

Source details

Journal of Pakistan Association of Dermatologists	CiteScore 2021 0.2	(i)
Scopus coverage years: from 1999 to 2022		
Publisher: Pakistan Association of Dermatologists		
ISSN: 1560-9014	sjr 2021 0.127	(j)
Subject area: Medicine: Dermatology	0.127	
Source type: Journal		
View all documents > Set document alert Save to source list Source Homepage	SNIP 2021 0.127	(j)

CiteScore CiteScore rank & trend Scopus content coverage

i Improved CiteScore methodology
 CiteScore 2021 counts the citations received in 2018-2021 to articles, reviews, conference papers, book chapters and data
 papers published in 2018-2021, and divides this by the number of publications published in 2018-2021. Learn more >

CiteScore 2021 \checkmark $0.2 = \frac{90 \text{ Citations } 2018 - 2021}{414 \text{ Documents } 2018 - 2021}$ Calculated on 05 May, 2022 CiteScore rank 2021 Category Rank Percentile

Medicine

— Dermatology #107/126 15th

View CiteScore methodology \succ CiteScore FAQ \succ Add CiteScore to your site ${}_{\mathcal{O}}^{\mathcal{O}}$

CiteScoreTracker 2022 ①

0.3 = 112 Citations to date 448 Documents to date

Last updated on 05 January, 2023 • Updated monthly

Q

About Scopus

- What is Scopus Content coverage
- Scopus blog
- Scopus API
- Privacy matters

Language

日本語版を表示する **查看简体中文版本** 查看繁體中文版本

Просмотр версии на русском языке

Customer Service

Help Tutorials Contact us

ELSEVIER

 $\label{eq:copyright} \textcircled{Copyright} \hline{Copyright} \textcircled{Copyright} \hline{Copyright} \hline Copyright \hline{Copyright} \hline{Copyright} \hline Copyrig$

RELX



Journal of Pakistan Association of Dermatologists <a>3

COUNTRY

Pakistan

<u>,</u>]]]]]

Universities and research institutions in Pakistan SUBJECT AREA AND CATEGORY

Medicine Dermatology

PUBLICATION TYPE

ISSN

Journals

15609014

SCOPE

Information not localized

 $\ensuremath{\bigcirc}$ Join the conversation about this journal

Quartiles

₿



1
Indian Journal of
Dermatology, Ven
IND

rmatology, Venereology IND

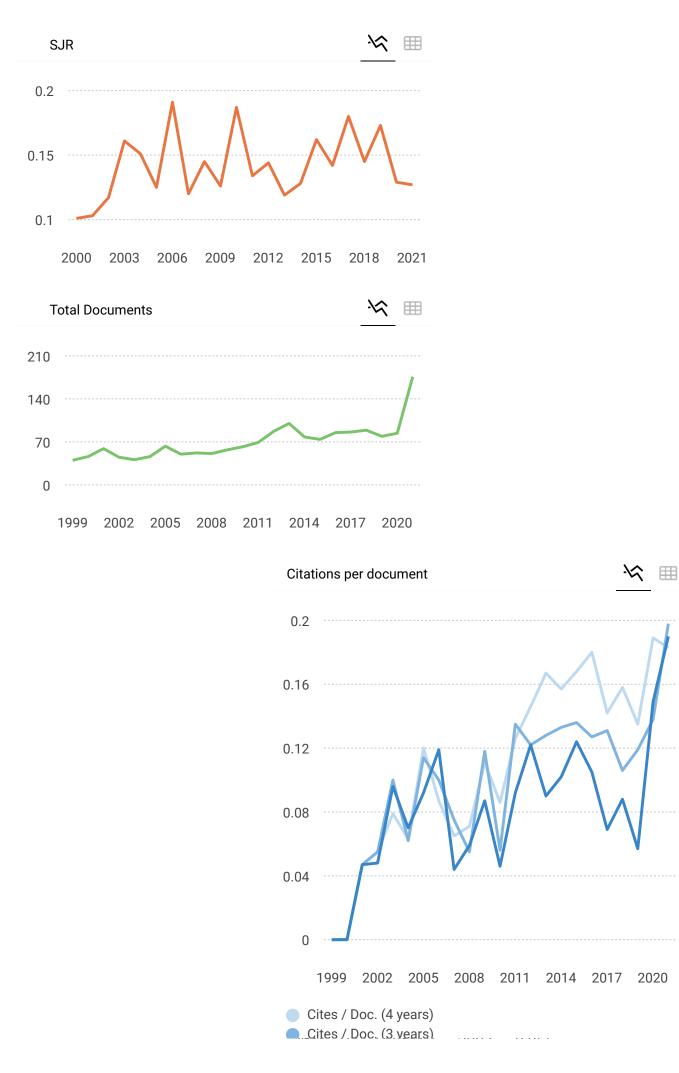


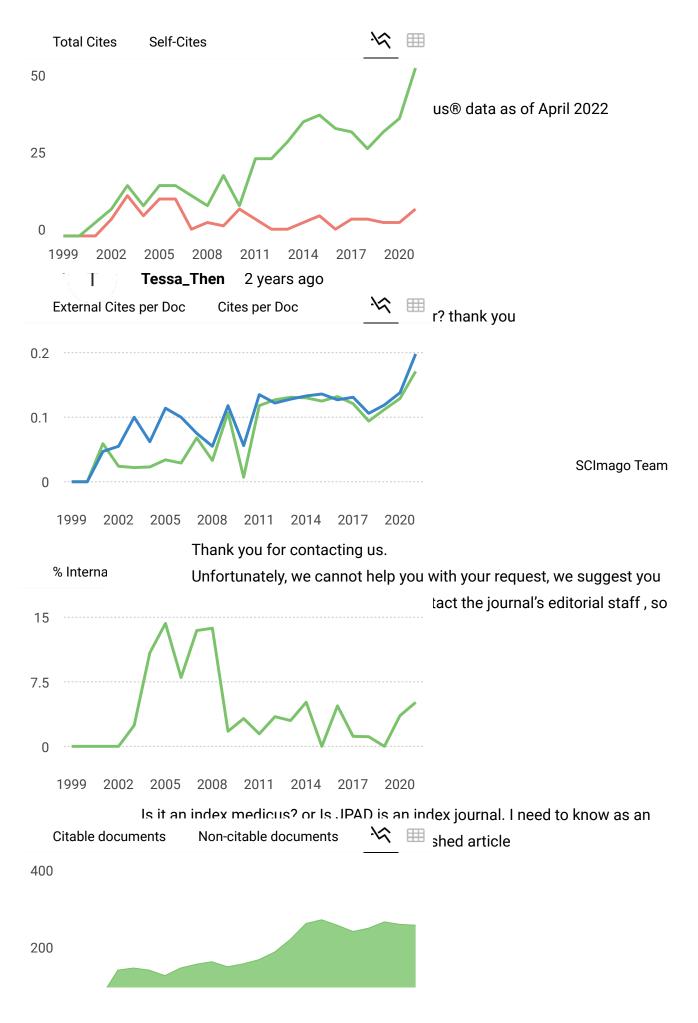
2 Indian Journal of Dermatology IND

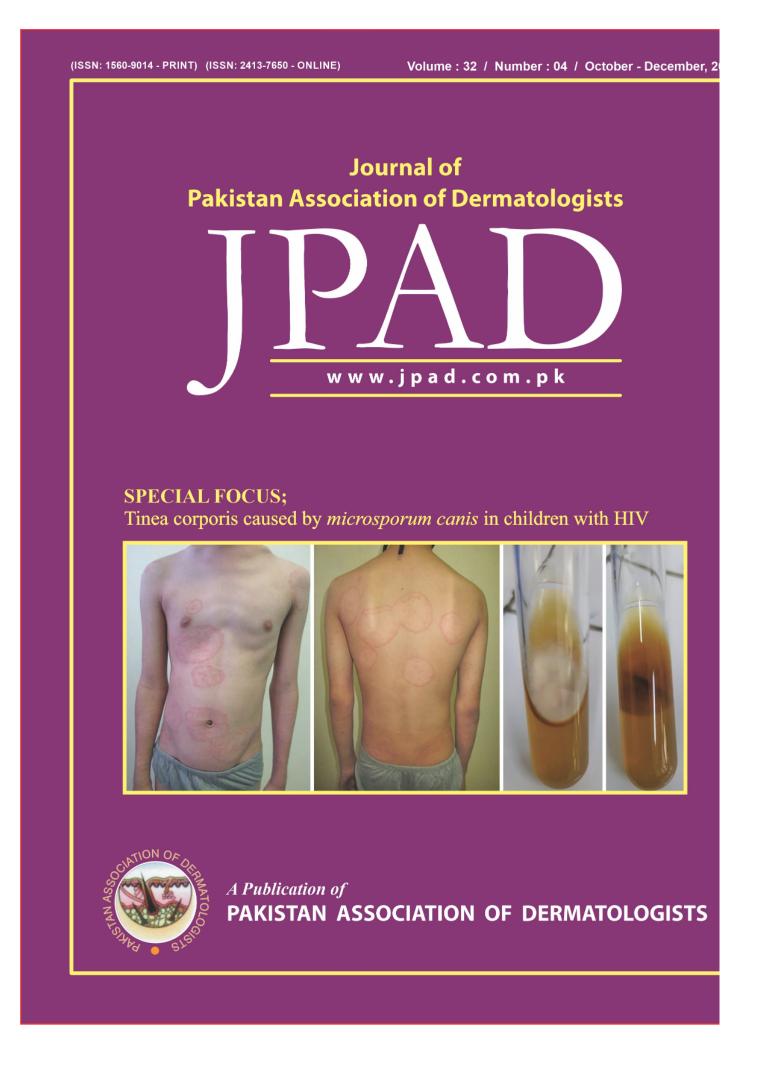
78%

3 Iranian Dermat IRN





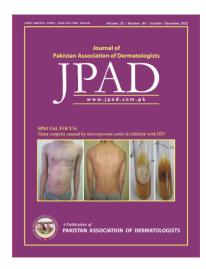




Journal of Pakistan Association of Dermatologists

Home / Archives / Vol. 32 No. 4 (2022): October - December

Vol. 32 No. 4 (2022): October - December



Published: 2022-12-08

Editorial

The importance of dermatological lesions in the agile diagnosis of Monkeypox

Gonzalo Dominguez Alvarado, Angie Marcela Arboleda Roca , Aymer Daniel Bolívar Cabrera , Gabrielle Yolmari Curicó Martínez , Laura Camila Millán Guerrero 662-664



Original Articles

The Spectrum of Dermatoses among Prisoners; A Retrospective Analysis

Hira Tariq 665-668



Basil Leaf Extract and Eugenol against Isolates of Candida sp. Causing Oral Candidiasis in HIV/AIDS

Dwi Murtiastutik, Emma Hidayati Sasmito, Diah Mira Indramaya, Sawitri Sawitri, Rahmadewi Rahmadewi, Afif Nurul Hidayati, Pepy Dwi Endraswari, Sudjarwo Sudjarwo, Budi Utomo 669-675



Can Ki-67 immunohistochemistry marker differentiate mycosis fungoides from cutaneous lichen planus?

Mazaher Ramezani, Kambiz Kamyab Hesari, Negar Ramezanizoorab, Masoud Sadeghi, Mansour Rezaei, Sedigheh Khazaei

676-682



Correlation between quality of life and clinical severity of melasma in Pakistani women

Maryam Qayyum, Saadiya Siddiqui, Mohsina, Mahmoona Ilyas, Atif Shahzad, Nadia Ali Zafar

Saadiya Siddiqui, Maryam Qayyum 683-689



Association of Pruritus Visual Analogue Scale and Risk Factors in Adolescence Pediculosis Capitis in Two Public Boarding Schools, West Java

Arlene Rainamira, Firman Parrol, Yari Castiliani, Githa Rahmayunita, Mochamad Helmi Aziz, Kusmarinah Bramono, Sandra Widaty

690-695



Scalp defects and their reconstruction in patients at Allama Iqbal Memorial Teaching Hospital, Sialkot

Sarfaraz Ahmad, Raffad, Zahida Rani, Atika Malik 696-700

🔁 PDF

Novel trends in cutaneous manifestations of geriatric dermatoses in a tertiary care hospital, South India

Latest trends in geriatric dermatosis

Hasini budeda, Indira B, Darsan S, V. V. V. Satyanarayana, Suruthi Purushothaman

701-706



Effect of Lactobacillus plantarum IS-10506 supplementation with oral metronidazole for the treatment of bacterial vaginosis : a randomized placebo-controlled clinical trial

Afif Nurul Hidayati, Ridha Ramadina Widiatma, Dwi Murtiastutik, Ingrid Suryanti Surono, Cita Rosita Sigit Prakoeswa, Lunardi Bintanjoyo

707-714



The Platelet Rich Plasma (PRP) Preparation and Standardization of time interval for scalp hair loss

Munir Alam 715-718

🖹 PDF

Review Articles

Zinc as an essential element for normal immune reactions and as a therapeutic agent for autoimmune diseases

Inas K. Sharquie, Khalifa E. Sharquie , Wasan W. Al-bassam 719-725

🖾 PDF

Review of treatment modalities in the management of Genital Warts

Review of treatment modalities in the management of Genital Warts

Jai Singh Solanki, Swagat Waghmare, Manjyot Gautam, Praneet P. Awake 726-733

🕒 PDF

Case Reports

Kaposi's sarcoma in a male with human immunodeficiency virus and condyloma acuminata: A case report

Minna Hasniah, Budi Eko Prasetyorini, Endra Yustin Ellistasari 734-740



Tinea Corporis Caused by Microsporum Canis in Children with HIV

Sri Mariyani

741-744



Recurrent herpes simplex in elderly with vitiligo vulgaris: A dilemmatic case

Novita Suprapto, dr. Lili Legiawati, dr. Rinadewi Astriningrum, dr. Shannaz Nadia Yusharyahya 745-747

🕒 PDF





Recognized by HEC



Indexed

Information

For Readers

For Authors

For Librarians

Open Journal Systems

Platform & workflow by OJS / PKP

Journal of Pakistan Association of Dermatologists

Home / About the Journal

About the Journal

Editor-in-Chief

Prof. Ijaz Hussain

Editor
Prof. Zahida Rani

Associate Editors

Prof. Faria Asad Prof. Shehla Shaukat

Assistant Editors

Dr. Ghazala Butt	Dr. Mahwish Zahoor	Dr. Amina Afzal
Dr. Bushra Bashir	Dr. Umara Siddique	Dr. Zareen Saqib
Dr. Sadia Jabeen	Dr. Wajieha Saeed	Dr. Nabeela Shehzadi
Dr. Rutaba Kiran		

JPAD, the official journal of Pakistan Association of Dermatologists is published quarterly, four issues per volume and one volume per year (ISSN 1560-9014, print; ISSN 2413-7650, online). The journal is recognized by Pakistan Medical Commission, Higher Education Commission of Pakistan and is indexed in College of Physicians and Surgeons Pakistan MEDLIP; Ulrich's International Periodical Directory, USA; ExtraMED, London; EMBASE/Excerpta Medica, PakMediNet, The Netherlands; and Index Medicus, WHO Alexandria, Egypt.

Make a Submission



Recognized by HEC



Indexed

Information

For Readers

For Authors

For Librarians

Open Journal Systems

Platform & workflow by OJS / PKP

Original Article

Basil leaf extract and eugenol against isolates of *Candida* sp. causing oral candidiasis in HIV/AIDS

Dwi Murtiastutik¹, Emma Hidayati Sasmito¹, Afif Nurul Hidayati^{1,2}, Rahmadewi¹, Sawitri¹, Budi Utomo³, Sudjarwo⁴, Pepy Dwi Endraswari^{5,6}, Diah Mira Indramaya¹

- ¹Department of Dermatology and Venereology, Dr. Soetomo General Academic Teaching Hospital, Surabaya, Indonesia.
- ²Department of Dermatology and Venereology, Universitas Airlangga Teaching Hospital, Surabaya, Indonesia.
- ³Department of Public Health and Preventive Medicine, Faculty of Medicine, Universitas Airlangga, Surabaya, Indonesia.
- ⁴Department of Pharmaceutical Science, Faculty of Pharmacy, Universitas Airlangga, Surabaya, Indonesia.
- ⁵Department of Clinical Microbiology, Faculty of Medicine, Universitas Airlangga, Dr. Soetomo General Academic Teaching Hospital, Surabaya, Indonesia.
- ⁶Department of Clinical Microbiology, Universitas Airlangga Teaching Hospital, Surabaya, Indonesia.
- **Abstract** *Background* Oral candidiasis is an opportunistic infection of the oral mucosa caused by *Candida* sp., frequently found in HIV/AIDS patients. Basil leaf extract (*Ocimum sanctum* Linn.), which contains eugenol, a compound believed to inhibit the growth of *Candida* sp.

Objective To evaluate the antifungal effect of basil leaf extract (*Ocimum sanctum* Linn.) and eugenol compared to fluconazole against *Candida* sp. isolates.

Methods Basil leaf extract (*Ocimum sanctum* Linn.) with doses equivalent to eugenol 800 μ g/mL and 400 μ g/mL, eugenol doses at 800 μ g/mL and 400 μ g/mL, which was then compared with fluconazole 25 μ g/mL against 40 *Candida* sp. isolated stored from the oral cavity of HIV/AIDS patients.

Results The average inhibition zone of fluconazole was 21.81 mm, the mean inhibition zone of eugenol with doses of 800 μ g/mL and 400 μ g/mL were 17.07 mm and 15.89 mm, and the mean inhibition zone of basil leaf extract (*Ocimum sanctum* Linn.) with doses equivalent to eugenol 800 μ g/mL and 400 μ g/mL were 14.87 mm and 14.01 mm (p = 0.001 and p < 0.05).

Conclusion Fluconazole had a significantly higher inhibition zone against *Candida albicans* and *Candida* non-*albicans* isolates than basil leaf extract (*Ocimum sanctum* Linn.) and eugenol.

Key words

Basil leaf extract (*Ocimum sanctum* Linn.), eugenol, fluconazole, antifungal susceptibility testing, oral candidiasis, HIV, AIDS.

Introduction

Oral candidiasis is a fungal infection of the oral mucosa caused by *Candida* sp.¹ *Candida* sp. is a commensal fungus in healthy people, but under certain conditions it can cause an opportunistic infection. Opportunistic infections can occur as a

consequence of infection with the human immunodeficiency virus (HIV) and the acquired immunodeficiency syndrome (AIDS). During the course of their illness, nearly 90% of HIV/AIDS patients develop oral candidiasis.^{2,3}

Oral candidiasis can be treated with the

antifungal drug nystatin oral suspension and/ or fluconazole as a systemic antifungal.^{4,5} In a test of *Candida* sp. resistance to fluconazole, isolates resistant to fluconazole were found to be 48.6%. *Candida* non-*albicans* was the most resistant isolate, accounting for 72.2% of the total.² Antifungal drug biofilm formation and resistance are two issues that can be addressed by seeking for alternative antifungal drugs.⁶

Basil (Ocimum sanctum Linn.) is a tropical plant that can be found in abundance in Indonesia.⁷ The main component of basil leaf (Ocimum sanctum Linn.) is eugenol. This compound is effective against the adaptive mechanism of Candida albicans biofilm resistance to fluconazole.⁸ Based on these findings, the researchers proposed to hold an in vitro test of basil leaf extract (Ocimum sanctum Linn.) and eugenol against stored isolates of Candida sp. and compare them to standard oral candidiasis treatment fluconazole, which was now showing signs of resistance.

Methods

The research design used in this study was an experimental laboratory with the aim of evaluating the comparison of the antifungal activity of basil leaf extract (*Ocimum sanctum* Linn.) with doses equivalent to eugenol 800 μ g/mL and 400 μ g/mL, eugenol at doses 800 μ g/mL and 400 μ g/mL was then compared with fluconazole 25 μ g/mL against 40 isolates stored *Candida* sp. which consisted of 20 *Candida albicans* and 20 *Candida* non-*albicans* isolated

Address for correspondence
Dwi Murtiastutik,
Department of Dermatology and Venerology
Faculty of Medicine, Universitas Airlangga/ Dr.
Soetomo General Academic Teaching Hospital/
Universitas Airlangga Teaching Hospital,
Surabaya, Jl. Mayjen Prof. Dr. Moestopo No. 6-8
Surabaya 60131, Indonesia.
Ph: (031) 5501609
Email: dwimurtiastutik@yahoo.co.id

from the oral cavity of HIV/AIDS patients who were hospitalized at the Infectious Disease Intermediate Treatment Unit (UPIPI) RSUD Dr. Soetomo Surabaya for the period April 2019 to July 2019 which was reactivated.

The antifungal activity was evaluated using the disk diffusion method with paper discs or blank discs. These data were entered into a data collection sheet and analysed with SPSS (Statistical Package for Social Sciences). This research has obtained ethical approval from the Ethics Committee of Dr. Soetomo General Academic Teaching Hospital Surabaya (0522/LOE/301.4.2/VII/2021).

Results

In this study, using the disc diffusion method, the average results of the inhibition zone test of basil leaf extract (*Ocimum sanctum* Linn.) was equivalent to 800 μ g/mL eugenol, basil leaf extract (*Ocimum sanctum* Linn.) was equivalent to 400 μ g/mL eugenol, eugenol 800 μ g/mL, and eugenol 400 μ g/mL will be tested and compared with the standard antifungal treatment fluconazole against *Candida* sp.

The average fluconazole zone for all *Candida* sp. was 21.81 mm, eugenol 800 μ g/mL was 17.07 mm, eugenol 400 μ g/mL was 15.89 mm, basil leaf extract (*Ocimum sanctum* Linn.) was equivalent to eugenol 800 μ g/mL was 14.87 mm, and basil leaf extract (*Ocimum sanctum* Linn.) was equivalent to eugenol 400 μ g/mL was 14.01 mm. **Table 1** shows a comparison of the average inhibition zone results for all *Candida* sp.

The average fluconazole inhibition zone in *Candida albicans* species was 22.75 mm, eugenol 800 μ g/mL was 16.76 mm, eugenol 400 μ g/mL was 15.52 mm, basil leaf extract (*Ocimum sanctum* Linn.) was equivalent to

cugent	of and fluconazole in all species.			
No.	Antifungal type	Number of Isolates	Average Inhibition Zone Diameter (mm)	P value
1	Fluconazole 25 µg/mL	40	21.81	
2	Eugenol 800 µg/mL	40	17.07	
3	Eugenol 400 µg/mL	40	15.89	0.001
4	Basil leaf extract equivalent to eugenol 800 µg/mL	40	14.87	
5	Basil leaf extract equivalent to eugenol 400 µg/mL	40	14.01	

Table 1 Comparison of the mean diameter of the inhibition zone of basil leaf extract (Ocimum sanctum Linn.), eugenol and fluconazole in all species.

Table 2 Comparison of the mean diameter of the inhibition zone of basil leaf extract (*Ocimum sanctum* Linn.), eugenol and fluconazole in *Candida albicans*.

No.	Antifungal type	Number of Isolates	Average Inhibition Zone Diameter (mm)	P value
1	Fluconazole 25 µg/mL	40	22.75	
2	Eugenol 800 µg/mL	40	16.76	
3	Eugenol 400 µg/mL	40	15.52	0.001
4	Basil leaf extract equivalent to eugenol 800 µg/mL	40	14.41	
5	Basil leaf extract equivalent to eugenol 400 µg/mL	40	13.62	

Table 3 Comparison of the mean diameter of the inhibition zone of basil leaf extract (*Ocimum sanctum* Linn.), eugenol and fluconazole in *Candida* non-albicans.

No.	Antifungal type	Number of	Average Inhibition	P value
		Isolates	Zone Diameter (mm)	P value
1	Fluconazole 25 µg/mL	40	20.86	
2	Eugenol 800 µg/mL	40	17.38	
3	Eugenol 400 µg/mL	40	16.25	0.001
4	Basil leaf extract equivalent to eugenol 800 µg/mL	40	15.34	
5	Basil leaf extract equivalent to eugenol 400 µg/mL	40	14.39	

eugenol 800 μ g/mL was 14.41 mm, and basil leaf extract (*Ocimum sanctum* Linn.) was equivalent to eugenol 400 μ g/mL was 13.62 mm. **Table 2** shows a comparison of the average inhibition zone results against *Candida albicans*.

In *Candida* non-*albicans*, the average fluconazole inhibition zone was 20.86 mm, eugenol 800 µg/mL was 17.38 mm, eugenol 400 µg/mL was 16.25 mm, basil leaf extract (*Ocimum sanctum* Linn.) was equivalent to eugenol 800 µg/mL was 15.34 mm, and basil leaf extract (*Ocimum sanctum* Linn.) was equivalent to eugenol 400 g/mL was 14.39 mm. The comparison of the average inhibition zone results against *Candida* non-*albicans* could be observed in **Table 3**.

Because the data were not normally distributed and homogeneous, non-parametric statistical methods (Mann Whitney Test) was used to analyze it. The results of the non-parametric statistical test showed that the data significance value was 0.001, so <0.05, which means that there was a significant difference between the mean inhibition zone of eugenol 800 μ g/mL, eugenol 400 μ g/mL, and basil leaf extract (*Ocimum sanctum* Linn.) equivalent to eugenol 800 μ g/mL, and basil leaf extract (*Ocimum sanctum* Linn.) equivalent to eugenol 400 μ g/mL compared to fluconazole which is the standard antifungal drug against the growth of *Candida* in all species, *Candida albicans* and *Candida* non-*albicans*.

Discussion

In this study, the average inhibition zone of fluconazole was greater than the average inhibition zone of eugenol and basil leaf extract (*Ocimum sanctum* Linn.). The inhibition zones of both eugenol and basil leaf extract (*Ocimum sanctum* Linn.) are not better than fluconazole as a standard antifungal drug, but both eugenol and basil leaf extract (*Ocimum sanctum* Linn.) have antifungal effects capable of inhibiting the growth of *Candida* sp., according to the mean comparison.

In a similar study published in 2010, Ahmad and colleagues discovered that fluconazole had a better antifungal effect than eugenol, but some Candida sp. Have become resistant to Subsequently study fluconazole. а was conducted with combining fluconazole and eugenol which concluded that a combination of the two were superior to inhibit Candida sp. than fluconazole alone.9 In a similar study published in 2021, Godil and colleagues discovered that basil leaf (Ocimum sanctum Linn.) had antifungal activity by inhibiting the growth of Candida sp., but in an in vitro test, the fluconazole inhibition zone was larger than that of basil (Ocimum sanctum Linn.) against Candida.¹⁰

In a study by Khan and colleagues in 2014, fluconazole revealed an MIC value of 256 μ g/mL, capable of inhibiting *Candida* sp., whereas MIC of eugenol were 400 μ g/mL, showing that MIC of fluconazole were inferior to MIC of eugenol. This indicates that fluconazole is more sensitive in inhibiting *Candida* sp.¹¹ Basil leaf (*Ocimum sanctum* Linn.) have been shown to inhibit *Candida* sp. in previous studies, but the antifungal activity of basil leaf (*Ocimum sanctum* Linn.) is considered small due to the high MIC. Eugenol is a constituent of basil leaf (*Ocimum sanctum* Linn.) that has been shown to be effective in inhibiting *Candida* sp.⁹

The fluconazole inhibition zone was also significantly higher than the eugenol and basil

leaf extract (*Ocimum sanctum* Linn.) inhibition zones. Fluconazole works by inhibiting the enzyme lanosterol 14-demethylase, a microsomal cytochrome P450 enzyme found in fungal cell membranes that prevents lanosterol from being converted to ergosterol. Disturbances in these enzymes cause the integrity of the fungal membrane to be disrupted, causing fungal growth to be inhibited and thus having a fungistatic effect.^{4,12}

In a study published in 2020 by Sharifzadeh and Shokri, it was discovered that eugenol can inhibit the growth of *Candida* sp.¹³ Eugenol has antifungal effect by inhibiting the an biosynthesis of ergosterol, a key component of fungal cell membranes, causing damage to the membranes and a reduction in function. Damage to cell membranes disrupts the transport of nutrients (compounds and ions) through the cell membrane, preventing fungal cells from growing and resulting in cell lysis. Eugenol also contains lipophilic compounds, which can penetrate the lipid bilayer membrane, which is made up of fatty acid chains, altering the fluidity and permeability of the cell membrane, causing the cell to lose its structure and function, resulting in cell lysis.¹⁴ Eugenol can also prevent the formation of pre-formed biofilms and Candida albicans biofilms. This substance inhibits Candida albicans biofilm resistance to fluconazole through an adaptive mechanism.¹⁵

In several studies investigating the antifungal effect of eugenol, the mechanism by which eugenol induces *Candida* cell death has not been fully understood. The inactivation of ergosterol synthesis and the production of free radicals, which can be an antifungal effect of eugenol, are the mechanisms of action of eugenol against *Candida*. However, the mechanism of action of eugenol is probably not related to the degradation of *Candida* fungal cell walls.¹⁵ According to Jawetz and colleagues, the

mechanism by which antifungal active substances inhibit fungal growth is through membrane damage. The integrity of the cellular components will be compromised if the cell membrane is damaged, and the fungal respiration process will be disrupted. In the end, there is insufficient energy for active substance transport, causing fungal growth to be disrupted.¹⁶

The results of this study showed that by giving basil leaf extract (*Ocimum sanctum* Linn.) to both *Candida albicans* and *Candida* non-*albicans*, all isolates showed inhibition zones for the growth of *Candida* sp. Basil leaf (*Ocimum sanctum* Linn.) are a tropical plant that can be found throughout Indonesia. Research by De Ornay *et al.* in 2017 and Desmara *et al.* in 2017, found antifungal activity from basil leaf extract (*Ocimum sanctum* Linn.) on the in vitro growth of *Candida albicans.*^{7,8} Eugenol is the main component of basil leaf (*Ocimum sanctum* Linn.). Several in vitro and in vivo studies show that eugenol inhibits the growth of the *Candida albicans*.¹⁷

Khan and colleagues found that basil leaf (*Ocimum sanctum* Linn.) inhibited the transition of yeast formation into hyphae, inhibited fungal proteinase enzymes, and inhibited the expression levels of HWP1, SAP1, and PLB2 genes, which are pathogenesis crucially expressed during a *Candida albicans* infection, in a study published in 2014. As a therapeutic effect of the content of basil leaf (*Ocimum sanctum* Linn.), the content of basil leaf (*Ocimum sanctum* Linn.) can cause *Candida* apoptosis by performing programmed cell death (PCD) on fungi.¹¹

The main component of eugenol, which is highly volatile, may have caused the inhibition zone of basil leaf extract (*Ocimum sanctum Linn*.) to be inferior to fluconazole in the study.¹⁸ The antifungal activity test method used can also have an impact; in this study, the diffusion method was used to test antifungal activity, which is a common method. The diffusion method has the advantage of being a simple and quick way to see the antifungal effect by measuring the diameter of the inhibition zone formed; however, the disc diffusion method has a disadvantage in that not all substances are absorbed on the paper disc, potentially affecting the inhibition zone formed.¹⁶

The concentration of the extract, the content of antifungal compounds, the type of fungus inhibited, and the diffusion power all affect the antifungal activity of a compound. The concentration of the extract can also influence the inhibition zone formed, with the higher the concentration, the larger the clear zone. The more active compounds present, the more focused the concentration, affecting the diameter of the inhibition zone formed on fungal growth.¹⁹

Several other studies produced very encouraging results. According to Ahmad and colleagues 2010 research, eugenol is an antifungal agent with antifungal activity in vitro against Candida albicans and Candida non-albicans, that are intrinsically resistant to fluconazole. The interaction of eugenol and fluconazole was found to have a high level of synergism. In the strains that were tested, no antagonistic interactions were found. Fluconazole in combination with eugenol improves efficacy while lowering the fluconazole minimum effective dose.9

In a study conducted by Jafri and colleagues in 2020, it was discovered that administering fluconazole and eugenol to *Candida* isolates worked synergistically, with eugenol disrupting cell membrane integrity and allowing the entry of standard fluconazole drugs into mold cells. Combining antifungal drugs with eugenol has

several benefits, including increased potency, reduced drug doses, and reduced toxicity, all of which help to inhibit or overcome biofilms and antifungal drug resistance.²⁰

Given the growing treatment failure and antifungal resistance in *Candida* sp., more in vivo studies are needed to assess the potential of these compounds for therapeutic applications and suggest ways to treat resistant *Candida* infections using a combination drug approach. The synergistic interactions between eugenol and antifungal drugs must also be evaluated, with additional studies in animal models needed to assess therapeutic efficacy, topical formulation and toxicity.^{9,20}

References

- Quindós G, Gil-Alonso S, Marcos-Arias C, Sevillano E, Mateo E, Jauregizar N, dan Eraso E. Therapeutic tools for oral candidiasis: Current and new antifungal drugs. *Med Oral Patol Oral Cir Bucal*. 2019;24(2):172-80.
- 2. Murtiastutik D, Maharani CS, Rahmadewi R, dan Listiawan MY. Nystatin profile on Candida species in HIV/AIDS patients with oral candidiasis: a phenomenology study. *J Pure Appl Microbiol*. 2019;**13**(**4**):2013-19.
- 3. Patil S, Majumdar B, Sarode SC, Sarode GS, dan Awan KH. Oropharyngeal candidosis in HIV-Infected patients—an update. *Front Microbiol.* 2018;1(1):1-9.
- Kundu RV, dan Garg A. 2012. 'Yeast Infections: Candidiasis, tinea (pityriasis) versicolor, and malassezia (pityrosporum) folliculitis'. In: Goldsmith LA, Katz SI, Gilchrest BA, Paller AS, Leffel DJ, Wolff K, editors. Fitzpatrick's dermatology in general medicine. 8th ed. New York: McGraw Hill. 2298-301.
- Lydiawati E, Listiawan MY, Murtiastutik D, Rahmadewi R, Prakoeswa CRS, Avanti C, Fitriani E, Astha ET, Astari L, dan Zulkarnain I. In vitro antifungal susceptibility testing of tea tree oil (TTO) 5% compared with nystatin against Candida sp. as important agent of oral candidiasis in HIV/AIDS patients. *BIKKK*. 2020;**32(3)**: 189-94.

- 6. Rauseo AM, Coler-Reilly A, Larson L, dan Spec A. Hope on the horizon: novel fungal treatments in development. *Open Forum Infect Dis.* 2020;7(2):5-23.
- Desmara S, Rezeki S, dan Sunnati S. Konsentrasi hambat minimum dan konsentrasi bunuh minimum ekstrak daun kemangi (O. sanctum L.) terhadap pertumbuhan Candida albicans. *J Caninus Denst.* 2017;2(1): 31-9.
- 8. De Ornay AK, Prehananto H, dan Dewi ASS. Growth inhibition of Candida albicans and power kill Candida albicans extract basil leave. *Jurnal Wiyata*. 2017;**4**(1):78-83.
- Ahmad A, Khan A, dan Yousuf S. Proton translocating ATPase mediated fungicidal activity of eugenol and thymol. *Fitoterapia*. 2010;81(1):1157-62.
- Godil AZ, Bhagat D, Das P, Kazi AI, Dugal, R, dan Satpute S. Fluconazole and Ocimum sanctum Oil in soft denture liners to treat biofilms of Candida albicans associated with denture stomatitis. *Dentristry*. 2021;9(1):1-11.
- Khan A, Ahmad A, Khan LA, dan Manzoor N. Ocimum sanctum (L.) essential oil and its lead molecules induce apoptosis in Candida albicans. *Res Microbiol*. 2014;165(6):411-9.
- Ghannoum M, Salem I, dan Christensen L. 2019. 'Antifungals'. In: Kang, S., Amagai, M., Bruckner, A.L., Enk, A.H., Margolis, D.J., McMichael, A.J., Orringers, J.S., editors. Fitzpatrick's dermatology. 9th ed. New York: McGraw Hill, 3436-45.
- 13. Sharifzadeh A, dan Shokri H. In vitro synergy of eugenol on the antifungal effects of voriconazole against Candida tropicalis and Candida krusei strains isolated from the genital tract of mares. *Equine Vet J*. 2020;**3**(**1**):94-101.
- 14. Brajawikalpa RS. dan Ramzy AN. Uji efektivitas antijamur minyak atsiri daun cengkeh (Syzygium aromaticum L.) terhadap pertumbuhan Malassezia furfur. *Jurnal Kedokteran dan Kesehatan*. 2018;**4**(1):49-51.
- 15. Silva ICG, Santos HBP, Cavalcanti YW, Nonaka CFW, Sousa SA, dan Castro RD. Antifungal activity of eugenol and its association with nystatin on Candida albicans. *Pesquisa Brasileira em Odontopediatria e Clínica Integrada*. 2017;**17(1)**:1-8.
- Jawetz, Melnick, dan Adelberg. 2007. 'Mikrobiologi Kedokteran'. 23rd ed. Jakarta: ISBN 978-979- 448-859-1. 273-5.

- Núñez IC, Arranz JC, Rivas CB, Mendonça PM, Perez K, Sánchez CD, Cortinhas LB, Silva CF, Carvalho M, dan Queiroz M. Chemical composition and toxicity of O. sanctum L. var. cubensis essential oil upgrowing in the Eastern of Cuba. *Phytopathology*. 2017;9(7):1021-8.
- 18. De Paula SB, Bartelli TF, Di Raimo V, Santos JP, Morey AT, Bosini MA, Nakamura CV, Yamauchi LM, Yamada-Ogatta SF. Effect of eugenol on cell surface hydrophobicity, adhesion, and biofilm of Candida tropicalis and Candida dubliniensis isolated from oral cavity of HIV-Infected

patients. *Evid Based Complement Alternat Med.* 2014;**1(1)**:1-8.

- 19. Andayani A, Susilowati A, dan Pangastuti A. AntiCandida minyak atsiri lengkuas putih (Alpinia galanga) terhadap Candida albicans penyebab candidiasis secara invitro. El Vivo. 2014;**2**(2):1-9.
- 20. Jafri H, Banerjee G, dan Khan MSA. Synergistic interaction of eugenol and antimicrobial drugs in eradication of single and mixed biofilms of Candida albicans and Streptococcus mutans. *AMB Expr.* 2020;**1**(**10**):185.