

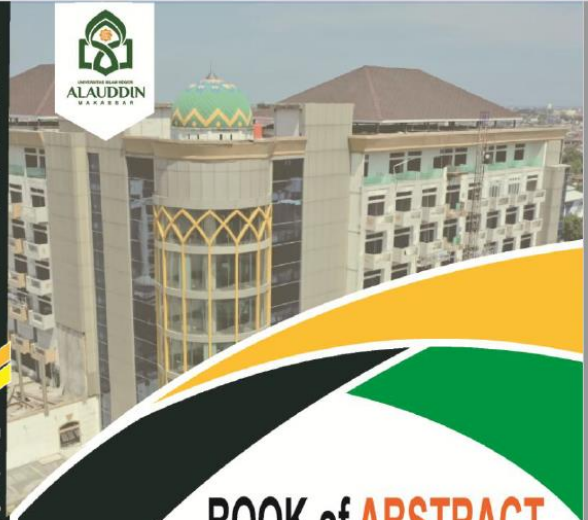


MEDICAL AND HEALTH SCIENCE FACULTY
MAKASSAR, 27th-28th AUGUST 2022



The 1st Alauddin Health and Medical International Conference

Book of Abstract Alhamdic 2022



BOOK of **ABSTRACT**

ALAUDDIN HEALTH AND MEDICAL
INTERNATIONAL CONFERENCE

NOVOTEL HOTEL, 27th-28th AUGUST 2022



UNIVERSITAS ISLAM NEGERI ALAUDDIN MAKASSAR
MEDICAL AND HEALTH SCIENCE FACULTY

Introduction

Dr. Syatirah Djalaluddin, SpA., M.Kes

Dean of Medical and Health Science Faculty, Universitas Islam Negeri
Alauddin Makassar

Medical and Health Science Faculty, Universitas Islam Negeri Alauddin Makassar hold the Alauddin Health and Medical International Conference (Alhamdic) 2022 with the theme *"Advance Technology in Health and Medicine in Improve Quality of Life"*. This activity aims to be a forum for socializing and exchanging ideas and experiences related to technological advances in health and medicine in improving the quality of life. In addition to this program, it is expected that stakeholders, industries, and researchers can establish business or research collaboration that focus on key issues in the future on a national and international scale.

The program activity will be in the form of a scientific seminar consisting of plenary lectures, oral presentations and poster presentations. The target participants are either practitioners or academics (lecturers, researchers and students) in the health and medical fields, from within and outside the country.

The main objective of this program is to socialize and increase knowledge about technological advances in health and medicine in improving the quality of life. Specifically, this activity aims to:

- As a medium for meeting and exchanging information between researchers and business partners
- As a means of socializing and identifying the level of understanding of the general public about technological advances in health and medicine in improving the quality of life
- Develop research networks and other collaborations in the field of health and medicine both on a national and international scale

Thank you for participating and see you at the next scientific event

Campus Map

Fakultas Kedokteran Dan Ilmu Kesehatan

UIN Alauddin Makassar

<https://www.google.com/maps/@-5.2055754,119.4986499,237m/data=!3m1!1e3>





PROGRAM AT A GLANCE OF ALHAMDI (ALAUDDIN HEALTH & MEDICAL INTERNATIONAL CONFERENCE)

MEDICAL & HEALTH SCIENCE FACULTY OF UIN ALAUDDIN MAKASSAR, INDONESIA



Time	AGENDA 1st Day (27th August 2022)	PIC	ROOM
07.30-08.30	Registration	Committee	Registration Desk
	Opening Ceremony		
08.30-08.32	Welcome Speech by MC		
08.32-08.42	Traditional Dance "Padduppa" By Rutikah		
08.42-08.45	National Anthem "Indonesia Raya" & Main UIN Alauddin Makassar		
08.45-08.50	Recitation of the Holy Quran By Gori Rahmat Hidayat, S.Sos		
08.50-08.55	Opening Speech: Dean of Medical and Health Science Faculty UIN Alauddin Makassar: Dr. dr. Syatrah, Sp.A, M.Kes	MC	
08.55-09.00	Opening Speech: Rector of UIN Alauddin Makassar: Prof. Hamdan Juhanis, Ph.D		
09.00-09.05	Opening Speech & Opening The Conference Officially: Governor of South Sulawesi Province, Indonesia: H. Andi Sudirman Sulaiman, S.T		
09.05-09.10	Recitation of Do'a By Dr. Abdul Hakim, Sp.d, M.Pd		
09.10-09.20	Coffee Break		
	Opening Symposium 1	MC	
	Symposium 1		
09.20-09.30	Traditional Dance "4 Etnisitas" By Rutikah		
09.30-10.00	Director of MST Malaysia: Madam Marlan Mohd Nasir, A.M.N, ETh		
10.00-10.30	Thammasat University Thailand: Rathaporn Aassutjarit, Ph.D		
10.30-11.00	Chairman of the SPUS Health Supervisory Board: Prof. dr. Abdul Kadir, Ph.D, Sp.THT-KL(2), MARS		
11.00-11.30	General Director of Islamic Education, Ministry of Religious Affairs: Prof. Dr. Muh Ali Ramdhan, S.TP., M.T		
11.30-12.00	Discussion		
12.00-13.00	Lunch (Product Presentation by Merka Venus)	MC	
	Symposium 2	MC	
13.00-13.30	UIN Alauddin Makassar, Indonesia: Dr. Hj. Siti Saleha, S.St, SKM, M.Keb		
13.30-14.00	UIN Alauddin Makassar, Indonesia: Ns. Ahmad Jamaluddin, M.Kep, Sp.Kep.MS, ETh		
14.00-14.30	UIN Alauddin Makassar, Indonesia: Dr. Andi Sutisawaty, S.St, M.Kes		
14.30-15.00	UIN Alauddin Makassar, Indonesia: Dr. apt. Karina Amir Yahr, M.St		
15.00-15.30	Discussion		
15.30-16.00	Coffee Break & Closing	MC	
Time	2nd Day (28th August 2022)	PIC	ROOM
07.30-08.30	Registration & Opening	Committee	Registration Desk
	Symposium 3		
08.30-09.00	West Virginia University, USA: Salman Khan, MD, Ph.D		
09.00-09.30	Xoenda, Comoliton, USA: Ryan Plano, MPH., Ph.D		
09.30-10.00	WHO: Mr. Stephan Himley, MD, MPH		
10.00-10.30	Discussion		
10.30-10.45	Coffee Break	MC	
10.45-12.00	Poster Presentation	Committee	Hall 7 (Hall Room Front Hall)
12.00-13.00	Lunch		
13.00-15.00	Oral Presentation	Moderator	Breakout Room
		Yusmaidah Jayadi, S.Gi, M.Kes	Hall 1 (Online)
		Ns. Andan Athiwiqaya, M.Kep	Hall 2 (Online)
		Mubrizah, S.S, M.S	Hall 3 (Offline)
		Titi Adhitya Kartini, MPH	Hall 4 (Offline)
		Rinaewati Aulia Inward Sadarang, MPH	Hall 5 (Offline)
		dr. Andri Faridillah, Sp.GC, M.Kes	Hall 6 (Offline)
15.00-16.00	Coffee Break	MC	
	Closing Ceremony		
16.00-16.30	Announcement: Best poster & oral presentation	MC	
	Closing Speech Officially by Dean of Medical and Health Science Faculty, UIN Alauddin Makassar		

List Of Content

	Page
Title Page	i
Introduction	ii
Conference Agenda	iii
List of Content	iv
Antiplasmodial activities test from non-chloroform fraction and its sub-fraction from <i>Hyrrios reticulatus</i> sponge extract	Mahfur Mahfur 1
A Narrative Inquiry Study On Parents About Children With Post Traumatic Stress Disorder After Disaster	Etik Rositasari 2
Bougenville flower (<i>Bougainvillea spectabilis</i> wild) active fraction test of ethanol extract (in vitro) as sunscreen	Ni Nyoman Yuliani 3
Coprocessed Lactosa-Sorbitol as a Filler Excipient in Formulation of Tablet Teophylline by Direct Compression	A. Mumthahanah Mursyid 4
Analysis of Chemical Compound, Antioxidant Activity, and Conventional and Molecular Characterization of Endophytic Fungi Isolated from <i>Muntingia calabura</i> L. Blossoms	St. Maryam 5
The Suitability of The Prescription of Non-Sterile Concoctions For Children at XXX Mother and Child Hospital Makassar : Compatibility and Stability Stud	Vina Purnamasari M 6
Effect of N1- Benzoiloxymethyl-5-Fluorouracil against The p53 Expression and Apoptosis Activity In Breast Cells of Female Mice (Sprague-Dawley) Induced with 7.12-Dimethylbenz(a)anthracene (DMBA)	Dewi yuliana 7
Anti inflammatory activities against LPS-induced J774.1 Cell and Total Flavonoid content of <i>Vitex coffasus</i>	Faradiba Abdul Rasyid 8
Radiotracer on Non-Small Cell Lung Cancer for EGFR L858R/T790M Double Mutations	Muammar Fawwaz 9
Effectiveness Of Moringa Honey On Hemoglobin Level In Pregnant Women With Anemia	Alifia Ayu Delima 10
How To Improve Maternal Health Care Quality: Scoping Review	Mahindria Vici Virahaju 11
The Effect Of Java Massage Towards The Quality Of Life Child With Asthma	Retno Puji Hastuti 12
The "Stopping the Heart" Technology Improves Quality of Life in South Sulawesi	Jayarasti Kusumanegara 13

Antiinflammatory activities against LPS-induced J774.1 Cell and Total Flavanoid content of *Vitex coffasus*

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Context: The secondary Metabolite compounds found in the vitex sp are terpenes, flavonoids, and lignans. Flavanoids have shown antiinflammatory, anti-oxidative, anti-mutagenic, and anti-carcinogenic properties. Virex coffasus (VCS) research was limited to cancer activities and larvicides compared to other vitex species.

Aim: This study aims to determine the anti-inflammatory activity against LPS-induced J774.1 cells. And the total flavonoid content of *Vitex coffasus*

Method: : Initially, the *Vitex coffasus* leaves were extracted by using methanol. Next, the extracts were divided into three concentrations 10, 30, and 100 µg/ml, and then evaluated Antiinflammatory by using J774.1 cells function as a macrophage and after being triggered by LPS from *Escherichia coli*, these macrophages could produce Nitric oxide (NO). Inhibition of NO was evaluated using Griess reagent. The qualitative analysis of flavonoids used TLC and visualization by spraying with AlCl₃. The determination of total flavonoids is conducted based on the AlCl₃ with total flavonoids expressed in QE (Quercetin equivalent) at the maximum wavelength of 431 nm.

Results: The extracts effectively reduced lipopolysaccharide (LPS)-induced NO levels in a 100 µg/ml concentration. The total flavonoid content of VC leaf extract is 1.0318 mg QE / g extract. was 0,003817 mgQE/g extract

Conclusion: From these results, we can conclude that the VCS leaves possess the antiinflammatory activity and total flavonoid content was 0,3817%

Keywords: Antiinflammatory, Flavanoid, *Vitex coffasus*

Introduction

Bitti wood (*Vitex cofassus*) is one of Sulawesi's indigenous plants known as gofasa. Bitti wood (*V. cofassus*) is utilized as a construction material and as a deodorizing agent by the inhabitants of South Sulawesi. Several investigations have shown that *Vitex coffasus* has anticancer properties¹. Secondary metabolites of flavonoid and alkaloid kinds found in bitti wood (*V. Coffasus*) have the capacity to inhibit *Staphylococcus aureus* germs.

According to previous work, VCS has anticancer properties. Several studies indicate that cancer has a relationship or association with other disorders, specifically inflammation²⁻⁴, which is one of the body's natural defense systems against outside incursions such as microbes and can manifest as rash, fever, and sore³. Cancer development has been linked to persistent and permanent inflammation. Its emergence might be connected to tumor-related inflammation^{4, 8-11}. Furthermore, infection and inflammation were linked to nearly 25% of all tumor incidents⁴. Nuraini's (2014)⁵ research

it was explained that the bark of the beetroot (*V. coffasus*) contains secondary metabolites, namely flavonoids which are proven to be anticancer and antiinflammation. Several medicinal plants containing flavonoids have antioxidant, antibacterial, antiviral, anti-inflammatory, allergy, and anticancer activities⁶. Flavonoid compounds are thought to be very useful in food because, in the form of phenolic compounds, these compounds are potent antioxidants. Many disease conditions are known to be exacerbated by the presence of free radicals such as superoxides and hydroxyl, and flavonoids can virtually eliminate these damaging oxidizing species⁷.

Based on the description above, this research was conducted to determine the anti-inflammatory and flavonoid levels of *Vitex cofassus* to increase scientific data from medicinal plants.

Methods

Preparing Powder From *Vitex coffasus* Leaves

The VCS (*Vitex cofassus*) leaf samples were collected, washed with running water, and dried in a 40-50°C drying cabinet. The sample is then ready for extraction after being ground into a powder using a blender.

The leaves of this plant were crushed into small chunks then macerated with methanol (1:20, v/v) at ambient temperature (28 ± 2 °C) for 3 days with periodic stirring. After filtering the extract, powdered marc was re-macerated in the same organic phase until the extraction was accomplished. The filtrate resulting from maceration is collected and then dried and use a rotary evaporator⁹.

Anti-inflammatory action against J774.1 cells stimulated by LPS

J774.1 murine macrophage cells were placed into 96-well plates containing 100 µL per well. These were then cultured for one day in the CO₂ incubator. On the following day, the samples were split up into two groups, one of which had LPS and the other of which had not. The plate was withdrawn from the incubation, and thus the old media was substituted with a sample that had earlier been prepared. The plate was incubated for another one day. On the next day, a Griess reagent was created by blending sulphanilamide and N-1-naphthylethylenediamine dichloride. Subsequently, it was dissolved in aqueous solution containing phosphoric acid, and then extra was added to it. Approximately 100 µL of Griess reagent was added to fresh 96-well plates.

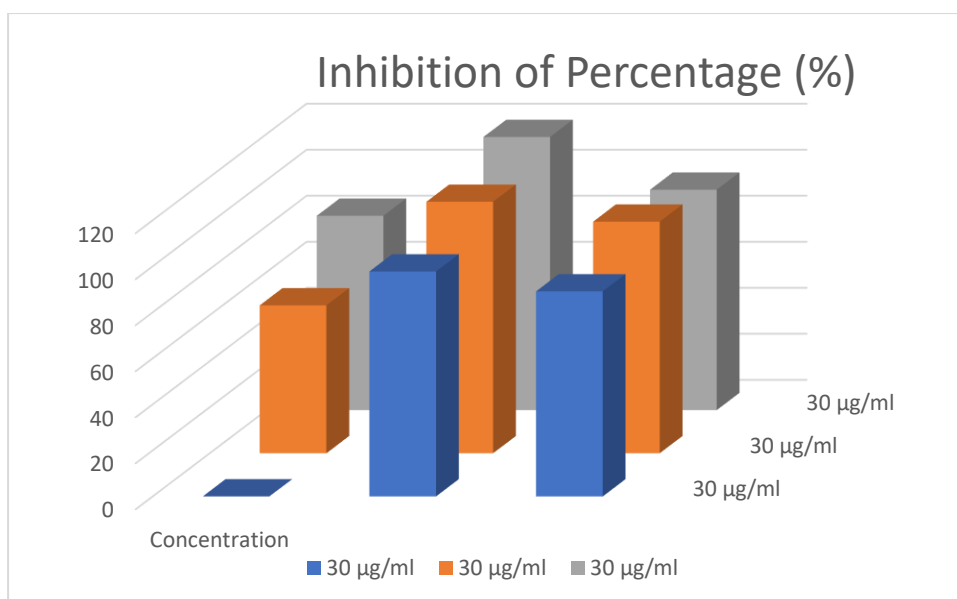
The cell plates were withdrawn from the incubator, placed in a cooler, and 100 μ l was placed to the Griess agent plate. The absorbents as from sample in the Griess solution plate were formed using a colorimetric method and then the percent of NO inhibition was determined. 200 g of VCS (*Vitex coffasus*) leaf extract was macerated in 2.75 L of methanol to yield a weight fraction of 10.063%. The outputs of extracting samples of VCS leaves (*Vitex coffasus*) are presented in the table below.

Table 1. Yield value of VCS (*Vitex coffasus*) leaves extract

Sample	Sample weight (g)	Extract result (g)	Yield value (%)
VCS Leaves	200	12,92	6,46

The result of anti-inflammatory of VCS leaves (*Vitex coffasus*) can be seen in the following table 2 :

Concentration (μ g/ml)			
Inhibition of Percentage (%)			
10	44,80	97,73	89,09
30	64,25	109,25	100,61
100	84,40	118,65	95,76



qualitative test was performed to determine the sample's chemical components using the NaOH 10 % reagent, .

Table 3. Qualitative test Results of VCS leaf (*Vitex coffasus*) extract flavonoid compound using TLC

Sample	Flavonoid test (NaOH 10 %)	Reference (Theodore, 2019)	Result
VCSLeaf	Brown-orange	Orange	+

Note : + (contains flavonoid)

Table 4. Results of Quercetin Absorbance Measurements at 431 nm

Concentration (ppm)	Absorbance
40	0,259
50	0,339
60	0,407
70	0,506
80	0,568

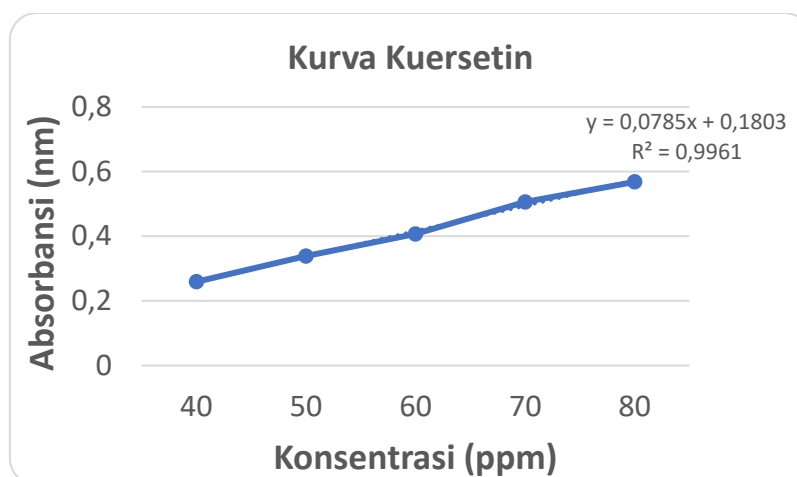


Figure 2. Quercetin Calibration curve at a maximum wavelength of 431 nm

Table 5. Determination of Total Flavonoid Levels of VCS(*Vitex coffasus*) leaf extract

Sample	Replication	Absorbance	Initial	Total	Total	Flavanoid
			Flavonoid	Flavonoid	Flavonoid	content
			Content	Content	Content	(5)
			(mg/ml)	(mgQE/ g	average	
				eks)	mgQE/ g	
					ekss	
VCSLeaves	1	0,473	0,003728	0,003728	0,003817	0,3817
	2	0,489	0,003932	0,003932		
	3	0,478	0,003792	0,003792		

Discussion

According to research published by Faradiba, a particular bioactive components in the vitex coffasus showed anti - cancer properties. The formation of cancer was related to chronic and prolonged inflammation. Its growth may be related with inflammation linked to tumors²⁻⁴. Moreover, roughly 25% of all reported cases were connected with inflammation and infection. 4. In this experiment, the initial anti-inflammatory VCS evaluation test was carried out. Throughout this study, the NO inhibiting test on LPS-induced J774.1 cells termed macrophage proved that these cells exhibited the same characteristics as typical macrophage. They can synthesize and discharge lysozyme, and then when they destroy cells, antibody recognizes the target.

These cells were derived from a BALB/c mouse who suffered from unpremeditated cancer. After being stimulated by LPS from Escherichia coli, the J774.1 cells, that functioned like macrophages, may produce NO. Suppression of NO was tested with griess reagent ¹⁰⁻¹¹

Griess reagents originally established in 1879, and then these assays were widely used to assess nitric oxide in biological materials including urine, plasma, and grown cells. The production of griess reagents is set up as follows: In the beginning, it will form short-term diazonium salt, which is made by a reaction combining nitrite and sufanilamide to which acid had earlier been applied, generally phosphoric acid. After adding N-naphthyl-ethylenediamine, the short-term salt converts to non-labile azo agents with a deep purple color. A plate reader is then utilized to quantify the absorbant or optical density of a strong purplish color to establish the NO content. The absorbent computation results are shown in Table 2 and Figure 1. the VCS t had a significant effect on NO inhibition effect of more than 80 % at 100 µgg/ml level. The VCS may be a prime source for some further anti-inflammatory action.

The plants possess anti-inflammatory action due of the herbal ingredient which is flavonoid ¹².Based on the faradiba's research which conclude that one of the compound found in the vitex coffasus had anticancer activity . The cancer development were subscribed from the chronic and permanent inflammation. its development could be associated with tumour-linked inflammation²⁻⁴. Furthermore, it is approximately that 25 % of all tumour incidents were correlated with infection and inflammation⁴. In this research, the preliminary assay was performed for the screening of anti-inflammatory's VCS. In this research, the NO inhibitory

assay on LPS-induced J774.1 cells as macrophage, it was discovered that these cells have similar properties as the regular macrophage. They can produce and liberate lysozyme and when they destroy cells, the object is determined by antibody. These cells were obtained from a BALB/c mouse which suffered from unpremeditated tumour. The J774.1 cells function as a macrophage and after being triggered by LPS from *Escherichia coli*, these macrophages could produce NO. Inhibition of NO was evaluated using griess reagent¹⁰⁻¹¹

Griess reagents were introduced in 1879, and these methods were widely used to examine nitric oxide in living specimens such as urine and blood, and also in a cell which is cultured in a medium. The process of griess reagents is established as follows: initially, it will produce short-term diazonium salt which is produced from a reaction between nitrite and sulfanilamide that is already added with acid, usually phosphoric acid. This short-term salt becomes non-labile azo agents which has a strong purple colour, after adding a *N*-naphthyl-ethylenediamine. Afterward, a microplate reader is used to measure the absorbant or optical density of high purple colour for NO concentration. Results of the calculations of absorbant can be seen in table 2 and Fig.1. the VCS t had a significant result on NO inhibitory activity of more than 80 % in 100 µgg/ml concentration. The VCS can be a potent source for continuing to develop as aanti-inflammatory activity .

The plants have aantiinflammatory activity because of the plants's compound which is flavonoid ¹². To assess the total flavonoid content of VCS (*Vitex coffasus*) leaf extract, a quantitative study of total flavonoid components was performed using UV-Vis spectrophotometry. UV-Vis spectrophotometry was used for flavonoid analysis because to the conjugated aromatic systems of flavonoids, which exhibit high absorption bands in the ultraviolet light spectrum and visible light spectrum. As a standard solution for measuring total flavonoid concentrations, quercetin was used in this work. Because quercetin is a kind of flavonoid, is often used as a standard for measuring quantities of flavonoids, quercetin is a type of flavonol that is abundant in plants; flavonols contain a keto group on C-4 and a hydroZyl group on C-3 or C-5 atoms, and they may form a complex with AlCl₃ ¹³

The total flavonoid content of VCS (*Vitex coffasus*) leaf extract was determined by dissolving 25 mg of VCS leaf extract in 10 ml p.a. methanol. Then, take 1 mL and add 3 ml of methanol pa, 0.2 mL of 10% AlCl₃ that might create a complex, causing a change in wavelength more toward the visible (visible) and noted with a solution generating a strongly

yellow hue, then add 0.02 mL of potassium acetate to retain the wavelength in the visible region, then add up to 10 mL of aquadestilleda. The combination is kept at ambient temperature for 30 minutes; it is incubated for 30 minutes to ensure that the reaction works precisely, resulting in a greater brightness (Azizah and Faramayuda 2014). They are then examined at 431 nm wavelength. The outcomes of the experiments are presented in table 6. In a concentration of 100 g/ml, the extracts considerably decreased lipopolysaccharide (LPS)-induced levels of nitric oxide (NO). The total flavonoid concentration of VC leaf extract is 1.0318 mg QE per gram of extract.

Conclusion

We may deduce from these data that VCS leaves contain anti-inflammatory potential while their total flavonoid concentration was 0.3817%.

Ethical approval

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Conflicting Interest

All authors declare no conflict of interest.

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