



KEMENTERIAN PENDIDIKAN DAN KEBUDAYAAN
UNIVERSITAS HASANUDDIN FAKULTAS KEDOKTERAN
KOMITE ETIK PENELITIAN KESEHATAN
RSPTN UNIVERSITAS HASANUDDIN

RSUP Dr. WAHIDIN SUDIROHUSODO MAKASSAR

Sekretariat : Lantai 2 Gedung Laboratorium Terpadu

JL. PERINTIS KEMERDEKAAN KAMPUS TAMALANREA EMLTO MAKASSAR 90245

Contact Person: dr. Agussalim Bukhari, M.Med., Ph.D., Sp.GK. TELP. 081241850858, 0411-5780103, Fax: 0411-581431



REKOMENDASI PERSETUJUAN ETIK

Nomor : 820/UN4.6.4.5.31/PP36/2020

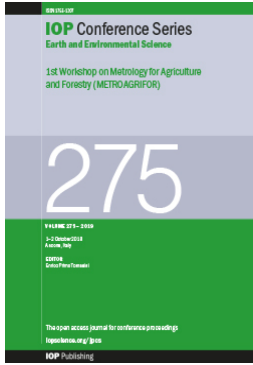
Tanggal: 23 Desember 2020

Dengan ini Menyatakan bahwa Protokol dan Dokumen yang Berhubungan Dengan Protokol berikut ini telah mendapatkan Persetujuan Etik:

No Protokol	UH20110627	No Sponsor Protokol	
Peneliti Utama	Sri Anggarini Rasyid, S.Si, MSI	Sponsor	
Judul Peneliti	Potensi Kerang Pokea (Batissa Violaceae Celebensis Marten 1897) Sebagai Imunostimulan Dan Anti Proliferasi Sel Kanker Secara In Vitro Dan In Vivo		
No Versi Protokol	2	Tanggal Versi	15 Desember 2020
No Versi PSP		Tanggal Versi	
Tempat Penelitian	RS Universitas Hasanuddin Makassar, Balai Besar Veteriner Maros, Laboratorium Biofarmaka dan Biofarmasi Prodi Farmasi STIKES Mandala Waluya Kendari		
Jenis Review	<input type="checkbox"/> Exempted <input checked="" type="checkbox"/> Expedited <input type="checkbox"/> Fullboard Tanggal	Masa Berlaku 23 Desember 2020 sampai 23 Desember 2021	Frekuensi review lanjutan
Ketua Komisi Etik Penelitian Kesehatan FKUH	Nama Prof. Dr. dr. Suryani As'ad, M.Sc., Sp.GK (K)	Tanda tangan	
Sekretaris Komisi Etik Penelitian Kesehatan FKUH	Nama dr. Agussalim Bukhari, M.Med., Ph.D., Sp.GK (K)	Tanda tangan	

Kewajiban Peneliti Utama:

- Menyerahkan Amandemen Protokol untuk persetujuan sebelum di implementasikan
- Menyerahkan Laporan SAE ke Komisi Etik dalam 24 jam dan dilengkapi dalam 7 hari dan Laporan SUSAR dalam 72 jam setelah Peneliti Utama menerima laporan
- Menyerahkan Laporan Kemajuan (progress report) setiap 6 bulan untuk penelitian resiko tinggi dan setiap setahun untuk penelitian resiko rendah
- Menyerahkan laporan akhir setelah Penelitian berakhir
- Melaporkan penyimpangan dari prokol yang disetujui (protocol deviation / violation)
- Mematuhi semua peraturan yang ditentukan



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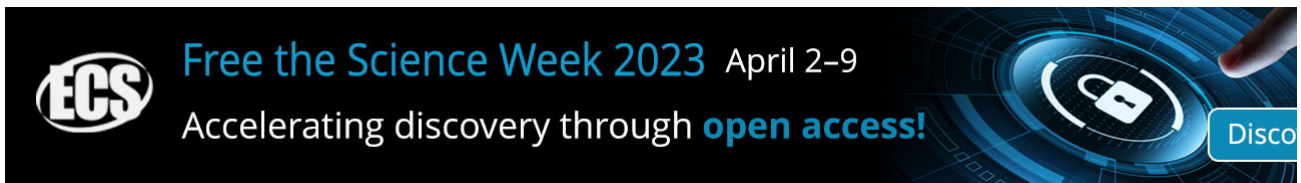
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2021

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7th International Conference on Environment and Renewable Energy 26-28 March 2021, Qingdao, China

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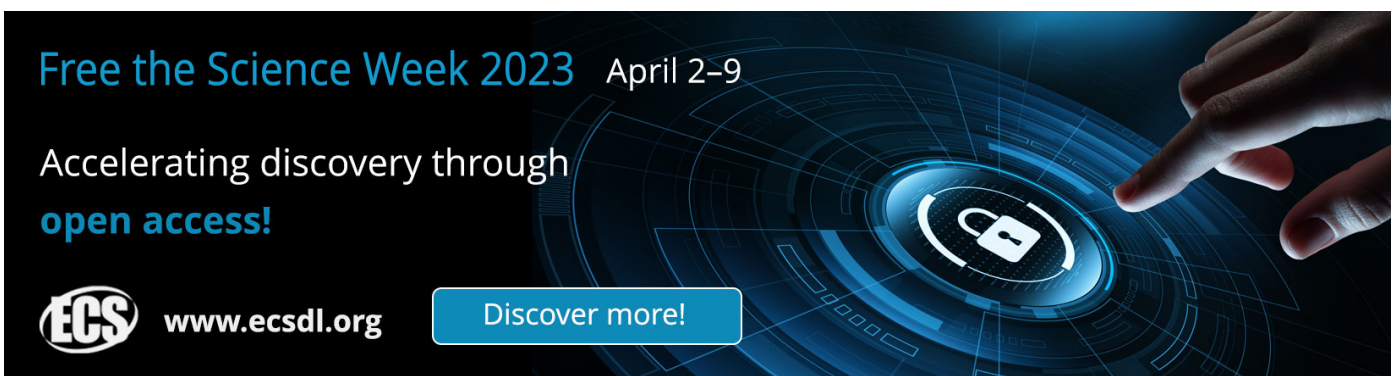
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Screening of Anti-MRSA metabolites in bacteria symbiotic with *Batissa violaceae celebensis* marten 1897

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Abstract. Currently, infectious diseases are still a serious problem in Indonesia, especially with the widespread resistance of microbes to antibiotics. *Staphylococcus aureus* as one of the common pathogens causing infection has experienced resistance to various classes of beta-lactam antibiotics known as Methicillin-resistant *Staphylococcus aureus* (MRSA). Therefore antibiotic alternatives sourced from nature are needed to treat the MRSA pathogenic infection. This study aims to explore the potential of bacteria in symbiosis with Pokea Shells (*Batissaviolaceacelebensis* Martens 1897) from Konawe Regency, Southeast Sulawesi Province to be used as the newest anti-MRSA. In this research, Pokea Shell's symbiont bacteria were isolated from Konawe Regency to obtain pure isolates. Pure isolates that have been produced from secondary metabolites to be tested for their potential in inhibiting the growth of pathogenic bacteria MRSA. The inhibition test was done qualitatively by using a paper disk. Based on the results five bacterial isolates were obtained with different abilities to inhibit MRSA bacteria. Based on the qualitative test obtained that the Bvc1 and Bvc3 bacterial isolates had the potency to be used as anti-MRSA agents. These isolates have an inhibition zone of 25 mm and 22 mm with a sensitive category according to CLSI standards (Sensitive ≥ 17 mm). Determination of the gram of these 2 isolates found as gram-positive bacteria in the form of bacilli so it can be concluded that the 2 isolates can be used as anti-MRSA agents.

Keywords: Anti-MRSA, *Batissa violacea celebensis* Martens 1897, Screening, Symbiont bacteria



1. Introduction

The danger of antibiotic resistance is recognized as an ongoing serious problem both in developed and developing countries[1]. In Indonesia, it exacerbates the serious threat of infectious diseases. *Staphylococcus aureus*, one of the common pathogens causing various infections, has evolved resistance to various classes of antibiotic beta-lactam, known as *Methicillin-resistant Staphylococcus aureus* (MRSA)[2]

Infections due to *Staphylococcus aureus* are usually treated with the administration of antibiotics[3]. However, in some cases, several strains of *Staphylococcus aureus*, such as *Methicillin-resistant Staphylococcus aureus* (MRSA), were found to be resistant to antibiotics[4]. One of the worst outcomes of MRSA infection is amputation. Apart from causing wound severity, MRSA is also strongly suspected to exacerbate pneumonia (lung infection) [5].

Indonesian waters are rich in invertebrates such as Mollusca, one of which is Bivalvia or shellfish [6]. Southeast Sulawesi Province is known for its biodiversity. The evidence of it can be found in the freshwater ecosystem on the Pohara River in Konawe District. The Pohara River holds potential biological resources such as freshwater clams known as *pokea* (*Batissa violacea celebensis* Martens 1897) by local people[7]. Several studies [8] mentioned that bivalves possess a plethora of benefits on human health. According to [9], *pokea* clam extract possesses the ability to inhibit the growth of *Staphylococcus aureus*. [10] reported that jackknife clam (*Solen* sp.) and windowpane oyster (*Placuna placenta*) extracts possess antibacterial activities against *S. aureus*. [11] reported that the extract of *Atactodeastriata* clam possesses antibacterial activities against *S. aureus* and *E. coli*. [13] reported that *Pokea* clam can empirically treat various diseases, one of which is an infectious disease caused by microorganisms such as bacteria. Therefore, this study was conducted in the hope to identify the anti-MRSA potential of bacterial isolates that are symbiotic with *Batissa violacea celebensis* Martens 1897.

2. Materials and Methods

As an explorative study, this study sought to find and identify the symbiont bacteria of *pokea* clam (*Batissa violacea celebensis* Martens 1897) that hide the potential as an anti-MRSA. This study consisted of several stages :

2.1. A sampling of *Pokea* Clams (*Batissa violacea celebensis* Marten 1897).

The source of bacterial isolates used in this study was *pokea* clams, which is a local endemic resource of Konawe District, Southeast Sulawesi Province. These clams were taken from the Pohara River and put into sterile plastic bags and stored in a cool box.

2.2. Isolation of Symbiont Bacteria

The symbiont bacteria were isolated using the pour plate method on nutrient agar media. The bacteria association were incubated overnight at 28 °C. *S. aureus* was inoculated on nutrient broth and was grown overnight at 27 °C according to the shaker condition. After the incubation period, colony morphological observations were made to determine which isolates were purified by the pour plate method. The bacterial isolates obtained were purified by re-isolation to ensure the purity of the culture. The purified isolates were then stored on agar slants as isolate stocks for further testing [13].

2.3. Production of Bacterial Secondary Metabolite Compounds

Secondary metabolite compounds were produced based on the phase of bacterial growth, which was the stationary phase. This process was performed by taking 10% (10^7 cells/mL) of bacterial culture to be inoculated into 50 mL of production medium, which was then shaker incubated in accordance with the optimum incubation time. Then, the metabolites were harvested and centrifugated at 1000 rpm for 10 minutes to obtain a metabolite compound form of supernatants (free cells) [13]

2.4. Testing of the Secondary Metabolite Compound Against MRSA Bacteria

The qualitative test of the antagonist test of the bacterial metabolite compound was carried out by testing the secondary metabolites of bacteria in the form of free cells. Disc paper with a diameter of 6 mm which has been inoculated with a bacterial supernatant with a cell density of 10^7 cfu/ml was placed on the surface of the agar medium that has been inoculated with MRSA bacteria with a density of 10^7 cfu / ml and then incubated at 37°C for 24 hours [14]. The diameter of the clear zone formed around the disc was calculated using a digital caliper.

3. Result and Discussion

The results of the test of anti-MRSA potential of bacterial isolates that were symbiotic with *Batissaviolaceacelebensis* Martens 1897 are as follows:

3.1. Results of Symbiont Bacteria Isolation

The isolation of symbiont bacteria was performed using the pour plate method. After the incubation period, the growing bacterial colonies were observed morphologically. The results are presented in Table 1.

Table 1. The Number of Bacterial Isolates That Are Symbiotic with *Batissa violacea celebensis* Martens 1897 based Differences in Colony Morphology

No.	Isolate Code	Colony Shape	Edge	Elevation	Colour	Inner Structure
1.	Bvc1	Circular	Entire	Low Convex	Beige	Transparent
2.	Bvc2	Irregular	Undulate	Low Convex	Beige	Transparent
3.	Bvc3	Circular	Entire	Low Convex	Beige	Transparent
4.	Bvc4	Irregular	Undulate	Low Convex	Beige	Transparent
5.	Bvc5	Circular	Entire	Low Convex	Beige	Transparent

3.2. Results of Inhibition Test against MRSA

The bacterial inhibition test was carried out by using the disc diffusion method. The results are presented in the following table 2 and figures 1 and 2:

Table 2. Inhibition Ability of Bacteria That Are Symbiotic with *Batissa violacea celebensis* Martens 1897

No.	Isolate Code	Inhibition Zone Area (mm)	Description*
1.	Bvc1	25	Sensitive
2.	Bvc2	-	
3.	Bvc3	22	Sensitive
4.	Bvc4	-	
5.	Bvc5	-	
6.	Control +	17	Sensitive
7.	Control -	-	

*Resistant (≤ 13 mm), Intermediate (14-16 mm), Sensitive (≥ 17 mm)
(Standard Clinical and Laboratory Standards Institute (CLSI))

Based on the results of a qualitative screening test, two bacterial isolates were produced which had inhibitory activity against MRSA pathogenic bacteria and the two isolates were used for the next test stage. The five isolates that possessed the highest inhibitory activity were Bvc1 (25 mm) and Bvc3 (22 mm). Isolate Bvc1 and Bvc3 had a wider inhibition zone than the positive control (penicillin). Penicillin is a methicillin class of antibiotics that are already resisted by *Staphylococcus aureus*. One form of antibiotic resistance by *Staphylococcus aureus* bacteria is resistant to penicillin class antibiotics[15].

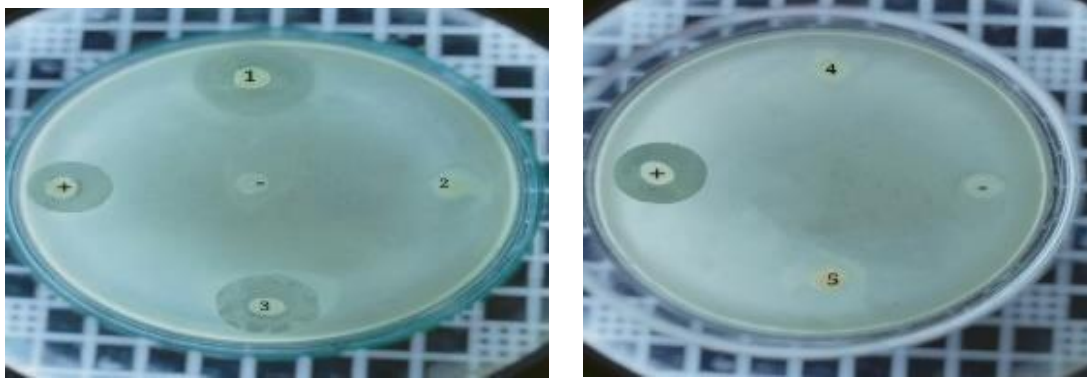


Figure 1. The results of the Test of Anti-MRSA Potential of Bacterial Isolates That Were Symbiotic with *Batissaviolaceacelebensis* Martens 1897

3.3. Characteristic Results of Bacterial Isolate Cells with Potential Anti-MRSA

Bacterial isolates that had inhibition zones were characterized by Gram staining. The characteristic results of bacterial isolate cells with potential anti-MRSA are shown in the following figure.

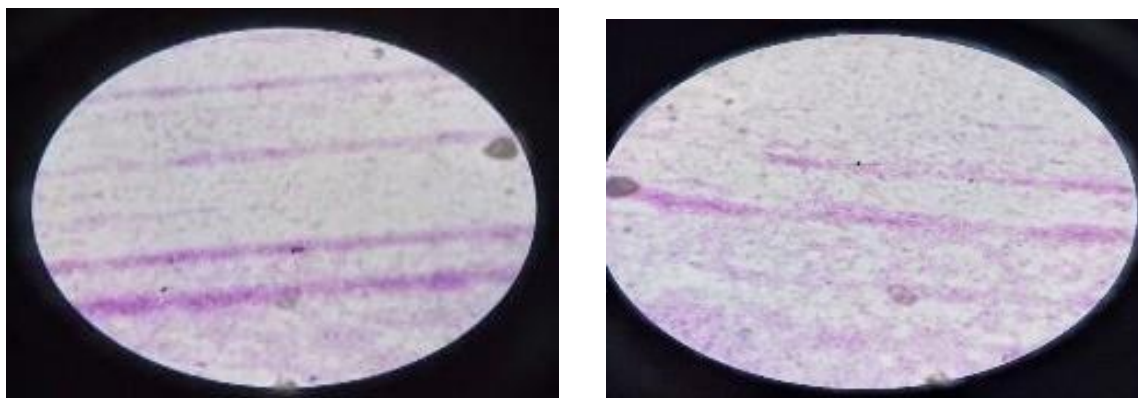


Figure 2. The Results of the Bacterial Isolate Staining Test That Were Symbiotic with *Batissaviolaceacelebensis* Martens 1897 and Had the Potential as an Anti-MRSA

4. Discussion

Based on the results of bacterial isolation using nutrient agar media and observation of colony morphological characters, 5 (five) bacterial isolates that were symbiotic with *Batissa violacea*

celebensis Martens 1897 obtained, which are Bvc1, Bvc2, Bvc3, Bvc4, and Bvc5 isolates (Table 1). After being incubated for 24 hours, colonies of 5 bacterial isolates on NA media were observed macroscopically. Characteristics observed included colony shape, edge, elevation, color, and structure in the colony. Bvc1, Bvc3, and Bvc5 showed circular colonies with entire edges, or smooth. Bvc2 and Bvc4 showed irregular colonies with undulate edges, or wavy. All isolates showed low convex elevations that resembled water droplets, beige-colored with transparent inner structure.

In Table 1, we can see that the colony morphology of the five isolates showed nearly identical characteristics. In general, colony morphological characteristics could be used as a basis for identification at the genus level, but only a basis in differentiating between isolates[16]. Bacterial isolates can be said to be different if they have different colony characteristics[17]

Purified bacterial isolates were tested for their activities against MRSA pathogens using the paper disc method. The use of a single disc for each antibiotic with good standardization could determine whether bacteria are sensitive or resistant by comparing the standard resistance zones of the same drug [18]. The results of the qualitative inhibition ability test of bacterial isolates that were symbiotic with *Batissa violacea celebensis* Martens 1897 are shown in Table 2.

Based on the qualitative screening test, the five bacterial isolates exhibited different inhibitory ability against MRSA pathogenic bacteria. As shown in the results of the potential test in figure 2, there were two bacterial isolates, Bvc1, and Bvc3, that showed more potent inhibitory ability than the positive control. According to [19], different degrees of activity shown by each bacterial isolate may be caused by differences in secondary metabolites produced by each isolate. The positive control used was penicillin 1 g diluted with 9 ml ddH₂O.

The inhibitory ability test conducted obtained two potential isolates as an anti-MRSA, namely Bvc1 and Bvc2. The activity of both isolates was more potent than control penicillin. The inhibitory ability of both isolates was in the sensitive category, in line with the Clinical and Laboratory Standards Institute (CLSI) stating that their inhibitory zones could be considered resistant (≤ 13 mm), intermediate (14-16 mm), sensitive (≥ 17 mm). Penicillin was also resistant (≤ 13 mm), intermediate (14-16 mm), sensitive (≥ 17 mm). The formation of inhibitory zone zones in BV1 and BV3 was a result of the antibacterial activities of resultant symbiont bacteria isolates. According to Yuliana (2014), symbiont bacteria can produce similar bioactive compounds as the host. In [7] study, *Batissa violacea celebensis* Martens 1897 contained bioactive compounds such as alkaloid, steroid, peptide, saponin, flavonoid, and phenol.

Studies on natural materials from the sea or rivers have also been carried out by several researchers, one of which was by [12] who used the symbiont bacteria in *Holothuriascabra* as anti-MRSA. The results of the study of 9 bacterial isolates found 6 isolates included in the sensitive category.

After the sensitive bacterial isolates have been identified, the process continued to the gram staining stage. Gram staining was performed to group the bacteria into 2, Gram-positive and Gram-negative bacteria. Based on Figure 2, it can be seen that the two isolates are Gram-positive with bacteria in the form of bacilli. Gram-positive bacteria have a cell wall structure with a thick peptidoglycan content, thus unaffected during alcohol decolorization. Gram-positive bacteria will retain the purple color of crystal violet that when observed under a microscope it will show a purple color[19]. So it can be concluded that both Bvc1 and Bvc3 isolates are classified as specific in inhibiting Gram-positive bacteria, this shows the potential for antibiotics that have specifications in inhibiting MRSA pathogenic target bacteria as shown by the two isolates to have high activity compared to penicillin in inhibiting the growth of MRSA bacteria.

5. Conclusion

Based on the results of the research, it is concluded that two bacterial isolates obtained from *Batissa violacea celebensis* Martens 1897 have the potential to be anti-MRSA.

6. Conflict of Interest:

The authors declare no conflict of interest.

Acknowledgement

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