through anti-inflammatory mechanisms and enhanced renal perfusion, contributing to normalized ion exchange. A defining pathological feature of *Candida* infection is robust pro-inflammatory cytokine activation. Elevated TNF- $\alpha$  and IL-6 in the negative control group reflect an activated innate immune response, while IL-4, an anti-inflammatory Th2 cytokine was suppressed. Ginger treatment shifted this balance: it significantly reduced IL-6 ( $p < 0.05$ ) and lowered TNF- $\alpha$ , while elevating IL-4. This pattern supports immune modulation, suppression of excessive pro-inflammatory signaling and enhancement of anti-inflammatory pathways. Mechanistically, ginger has been shown to inhibit nuclear factor- $\kappa$ B (NF- $\kappa$ B) activation, resulting in reduced expression of TNF- $\alpha$ , IL-6, and other inflammatory mediators. The increase in IL-4 further suggests that ginger may steer immune responses toward regulatory profiles, diminishing tissue damage while preserving antifungal defense.

## REFERENCES

1. Abdallah, E. M., Alhatlani, B. Y., de Paula Menezes, R., & Martins, C. H. G. (2023). Back to Nature: Medicinal plants as promising sources for antibacterial drugs in the post-antibiotic era. *Plants*, 12(17), 3077.
2. Agi, V. N., & Azike C. A. (2019). Antifungal action of garlic (*Allium sativum*) and ginger (*Zingiber officinale*) on some pathogenic fungi. *Asian Journal of Research in Biochemistry*, 4(4), 1-6.

3. Al-Gbouri and A. G. Hamza. (2018). Evaluation of phyllanthus mplica extract as antibacterial and antibiofilm against biofilm formation bacteria. Iraqi Journal of Agricultural Science.49(1):24-30.
4. Altman, D. G. (1995). Absence of evidence is not evidence of absence. BMJ, 311(7003), 485.
5. Anwer, N., Waqar, M. A., Iqbal, M., Mushtaq, M. and Sobia, A. (2017). Phytochemical analysis, free radical scavenging capacity and antimicrobial properties of Impatiens bicolor plant. International Food Research Journal, 20(1):3-9.
6. Azene, T. (2021). Revealing the Therapeutic Uses of Garlic (*Allium sativum*) and Its Potential for Drug Discovery. Hindawi the Scientific World Journal, 7(3):2-6.
7. Banerjee, S., Denning, D. W., & Chakrabarti, A. (2021). One Health aspects & priority roadmap for fungal diseases: A mini-review. Indian Journal of Medical Research, 153(3), 311-319.
8. Cerami, A., & Beutler, B. (1989). TNF and TNF receptors. Annual Review of Immunology, 7, 425–452.
9. Cerami, A., & Beutler, B. (1989). The role of tumor necrosis factor in the pathogenesis of disease. Annual Review of Immunology, 7, 625–655.
10. Chen, Y., Bruning, E., Rubino, J., & Eder, S. E. (2017). Role of female intimate hygiene in vulvovaginal health: Global hygiene practices and product usage. Women's Health, 13(3), 58-67.
11. Dinarello, C. A. (2000). Proinflammatory cytokines. Chest, 118(2), 503–508.
12. Doughari, J. H. (2016). Phytochemicals: extraction methods, basic structures and mode of action as potential chemotherapeutic agents. Phytochemicals-A Global Perspective of their Role in Nutrition and health. PLoS Pathogens, 3:1-27.
13. Erb, M., & Kliebenstein, D. J. (2020). Plant secondary metabolites as defenses, regulators, and primary metabolites: The blurred functional trichotomy. Plant Physiology, 184(3), 39-43
14. Ghasemzadeh A., Jaafar H. Z. E., Rahmat A., Wahab P. E. M. and Halim M. R. A. (2010): Effect of Different Light Intensities on Total Phenolics and Flavonoids Synthesis and Anti-oxidant Activities in Young Ginger Varieties (*Zingiber officinale* Roscoe) Int. J. Mol. Sci. 11, 3885-3897; Doi:10.3390/ijms11103885.
15. Gómez-Gaviria, M., & Mora-Montes, H. M. (2020). Current aspects in the biology, pathogeny, and treatment of *Candida krusei*, a neglected fungal pathogen. Infection and drug resistance, 1673-1689.
16. Gonzalez, M.W., Kann, M.G., (2012). Chapter 4: Protein interactions and disease. J. PLoS Comput. Biol. 8, e1002819.
17. Kraut, J. A., & Madias, N. E. (2012). Metabolic acidosis: pathophysiology, diagnosis and management. Nature Reviews Nephrology, 8(10), 589–601.
18. Kishimoto, T. (2005). Interleukin-6: From basic science to medicine. Annual Review of Immunology, 23, 1–21.
19. Kishimoto, T. (2005). Interleukin-6: Discovery of a pleiotropic cytokine. Arthritis Research & Therapy, 7(Suppl 2), S2.
20. Liamis, G., Milionis, H. J., & Elisaf, M. (2010). Electrolyte disorders associated with infectious diseases. Clinical Microbiology and Infection, 16(7), 868–874.
21. McInnes, I. B., & Schett, G. (2007). Cytokines in the pathogenesis of rheumatoid arthritis. Nature Reviews Immunology, 7(6), 429–442.
22. McKay DL, Chen CY, Zampariello CA, and Blumberg JB (2015): Flavonoids and Phenolic Acids from Cranberry Juice are Bioavailable and Bioactive in Healthy Older Adults. Food Chem 168: 233—240.
23. Moiketsi, B. N., Makale, K. P., Rantong, G., Rahube, T. O., & Makhzoum, A. (2023). Potential of selected African medicinal plants as alternative therapeutics against multi-drug-resistant bacteria. Biomedicines, 11(10), 2605.
24. Nile, S. H., & Park, S. W. (2015). Chromatographic analysis, antioxidant, anti inflammatory, and xanthine oxidase inhibitory activities of ginger extracts. Industrial Crops and Products, 70, 238–244.
25. O'Garra, A., & Vieira, P. (2007). T(H)1 cells control themselves by producing interleukin-10 and interleukin-4. Nature Reviews Immunology, 7(6), 425–428.
26. O'Garra, A., & Vieira, P. (2007). T cell subsets and regulation of Th1/Th2 balance. Microbes and Infection, 9(6), 819–826.
27. Ostrosky-Zeichner, L., & Sobel, J. D. (2023). Candidiasis. In Diagnosis and Treatment of Fungal Infections (pp. 151-166). Cham: Springer International Publishing.
28. Palmer, B. F., & Clegg, D. J. (2016). Physiology and pathophysiology of potassium homeostasis. Advances in Physiology Education, 40(4), 480–490.

29. Paul, W. E. (1991). Interleukin-4: A prototypic immunoregulatory lymphokine. *Blood*, 77(9), 1859–1870.
30. Rahmani, A. H., Al Shabrmi, F. M., & Aly, S. M. (2014). Active components of ginger and their biological effects. *International Journal of Physiology, Pathophysiology and Pharmacology*, 6(2), 125–136.
31. Riyazi A, Hensel A, Bauer K, Geissler N, Schaaf S, Verspohl E. (2007). The effects of the volatile oil from ginger rhizomes (*Zingiber officinale*), its fractions and isolated compounds on the 5-HT<sub>3</sub> receptor complex and the serotonergic system of the rat. *Plant Med.*
32. Rodriguez-Morales, A. J., Garbacz, K., & Jarzembowski, T. (2023). *Bacterial Infectious Diseases Annual Volume 2023*. BoD–Books on Demand.
33. Yadufashije, C., Adolyne N., Emanuel M., Sibomana M., and Joseph M. (2020). Antibacterial activity of ginger extracts on bacteria isolated from digestive tract infection patients attended Muhoza Health Center. *Asian Journal of Medical Sciences*. 11(2), 35-41.
34. Zapata, M. R. (2023). Diagnosis and Treatment of Vulvovaginitis. In *Handbook of Gynecology* (pp. 297-317). Cham: Springer International Publishing.