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Comparative Study on the Extraction of Crude Fucoïdan from Brown Seaweed Using Conventional, Microwave and Ultrasound-Assisted Methods

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ABSTRACT

Brown seaweeds are rich in fucoidan, a sulfated polysaccharide with antimicrobial, immunomodulatory, antioxidant, and anticancer properties. The brown algae *Sargassum* sp. has not been thoroughly investigated for fucoidan extraction using various techniques and evaluations of their effects on extraction yield and its structural properties. The purpose of this study was to compare the structural characteristics and extraction yield of fucoidan from *Sargassum* sp. using conventional, microwave, and ultrasonic-assisted extraction methods. The results showed that a slightly higher yield was obtained by using the ultrasonic-assisted extraction method (2.772%) followed by the microwave-assisted extraction method (2.494%) and conventional extraction method (2.399%). However, the IC₅₀ values for antioxidants were found to be lower (less value is preferable) for crude fucoidan obtained by microwave-assisted extraction method (175 µg/ml) than for conventional (195 µg/ml) and ultrasound-assisted extraction methods (230 µg/ml). The crude fucoidan obtained from the three different extraction methods showed moderate antioxidant strength

Contribution to Sustainable Development Goals (SDGs)

SDG 3: Good Health and Well-being

SDG 9: Industry, Innovation, and Infrastructure

SDG 12: Responsible Consumption and Production

DG 14: Life Below Water

1. INTRODUCTION

1.1. Research Background

Brown seaweed's cell walls contain a complex sulfated polysaccharide called fucoidan. Usually, hot water or acid is used as an extractant to extract this compound. However, important factors that can significantly affect the yield of fucoidan include the solid-to-solvent ratio, extraction duration, pH, and temperature [1]. Conventional extraction methods have several

drawbacks, including longer operating durations, higher solvent and power consumptions, lower yields, and lower selectivity. Eco-friendly substitutes that employ high pressure, ultrasound, microwave, and enzymes to extract fucoidan from different algal sources have been developed to overcome these problems [2].

1.2. Literature Review

Based on the color of their pigments, seaweeds are divided into three taxonomic groups, namely green (*Chlorophyceae*), red (*Rhodophyceae*), and brown (*Phaeophyceae*) [1]. Fucoidan, an anionic water-soluble sulfated polysaccharide especially



prevalent in L-fucose, is found in brown seaweed [3]. Numerous beneficial biological qualities are known to be exhibited by this compound, which was isolated from different species of *Sargassum*. For instance, research has demonstrated that fucoidan has antibacterial, anticancer, antilipogenic, antithrombin, antiangiogenic, antioxidant, and antidiabetic properties that are associated with its chemical structure [1]. Furthermore, because of its emulsifying properties, fucoidan is considered as valuable ingredient in the food industry [4].

Research on fucoidan extraction from *Sargassum siliquosum*, *Sargassum mcclurei*, *Sargassum wightii*, and *Sargassum muticum* has demonstrated the potential of ultrasound-assisted extraction as a faster and more effective substitute for other methods [5-8]. These investigations showed that fucoidan yield can be considerably increased by ultrasound-assisted extraction, while extraction time can be decreased. On the other hand, another study on fucoidan extraction from *Ascophyllum nodosum* [9] and *Fucus vesiculosus* [10] found that when compared to the conventional extraction method, microwave-assisted extraction method resulted in a significantly shorter amount of extraction time.

1.3. Research Objective

The aim of this research is to compare the effect of different extraction methods namely conventional, microwave, and ultrasound-assisted methods on the yield of extraction of fucoidan from brown seaweed *Sargassum sp.* The chemical structure of crude fucoidan extract and its antioxidant activities were also investigated.

2. MATERIALS AND METHODS

2.1. Materials

The dried brown seaweed *Sargassum sp* was obtained from Sepanjang Beach, Gunung Kidul, Special Region of Yogyakarta, Indonesia. All chemicals used in extraction such as ethanol, calcium chloride, and demineralized water were purchased from a local chemical store in Yogyakarta.

2.2. Depigmentation of brown seaweed

The dried brown seaweed was grinded using a grinder to obtain fine powders. The powdered materials were passed through a 30-mesh sieve, which allows particles smaller than 500 μm to pass through. Dried *Sargassum sp* was depigmented to remove the pigments from the seaweed sample. The procedure involved treating 100 g of seaweed powder with 1 liter of ethanol for 24 hours at room temperature. The depigmented seaweed residue was filtered through a paper filter, dried, and then stored in airtight containers at 25°C until further use.

2.3. Conventional extraction method

Depigmented seaweed powder was mixed with deionized water at a ratio of 1:20 and heated to 65°C, for 60 minutes at 130 rpm using a water bath shaker (B-One SWB-30 Shaker Water Bath). Following extraction, the resulting mixture was filtered using filter paper to separate the residue from the supernatant. The residue was then allowed to dry in the shade. To precipitate the calcium alginate, the supernatant was treated with 1% calcium chloride and then kept in cold storage overnight. Centrifugation was used for 15 minutes at 5,000 rpm to remove the resultant

precipitate. Crude fucoidan was precipitated by mixing the remaining supernatant with twice as much ethanol and incubated in a cold environment. The precipitated fucoidan could be separated from the supernatant by the centrifugation process that followed. The dried fucoidan's weight was calculated after freeze dried at -55 °C using Freeze Dryer (VirTis Benchtop K).

2.4. Microwave assisted extraction method

Fucoidan was extracted from depigmented seaweed powder using a microwave reactor with power of 300 watts for 15 minutes at 30°C and a solid solvent ratio of 1:20. Following filtration, the seaweed residue was removed from the extracts. The process of gradually removing alginate and recovering fucoidan was performed as outlined in Section 2.3.

2.5. Ultrasound-assisted extraction method

Fucoidan was extracted from depigmented seaweed powder by using an ultrasonic processor (Ultrasonic Processor FS-300N) with a frequency of 40 kHz and a power of 300 W, for 15 minutes and a solid solvent ratio of 1:20. Following filtration, the seaweed residue was removed from the extracts. Alginate was gradually removed, and fucoidan was then eventually recovered following the procedure as outlined in Section 2.3.

2.6. Fucoidan yield

The fucoidan yield % was determined using the following formula [1].

$$\text{Fucoidan yield \% (w/w)} = \frac{\text{Weight of dried fucoidan (g)}}{\text{Weight of depigmented algal powder (g)}} \times 100$$

2.7. Fucoidan characterization

To identify the extracted fucoidan's functional groups, Fourier transform infrared (FTIR) spectroscopy was carried out using a PerkinElmer FTIR. The range of the FTIR spectral scans was 4000 cm^{-1} to 400 cm^{-1} .

2.8. Analysis of antioxidant activity

Antioxidant activity was assessed using the 2,2-Diphenyl-1-picrylhydrazyl (DPPH) free radical inhibition assay. Several concentrations of fucoidan (0, 60, 120, 180, 240, and 300 $\mu\text{g/mL}$) were prepared for the sample solution. The absorbance was measured at a wavelength of 517 nm. The following formula was used to determine the inhibition percentage.

$$\% \text{inhibition} = \frac{(A_{\text{blank}} - A_{\text{sample}})}{A_{\text{blank}}} \times 100\%$$

and then the linear plot of %inhibition versus concentration was analyzed using the following equation.

$$y = a + bx$$

where x is the concentration of the measured substance and y is the % inhibition. Meanwhile, the IC_{50} value was determined as the x value of this equation when y was equal to 50%.

3. RESULT AND DISCUSSION

3.1. Fucoidan yield

The cell wall of brown seaweed is primarily composed of sulfated polysaccharides containing fucose and polyanionic polysaccharides called alginates [11]. About 25–30% of the dry weight of brown seaweeds is made up of fucoidan, a sulfated polysaccharide [1].

The yields of fucoidan extracted using conventional, microwave, and ultrasound-assisted extraction methods are presented in Table 1. The results from Table 1 showed that using microwave and ultrasound could increase the yield of fucoidan. The highest yield in this case was obtained by using ultrasound-assisted method. A similar finding was also obtained by Hanjabam et al. [7] for the extraction of crude fucoidan from *Sargassum wightii* using conventional and ultra-sonication extraction methods. They concluded that the ultra-sonication extraction method gave higher yield than the hot water extraction method. Similar conclusion was also reported by Antonisamy et al. [1] for the extraction of fucoidan from *Sargassum ilicifolium*. They reported that the ultrasonic bath method produced a 2.5-fold higher yield of fucoidan than the conventional method. This is because ultrasonic waves have a cavitation effect that improves cell wall lysis and solvent penetration by reducing mass transfer resistance and increasing polysaccharide release [1].

Ref. [12] reported the published data of fucoidan content from brown macroalgae of 102 different algae species. The crude fucoidan content from *Sargassum* genera from [12] with some additional data is summarized in Table 2. From Table 2, it can be concluded that the yield of fucoidan was varied depended on the algae species and the method of extraction. The result of the current study was similar to that of Sinurat, E. and Kusumawati, R. [42] for the extraction of fucoidan from brown seaweed *Sargassum binderi* Sonder using conventional method by extraction with calcium chloride at 85 °C for 4 hours. At higher temperature and longer extraction duration, the yield of fucoidan was 2.57% [42], whereas in the current study, the yield of 2.494 – 2.772% could be obtained for the extraction duration of only 15 minutes with the assistance of microwave and ultrasound. Thus, it is clear that extraction by adding microwave as well as ultrasound could produce fucoidan with higher yield at a much shorter extraction duration. To obtain a higher yield of fucoidan, more studies should be conducted for the extraction with the assistance of microwave and ultrasound but at higher temperatures (i.e. 85 °C) and longer duration (i.e. up to 1 hour).

Table 1. Fucoidan extraction yields obtained using the conventional, microwave and ultrasound-assisted extraction methods

Parameter	Conventional	Microwave	Ultrasound
Extraction yield (%)	2.399±0.248	2.494±0.004	2.772±0.154

3.2. Functional group analysis

The FTIR analysis for all the samples is shown in Fig. 1. The broad bands at 3350–3325 cm⁻¹ for all the samples is explained by O-H stretching vibrations, which are present in all polysaccharides [1]. The bands at 1029–1027 cm⁻¹ show the S=O stretching, O-H bending (1412 cm⁻¹), and the C=C stretching (1595–1594 cm⁻¹). The spectra at 1412 cm⁻¹ and 1595–1594 cm⁻¹, are ascribed to symmetric and asymmetric stretching vibrations of carboxylate groups, respectively, which may indicate the presence of uronic acid [46]. The area of polysaccharides of about 1029–1027 cm⁻¹ shows the glucose and galactose that are typical of a pyranose ring [46]. The characteristic of fucoidan seen from the FTIR spectra is proven by the presence of absorption at 1217–1215 cm⁻¹ which indicates sulfated polysaccharides [42].

Table 2. Summary of crude fucoidan content reported for several algal species from *Sargassum* genera

Species	Location	Crude content (%DW)	Ref.
<i>Sargassum binderi</i>	Malaysia	6.2	[13]
<i>Sargassum duplicatum</i>	Vietnam	0.32	[14]
<i>Sargassum feldmannii</i>	Vietnam	0.46	[14]
<i>Sargassum filipendula</i>	Indonesia	4.5 – 6.1	[15]
<i>Sargassum fusiforme</i>	China	3.94–11.24	[16]
<i>Sargassum glaucescens</i>	China	4.20	[17]
<i>Sargassum hemiphylum</i>	China	2.72 ± 0.18	[18]
<i>Sargassum hemiphylum</i>	China	4.69 ± 1.05	[5]
<i>Sargassum henslowianum</i>	China	6.25	[19]
<i>Sargassum horneri</i>	China	4.80	[17]
<i>Sargassum mcclurei</i>	Vietnam	2.7	[20]
<i>Sargassum pallidum</i>	Japan	1.86 – 6.31	[21]
<i>Sargassum polycystum</i>	China	3.41 ± 0.77	[5]
<i>Sargassum ringgoldianum</i>	China	0.95 ± 0.02	[22]
<i>Sargassum siliquosum</i>	China	5.08 ± 1.17	[5]
<i>Sargassum stenophyllum</i>	Brazil	0.4	[23]
<i>Sargassum tenerrimum</i>	India	3.1	[24]
<i>Sargassum vachellianum</i>	China	5.5 ± 0.25	[25]
<i>Sargassum ilicifolium</i>	India	2.68 – 6.74	[1]
<i>Sargassum sp</i>	Egypt	5.99–36.79	[4]
<i>Sargassum wightii</i>	India	10.59-14.61	[7]
<i>Sargassum cinereum</i>	Egypt	21.79±4.99	[26]
<i>Sargassum thunbergii</i>	China	2.85 ± 0.08	[27]
<i>Sargassum polycystum</i>	Bangladesh	4.1 – 6.1	[28]
<i>Sargassum horneri</i>	China	2.91±0.17	[29]
<i>Sargassum Zhangii</i>	China	2.85±0.35	[30]
<i>Sargassum hemiphylum</i>	Taiwan	4.02 ± 0.27	[31]
<i>Sargassum fusiforme</i>	China	1.56	[32]
<i>Sargassum polycystum</i>	Malaysia	1.16 ± 0.07	[33]
<i>Sargassum wightii</i>	India	0.7–1	[34]
<i>Sargassum cinereum</i>	India	9.4 ± 1.9	[35]
<i>Sargassum fluitans</i>	Mexico	0.84	[36]
Borgesen			
<i>Sargassum swartzii</i>	India	5.96	[37]
<i>Sargassum fusiforme</i>	China	7.3	[38]
<i>Sargassum polycystum</i>	India	4.51±0.24	[39]
<i>Sargassum fusiforme</i>	China	6.02	[40]
<i>Sargassum latifolium</i>	Egypt	10.1±0.2	[41]
<i>Sargassum binderi</i>	Indonesia	2.57 – 6.00	[42]
Sonder			
<i>Sargassum sp</i>	Indonesia	1.04 – 4.1	[43]
<i>Sargassum binderi</i>	Indonesia	4.02±0.27	[44]
Sonder			
<i>Sargassum filipendula</i>	Indonesia	5.8 – 8.3	[45]

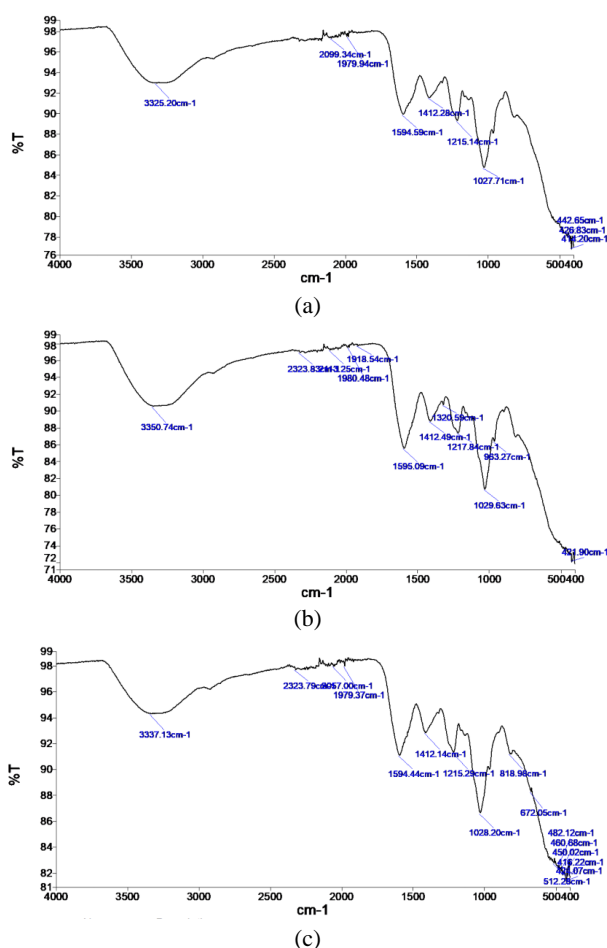


Fig. 1. FTIR spectra of fucoidan extracted from *Sargassum sp* using extraction method: (a) conventional, (b) microwave, (c) ultrasound

3.3. Antioxidant activity

DPPH test (2,2-diphenyl-1-picrylhydrazyl) method was used to obtain the antioxidant activity. IC₅₀ (µg/ml) is a measure of antioxidant capacity that is used to determine antioxidant activity. The amount of test compound concentration that can inhibit free radicals by up to 50% is known as the IC₅₀ value. The free radical reduction activity increases with a smaller IC₅₀ value. The IC₅₀ value category is very strong if the IC₅₀ value <50 µg/ml, strong if the IC₅₀ value is between 50 and 100 µg/ml, moderate if the IC₅₀ value is between 100 and 250 µg/ml, and weak if the IC₅₀ value is between 250 and 500 µg/ml [47].

The IC₅₀ values for crude fucoidan obtained from the current study are presented in Table 3. It can be seen from Table 3 that the level of antioxidant strength for all samples were moderate. Among the three extraction methods used, fucoidan extracted using microwave assisted method resulted in the highest antioxidant activity (the lowest value is preferable). Other studies reported the lower values of IC₅₀ suggesting the strong antioxidant activity for fucoidan extracted from different seaweed such as *Sargassum cinereum* with IC₅₀ value of 75 µg/ml [35], and 69 – 99 µg/ml for *Sargassum fusiforme* [16].

Table 3. Average IC₅₀ values of fucoidan obtained using the conventional, microwave, and ultrasound-assisted extraction methods

Parameter	Conventional	Microwave	Ultrasound
Average IC ₅₀ values (µg/ml)	195.58±5.40 (moderate)	175.62±2.80 (moderate)	230.21±21.29 (moderate)

4. CONCLUSION

This study offers a comparative assessment of the effects of conventional, ultrasound, and microwave-assisted extraction methods on the yield and functional characteristics of the crude fucoidan that was extracted from brown seaweed *Sargassum sp*. The yield of fucoidan extracted using an ultrasonic probe was slightly higher than those of conventional and microwave-assisted extracted fucoidan. The results also showed that the fucoidan obtained by all three methods exhibited moderate antioxidant activity. The IC₅₀ values of fucoidan for all samples were between 175–230 µg/ml. The results of this investigation indicate that crude fucoidan derived from *Sargassum sp*. may prove to be a useful resource for the food industry as well as for medicinal purposes.

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