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Potentials of Omega-3 Rich Microalgae from Kulonprogo Mangrove Forest Yogyakarta for Nutraceuticals and Pharmaceuticals Products

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Abstract. The Covid-19 pandemic has increased human needs for nutrition including functional food and nutraceutical products in line with the need to boost immunity to resist viral infection, maintain a healthy life, and limit fatalities. In this regard, the choice to use functional food and nutraceuticals seems to be a promising panacea. This paper aims to present the potential of the microalgae species Aurantiochytrium, which is commonly found in the mangrove ecosystem. As a maritime country with the largest mangrove forest worldwide, Indonesia is known having abundant mangrove's biodiversity. One of the vital biodiversity in a mangrove ecosystem is microalgae, which have been intensively studied due to promising beneficial human iffe products. The high DHA content in microalgae has the potential to develop nutraceutical and pharmaceutical products. However, it is unfortunate that the potential of this microalga from Indonesian mangroves is rarely observed. Therefore, this study will share blodiscovery experience in isolating microalgae from the Indonesian mangrove forest. The focus of this study shall isolate microalgae from samples collected from Kulonprogro's Mangrove forest. The isolation results will then be identified to find out the morphological characteristics, and the found microalgae will be analyzed further in terms of valuable compounds contained in the selected strains. It is expected to trigger a further intensive exploration of microalgae in other mangrove forests and will grow interested in biodiversity research of Indonesian mangrove forests. The isolation of DHA producing microalgae has been successfully done using direct planting methods. The results indicate that the investigating microalgae can be cultured in the media prepared in this experiment.

Bioprocess Potentials of *Aurantiochytrium* Microalgae from Kulonprogo Mangrove Forest Yogyakarta, Indonesia

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Aurantiochytrium sp. is a marine protist that is highly ecologically relevant in the mangrove environment. This species of microalgae has long been known to produce high concentration of Docohexanoic acid (DHA, omega-3, 22:6n-3). DHA has a beneficial role for maintenance of normal brain function, visual acuity as well as prevention of cardiovascular and neurological diseases in humans. Although Indonesia is known to have the largest mangrove forest in the world, unfortunately, research on the isolation, potential observation and use of Aurantiockytrium microalgae has been still rarely published. Therefore, this research aims to promote the importance of research field using microalgae. Aurantiockytrium from Indonesian strain for future biochemical production processes. In this work, the samples of targeting microalgae were isolated from Kulonprogo mangrove, Yogyakarta using the direct plating method. The isolation medium contained 50% (v/v) seawater, 1% (w/v) agar, 0.5% (w/v) glucose, 0.1% (w/v) peptone, and 0.1% (w/v) yeast extract, without the addition of streptomycin. and penicillin. Morphological characteristics were observed. The observation results confirmed the availability of Aurantiochytrium isolates in prepared medium. Therefore, future isolation of *Aurantiockytrium* microalgae from other Indonesian mangrove forests can use the direct planting methods presented in this paper. In addition, this paper presented future important applicability of Indonesian Aurantiochytrium strains for uses in strategic industrial sectors, including pharmaceuticals, nutraceuticals and biofuel companies.

Keywords: Aurantiochytrium, docosahexaenoic acid, mangrove, microalgae, omega-3.

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INTRODUCTION

Conventional of source docosahexaenoic acid (DHA omega-3 22:6 n-3) for human nutrition, nutraceuticals and pharmaceutical products is marine fatty fish such as salmon and mackerel [1]. However, omega-3 extraction from fish oil arises some threatening problems such as contamination. heavy metals environmental issues related to illegal fishing and overfishing of the oceans. In addition, fish oil is not suitable for vegetarians and the odor makes it unattractive.

Therefore, recent trend of omega-3 sources shifts into omega-3 producing microorganisms [2]. Among the alternatives of marine microorganisms, Aurantiochytrium species is the most promising candidate for producing omega-3 PUFAs [3], [4]. This microalgae species has been chosen as the raw material to produce DHA-containing oil by Martek Biosciences Corporation (now part of the Royal DSM (Dutch State Mines) [2]. The superiorities of Aurantiochytrium species, among others, are high oil contents of 50-77% (dry weight), high DHA productivity with more than 30% DHA within the total oil produced and shorter cultivation time [5]. A comparison shows that Aurantiochytrium strains can

produce high amount of DHA and biomass in a relatively short cultivation time (Table 1). In addition, tabel 1 also shows that *Aurantiochytrium* sp. can achieve DHA production of 5,5 g/L until 20,5 g/L and the productivity rate of those strains is more than 100 g/L / day within 4 days.

Aurantiochytrium sp. is ubiquitous in marine and estuarine environments, in both tropical and sub-tropical areas and is also reported to be associated with mangrove habitats. Followings description are taxonomy of Aurantiochytrium microalgae [6]:

Domain : *Eukaryota* Kingdom : *Chromophyta* Phylum : *Heterokonta*

Family : Throustochytrioceae Order : Throustochytriales Genera : Aurontiochytrium

In addition to the potentials of high lipid production, *Aurantiochytrium* microalgae has also potentials usage in producing astaxanthin [7]–[9]. Recent potential of Aurantiochytrium is to fulfill a great demand by industry on squalene supply for adjuvant vaccine covid-19 [10], [11]. It is obvious that further study concerning of Aurantiochytrium microalgae shall be of importance in the near future.

Table 1. Examples of Aurantiochytrium strains for DHA production.

Strain name	DHA content (gL-1)	Cultivatio n time (h)	DHA productivity (mg/Lh-1)	Reference
Aurantiochytrium limacinum SR21	13.3	96	138	[12]
Aurantiochytrium mangrovei Sk-02	5.5	48	115	[13]
Schizochytrium sp. G13/2S	6.1	49	125	[14]
Aurantiochytrium sp. HX-308	17.7	160	111	[15]
Aurantiochytrium sp. SD116	17.4	110	158.4	[16]
Aurantiochytrium sp. BUCHHT 093	20.7	96	215.6	[17]

With more than 3 million hectares of mangrove forests, Indonesia is recognized as the richest mangrove country worldwide [18]. Fallen mangrove leaves might be colonized by a plentiful Aurantiochytrium microalgae, as Aurantiochytrium feed on the leaves resulting in their decay [19]. In addition, Aurantiochytrium microalgae inhabit on the above ground roots, free living on the mud and pelagic water column.

Despite the widest mangrove forest as abundant resources of Aurantiochytrium ecosystem. scientific publications concerning experimental work of isolation, characterization, screening and utilization Indonesian Aurantiochytrium microalgae have been rarely observed and published. Hutari et.al [20] isolated, characterized and did some cultivation work of Aurantiochytrium sp. from Lampung mangrove forest There are two review journals on the application of Aurantiochytrium microalgae [21], [22]. However, there is a lack of experimental studies in isolating, screening cultivating of Aurantiochytrium sp. from Indonesian mangrove.

Therefore, this paper presents our biodiscovery experience in isolating microalgae from a mangrove forest in Kulonprogo, Yogyakarta. This research is expected to promote the potentials of Aurantiochytrium microalgae as sources of valuable products. Following on this the modified method in the isolation of Aurantiochytrium microalgae shall be useful for further work in this field.

RESEARCH METHODOLOGY

Sampling Location

Targeting microalgae are isolated from a mangrove area in Kulonprogo, Yogyakarta. Isolation work was done on July 1st, 2019. The chosen mangrove areas are located close to the Hindia Ocean. The samples were cultured in media in the Laboratory of Pharmacy, Ahmad Dahlan University. Their morphological characteristics will be investigated afterward.

Bolation of Microalgae

Isolation of microalgae was carried out with direct plating method [23]. Fallen mangrove leaves of yellow and brownblack color were collected from the selected mangrove area. The illustration of isolation work was visualized in our education video [24]. Each leaf was cut into pieces of app. 1 x 1 cm size and placed directly on By+ agar medium and then incubated at 23°C for two days. The medium contained 50% (v/v) seawater, 1% (w/v) agar, 0.5% (w/v) glucose, 0.1% (w/v) peptone, and 0.1% (w/v) yeast extract. The addition of streptomycin and penicillin is excluded in this preliminary research.

After incubation, colonies were picked and transferred into 50 mL flasks containing 20 mL liquid By+ broth medium (without agar) and incubated for 3 days at 23°C. Ten microliters of these liquid cultures were spread on By+ agar medium and incubated for 3 days. Streak plate technique was applied for obtaining pure isolates of each colony without contaminants.

Morphological Characterization

Morphological characterization was carried out during isolation of pure isolates to distinguish *Aurantiochytrium* colonies from other microorganisms. The selected isolates were observed under a light microscope and the figures were captured by digital camera.

RESULTS AND DISCUSSION

Figure 1 represents an insight of the recent research project. Figure 1A shows the sampling work located in a mangrove area of Kulonprogo, Yogyakarta, whereas Figure 1B shows some isolates inoculated from the collected samples. Different colonies were identified by their morphology as the microbes grew. Spherical, uneven, and slimy producing colonies were taken in cultivating media, as shown in Figure 1C, for further analysis to obtain pure colonies.

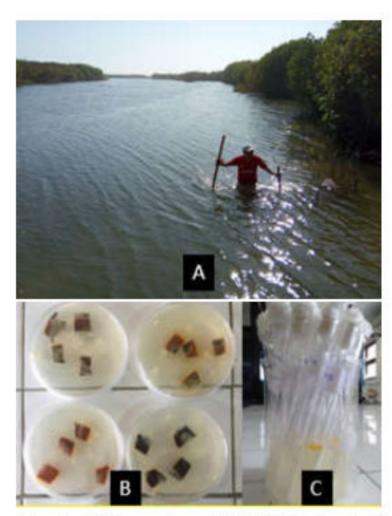
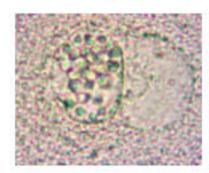


Fig. 1: An overview of isolation steps of targeting microalgae from a mangrove forest in Kulonprogo, Yogyakarta, Indonesia. A. situation of the mangrove area, B. isolates inoculated from the collected samples, and C. colonies were taken in agar media

Figure 3 depicted the micrograph of isolated microalgae from Kulonprogo mangrove forest. The figure shows typical

morphological characteristics of Aurantiocythrium's cell as compared with Aurantiochytium LR 52 [21].



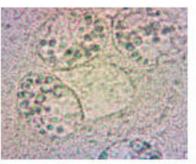


Fig. 3: Micrograph of discovered microalgae from Kulonprogo mangrove forest, Yogyakarta, Indonesia.

DISCUSSION

The work presented in this paper that observed Aurantiochytrium microalgae has been successfully isolated fallen of Kulonprogo from leaves mangrove forest. It is suggested that direct plating method presented in this paper shall be suitable for experimental work in isolating Aurantiochytrium sp. from other Indonesian mangrove forests. This method was also applicable with the results obtained from previous results [20], [25]-[27].

The genus Aurantiochytrium is attributed to a DHA content of about 80% of the polyunsaturated fatty acids [28]–[30]. For example, fatty acid profile of Aurantiochytrium LR52 reveals a simple fatty acid profile with DHA content of more than 80% of PUFA and other fatty acids such as eicosapentaenoic acid (EPA; 20:5, n-3) and arachidonic acid (ARA; 20:4 n-6) [31].

As mentioned previously, the DHA from Aurantiochytrium microalgae becomes an emerging important sustainable resource to replace fish oil as a sole omega-3 resource. Recent industrial application also convinces that strains of Aurantiochytrium species are the most productive microbial omega-3 producers [32].

Other main interest of the usage of Aurantiochytrium microalgae is in the field nutritional, pharmaceutical biochemical products (e.g., proximate nutrients, amino acids, and fatty acid composition). Among others, fatty acid palmitic prcfiles cf acid (C16:0),pentadecylic acid (C15:0), heptadecanoic acid (C17:0 n-6) has been of interest to be observed in previous publications [33]-[35].

With the continuous emerging observation of promising benefits from Aurantiocytrium as an alternative omega-3 resource, however, there have been so far very few biodiscovery and isolation work of valuable strains of Aurantiocytrium from Indonesia, as the country with the world's largest mangrove area. Therefore, this describing Aurantiochytrium research isolation work from one selected mangrove areas in Indonesia is relevant and important for future research field of biochemicals production.

CONCLUSIONS

Experimental work in this paper shows that the isolation of Aurantiochytrium microalgae has been successfully done using direct plating method. The results indicate that the investigating microalgae can be cultured in the media prepared in this experiment. Further work in in this field shall explore characterization of indigenous Indonesian Aurantiochytrium in terms of their fatty-acid profiles and biochemical compositions. In addition, cultivation shall be conducted to analyze growth rate and high added value components from isolated Aurantiochytrium species. Following this successfully isolation work, it is expected that this result can contribute to further thorough observation in the field of producing valuable nutritional compounds from Aurantiochytrium microalgae isolated in other Indonesian mangrove forests.

Further study is required to demonstrate the important applicability of Indonesian Aurantiochytrium strains in strategic industrial sectors, including pharmaceuticals, nutraceuticals and biofuel companies

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CERTIFICATE





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