Efficiency of Plant Active Compounds as Anti-Adhesion Factors for Clinical Candida albicans Isolates in vitro

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Abstract
The aim of this study is the determination of the anti-adhesion activity of active compounds extracted from plants in the inhibition of adhesion ability of C. albicans that is used as a virulence factor for infection. Adherence to host surfaces is a primary factor in the colonization of human tissues by fungi, which can also adhere to the surfaces of medical devices and form biofilms. Medicinal plants possess therapeutic properties or beneficial pharmacological effects on the body. A total of one hundred samples were collected from female patients with vaginal infection in two hospitals in Baghdad city for three months. The fungi were isolated and identified by microscopic morphology, morphological features on culture media, and Vital Index of Traditional Environmental Knowledge (VITEK 2) System. Extraction and purification of plant active compounds, vasicine from Adhatoda vasica and gallic acid from Punica granatum L., were performed. The active compounds were used as anti-adhesion factors and the minimum inhibition concentration (MIC) against Candida albicans was determined. We also used the antifungal drug fluconazole for comparison as a control. The results of vaginal discharge characterization of clinical samples showed that 30% of the infections are caused by fungi (100% were Candida albicans), 47% by Gram negative bacteria, and 23% by Gram positive bacteria. The results of MIC assays showed that the plant extract and the antifungal drug do not seem to have fungicidal effects on the isolates of C. albicans tested. The ability of C. albicans isolates for adhesion to the epithelial cells was detected by the calculation of the number of adherent C. albicans cells under light microscope. The activity of vasicin and gallic acid appeared to be approximately equal on the number of C. albicans adherent on epithelial cells. Vasicin concentration activity used in mg/ ml was higher than that used in µg/ ml, with less numbers of C. albicans adhered on epithelial cell. However, treatment with fluconazole showed that more than half of the number of C. albicans were non-adherent on epithelial cells when added to the suspension, in contrast to the control (C. albicans without the extracts or antifungal drug).

Keywords: Efficiency: Candida albicans: Vasicine: Gallic acid; anti-adhesions.

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Introduction

*Candida* infection is the main cause of nosocomial bloodstream infections (BSIs) in tertiary care hospitals worldwide. These organisms are the most common cause of superficial vaginal or mucocutaneous oral infections and may under propitious conditions enter the bloodstream leading to deep-tissue infections [1]. It is generally a harmless commensal fungus that can turn into an opportunistic organism in immunocompromised or immunologically deficient individuals. Abnormal growth of *C. albicans* due to environmental imbalance such as the reduction of pH may result in candidiasis [2]. Different morphologies of *C. albicans* undergo certain interactions with host cells during an infection. It was suggested that epithelial mechanisms drive mucosal tissues to dissociate the pathogenic and commensal states of *C. albicans* [3]. For example, *C. albicans* cells treated with a medicinal plant extract showed a declined adherence to surfaces, thereby inhibiting virulence factors of *C. albicans* [4].

The primary factor in the fungal colonization of human tissues is adherence to host surfaces; this process is controlled and induced by several cell signaling cascades in both the fungus and the environment. Also, *Candida* can adhere to the surfaces of medical devices and form biofilms [5]. The difficulty to eradicate *Candida* infections is owing to its unique switch between yeast and hyphae forms and more likely to biofilm formations that render resistance to antifungal therapy.

Plants are known sources of natural medicines [6]. Medicinal plants possess therapeutic properties and beneficial pharmacological effects on the body. A certain plant can be declared as medicinal when one or more of its organs contain a necessary substance that can be used for therapeutic purposes or used as a precursor for the synthesis of useful drugs [7].

Plants synthesize hundreds of chemical compounds for functions that include defense against insects, fungi, diseases, and herbivorous mammals. Since a single plant contains widely diverse phytochemicals, the effects of using a whole plant as medicine are uncertain. Further, the phytochemical content and pharmacological actions, if any, of many plants having medicinal potential remain unassessed by rigorous scientific research to define efficacy and safety [7].

Therefore, this study investigates the effects of gallic acid and vasicin as anti-adhesions against *C. albicans* adhesion during epithelial cell infection of vagina.

Materials and methods

Samples Collection and Fungal isolation

Vaginal swabs were collected from women referred to Al-Yarmouk Hospital, Alwiya Hospital and Gynecological Private Clinic in Baghdad, Iraq, during the period extended from September to
December 2018. The samples were collected in sterile swabs containing 5 ml of normal saline from women patients with vaginal discharge after being subjected to CLS for diagnosis. All samples were cultured on Sabouraud Dextrose agar (SDA) as slants, tightly wrapped with parafilm and stored at 4 °C [8].

Identification of candida isolates
The isolates were identified as C. albicans depending on the microscopic morphology, morphological features on culture media, and VITEK 2 System.

Plant extracts
Vasicine alkaloid was isolated from the concentrated leaves extract of A. vasia by the acid-base method [9], while Gallic acid was isolated from Punica granatum L. using ethyl acetate (organic solvent) by a separator funnel [10].

Characterization of plant extracts
Vasicine alkaloid was characterized by the melting point of the partially purified sample according an earlier study [11]. The chemical characteristics were investigated using the modified Dragendorff’s reagent and high performance liquid chromatography (HPLC), as previously described [12]. Gallic acid was characterized by phytochemical screening, using few drops of ferric chloride solution with change in color indicating to the presence of phenols, while HPLC was also used for the same purpose [13].

Determination of MIC Concentration for active plant compounds and antifungal drugs
The MIC of the plant active materials (vasicine and gallic acid) and fluconazole drug was determined by the agar dilution technique. The test was used to ensure that the highest concentration of these compounds that did not seem to have a fungicidal effect against C. albicans growth, by mixing with Muller Hinton agar and incubation for 24h at 37 °C [14].

Preparation of C. albicans suspension
Ten milliliters of Sabouraud Dextrose broth medium was inoculated with C. albicans growth culture and incubated at 37°C for 24 hours. The yeast culture was washed twice with Phosphate Buffer Saline (PBS) concentrated by centrifugation at 1000 rpm for 20 min, and re-suspended in PBS [15, 16].

Preparation of epithelial cells
Uroepithelial cells were isolated from the urine of some healthy females from September 2018 until December 2019 by centrifugation at 6000 rpm for 5 min, then washed three times with PBS and re-centrifuged at 6000 rpm for 10 min before re-suspension in PBS [14].

Fixative solution
The fixative was prepared by mixing 30ml of methanol (99.8%) with 10 ml of acetic acid (99.8%). It was used for the fixation of yeast and uroepithelial cells during staining by methylene blue [17].

Candida adhesion test
The adhesion test was achieved as follows [18]:
● A mixture of 0.1 ml of the yeast suspension, 0.1ml of the epithelial cells suspension and 0.05 of PBS were incubated at 37°C for 2 hrs.
● Unattached yeast cells to uroepithelial cells were removed by centrifugation in PBS at 6000 rpm for 10 min.
● The pellet was re-suspended in PBS. A drop of the suspension was put on a microscopic slide, air-dried, fixed with methanol: acetic acid (3:1) and stained with methylene blue.
● The adhered yeast cells to epithelial cells were observed by the compound light microscope.
● The control contained only epithelial cells.

Anti-adhesion test using Vasicin, Gallic acid and Fluconazole
After the activation of Candida for 24 hours, 500 microliters of the activated inoculum was taken and concentrated using centrifuge at 6000 rpm for 5 min, then 50 μl of PBS was added and mixed with 500 microliters of epithelial cells [18]. After completing this step, the experience was divided into two lines:
In the first line, the mixture from the previous step was incubated at 37°C for 2 hours, then 50 microliters of the vasicin extract was added with different concentrations (200, 400, 600, 800,1000,1200,1400,1600,1800 and 2000 µg/ml) and (1, 2, 4, 8, 16, 32, 64, 128, 256 and 516 mg/ml ), and stained for examination under light microscope. In the second line, 50 microliters of the vasicin extract with different concentrations (200, 400, 600, 800, 1000, 1200,1400,1600,1800 and 2000
μg/ml) and (1, 2, 4, 8, 16, 32, 64, 128, 256 and 516 mg/ml) were added to the mixture. The samples were then incubated at 37°C for 2 hours, stained, and examination under a light microscope. As for gallic acid extract, the previous experiment repeated with various concentrations (200, 400, 600, 800, 1000, 1200, 1400, 1600, 1800, 2000 μg/ml). Fluconazole antifungal drug is commonly used to treat vaginal yeast infections caused by Candida. This antifungal drug was treated in the same way as the vasicin and gallic acid extracts in the previous experiments, with different concentrations (200, 400, 600, 800, 1000, 1200, 1400, 1600, 1800, 2000 μg/ml).

Results and discussion

Isolation and Identification

One hundred samples of high vaginal swabs [HVS] were collected from women with symptoms of vaginal infection who visited Gynecological private clinics and educational hospitals (Al-Yarmouk and Al-Alwiya). The collection was performed between September and December 2018. Half of the samples was collected from pregnant whereas the other half was from non-pregnant women. Samples were tested by microscopic examination and culture on SDA.

Colonial morphology, gram staining, and VITEK2 test were carried out for the identification of the isolated organisms. Thirty samples showed positive results of Candida albicans. The characterization of the samples of vaginal discharge showed that 30 (30%) of infections were caused by Fungi (100% were Candida albicans), 47 by Gram negative bacteria, and 23 by Gram positive bacteria.

These results approximately agree with those reported by other authors [19], who showed that Candida albicans was detected in 20% of the subject samples. Onyia revealed that about 90% of this infection is caused by Candida albicans and 10% by other species of Candida [20]. This infection has been reported as the common cause of vaginitis, second to bacterial vaginosis and, generally, the infections occur when there is an imbalance in the pH of the vagina [21].

The over-growth of this fungus in the vagina leads to a burning sensation in the vagina, the production of heavy white/yellow curd-like discharge and/or an itchy vulva, pruritus, dyspareunia, dysuria, irritation, soreness of the vulva and other discomforting symptoms that will ensure frequent hospital visits [20]. The present results showed that C. albicans infections were more in pregnant (21 cases) than non-pregnant women (9 cases), which agrees with the results reported earlier [22].

Determination of MIC for active plant compounds and antifungal drug

The effects of the active compounds of plant extracts and the antifungal drug against C. albicans were tested by using MIC with four dilutions which showed resistance to fluconazole. The results are shown in Tables- 1 and 2. The MIC assays showed that the plant extracts and the antifungal drug do not appear to have fungicidal effects on the isolates of C. albicans tested at these concentrations.

**Table 1-** Minimum inhibitor concentration (MIC) of active compound of plant extract against C. albicans.

| MIC | C. albicans Plant extract | CA. 2 | CA. 8 | CA. 20 | Stock of plant extract |
|-----|---------------------------|-------|-------|-------|-----------------------|
| Vasin (mg/ ml) | >516 mg/ml | >516 mg/ml | >516 mg/ml | 516 mg/ml |
| Vasin (μg/ ml) | >2000 μg/ml | >2000 μg/ml | >2000 μg/ml | 2000 μg/ml |
| Gallic acid (μg/ ml) | >2000 μg/ml | >2000 μg/ml | >2000 μg/ml | 2000 μg/ml |

**Table 2-** Minimum inhibitor concentration (MIC) of the antifungal drug (Fluconazole) against C. albicans.

| MIC | C. albicans Fluconazole | CA. 2 | CA. 8 | CA. 20 | Stock of Fluconazole |
|-----|-------------------------|-------|-------|-------|---------------------|
| Fluconazole (μg/ ml) | >2000 μg/ml | >2000 μg/ml | >2000 μg/ml | 2000 μg/ml |

Fluconazole is widely used clinically for treating pathogenic fungal infections, specifically those caused by C. albicans [23]. Recently, studies from many countries have shown an increased prevalence of fluconazole-resistant strains of C. albicans [24].
The antibacterial as well as antifungal activities of any plant depend on the extraction conditions, such as type and concentration of the solvent and time and temperature for the extraction process. All these factors affect the type and the amount of the active material that is extracted [25]. The mode of action of these compounds, such as polyphenols, is generally attributed to polyphenol-protein interactions, although different mechanisms have been suggested, including inhibition of microbial enzymes, actions on membranes, or deprivation of substrates required for microbial growth [26].

They have many roles in eukaryotic membranes, including establishing membrane fluidity, regulating membrane-bound enzymes and maintaining membrane permeability. Ergosterol is found in fungal cell membranes, and the lack of detectable ergosterol in fungal cell membranes after gallic Acid treatment suggests damage to the sterol biosynthetic pathway and subsequent reduction in the viability of *C. albicans* [27, 28].

Phenolic compounds (e.g. gallic acid), initially affect cell membrane, changing permeability and causing the leakage of cellular content, or interfere with membrane proteins resulting in structure disrupting. In addition, they disrupt cell wall and cell membrane synthesis [29]. Polyphenols may affect the bacterial cell wall, inhibit enzymes by oxidized agents, interact with proteins and disturb the co-aggregation of microorganisms [30]. Pomegranate extracts were reported to have broad bioactivities including antimicrobial, anti-inflammatory and antitumor effects, and have been used in traditional Uyghur medicine for many years [31].

**Effects of the plants active compounds and antifungal drug on adhesion activity**

The ability of *C. albicans* isolates to adhere to the epithelial cells was detected by calculations of the number of adherent *C. albicans* cells under the light microscope. Forty cells of *C. albicans* adhered to the assayed epithelial cells. These data are regarded as a control for the detection of the effects of the MIC of plants active compounds and the antifungal drug on the adhesion ability of *C. albicans* (Figure-1).

![Figure 1](image.png)

*Figure 1*- The adhesion of *C. albicans* isolates on epithelial cells.

**Effect of vasicin on adhesion activity of C. albicans**

No *C. albicans* cells were adhered to epithelial cells before and after the incubation with vasicin with concentrations of 128-516 mg/ml. Also, at treatment with 64 mg/ml of vasicin, no *C. albicans* cells were adhered; but before incubation at the same concentration, four cells of *C. albicans* were found adhered on epithelial cells. The other concentration showed a variable number of *C. albicans* adhered to epithelial cells, in contrast to the control (*C. albicans* without the extracts or antifungal drug), as shown in Figure-2.
As shown in Figure-3, no *C. albicans* cells were adhered to epithelial cells, only when vasicin was added with a concentration of 1800-2000 µg/ ml. The other concentrations showed a variable number of *C. albicans* adhered to the epithelial cells.

**Figure 2-** Anti-adhesion activity of Vasicin (mg/ ml) against *C. albicans*.

The activity of vasicin concentration in the mg/ ml was higher than that in µg/ ml, which showed less number of *C. albicans* adhered on epithelial cells.

**Effects of Gallic Acid on the adhesion activity of C. albicans**

*C. albicans* showed no adhered cells to epithelial cells when gallic acid was added with concentrations of 1800-2000 µg/ ml. Only the concentration of 2000 µg/ ml of gallic acid showed no adhered cells before incubation in the suspension. The other concentrations showed a variable number of *C. albicans* adhesion to epithelial cells in contrast to the control (*C. albicans* without the extracts), as shown in Figure-4.

**Figure 3-** Anti-adhesion activity of Vasicin (µg/ ml) against *C. albicans*.
Figure 4. Anti-adhesion activity of Gallic Acid (µg/ml) against C. albicans.

Vasicin and gallic acid activities when used the concentrations of µg/ml showed approximately equal effects on the number of C. albicans adhesion on epithelial cells. The radiolabeled adhesion and the CFU counting assays suggested that vasicin and gallic acid have anti-adhesive effects.

Plant extracts and surfactants were evaluated for their anti-candidal-adhesive properties both on epithelial cells and on inert surfaces by some researchers [32]. Patel et al. showed that Dodonaea viscosa var. angustifolia (Sand Olive plant) crude extract could inhibit the adhesion of C. albicans isolated from HIV-seropositive and HIV-seronegative individuals to oral epithelial cells [33]. The date extract has also been proven to decrease the adhesion of three different species of candida to epithelial cells [40]. Previous studies have been carried out using various substances to assess their effect on C. albicans adhesion, for example, some plant extracts and surfactants, including bio-surfactants, etc. [34]. A recent study in Iraq detected the effects of the MIC of pomegranate peel extracts on the adhesion ability of bacteria and showed a clear decline in the adhesion activity when the bacteria was treated with peel extracts [35].

The relation between candidal adherence to host epithelial cells or acrylic denture surfaces and the ability of Candida to colonize and cause disease has been investigated in a number of studies. Colonization of the oral cavity is facilitated by several specific (receptor-ligand) and non-specific (electrostatic and electrodynamic) interactions between Candida and oral surfaces [36]. The adhesion process of Candida is complex and involves biological and non-biological factors. The adhesion capacity of Candida seems to influence its infection and colonization potential. Adhesion is not only affected by yeast conditions; variations in the epithelial surface also have a critical role in the adhesion phenomenon [37].

Cell surface hydrophobicity, which contributes to the interaction between the cells and the surfaces, seems to be an important factor in the adhesion of C. albicans. There are statistically significant correlations between cell hydrophobicity and the adhesion of C. albicans to the epithelial cells of the mouth and acrylic resin surfaces. The hydrophobic nature of C. albicans’ cell surface is thought to be associated with the adherence of the yeast to epithelial cells and plastic devices. A correlation between the hydrophobicity of C. albicans and its adherence to HeLa cells has been reported. Polyenes bind avidly to ergosterol in fungal membranes, forming channels that allow K⁺ and Mg²⁺ to leak out of the cell. The limited exposure of oral C. albicans isolates to polyenes reduces the relative cell surface hydrophobicity and candidal adherence to epithelial cells [38]. The adhesion occurs because C. albicans forms the fibrous layer composed of multiple sugars on the surfaces of cells during the stationary phase of growth [39].

Receptor analogs, adhesion analogs, and surface-modifying agents were used to inhibit the adhesion of microbes onto host cells. The number of organisms developing resistance to antiadhesive
substances seems to be less compared with antimicrobial agents. However, the limitation of this kind of therapy is the requirement of multiple anti-adhesion agents to counter each type of adhesion of infecting pathogens [40].

The effect of the plant extracts is attributed to their ability to inhibit cell attachment. The cell attachment is the initial stage in biofilm formation following surface conditioning which creates a favorable environment for bacterial attachment. Surface conditioning is achieved by the adsorption of substances that include nutrients, organic, and inorganic molecules that are important for the growth of the cells, which in turn promotes cell adhesion and biofilm formation. Therefore, it can be postulated that pretreatment of the surface with plant extracts produces an unfavorable film that promotes detachment, thereby reducing the surface adhesion [41]. Janecki and Kolodziej explained the activity of plants active compounds that could be used as anti-adhesive agents against pathogenic microbes and showed capabilities of these compounds of interacting with macromolecules, including carbohydrates and proteins, which made these compounds as promising anti-adhesive and anti-biofilm agents [26]. The success of plant extracts in inhibiting cell attachment and reducing microbial colonization on surfaces of epithelial mucosa, which lead to infection, represented the important steps that achieved the growth of an already established biofilm [41].

**Effects of Fluconazole on adhesion activity of C. albicans**

*C. albicans* showed no cells to be adhered to epithelial cells when fluconazole was added with concentrations of 1000-2000 µg/ ml. Only a concentration of 2000 µg/ ml of fluconazole showed no adhered cells before incubation with the suspension. The other concentration demonstrated a variable number of *C. albicans* adhered to epithelial cells, in contrast to the control (*C. albicans* without the extracts), as shown in Figure-5.

![Figure 5](image_url)

**Figure 5**- Anti-adhesion activity of Fluconazole (µg/ ml) against *C. albicans*.

Vasicin and gallic acid activities in the µg/ ml concentration showed less effects on the number of *C. albicans* adhered on the epithelial cells than that of fluconazole, which showed more than the half of the number of *C. albicans* adhered on epithelial cells when added to the suspension.

A previous study showed that cells pre-treated with fluconazole had a reduction in the adhesion of *C. albicans* cells to the epithelial cells, which ranged from 0 to 72.9%. [42]. Also, another study used low doses of fluconazole and showed a reduction in the adhesion of *C. albicans* [43].

Another study found that incubation of HeLa cells with *Candida* spp., in the presence of amphotericin B, nystatin or natamycin, reduced the candidal adherence to these cells. When compared to amphotericin B, nystatin and natamycin suppressed adherence less effectively and these differences were statistically significant. The MIC and sublethal concentrations of nystatin reduced the adherence of *C. albicans* to cell monolayers by 35 and 47%, respectively. Interestingly, following the “limited exposure prior to adhesion” procedure, nystatin reduced the adherence of *C. albicans* to the epithelial cells by 54 and 73% at the MIC and sublethal concentration, respectively [15].
The effects of antifungal drugs on candidal adhesion to epithelial cell has been examined following (i) pre-incubation of the isolates with the drug, (ii) pre-treatment of BECs with the drug, and (iii) limited exposure (1 h) of the isolates to sub lethal concentration of the drug, and subsequent removal of the drug. This study have indicated that pre-exposure of either the target cell surface or the yeasts to antifungal drugs results in reduced adhesion, which in vivo may enhance the effect of the topical agent [15].

Conclusions
From 100 samples of infections, 30% were caused by C. albicans. The maximum concentration used for vasicin (516 mg/ml and 2000 µg/ml), Gallic acid (2000 µg/ml) and Fluconazole (2000 µg/ml) did not seem to have a fungicidal effect on the isolates of C. albicans tested. Both the alkaloid (vasicine), the phenol (gallic acid) and fluconazole used in this study were able to show anti-adhesion activity in vitro. Vasicin and gallic acid activities exhibited approximately equal effects on the number of C. albicans adhered to epithelial cells, while fluconazole showed more effects on C. albicans adhesion.

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