

# Chemical and sensory characteristics of low molecular weight fractions obtained from three types of Japanese soy sauce (shoyu) – Koikuchi, tamari and shiro shoyu

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## Abstract

Three typical forms of Japanese soy sauce (shoyu), koikuchi, tamari and shiro shoyu, can be differentiated, primarily due to their different compositions of soybeans and wheat used for their productions. To evaluate and compare the chemical characteristics of the low molecular weight (MW) fractions of three types of shoyu with their sensory data, gel filtration fractions of ultrafiltration products with MW less than 500 Da (F-500) were subjected to chemical and sensory analyses. The results showed that salty and umami tastes were characteristic of all F-500 fractions, however, the umami taste intensities of those of koikuchi and tamari shoyu were found to be twice as large as that of shiro shoyu. After separation by gel filtration, it was found that the tastiest fractions of the three types of shoyu were those containing sodium salt, free l-glutamic acid and most other free amino acids, especially sweet taste-eliciting amino acids, at concentrations above their thresholds. In some umami fractions of koikuchi and shiro shoyu, that predominantly contained salt and phenylalanine but had a relatively low free l-glutamic acid content, a potential synergistic effect among free l-glutamic acid, salt and phenylalanine was obvious. This first report offers new insights into soy sauce research.

**Keywords:** Ultrafiltration; Gel filtration chromatography; Koikuchi, tamari and shiro shoyu; Soy sauce; Umami taste

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## 1. Introduction

Shoyu is a traditional Japanese soy sauce that is made from a mixture of soybeans and wheat using a well-established two-step fermentation process, which involves koji-mold fermentation to yield koji and brine fermentation to yield moromi. The Japanese people recognize five distinct types of shoyu, based on specific physical and aroma characteristics, that result from differences in the soybean and wheat composition as well as differences in the fermentation processes that are used in their production ([Fukushima, 1981](#) and [Fukushima, 1989](#)). Among these, three typical forms of shoyu – koikuchi, tamari and shiro shoyu – can be differentiated, primarily due to their different raw material compositions. These are thought to contribute to the different types and compositions of breakdown products that are released during fermentation, which consequently impart the characteristic tastes of these three types of shoyu.

Koikuchi shoyu, which has a strong aroma and a deep brown colour, is produced from equal amounts of soybeans and wheat. Tamari shoyu, which is characterized by greater viscosity and less aroma than koikuchi shoyu but has a darker brown colour, is produced using soybeans as the main ingredient with a relatively small amount of wheat. Shiro shoyu is made using a relatively high ratio of wheat to soybeans and has a light yellow to tan colour ([Fukushima, 1981](#), [Fukushima, 1989](#) and [Yokotsuka, 1986](#)).

In shoyu fermentation, which commonly takes at least 6–8 months, ~70–90% of the total soybean and wheat proteins are degraded into free amino acids ([Chou and Ling, 1998](#), [Fukushima, 1989](#) and [Takeuchi et al., 1962](#)), the most abundant of which are glutamic acid, aspartic acid and leucine ([Chou and Ling, 1998](#), [Fujiwara, Tanaka, et al., 1962](#) and [Fujiwara, Tokuda, et al., 1962](#)). Short peptides, with an average length of 2 or 3 residues, are also present in both koikuchi and tamari shoyu at concentrations ranging from 10% to 20% of the total nitrogenous compounds ([Takeuchi et al., 1962](#)). Therefore, during shoyu fermentation, it is likely that most of the proteins that are contained in soybeans and wheat are primarily hydrolyzed into free amino acids and short peptides with a molecular weight (MW) of less than 500 Da. Fractions of another soy sauce, which is known as Indonesian soy sauce, within this MW range were reported to contain the majority of the taste compounds ([Apriyantono et al., 2004](#) and [Lioe et al., 2004](#)). However, to our knowledge, the characteristic tastes and chemistry of the low MW fractions of the three types of Japanese soy sauce have not been investigated previously. The last taste study of Japanese soy sauce was even reported more than 30-years-ago ([Oka and Nagata,](#)

[1974a](#) and [Oka and Nagata, 1974b](#)). Therefore, attempts were made to characterize shoyu fractions with a MW of less than 500 Da obtained by ultrafiltration, chemically and sensorially. These fractions were then separated further by gel filtration chromatography to characterize their taste-active fractions through evaluations of their chemical compositions in relation to their taste characteristics.

## **2. Materials and methods**

### **2.1. Materials**

Three types of shoyu – that is koikuchi, tamari and shiro shoyu – were purchased at a supermarket in Okinawa, Japan, and immediately stored in a refrigerator at 5–7 °C prior to use. Milli-Q water, which was used throughout this research, was obtained from a Milli-Q SP reagent water system (Millipore Corp., Bedford, MA, USA). All of the chemicals used in the analysis were of analytical grade.

### **2.2. Methods**

#### **2.2.1. Stepwise ultrafiltration**

Each original shoyu sample was diluted 1:1 with Milli-Q water to enable it to be ultrafiltered under the conditions described below. Each diluted shoyu sample was then submitted to an ultrafiltration cell, using an Amicon model 202 ultrafiltration unit (Amicon Inc., Beverly, MA, USA) at 4 °C under 1.5–2.0 bar N<sub>2</sub> pressure. Stepwise ultrafiltration was carried out to obtain fractions with a MW of less than 500 Da (F-500) for each type of shoyu using a 0.45 µm mixed cellulose ester membrane (Advantec, Toyo Roshi Kaisha Ltd., Japan), followed by Q0100 (MWCO 10,000 Da, Advantec), YM3 (MWCO 3000 Da, Millipore), and YC05 (MWCO 500 Da, Millipore) membranes. Each F-500 fraction was collected, freeze-dried and reconstituted to its original volume using Milli-Q water.

#### **2.2.2. Gel filtration chromatography**

Samples (5 ml) of the non-diluted F-500 fractions of koikuchi and shiro shoyu, and the 1 + 1 diluted F-500 fraction of tamari shoyu, were chromatographed on a Sephadex G-25 superfine (SF) column (2.6 × 90 cm; Pharmacia, Uppsala, Sweden) at 4 °C and a flow rate of 26 ml/h with Milli-Q water as the eluant. The different application for tamari shoyu was chosen to obtain a better separation. One-hundred and fifty aliquots (7.5 ml each) of the eluate were collected using an Iwaki FRC-2120 fraction collector (Iwaki Glass Co. Ltd., Tokyo, Japan). The UV absorbance of each tube was measured at 214 and 280 nm, using a Shimadzu UV-160 spectrophotometer (Shimadzu Corp., Kyoto, Japan). To detect the salt peak, the sodium chloride (NaCl) concentration was also analyzed by the Mohr method ([Fischer & Peters, 1968](#)). The absorbance and salt concentration were plotted against the tube number. The collected fractions were then freeze-dried. The corresponding fractions from two chromatographic runs of the F-500 fractions of koikuchi and shiro shoyu, as well as those from four runs for tamari shoyu, were combined and reconstituted to the total initial volume (that is 10.0 ml) with Milli-Q water. Finally, the resulting fractions were subjected to several analyses as described below.

### 2.2.3. Chemical analysis

Analyses of dry matters, total peptides, NaCl concentration, total sugars, total acids, pH and free l-glutamic acid concentration were performed for the three F-500 fractions and their resulting gel filtration fractions in duplicate. Dry matters of samples were analyzed by a gravimetric method and using a freeze dryer TAITEC model VD-80 (Taitec Corp., Saitama, Japan). Total peptides were determined by the Lowry method ([Lowry, Rosebrough, Farr, & Randall, 1951](#)) using a DC protein assay reagent kit (Bio-Rad Laboratories, CA, USA) and a Benchmark Plus Bio-Rad microplate spectrophotometer. The salt concentration was quantified by the Mohr method ([Fischer & Peters, 1968](#)). Total sugars were analyzed by the anthrone method ([Clegg, 1956](#)) using the anthrone reagent (Nacalai Tesque, Kyoto, Japan), d-glucose standard (Cica, Kanto Chemical Co., Tokyo, Japan) and a Shimadzu UV-160 spectrophotometer. Total sugars were measured as total glucose, because this monosaccharide constitutes 85% of the total carbohydrates in soy sauce ([Fukushima, 1989](#)). Total acids were determined by titrating 200  $\mu$ l of the sample, which was combined with 20 ml of deionized water, with 0.10 N NaOH and using 0.1% phenolphthalein in ethanol as an indicator, according to the method described by the Association of Official Analytical Chemists ([AOAC, 1990](#)). The total acids were calculated, based on lactic acid, because it was abundant in shoyu, that is 70% of the total organic acids in soy sauce ([Fukushima, 1989](#)). pH was measured using a pH meter, model F.8L (Horiba, Japan) and Advantec pH test paper (that is, PB with a pH range of 3.2–5.6 and MR-BTB No. 20 with a pH range of 5.0–8.0). The free l-glutamic acid concentration was quantified using a Böehringer-Mannheim/R-Biopharm reagent kit (Böehringer Mannheim, Mannheim, Germany), following the enzymatic method described by [Böehringer Mannheim \(1995\)](#).

The total amino acid compositions of the F-500 fractions and their resulting gel filtration fractions were determined by HPLC-OPA precolumn derivatization ([Harris, 1988](#)) after hydrolysis with 6 N HCl (in a vacuum at 110 °C for 20 h). This analysis was performed on a Shimadzu LC-10A amino acid analyzer system using a Shim-pack Amino-Na column (i.d. 6.0 mm  $\times$  10 cm), OPA fluorescence detector and Na-form buffer solvents of amino acid analytical grade (Shimadzu).

### 2.2.4. Capillary zone electrophoresis (CZE) profiles

CZE profiles were obtained for the F-500 fractions of the three types of shoyu, in order to recognize their characteristic patterns. The profiles were obtained using a Photal CAPI-3300 System (Otsuka Electronics Co., Japan) on an uncoated fused silica column (i.d., 50  $\mu$ m, *l* 50 cm, *l*<sub>eff</sub>, 37.8 cm) under the following conditions: voltage, 15 kV; hydrodynamic injection time, 30 s; running time, 25 min; room temperature; buffer, phosphate 50 mM (pH 2.5); and detection at 190–400 nm (photodiode array detector/PDAD). Sample solutions were filtered through a 0.45  $\mu$ m membrane (Advantec) prior to analysis.

### 2.2.5. Taste dilution factors and taste descriptions

#### 2.2.5.1. Training of the panellists

Eight of a total of 16 subjects from the Department of Bioscience and Biotechnology of the University of the Ryukyus, Okinawa, Japan, were selected as panellists and trained using a

triangle test, according to the method of [Carpenter, Lyon, and Hasdell \(2000\)](#), to respond to the five basic tastes with the following taste solutions: NaCl (12 mM) for a salty taste; monosodium glutamate (MSG; 4 mM; pH 5.6) for an umami taste; saccharose (40 mM) for a sweet taste; lactic acid (10 mM) for a sour taste; and caffeine (1.5 mM) for a bitter taste. Each panellist examined three sets of each taste solution per session, which were served in a triangle test with Milli-Q water. The training was conducted through two separate sessions. In this training and the subsequent evaluation of samples, retasting was not allowed. Sensory evaluation was done in a room at a temperature of 25 °C.

#### 2.2.5.2. Determination of taste dilution factors and taste descriptions

In order to determine the taste dilution factors that best described the taste intensities of the F-500 fractions and their gel filtration fractions, taste dilution analysis was conducted, following the procedures of [Frank, Ottinger, and Hofmann \(2001\)](#), as well as [Ottinger and Hofmann \(2003\)](#). Exactly 1.0 ml of each of the shoyu fractions was diluted stepwise 1:1 with Milli-Q water (pH 5.6). Each 0.5 ml dilution was then evaluated by the eight trained panellists in a triangle test, in order of increasing concentration. The dilution at which a taste difference between the diluted fraction and two blanks of Milli-Q water could just be detected was determined as the taste dilution (TD) factor. Once a taste could be detected, the panellists were asked to describe it in terms of the five basic tastes that had been examined in the training sessions.

### 3. Results and discussion

#### 3.1. Characteristics of F-500 fractions from the three types of shoyu

The chemical compositions and sensory data of the ultrafiltration fractions with a MW of less than 500 Da (F-500) from koikuchi, tamari and shiro shoyu are shown in [Table 1](#). It is clear that salty and umami tastes were the dominant tastes of the three F-500 fractions. Comparing to shiro shoyu, umami taste occurred at higher intensities in both koikuchi and tamari shoyu, even though the free l-glutamic acid contents, the well-known key compound of umami taste, of the three F-500 fractions, were considerably similar. It is noted that the major components quantified in the three fractions were 84% to 94% of the total dry matters in F-500 fractions or 68% to 77% of those in original shoyu ([Table 1](#)). This indicated that most of the Japanese soy sauce components have been quantified. Among the major components that were observed in the three shoyu samples ([Table 1](#)), free amino acids seem to be the most responsible for the intense umami tastes of both koikuchi and tamari shoyu, because their concentrations in these two samples were much higher than that observed in shiro shoyu.

Table 1.

Chemical composition, taste dilution (TD) factor and sensory analysis results of the ultrafiltration fractions with MW less than 500 Da (F-500) obtained from three types of shoyu

Fraction F-500 (% w/v)	Koikuchi	Tamari	Shiro	Threshold value	Taste quality
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Fraction F-500 (% w/v)	Koikuchi	Tamari	Shiro	Threshold value	Taste quality
Dry matters	35.18	37.08	41.59		
NaCl	15.1	14.0	17.4	0.06 <sup>a</sup>	Salty
Total peptides	2.39	4.69	1.13		
Free AA <sup>b</sup>	6.03	6.86	1.82		
Free l-Glu	1.19	1.10	1.00	0.03 <sup>c</sup>	Umami
Total sugars (as glucose)	2.30	2.37	17.12	0.86 <sup>a</sup>	Sweet
Total acids (as lactic acid)	2.70	3.07	0.79	0.01 <sup>d</sup>	Sour
pH	4.8	4.7	5.2		
Taste quality (TD factor) <sup>e</sup>	Umami (128)	Umami (128)	Salty (128)		
	Salty (128)	Salty (128)	Umami (64)		
	Sour (16)	Sour (32)	Sweet (8)		
	Sweet (2)				
Dry matters of original shoyu	41.64	47.11	51.28		

<sup>a</sup> [Soldo et al. \(2003\)](#).

<sup>b</sup> Free amino acids calculated by difference between the total amino acids (determined by HPLC-OPA derivatization after acid hydrolysis) and the total peptides (Lowry method).

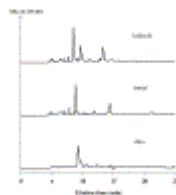
<sup>c</sup> Determined as l-Glu(Na) by [Kato et al. \(1989\)](#).

<sup>d</sup> Stahl (1978) cited by [Salles et al. \(2000\)](#).

<sup>e</sup> Following the methods of [Frank et al. \(2001\)](#) as well as [Ottinger and Hofmann \(2003\)](#).

A slight note of sweetness was found in koikuchi and shiro shoyu, whereas sourness was detected in koikuchi and tamari shoyu. The sweetness in shiro shoyu and the sourness in tamari shoyu can be easily explained by their respective high and low ratios of sugar to acid concentrations, although the presence of sugar and acid in the three shoyu fractions was considered above their thresholds ([Table 1](#)). In several reports on Japanese soy sauce, it is found that glucose and lactic acid impart 80–85% and 70–80% of the total sugars and organic acids, respectively ([Fukushima, 1985](#), [Fukushima, 1989](#) and [Yokotsuka, 1983](#)).

[Fig. 1](#) shows that the three types of shoyu had different CZE profiles of their F-500 fractions. As these profiles were observed at 214 nm – that is, at the high absorption range of most peptides and some free amino acids ([Belitz and Grosch, 1999](#), [Cliffe et al., 1993](#), [Frazier, 2001](#) and [Frazier et al., 1999](#)) – these conditions indicate that the three shoyu samples contained different varieties of low MW nitrogenous compounds, and this is assumed to be a result of the different ratios of soybeans to wheat, as explained above. Those varieties are thought to correlate to the taste differences among the three shoyu samples and led to further characterization below.

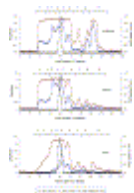


[Full-size image](#) (18K)

Fig. 1. CZE electropherograms of the ultrafiltration fractions with a MW of less than 500 Da (F-500) obtained from the three types of shoyu, detected at 214 nm. CZE was performed on an uncoated fused silica column,  $l_{\text{eff}}$ , 37.8 cm, at room temperature using 50 mM phosphate buffer at pH 2.5 as an eluting buffer and a photodiode array detector.

### 3.2. Characteristics of gel filtration fractions from the three F-500 fractions

To more precisely evaluate the above-mentioned characteristics of the low MW fractions from the three types of shoyu, further separations were performed by Sephadex G-25 SF gel filtration chromatography. The separation profiles of the three F-500 fractions are shown in [Fig. 2](#). Each F-500 fraction was separated further into nine fractions (fractions 1–9). In the fractionations of koikuchi and tamari shoyu, the profiles of the earlier runs (as shown by fractions 1–6) were similar, but were notably different from that of shiro shoyu, as illustrated by the 214 and 280 nm absorbance profiles. However, all of the separations had a similar pattern of salt peak; that is, the sodium salt that was contained in each F-500 fraction was separated into only one peak. It should also be noted that all of the salt fractions were collected in fractions 4 and 5.



[Full-size image](#) (116K)



Fig. 2. Gel filtration chromatograms of the F-500 fractions of the three types of shoyu. Fractionation was performed on a Sephadex G-25 SF column (2.6 × 90 cm) at 4 °C using Milli-Q water as the eluant.

The chemical compositions of the gel filtration fractions from the three types of shoyu are presented in [Table 2](#), [Table 3](#) and [Table 4](#), and their total amino acid compositions are shown in [Table 5](#), [Table 6](#) and [Table 7](#). In the total amino acid analysis, tryptophan and cysteine could not be measured, because the former was destroyed during acid hydrolysis and the concentration of the latter was too low to be detected as it was partially oxidized ([Harris, 1988](#)).

Table 2.

Chemical compositions of the Sephadex G-25 SF gel filtration fractions obtained from the F-500 fraction of koikuchi shoyu

Fractions	Content, % w/v							pH
	Dry matters	NaCl	Total peptides	Free amino acids <sup>a</sup>	Free l-Glu	Total sugars	Total acids	
1	0.34		0.12			0.03		5.6
2	0.29		0.11	0.03		0.05		5.6
3	6.10		0.94	1.37	0.32	0.61	0.68	5.2
4	15.41	5.15	0.51	3.76	0.69	0.85	1.46	5.2
5	11.60	8.98	0.24	0.88	0.10	0.11	0.56	4.8
6	0.12		0.11					5.6
7	0.03		0.03					5.8
8	0.03		0.03					5.8
9	0.08		0.07					5.8

<sup>a</sup> Calculated by difference between the total amino acids (determined by HPLC-OPA derivatization after acid hydrolysis) and the total peptides (Lowry method).

Table 3.



Chemical compositions of the Sephadex G-25 SF gel filtration fractions obtained from the F-500 fraction of tamari shoyu

Fractions	Content, % w/v							pH
	Dry matters	NaCl	Total peptides	Free amino acids <sup>a</sup>	Free l-Glu	Total sugars	Total acids	
1	0.80		0.29	0.04		0.04		5.6
2	0.63		0.22	0.04		0.05		5.4
3	8.93		1.74	1.48	0.38	0.57	1.46	4.8
4	22.35	11.2	1.16	5.10	0.64	0.47	2.48	5.2
5	3.71	1.86	0.63		tr <sup>b</sup>	0.01	0.23	5.4
6	0.26		0.21					5.8
7	0.12		0.03					5.8
8	0.10		0.06					5.8
9	0.10		0.03					5.8

<sup>a</sup> Calculated by difference between the total amino acids (determined by HPLC-OPA derivatization after acid hydrolysis) and the total peptides (Lowry method).

<sup>b</sup> Trace (<0.005% w/v).

Table 4.

Chemical compositions of the Sephadex G-25 SF gel filtration fractions obtained from the F-500 fraction of shiro shoyu

Fractions	Content, % w/v							pH
	Dry matters	NaCl	Total peptides	Free amino acids <sup>a</sup>	Free l-Glu	Total sugars	Total acids	
1	0.05		0.01					5.4
2	0.21		0.03	0.01		0.05		5.6
3	6.93		0.24	0.49	0.23	5.84	0.23	4.6
4	23.30	8.22	0.35	1.38	0.45	12.03	0.68	4.8

Fractions	Content, % w/v							pH
	Dry matters	NaCl	Total peptides	Free amino acids <sup>a</sup>	Free l-Glu	Total sugars	Total acids	
5	10.05	7.51	0.35		0.06	1.93	0.23	4.8
6	0.10		0.08			0.01		5.6
7	0.09		0.03					5.6
8	0.03		0.01					5.6
9	0.05		0.02					5.6

<sup>a</sup> Calculated as the difference between the total amino acids (determined by HPLC-OPA derivatization after acid hydrolysis) and the total peptides (Lowry method).

Table 5.

Total amino acid compositions of the gel filtration fractions of the F-500 fraction of koikuchi shoyu

Amino acids (AA) (%w/v)	Fractions					Threshold value of free AA <sup>a</sup>	Taste of free AA <sup>a</sup>
	1	2	3	4	5		
Asp	0.04	0.04	0.14	0.20	0.05	0.10	Umami
Glu	0.04	0.06	0.59	1.19	0.19	0.03	Umami
Ala			0.09	0.46	0.06	0.06	Sweet
His			0.07	0.11		0.02	Bitter
Lys		0.01	0.36	0.13	0.02	0.05	Sweet, bitter
Arg			0.10	0.26	0.07	0.05	Bitter
Gly	0.01	0.01	0.07	0.20	0.08	0.13	Sweet
Thr	0.01		0.06	0.22	0.01	0.26	Sweet
Ser	0.01		0.05	0.29	0.10	0.15	Sweet
Pro		0.01	0.21	0.29	0.04	0.30	Sweet, bitter

Amino acids (AA) (%w/v)	Fractions					Threshold value of free AA <sup>a</sup>	Taste of free AA <sup>a</sup>
	1	2	3	4	5		
Met			0.03	0.07	0.03	0.03	Bitter
Val			0.20	0.24	0.03	0.04	Bitter
Tyr					0.01	nd <sup>b</sup>	
Phe	0.01	0.01	0.03	0.05	0.34	0.09	Bitter
Ile			0.15	0.22	0.03	0.09	Bitter
Leu			0.16	0.34	0.06	0.19	Bitter

<sup>a</sup> [Kato et al. \(1989\)](#), Glu and Asp were determined as their sodium salts.

<sup>b</sup> Not determined.

Table 6.

Total amino acid compositions of the gel filtration fractions of the F-500 fraction of tamari shoyu

Amino acids (AA) (%w/v)	Fractions					Threshold value of free AA <sup>a</sup>	Taste of free AA <sup>a</sup>
	1	2	3	4	5		
Asp	0.12	0.08	0.33	0.79	0.02	0.10	Umami
Glu	0.09	0.08	0.73	1.56	0.01	0.03	Umami
Ala	0.01	0.01	0.09	0.38		0.06	Sweet
His	0.01		0.21	0.20		0.02	Bitter
Lys	0.01	0.01	0.47	0.16	0.01	0.05	Sweet, bitter
Arg	0.01		0.09	0.49	0.06	0.05	Bitter
Gly	0.01	0.01	0.10	0.30	0.01	0.13	Sweet
Thr	0.01	0.01	0.04	0.36		0.26	Sweet
Ser	0.01	0.01	0.08	0.48	0.01	0.15	Sweet
Pro	0.01	0.01	0.26	0.38	0.01	0.30	Sweet, bitter

Amino acids (AA) (%w/v)	Fractions					Threshold value of free AA <sup>a</sup>	Taste of free AA <sup>a</sup>
	1	2	3	4	5		
Met			0.04	0.09		0.03	Bitter
Val	0.01	0.01	0.31	0.25		0.04	Bitter
Tyr					0.03	nd <sup>b</sup>	
Phe	0.01	0.01		0.21	0.36	0.09	Bitter
Ile	0.01	0.01	0.22	0.25		0.09	Bitter
Leu	0.01	0.01	0.25	0.36		0.19	Bitter

<sup>a</sup> [Kato et al. \(1989\)](#), Glu and Asp were determined as their sodium salts.

<sup>b</sup> Not determined.

Table 7.

Total amino acid compositions of the gel filtration fractions of the F-500 fraction of shiro shoyu

Amino acids (AA) (%w/v)	Fractions					Threshold value of free AA <sup>a</sup>	Taste of free AA <sup>a</sup>
	1	2	3	4	5		
Asp			0.02	0.11	0.01	0.10	Umami
Glu	0.01	0.03	0.42	0.94	0.09	0.03	Umami
Ala			0.01	0.07	0.01	0.06	Sweet
His			0.01	0.04		0.02	Bitter
Lys			0.05	0.03		0.05	Sweet, bitter
Arg			0.02	0.05	0.01	0.05	Bitter
Gly			0.01	0.06	0.02	0.13	Sweet
Thr			0.01	0.04	0.01	0.26	Sweet
Ser			0.01	0.07	0.02	0.15	Sweet
Pro		0.01	0.06	0.12	0.02	0.30	Sweet, bitter

Amino acids (AA) (%w/v)	Fractions					Threshold value of free AA <sup>a</sup>	Taste of free AA <sup>a</sup>
	1	2	3	4	5		
Met				0.01		0.03	Bitter
Val			0.04	0.05	0.01	0.04	Bitter
Tyr						nd <sup>b</sup>	
Phe			0.01	0.02	0.09	0.09	Bitter
Ile			0.02	0.04		0.09	Bitter
Leu			0.04	0.08	0.01	0.19	Bitter

<sup>a</sup> [Kato et al. \(1989\)](#), Glu and Asp were determined as their sodium salts.

<sup>b</sup> Not determined.

The fractionation results shown in [Table 2](#), [Table 3](#) and [Table 4](#) indicate that fractions 3–5 were the major F-500 fractions, which contained notably more components than the other fractions. Their taste intensities, which are described by the value of the TD factors, as well as their taste descriptions, are shown in [Table 8](#). The results revealed that the umami taste was predominant in all of the major fractions at various taste intensities. The highest umami taste intensity was found in fraction 4 of koikuchi and tamari shoyu, which contained sodium salt, free l-glutamic acid and total free amino acids at relatively high concentrations ([Table 2](#) and [Table 3](#)). The concentrations of free l-glutamic acid in these fractions were known to be 21–23 times higher than the umami taste threshold value ([Kato, Rhue, & Nishimura, 1989](#)). Because the tastiest fractions contained sodium salt and it is known that the acidic amino acid can exhibit a synergistic effect with the salt ([Lioe, Apriyantono, Takara, Wada, & Yasuda, 2005](#)), the potential interaction between free l-glutamic acid and salt in the fractions is thought to play an important role in the intense umami taste. In addition, the high concentrations of other umami taste-eliciting amino acids, such as aspartic acid, along with sweet taste-eliciting amino acids, such as alanine, serine and glycine, above their threshold values ([Kato et al., 1989](#)), that were present in fraction 4 of koikuchi and tamari shoyu ([Table 5](#) and [Table 6](#)) might also contribute to the intense umami taste of the fractions and, certainly, their F-500 fractions because of the possible interaction between umami and sweet stimuli ([Heyer, Taylor-Burds, Tran, & Delay, 2003](#)). This fact is consistent with the result reported previously in the umami fractions of Indonesian soy sauce ([Lioe et al., 2004](#)).

Table 8.

TD factors and taste description of the gel filtration fractions from the F-500 fractions of the three types of shoyu<sup>a</sup>

Fractions	Koikuchi shoyu		Tamari shoyu		Shiro shoyu	
	Taste description	TD factor	Taste description	TD factor	Taste description	TD factor
1	– <sup>b</sup>	<1	–	<1	–	<1
2	–	<1	–	<1	–	<1
3	Umami	8	Umami	32	Umami	8
	Sour	8	Sour	16	Sour	4
	Sweet	2			Sweet	2
4	Umami	64	Umami	64	Umami	32
	Salty	32	Salty	64	Salty	32
	Sour	16	Sour	32	Sweet	8
5	Salty	64	Salty	16	Salty	64
	Umami	64	Umami	16	Umami	32
					Sweet	1
6	–	<1	Bitter	1	–	<1
7	–	<1	–	<1	–	<1
8	–	<1	–	<1	–	<1
9	Bitter	1	–	<1	–	<1

<sup>a</sup> Following the methods of [Frank et al. \(2001\)](#) as well as [Ottinger and Hofmann \(2003\)](#).

<sup>b</sup> No perceivable taste.

An exception was found in the other salt fraction of koikuchi shoyu (fraction 5), which contained much lower free l-glutamic acid and total free amino acid concentrations ([Table 2](#)), but also had an intense umami taste ([Table 8](#)). This fraction contained phenylalanine as a dominant amino acid ([Table 5](#)). This so-called bitter amino acid ([Solms, 1969](#)) was present in fraction 5 at concentrations above its bitter taste threshold value ([Kato et al., 1989](#)). Nevertheless, no bitter taste was perceived in the fraction ([Table 8](#)). This could be explained by the recent findings in our previous studies ([Lioe et al., 2004](#) and [Lioe et al., 2005](#)), which revealed a significant interaction among phenylalanine, sodium salt and free l-glutamic acid/MSG that elicited an intense umami taste. For a comparison, this interaction could not be found in fraction 5 of tamari shoyu, which also contained high levels of phenylalanine but a lack of free l-glutamic acid

([Table 3](#) and [Table 6](#)), and therefore had a low intensity of umami taste ([Table 8](#)). However, the interaction was also likely to be present in fraction 5 of shiro shoyu and strongly considered to contribute to the relatively high umami taste intensity of this fraction ([Table 8](#)). Therefore, the same interaction as found in Indonesian soy sauce was also present in this work. On the basis of our knowledge, the possible occurrence of the significant interaction among the three major components in shoyu has not been reported previously in a number of Japanese soy sauce studies; therefore it represents a new insight in the current study.

Free l-glutamic acid, which is as a umami taste-contributing amino acid, was also found to be abundant in fraction 3 of the three types of shoyu, i.e., the content is around half that of the corresponding fraction 4, in contrast, most of these fractions had very low umami taste intensities ([Table 8](#)). This effect was thought to be caused by the dominant sour taste of free l-glutamic acid ([Kato et al., 1989](#)) in the absence of sodium salt ([Table 2](#), [Table 3](#) and [Table 4](#)), even though the fractions had relatively high concentrations of lysine ([Table 5](#), [Table 6](#) and [Table 7](#)), which can also taste salty, as found in a cheese fraction reported by [Molina, Ramos, Alonso, and López-Fandiño \(1999\)](#). In addition, the presence of sweet taste-eliciting amino acids in fraction 3, which could contribute to the perceived umami taste, was mostly lower than their thresholds ([Table 5](#), [Table 6](#) and [Table 7](#)). Therefore, it can be inferred from these results that the presence of the other major components, such as sodium salt and sweet amino acids, together raised synergistic interactions that were important for the tastiness of shoyu.

Finally, in this study, peptide fractions were found in the early (fractions 1 and 2) and late (fractions 6–9) gel filtration runs ([Fig. 1](#) and [Table 2](#), [Table 3](#) and [Table 4](#)). However, these either had no taste or tasted bitter with low taste intensities or TD factors ([Table 8](#)). These results imply that the peptide fractions made no significant contribution to the umami taste of shoyu. Nevertheless, peptides were also present comparably in some umami fractions, i.e., fraction 4 of koikuchi shoyu ([Table 2](#)) and fractions 3 and 4 of tamari shoyu ([Table 3](#)). The precise contribution of these peptides has been investigated further by screening the peptide subfractions using the RP-HPLC/taste dilution analysis method developed by [Frank et al., 2001](#) and [Ottinger and Hofmann, 2003](#). The results showed that there was no significant contribution of peptides to the umami taste of any shoyu. Also, in this investigation, it was clearly found that free glutamic acid, as well as several sweet amino acids and sodium salt, were the key contributors of the high intensity of umami taste perceived in fraction 4 of koikuchi and tamari shoyu, which is consistent with that explained above.

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