

## Note

# Mineral Contents of Indonesian Seaweeds and Mineral Solubility Affected by Basic Cooking

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**This experiment was carried out to study the mineral contents and profiles of several Indonesian green, brown, and red seaweeds, and to evaluate the solubilities of Ca and Mg as affected by boiling in different solutions (water, 1% sodium chloride and 0.5% acetic acid). The macromineral profiles were dominated by Ca, K, Na and Mg. The trace minerals Cu, Fe and Zn were found at low concentrations. Boiling in water and 0.5% acetic acid significantly increased the solubilities of Ca and Mg ( $p < 0.05$ ), whereas boiling in 1% sodium chloride resulted in varying solubilities of Ca and Mg. Mainly soluble Ca was found in both low (MW < 10,000) and high (MW > 200,000) molecular weight fractions, whereas soluble Mg was found in the high (MW > 200,000) molecular weight fraction.**

Keywords: Ca, boiling, Indonesia, Mg, mineral, seaweed, solubility, ultrafiltration

## Introduction

From foodstuff standpoint, seaweeds have a long history and even today, they are an important part of the diet of many Asian countries, particularly Japan, China and Korea. Among these three, Japan, where seaweeds (sea vegetables) have a considerable market value, is the most important seaweed consumer (Arasaki and Arasaki, 1983; Nisizawa *et al.*, 1987; Nisizawa, 2002).

Seaweeds are the richest source of minerals (Japan Society for Research of Food Composition, 1985; Ito and Hori, 1989; Ruperez, 2000) and dietary fibers (Suzuki *et al.*, 1993<sup>3</sup>; Suzuki *et al.*, 1996; Wong and Cheung, 2000; Santoso, Yoshie and Suzuki, 2002). Moreover, seaweeds contain fats and proteins, although at low concentrations (Wong and Cheung, 2001; Sanchez-Machado *et al.*, 2004).

The most common minerals found in seafood including seaweeds are iodine, magnesium, sodium, calcium, phosphorus, iron, potassium, copper and fluorine (Ensminger *et al.*, 1995). Minerals are very important for biochemical reactions in the body as cofactors of enzymes. Mineral deficiency can lead to severe health impairment.

Mineral concentration is only one function of their physiological worth, since they should also be bioavailable. Among functional food components, minerals play an essential role in living a healthy life. Bioavailability is a term used to describe the proportion of a nutrient in food that can be utilized for normal body functions (Watzke, 1998; O'Dell, 1984). For minerals to be bioavailable they should to be in a soluble form, however, not all

soluble minerals are bioavailable (Clydesdale, 1988). The bioavailability of minerals and trace elements in humans is best studied using an *in vivo* model; however, experiments using such a model are often complex, time-consuming and costly, and yield nonreproducible results. The first steps in the study of mineral bioavailability are to measure the concentrations of minerals in foodstuff, in the form in which they exist, as well as the solubility of these minerals.

There are few reports on the mineral content of seaweeds. Studies of minerals in seaweeds mainly focused on sub-tropical and temperate seaweeds such as Japanese seaweeds (Ito and Hori, 1989; Yoshie *et al.*, 1994; Yoshie *et al.*, 1999), Korean seaweeds (Yoshie *et al.*, 1993), and Spanish seaweeds (Ruperez, 2000). There is little information available on mineral compounds in tropical seaweeds, including those in Indonesia.

Since there is no information or data available yet on the mineral profile of Indonesian seaweeds, it is a fertile area for research. Compared to their availability, seaweeds still have a low worldwide consumption and are consumed only in local areas. Seaweed utilization at present is limited to a few genera such as *Gracilaria*, *Gellidium*, *Euclidean*, *Turbinaria*, and *Sargassum*. Almost all collected seaweeds are exported as raw materials for industry. They are unprocessed materials in dry form. Therefore the first objective of the research is to measure the mineral profiles and contents of several Indonesian seaweeds.

Almost all seaweeds are eaten cooked except for the ones used in salads. The most common method of cooking seaweeds is boiling, where sodium chloride or acetic

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acid is usually added as seasonings. Therefore, the second objective of this experiment is to evaluate the effects of boiling in different solutions (*i.e.*, water, sodium chloride and acetic acid) on the solubilities of minerals (*i.e.*, calcium and magnesium) in several Indonesian seaweeds.

## Materials and Methods

**Materials** Eight species of Indonesian seaweeds were used in this experiment: five green algae (*Caulerpa racemosa*, *Caulerpa sertularioides*, *Cladophoropsis vaucheriaeformis* and *Ulva reticulata*), three brown algae (*Padina australis*, *Sargassum polycystum* and *Turbinaria conoides*), and one red alga (*Kappaphycus alvarezii*), all collected from the Seribu Islands, Jakarta Province.

**Sample preparation** After removing sand, the seaweed samples were washed with clean seawater and transported to the laboratory under refrigeration. The samples were washed with tap water, wiped with paper towels, then minced with a food processor (MK-K75; Matsushita Electric Corp., Osaka, Japan), and stored at  $-20^{\circ}\text{C}$  until use.

**Proximate analysis and dietary fiber analysis** Moisture, ash and protein contents were determined according to the AOAC method (Association of Official Analytical Chemist, 1990). Dietary fiber content was determined according to an enzymatic-gravimetric method (Prosky *et al.*, 1988).

**Total minerals** Each sample (2 g wet weight) was weighed in a Kjeldahl flask. Twenty milliliters of concentrated nitric acid was added to each sample and the flask was left to stand overnight. Five milliliters of concentrated perchloric acid and 0.5 mL of concentrated sulfuric acid were added, and the flask was then heated until no white smoke was emitted. The samples were dissolved in 2% of hydrochloric acid and transferred into a volumetric flask, then analyzed using an atomic absorption spectrophotometer (Model AA-600, Shimadzu Co., Kyoto, Japan) with acetylene flame, a single-slot head, and a Pt-Rh corrosion resistant nebulizer for measuring the total magnesium (Mg), calcium (Ca), potassium (K) and sodium (Na) contents. Copper (Cu), zinc (Zn) and iron (Fe) contents were measured using an inductively coupled argon plasma emission spectrometer (ICP Nippon Jarrel-Ash ICAP-757V, Kyoto, Japan).

All the reagents used were of analytical grade and their solutions were prepared using double-distilled deionized water. All the glassware and plastic bottles used were dipped in 2% Contaminon-L (Wako Pure Chemical Industries, Ltd., Osaka, Japan) for at least 2 h, and then rinsed with double-distilled deionized water to remove contaminants.

**Soluble minerals** The samples (10 g) were blended in a tube with water or 1% sodium chloride or 0.5% acetic acid (40 mL) at 5,000–10,000 rpm for 2 min using a blender (Ultra-Turrax T-25; Janke and Kunkel, IKA-Labortechnik, GmbH Co., Staufen, Germany) to produce a water-soluble fraction (0-min-treated sample). The tubes were then boiled in a water bath (Eyela Uni Thermo Shaker

NTS-3000, Tokyo Rikakikai Co., Ltd., Japan) at  $100^{\circ}\text{C}$  for 20 min (20-min-treated sample). Next, the samples were centrifuged at 10,000  $\times g$ ,  $2^{\circ}\text{C}$  for 10 min, and filtered through an Advantec filter paper No. 101 (Toyo Roshi Kaisha, Ltd., Tokyo, Japan). Nonboiled samples were directly centrifuged and filtered. The mineral concentration of the filtrate was measured using an atomic absorption spectrophotometer (Model AA-600, Shimadzu Co., Kyoto, Japan) and calculated as percentage relative the total mineral content. The boiling conditions of  $100^{\circ}\text{C}$  for 20 min were chosen in accordance with the previous method of Suzuki *et al.* (2000). The solubility of each mineral was calculated using  $\text{Solubility (\%)} = (\text{Soluble mineral mg/g}) / (\text{Total mineral mg/g}) \times 100$ .

**Soluble minerals fractionated by molecular weight** The filtered supernatant (2.0 mL) was completely passed through Advantec ultra-filter units USY-10 or USY-200 (Toyo Roshi Co.Ltd., Tokyo, Japan) with MW cut-off 10,000 or 200,000, respectively to obtain low, mid and high molecular weight fractions of the soluble minerals. The concentration of minerals (Ca and Mg) in the low (MW < 10,000), mid (MW = 10,000–200,000), and high (MW > 200,000) molecular weight fractions were measured using an atomic absorption spectrophotometer (Model AA-600, Shimadzu Co., Kyoto, Japan), and the results were expressed as percentage of the soluble mineral fraction (MW < 10,000, MW = 10,000–200,000 and MW > 200,000) relative to the total soluble mineral content.

**Statistical analysis** All experiments were performed in 3–5 replicates. Results are expressed as mean  $\pm$  standard deviation ( $n=3-5$ ). ANOVA was used to calculate the significance of differences using  $p < 0.05$ .

## Results and Discussion

**Proximate composition and dietary fiber content** Table 1 shows the moisture, ash, fat, crude protein, and dietary fiber contents in several Indonesian seaweed samples. The samples had ash, protein and dietary fiber contents of 2.1–5.5 g/100 g wet weight, 0.7–3.1 g/100 g wet weight and 7.1–11.6 g/100 g wet weight, respectively. After calculation, the dietary fiber and ash contents per 100 g dry matter samples were shown to be high.

**Mineral composition** Table 2 shows the macromineral profiles and contents of the several Indonesian seaweed samples. High Na concentrations were found in *Ulva reticulata* (26.4 mg/g dry weight), *Caulerpa sertularioides* (25.7 mg/g dry weight), and *Cladophoropsis vaucheriaeformis* (23.9 mg/g dry weight), whereas high K concentrations were found in *Kappaphycus alvarezii* (87.1 mg/g dry weight), *Turbinaria conoides* (27.9 mg/g dry weight), and *Sargassum polycystum* (17.5 mg/g dry weight). The ranges of Ca and Mg contents were 2.8–28.3 mg/g and 2.9–21.5 mg/g dry weight, respectively. The highest and the lowest Ca concentrations were found in *Padina australis* and *Kappaphycus alvarezii*, respectively. *Ulva reticulata* contained the highest Mg content, and *Kappaphycus alvarezii* contained the lowest Mg content.

The macromineral profiles and contents of the Indonesian seaweed samples were very similar to those of Japa-

**Table 1.** Moisture, ash, fat, protein, and dietary fiber contents of several Indonesian seaweed samples<sup>a</sup>.

Seaweed Sample	Moisture	Ash	Fat	Protein	Dietary fiber
<u>Green algae</u>					
<i>Caulerpa racemosa</i>	88.8 ± 0.5 <sup>d</sup>	2.1 ± 0.2 <sup>a</sup>	0.5 ± 0.1 <sup>c</sup>	1.5 ± 0.2 <sup>c</sup>	7.3 ± 0.5 <sup>cd</sup>
<i>Caulerpa sertularioides</i>	82.4 ± 0.6 <sup>a</sup>	2.9 ± 0.2 <sup>b</sup>	2.3 ± 0.1 <sup>e</sup>	3.1 ± 0.2 <sup>e</sup>	10.9 ± 0.2 <sup>bc</sup>
<i>Cladophoropsis vaucheriaeformis</i>	84.9 ± 0.7 <sup>c</sup>	4.0 ± 0.1 <sup>d</sup>	0.5 ± 0.1 <sup>bc</sup>	2.5 ± 0.5 <sup>d</sup>	7.1 ± 0.7 <sup>cd</sup>
<i>Ulva reticulata</i>	83.9 ± 0.3 <sup>bc</sup>	2.9 ± 0.1 <sup>b</sup>	1.2 ± 0.2 <sup>e</sup>	1.2 ± 0.2 <sup>bc</sup>	10.6 ± 0.1 <sup>ab</sup>
<u>Brown algae</u>					
<i>Padina australis</i>	83.1 ± 1.0 <sup>ab</sup>	5.5 ± 0.4 <sup>c</sup>	0.8 ± 0.1 <sup>d</sup>	1.5 ± 0.1 <sup>c</sup>	9.6 ± 0.6 <sup>a</sup>
<i>Sargassum polycystum</i>	84.7 ± 0.4 <sup>c</sup>	3.8 ± 0.3 <sup>d</sup>	0.3 ± 0.1 <sup>b</sup>	0.9 ± 0.1 <sup>a</sup>	10.1 ± 0.1 <sup>cd</sup>
<i>Turbinaria conoides</i>	85.1 ± 0.3 <sup>c</sup>	2.5 ± 0.1 <sup>ab</sup>	0.8 ± 0.1 <sup>d</sup>	1.0 ± 0.1 <sup>ab</sup>	9.5 ± 0.5 <sup>c</sup>
<u>Red alga</u>					
<i>Kappaphycus alvarezii</i>	83.3 ± 0.7 <sup>ab</sup>	3.4 ± 0.1 <sup>c</sup>	0.2 ± 0.1 <sup>a</sup>	0.7 ± 0.1 <sup>a</sup>	11.6 ± 0.3 <sup>d</sup>

<sup>a</sup> Data are means of three determinations ± SD (g/100g wet weight). Means in columns followed by different superscript letters (a-f) are significantly different ( $p < 0.05$ ).

**Table 2.** Macromineral contents of several Indonesian seaweed samples<sup>a</sup>.

Seaweed Samples	Mg	Ca	K	Na
<u>Green algae</u>				
<i>Caulerpa racemosa</i>	3.8 ± 0.3 <sup>a</sup>	18.5 ± 5.3 <sup>c</sup>	3.2 ± 0.2 <sup>b</sup>	25.7 ± 1.2 <sup>c</sup>
<i>Caulerpa sertularioides</i>	3.7 ± 1.0 <sup>a</sup>	12.0 ± 4.4 <sup>b</sup>	0.3 ± 0.0 <sup>a</sup>	0.7 ± 0.4 <sup>a</sup>
<i>Cladophoropsis vaucheriaeformis</i>	7.1 ± 0.6 <sup>b</sup>	22.3 ± 3.3 <sup>d</sup>	9.9 ± 0.4 <sup>c</sup>	23.9 ± 1.2 <sup>c</sup>
<i>Ulva reticulata</i>	21.5 ± 2.8 <sup>c</sup>	17.9 ± 5.3 <sup>c</sup>	12.6 ± 0.3 <sup>c</sup>	26.4 ± 0.8 <sup>c</sup>
<u>Brown algae</u>				
<i>Padina australis</i>	4.0 ± 1.6 <sup>ab</sup>	28.3 ± 4.3 <sup>e</sup>	0.5 ± 0.2 <sup>a</sup>	1.0 ± 0.9 <sup>a</sup>
<i>Sargassum polycystum</i>	5.7 ± 0.7 <sup>ab</sup>	18.7 ± 1.4 <sup>c</sup>	17.5 ± 1.4 <sup>d</sup>	9.7 ± 1.4 <sup>b</sup>
<i>Turbinaria conoides</i>	5.7 ± 0.3 <sup>ab</sup>	14.8 ± 2.2 <sup>b</sup>	27.9 ± 1.1 <sup>e</sup>	11.5 ± 0.5 <sup>b</sup>
<u>Red alga</u>				
<i>Kappaphycus alvarezii</i>	2.9 ± 0.3 <sup>a</sup>	2.8 ± 0.3 <sup>a</sup>	87.1 ± 5.8 <sup>f</sup>	11.9 ± 2.5 <sup>c</sup>

<sup>a</sup> Data are means of three determinations ± SD (mg/g dry weight). Means in columns followed by different superscript letters (a-f) are significantly different ( $p < 0.05$ ).

nese and Spanish seaweeds, with Na, Ca, Mg and K being the major minerals. In the raw Japanese brown alga *Laminaria japonica* (ma-kombu in Japanese) the Na, Ca, Mg and K contents were 590, 75, 120 and 42 mg/100 g of edible portion (Resources Council, Science and Technology Agency, 1991). In other brown algae (*Fucus vesiculosus*, *Laminaria digitata*, and *Undaria pinnatifida*) and red algae (*Chondrus crispus* and *Porphyra tennera*) grown in Spain, the ranges of Na, Ca, Mg and K contents were 36.3–70.6, 3.9–10.1, 5.7–11.8, and 31.8–115.8 mg/g dry weight, respectively (Ruperez, 2002). Furthermore, the Ca con-

tent of *Gracillaria changii* grown in Malaysia was 651 mg/100 g wet weight (Norziah and Ching, 2000).

The trace-mineral profiles and contents of several Indonesian seaweed samples are presented at Table 3. The range of Cu contents was from 0.002 mg/g dry weight (*Sargassum polycystum*) to 0.251 mg/g dry weight (*Caulerpa sertularioides*). The ranges of Fe and Zn contents were 0.041–0.813 and 0.003–0.227 mg/g dry weight, respectively. The highest Fe content was found in *Caulerpa racemosa*, and the highest Zn content was found in *Cladophoropsis vaucheriaeformis*.

**Table 3.** Micromineral contents of several Indonesian seaweed samples<sup>a</sup>.

Seaweed Samples	Cu	Zn	Fe
<u>Green algae</u>			
<i>Caulerpa racemosa</i>	0.008 ± 0.003	0.010 ± 0.002	0.813 ± 0.237
<i>Caulerpa sertularioides</i>	0.251 ± 0.062	0.003 ± 0.000	0.041 ± 0.010
<i>Cladophoropsis vaucheriaeformis</i>	0.227 ± 0.003	0.227 ± 0.003	0.111 ± 0.056
<i>Ulva reticulata</i>	0.179 ± 0.001	0.017 ± 0.001	0.280 ± 0.060
<u>Brown algae</u>			
<i>Padina australis</i>	0.005 ± 0.001	0.013 ± 0.003	0.446 ± 0.016
<i>Sargassum polycystum</i>	0.002 ± 0.001	0.004 ± 0.001	0.277 ± 0.214
<i>Turbinaria conoides</i>	0.003 ± 0.001	0.006 ± 0.003	0.062 ± 0.017
<u>Red alga</u>			
<i>Kappaphycus alvarezii</i>	0.005 ± 0.000	0.018 ± 0.004	0.070 ± 0.033

<sup>a</sup> Data are means of three determinations ± SD (mg/g dry weight).

Compared with the Japanese seaweeds *Hizikia fusiformis* (hijiki in Japanese), *Porphyra yezoensis* (susabi-nori in Japanese) and *Enteromorpha intestinalis* (ao-nori in Japanese), the Indonesian seaweed samples had slightly lower Fe and Zn contents with ranges of 9.12–54.4 and 0.82–5.24 mg/100 g dry weight, respectively (Yoshie *et al.*, 1999). Furthermore, the Fe and Zn contents of brown algae (*Fucus vesiculosus*, *Laminaria digitata*, and *Undaria pinnatifida*) and red algae (*Chondrus crispus* and *Porphyra tennera*) grown in Spain were almost the same, with ranges of 0.033–0.103 and 0.017–0.071 mg/g dry weight, respectively (Ruperez, 2002). The red alga *Gracillaria changii* had an Fe content of 95.6 mg/100 g wet weight (Norziah and Ching, 2000).

The dried products of common Japanese seaweeds (*Hizikia fusiformis*, *Porphyra yezoensis*, and *Undaria pinnatifida*) also had low Cu contents (0.08–0.62 mg/100 g edible portion, equal to 0.0009–0.007 mg/g dry weight) (Resources Council, Science and Technology Agency, 2001), and the Spanish brown and red algae had a low Cu content less than 0.005 mg/g dry weight (Ruperez, 2002).

The wide range of mineral contents, not found in edible land plants, is associated with factors such as phylum, geographical origin, and seasonal, environmental and physiological variations of the seaweeds (Mabeau and Fluerence, 1993). Mineral content also depends on the type of seaweed processing (Nisizawa *et al.*, 1987; Yoshie *et al.*, 1994) and the mineralization methods used (Fluerence and Le Coeur, 1993).

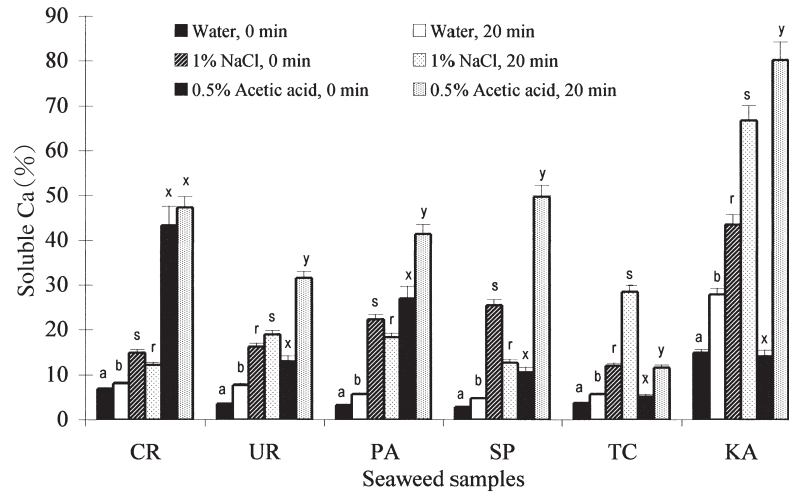
**Solubility of minerals** Six species of locally consumed Indonesian seaweeds were selected in order to evaluate the solubilities of Mg and Ca as affected by boiling in different solutions: two green algae (*Caulerpa racemosa* and *Ulva reticulata*), three brown algae (*Padina australis*,

*Sargassum polycystum* and *Turbinaria conoides*), and one red alga (*Kappaphycus alvarezii*).

The solubilities of Ca and Mg as affected by boiling in different solutions are shown in Figs. 1 and 2, respectively. Boiling in water or 0.5% acetic acid at 100°C for 20 min significantly increased the solubility of Ca in all seaweed samples, except for boiling *Caulerpa racemosa* in 0.5% acetic acid, which did not show a significant increase. Boiling in 1% sodium chloride significantly increased Ca solubility in *Ulva reticulata*, *Turbinaria conoides* and *Kappaphycus alvarezii*, whereas it significantly decreased Ca solubility in the other three seaweed samples. The percentages of soluble Ca in water after boiling were from 4.7% (*Sargassum polycystum*) to 27.9% (*Kappaphycus alvarezii*). After boiling in 0.5% acetic acid, the lowest and the highest Ca solubilities were found in *Turbinaria conoides* (11.6%) and *Kappaphycus alvarezii* (80.2%), respectively.

Mg solubility also significantly increased after boiling in water in all the species except *Kappaphycus alvarezii*. Boiling in 0.5% acetic acid, significantly increased the Mg solubility in *Ulva reticulata*, *Sargassum polycystum*, *Turbinaria conoides* and *Kappaphycus alvarezii*, whereas the increase was not significant in the other seaweed samples. Boiling in 1% sodium chloride significantly increased Mg solubility only in *Ulva reticulata* and *Turbinaria conoides*; it either had no effect or it decreased Mg solubility in the other seaweed samples.

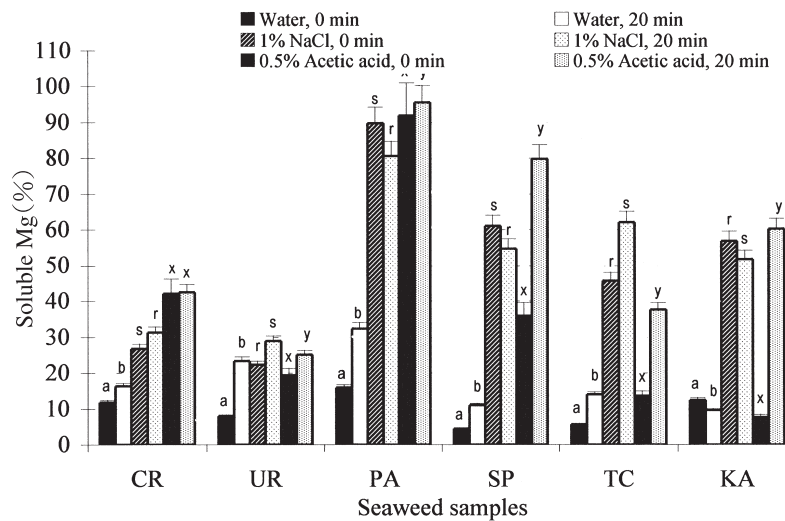
Suzuki *et al.* (2000) studied the effect of salt water heating on solubilities of minerals in shellfish. They reported that the percentage of soluble Mg sometimes increased after heating in water and 1% sodium chloride, whereas Ca solubility was not significantly affected by heating in either water or sodium chloride solutions.



**Fig. 1.** Percent solubility of calcium as affected by boiling in different solutions.

CR: *Caulerpa sertularioides*, UR: *Ulva reticulata*, PA: *Padina australis*, SP: *Sargassum polycystum*, TC: *Tubinaria conoides*, KA: *Kappaphycus alvarezii*.

Values in columns followed by the different superscript letters in the same solution <sup>a,b</sup> or <sup>r,s</sup> or <sup>x,y</sup> are significantly different ( $p < 0.05$ ).



**Fig. 2.** Percent solubility of magnesium as affected by boiling in different solutions.

CR: *Caulerpa sertularioides*, UR: *Ulva reticulata*, PA: *Padina australis*, SP: *Sargassum polycystum*, TC: *Tubinaria conoides*, KA: *Kappaphycus alvarezii*.

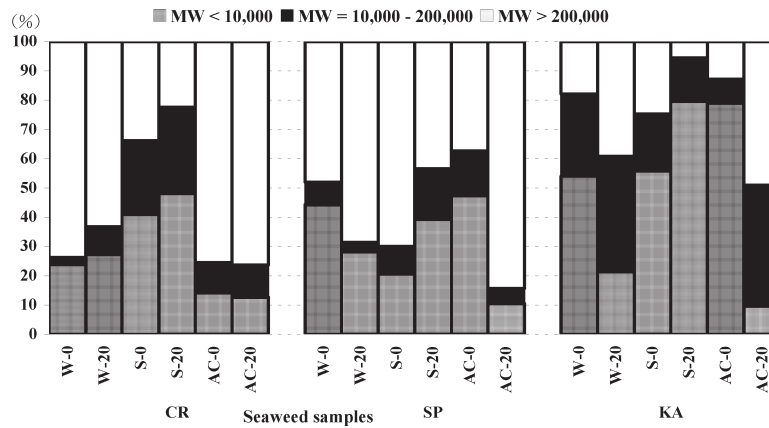
Values in columns followed by the different superscript letters in the same solution <sup>a,b</sup> or <sup>r,s</sup> or <sup>x,y</sup> are significantly different ( $p < 0.05$ ).

These results differ from our findings in Indonesian seaweeds, where the solubility of Ca significantly increases after boiling in water and 0.5% acetic acid. However, in case of soluble Mg, the results are very similar.

Minerals in food may change their chemical form during and/or after food processing, or interaction with other compounds. Thus their solubilities can increase or decrease depending on the type of processing method. Heating induces the protein denaturation, which may make minerals insoluble. However, since seaweeds have low protein content, the interaction between minerals and proteins may not occur.

Mineral solubility is also influenced by pH. The per-

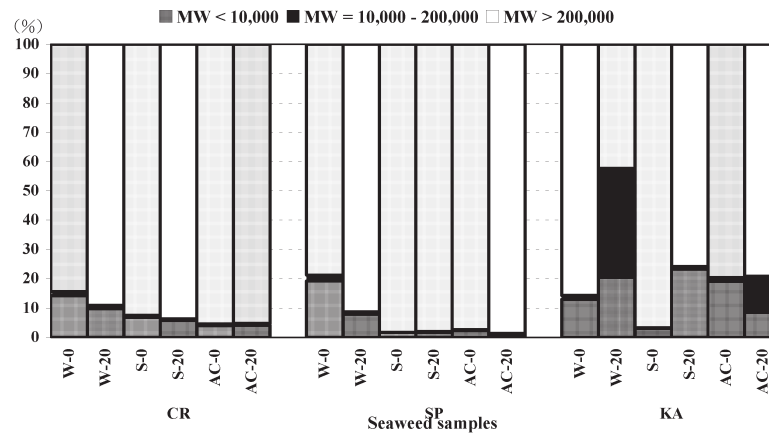
cent solubilities of Fe in three species of Japanese seaweeds *Porphyra yezoensis* (susabi-nori in Japanese), *Enteromorpha intestinalis* (ao-nori in Japanese) and *Hizikia fusiformis* (hijiki in Japanese) at pH 2 were higher than those at pH 6 (Yoshie *et al.*, 1999). The solubilities of Fe in cod, scallop and prawn decreased with increasing pH (Yoshie *et al.*, 1997), and the percentage of soluble Fe obtained at an endogenous pH (2.5-3.1) was higher than that at pH 5.5 in model systems containing organic acids and lignin (Suzuki, *et al.*, 1992). The percentages of soluble Ca in white wine pomace and apple pomace also increased with decreasing pH (Torre *et al.*, 1995). Furthermore, Ekhlom *et al.* (2003) demonstrated that the



**Fig. 3.** Molecular weight fraction of soluble calcium as affected by boiling in different solutions.

CR: *Caulerpa sertularioides*, SP: *Sargassum polycystum*, KA: *Kappaphycus alvarezii*

W-0: Water, 0- min boiling; W-20: Water, 20- min boiling; S-0: 1% sodium chloride, 0- min boiling; S-20: 1% sodium chloride, 20- min boiling; AC-0: 0.5% acetic acid, 0- min boiling; AC-20: 0.5% acetic acid, 20- min boiling.



**Fig. 4.** Molecular weight fraction of soluble magnesium as affected by boiling in different solutions.

CR: *Caulerpa sertularioides*, SP: *Sargassum polycystum*, KA: *Kappaphycus alvarezii*.

W-0: Water, 0- min boiling; W-20: Water, 20- min boiling; S-0: 1% sodium chloride, 0- min boiling; S-20: 1% sodium chloride, 20- min boiling; AC-0: 0.5% acetic acid, 0- min boiling; AC-20: 0.5% acetic acid, 20- min boiling.

solubilities of Ca, Mg and K in oat bran increased with citric acid concentration. In this study, the solubilities of Ca and Mg in Indonesian seaweed samples were also influenced by pH. By heating in 0.5% acetic acid, the solubilities of Ca and Mg increased compared with those in the case of heating in water or 1% sodium chloride.

Seaweeds contain a high dietary fiber content which is associated with minerals and influences their solubility. Fiber components, such as cellulose, hemicellulose, pectin, other polysaccharides and lignin, may form insoluble complexes with elemental minerals, and thus reduce the bioavailability of these minerals (Eklom *et al.*, 2003). In the case of seaweeds, heating was effective for extracting carbohydrates from matrixes and water-soluble dietary fiber content in all seaweeds steadily increased during heating (Suzuki *et al.*, 1993<sup>b</sup>; Suzuki *et al.*, 1993<sup>c</sup>).

The soluble dietary fiber contents of *Kappaphycus alvarezii* and *Padina australis* were higher than those in the other seaweed samples (Santoso *et al.*, 2002), and solubilities of Ca and Mg in both of these seaweeds was

expected to increase, particularly after boiling in 0.5% acetic acid. These results indicate that soluble dietary fibers are enhancers, and thus they could interact with magnesium and calcium and induce increases in their solubility. Clydesdale (1988) defined an enhancer as a molecular species in food that forms a compound with minerals which is soluble and can be absorbed by mucosal cells.

*Soluble minerals fractionated by molecular weight*  
Seaweeds contain a large amount of dietary fibers. During boiling dietary fibers can be extracted from matrix, particularly soluble dietary fibers can interact with other components including minerals. The interaction between soluble minerals and soluble dietary fibers produces mineral-dietary fiber complexes in different molecular weight fractions. In this experiment, we separated soluble minerals by molecular weight in three categories: low (MW < 10,000), mid (MW = 10,000–200,000), and high (MW > 200,000) molecular weight fractions by ultrafiltration. Three species of Indonesian seaweeds were se-

lected, namely *Caulerpa racemosa*, *Sargassum polycystum*, and *Kappaphycus alvarezii*, representative of green, brown and red algae groups, respectively.

The solubilities of Ca and Mg in the low, mid, and high molecular weight fractions as affected by boiling in different solutions are presented in Figs. 3 and 4, respectively. Boiling in 1% sodium chloride increased the solubility of Ca in the low and mid molecular weight fractions, but not in the mid molecular weight fraction in *Kappaphycus alvarezii*. However, in the high molecular weight fraction (MW > 200,000), its solubility decreased after boiling. The largest decrease was found in *Sargassum polycystum* followed by *Kappaphycus alvarezii*, where Ca solubility decreased from 70.0 to 43.4% and 24.7 to 5.5%, respectively.

Boiling in the other solutions gave various solubilities of Ca in the low, mid and high molecular weight fractions. Boiling in either water or 0.5% acetic acid sometimes increased and sometimes decreased Ca solubility. Soluble Ca was mainly in low and high molecular weight fractions. However, in the case of *Kappaphycus alvarezii* after boiling in water, the Ca solubility was also high (39.4%) in the mid molecular weight fraction.

The result suggests that the interaction between soluble Ca and low and high molecular weight compounds is stronger than that between soluble Ca and mid molecular weight compounds. Since dietary fibers are the main components of seaweeds, they can interact with other compounds including minerals. These low and high molecular weight fractions of soluble dietary fibers can bind to soluble Ca.

The solubility of Mg in the low, mid, and high molecular weight fractions was varyingly affected by boiling in different solutions and did not show a distinct pattern. More than 75% soluble Mg was found in the high molecular weight fraction (MW > 200,000), and boiling in 0.5% acetic acid produced the highest percentage (98.7%) of soluble Mg in *Sargassum polycystum*.

The percentages of soluble Mg in the low molecular weight fractions ranged from 1.6% (*Sargassum polycystum* in 1% sodium chloride, before boiling) to 23.2% (*Kappaphycus alvarezii* in 1% sodium chloride, after boiling), respectively. Generally, a moderate percentage of soluble Mg in the low molecular weight fraction was found in water without boiling, whereas in the other solutions the percentage was not so high, except in 1% sodium chloride after boiling and in 0.5% acetic acid without boiling in *Kappaphycus alvarezii*.

The percentages of soluble Mg in the mid molecular weight fraction was low ( $\leq 2.0\%$ ), except in *Kappaphycus alvarezii* after boiling in water (37.4%) and 0.5% acetic acid (12.2%).

Results obtained by Suzuki *et al.* (2000) showed that more than 50% of soluble Mg in low molecular weight fraction was found in shellfish, and only a low percentage of soluble Mg was found in the high molecular weight fraction in short-necked and hard clam. Shellfish have high protein and low fiber contents, different from Indonesian seaweeds, which contain a high fiber content and a low protein content (Santoso *et al.*, 2002). This suggests

that most soluble Mg interacts with high molecular weight compounds (*i.e.*, high molecular weight soluble dietary fibers).

From these results, it is concluded that Indonesian seaweeds contain high macromineral (*i.e.*, Na, K, Ca and Mg) contents, but low trace-mineral (*i.e.*, Cu, Zn and Fe) contents. The solubilities of Ca and Mg are significantly increased by boiling, particularly in acid solutions (0.5% acetic acid).

In this experiment, the sample collection was carried out only in the dry season. Therefore, further research is necessary to precisely determine the effects of different collection or harvest times (*i.e.*, wet season) on the level of minerals including heavy metals as contaminant indicators.

Since people consume food not in a single form but in mixed forms, our study of the interaction between food components and minerals as enhancers or inhibitors is ongoing.

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