



HOKKAIDO UNIVERSITY

Title	Molecular Species of Fish Muscle Lecithin
Author(s)	Takahashi, Koretaro; Hirano, Tsugihiko; Takama, Kozo et al.
Citation	日本水産学会誌, 48(12), 1803-1814 https://doi.org/10.2331/suisan.48.1803
Issue Date	1982
Doc URL	https://hdl.handle.net/2115/49848
Rights	© 1982 公益社団法人日本水産学会
Type	journal article
File Information	BJSSF48_1803.pdf



Molecular Species of Fish Muscle Lecithin^{*1,*2}Koretarō TAKAHASHI,^{*3} Tsugihiko HIRANO,^{*4} Kōzō TAKAMA^{*3} and
Kōichi ZAMA^{*3}

(Received May 19, 1982)

A satisfactory separation of the molecular species of fish muscle lecithin was made by modifying the lecithin into diglyceride acetate for high performance reversed phase liquid chromatography (HPLC). The molecular species of each chromatographic peak was determined by fatty acid composition and by total acyl carbon number analysis subsequent to separation, using a thin layer chromatographic technique with silver nitrate impregnated silica gel plates (Ag⁺-TLC). After plotting the relative retention time (RRT) of each molecular species from soybean, egg yolk, chum salmon, big-eyed tuna, Alaska pollack, and carp muscle lecithins semilogarithmically against the total acyl carbon number or the number of total double bonds of each molecular species, a set of parallel straight oblique lines was obtained. Thus we can express the molecular species in matrix relation by giving a variable integer x for the acyl carbon number and a variable integer y for the number of double bonds of each fatty acid in the molecular species. By using the correlations between the RRTs and the corresponding molecular species on semilogarithmic plots, it was concluded that an identification of molecular species from fish lipids could be done by RRTs from HPLC.

Lecithin has been studied by many workers concerning lipid changes during storage or processing of fish products.¹⁾ However, these works were done mainly by analyzing the lipid composition or fatty acid composition, which might not be enough to elucidate the mechanisms of deterioration of foods caused by the changes in lipid. Therefore, a new method for determining molecular species of lecithin is necessary not only in the field of biochemistry, but also in the field of food chemistry.

Recently, HPLC method has been introduced by ŌSHIMA *et al.*^{*5} in order to analyze fish phospholipids which might have the most complicated molecular species composition among biological sources.

The purpose of this study was to develop the method of ŌSHIMA *et al.*^{*5} to determine the individual molecular species of fish phospholipid.

Experimental

Preparation of Lecithin

Total lipids were obtained from the fish muscle tabulated in Table 1, according to the method of BLIGH & DYER. Soybean lecithin was purchased from Wako Pure Chemical Industries, Ltd., Osaka, and egg yolk lecithin was kindly supplied by Asahi Chemical Industry Ltd., Tokyo. These total lipid and crude lecithin were subjected to a silica gel column chromatography which has been done successfully by LANDS *et al.*²⁾ for the purification of lecithin. Eluates were monitored by TLC. And lecithin of more than 95% purity were collected.

Preparation of Diglyceride Acetate from Lecithin

Pure lecithin was dephosphorylized with phospholipase C (*Clostridium perfringens*), according to the method of RENKONEN.³⁾ Diglyceride was prepared by preparative TLC from the dephosphorylized lipid. The developing solvent was *n*-hexane/diethyl ether (1:1, v/v).

Acetylation was performed by adding an appropriate amount of acetic anhydride to the solution of diglyceride in pyridine, and by standing it for 12 h

*1 Molecular Species of Marine Animal Lipid-I.

*2 Presented at the Japanese Society of Scientific Fisheries Meeting, Tokyo, April, 1982.

*3 Laboratory of Food Chemistry I, Faculty of Fisheries, Hokkaido University, Hakodate, Hokkaido 041, Japan (高橋是太郎・高間浩蔵・座間宏一: 北海道大学水産学部).

*4 Nikkan Denshi Ltd., Hakodate, Hokkaido 041, Japan (平野次彦: 日函電子株式会社).

*5 T. ŌSHIMA, S. WADA, and C. KOIZUMI: Presented at the annual meeting of the Japanese Society of Scientific Fisheries, Tsu, Japan, October, 1981.

Table 1. Fish examined.

Species	Mean body length and weight	Locality of catch	Date of catch
Chum salmon (Summer)* ¹ <i>Oncorhynchus keta</i>	65 cm, 3.5 kg, (1)* ²	The offing of Akkeshi, Hokkaido	June 1980
Chum salmon (Fall)* ¹ <i>Oncorhynchus keta</i>	72 cm, 4.5 kg, (1)	The Moheji River, Hokkaido	Nov. 1981
Big-eyed tuna <i>Parathunnus obesus</i>	110 cm, 20 kg, (1)	Purchased from the Market.	—
Alaska pollack <i>Theragra chalcogramma</i>	44 cm, 610 g, (10)	The Uchiura Bay, Hokkaido	Dec. 1981
Carp <i>Cyprinus carpio</i>	23 cm, 175 g, (5)	Cultured	Sep. 1980

*¹ Male. *² Nos. of individual used.

at room temperature.⁴⁾ The resulting diglyceride acetates were purified by the method of preparative TLC by using the solvent *n*-hexane/diethyl ether (75:25, v/v). Finally, they were filtered through a 0.45 μ type FP-45 Fluoropore filter (Sumitomo Electric Industry, Ltd., Osaka) and subjected to HPLC.

HPLC Fractionation of the Molecular Species of Diglyceride Acetate Derived from Lecithin

The diglyceride acetates were fractionated into major molecular species on twin 8 $\times$ 250 mm LiChrosorb RP-18 columns which were connected in series. A Hitachi Liquid Chromatograph Model 638-50 (Hitachi Ltd., Tokyo) equipped with a Shodex RI detector Model SE-11 (Showa Denko Ltd., Tokyo) was used. The eluting solvent used was isopropanol/acetone/methanol/acetonitrile (1:1:3:4, v/v). Diglyceride acetates were dissolved into 5 volumes of tetrahydrofuran, and 25 μ l of these solutions were applied to the column under room temperature (lower the better) and at a flow rate of 1.5 ml/min.

Identification of Molecular Species of Each Peak on HPLC

Peaks on HPLC chromatograms were numbered in sequence of elution. The fatty acid composition of each collected predominant peak was analyzed by gas chromatography. The analytical conditions for fatty acid were as follows.

Gas chromatograph: Hitachi 063, Column: Unisole 3000 (Gasukuro Kogyo Ltd., Tokyo), glass column 0.2 $\times$ 200 cm, Column temp.: 220°C, Detector: FID, Detector temp.: 250°C, Injection temp.: 280°C, Carrier gas: N₂, flow rate: 20 ml/min.

Methyl esters of fatty acids were prepared ac-

cording to the method of CHRISTOPHER and GLASS described by PREVOT and MORDRET.⁵⁾ An aliquot amount (less than 20 mg) of lipid was dissolved in 1 ml *n*-hexane and 0.2 ml of methanolic 2N-NaOH solution was added. After shaking this mixture, it was stand for 20 s under 50°C and then 0.2 ml of methanolic 2N-HCl solution was added. The *n*-hexane layer was collected and then concentrated. Methyl esters prepared were subjected to a gas liquid chromatographic analysis.

The small peaks which have critical pairs were first subjected to Ag⁺-TLC. The developing solvent used was benzene/diethyl ether (8:2, v/v). The band obtained by Ag⁺-TLC were then eluted with diethyl ether containing dotriacontane which was used as an internal standard, and then applied to fatty acid analysis and total acyl carbon number analysis. The analytical conditions for total acyl carbon number analysis were as follows.

Gas chromatograph: Hitachi 063, Column: OV-101 (Gasukuro Kogyo Ltd., Tokyo), steel column 0.3 $\times$ 50 cm, Column temp.: 300–330°C, programed as 1°C/min, Detector: FID, Detector temp.: 340°C, Injection temp.: 345°C, Carrier gas: N₂, flow rate: 60 ml/min.

Hydrolysis of Diglyceride Acetate Derived from Lecithin by Pancreatic Lipase

Fraction of the molecular species of 16:0 in position 1 and 22:6 in position 2 was collected by HPLC. This lipid (less than 5 mg) was then suspended by shaking vigorously in a mixture of 1 M tris-HCl buffer, pH 8 (1 ml), 2.2% CaCl₂ (0.1 ml), and 0.05% taurocholic acid sodium salt (0.25 ml) at 40°C for 1 min. Then, 40 mg of pancreatic lipase (Calbiochem, San Diego, Calif. 92112) was added to the mixture and the reaction was proceeded for 4 min at 40°C by shaking it

virgorously. The reaction was stopped by adding 1 ml of ethanol and 1 ml of 6 N-HCl. The hydrolysate was extracted with diethyl ether and purified by using a preparative TLC with *n*-hexane/diethyl ether/formic acid (80:20:2, v/v) as developing solvent.

Results

Figs. 1-3 show chromatograms of diglyceride acetate derived from each lecithin sources on HPLC. Chromatography was regularly completed in about 2 h. The diglyceride composition of each collected predominant peak was easily determined by fatty acid analysis, as shown in Table 2. This table shows the results on big-eyed tuna lecithin as an example. Peak number 7 in Fig. 3-E is obviously ascribed to the diglyceride acetate composed of 16:0 and 22:6. We have considered that 22:6 is bound in position 2 of the molecule since this peak was the most predom-

Table 2. Determination of lecithin molecular species of major component*

Fatty acid	Peak number		
	5	6	7
15:0			trace
16:0	25.6	48.4	53.8
17:0			trace
18:1	25.9	4.1	trace
20:4			trace
20:5	23.3	3.7	trace
22:4			trace
22:6	25.1	43.8	46.2
Molecular species	$\begin{bmatrix} 18:1 \\ 20:5 \end{bmatrix}$	$\begin{bmatrix} 22:6 \\ 16:0 \end{bmatrix}$	$\begin{bmatrix} 16:0 \\ 22:6 \end{bmatrix}$

* Example of big-eyed tuna.

ant, and it is said that highly unsaturated fatty acids such as 20:5 or 22:6 are usually dominantly bound in position 2.⁸⁻¹⁰⁾ In addition, after pancreatic lipase hydrolysis, only a trace amount

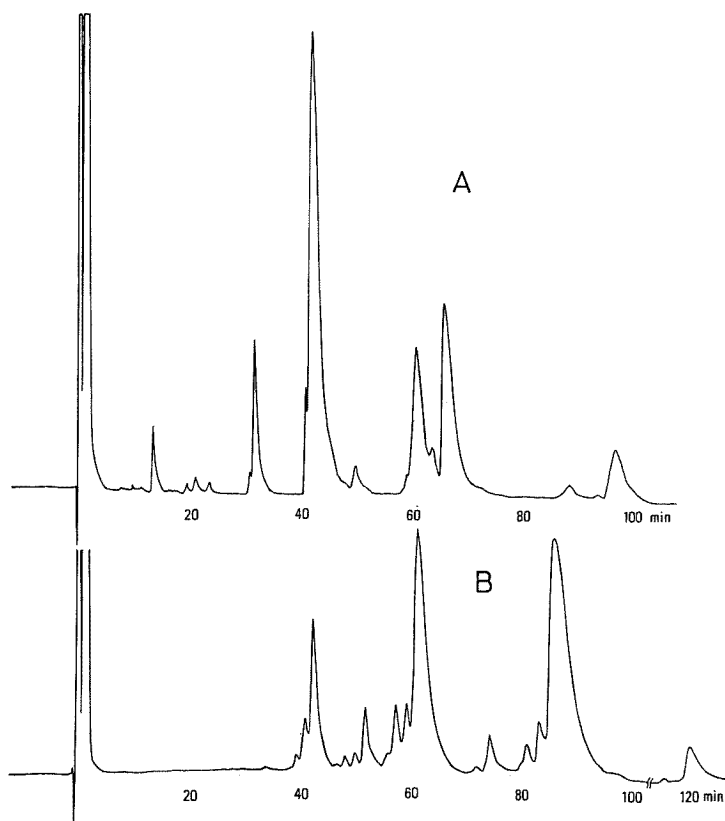


Fig. 1. HPLC chromatograms of soybean and egg yolk lecithin.
A: Soybean. B: Egg yolk.

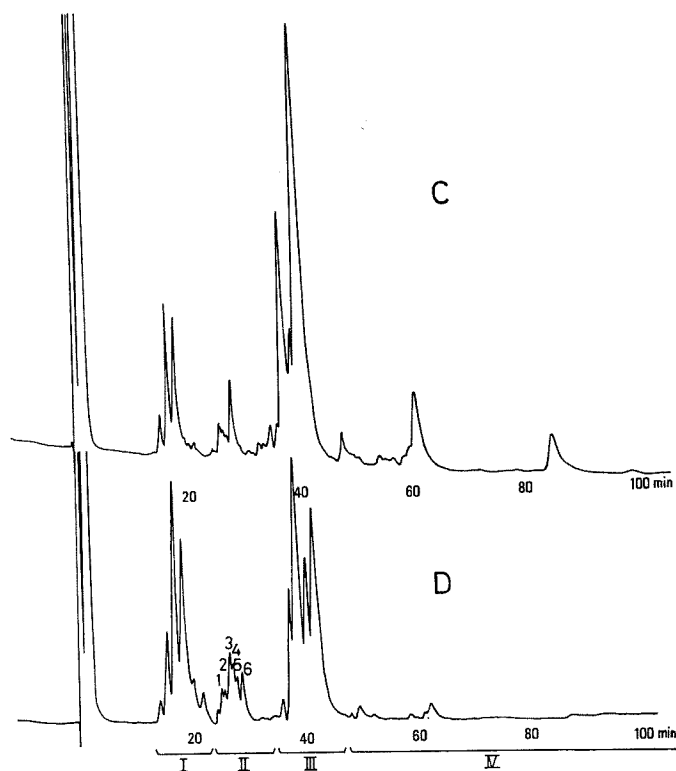


Fig. 2. HPLC chromatograms of chum salmon muscle lecithin.
C: Captured in summer. D: Captured in fall.

of 22: 6 was detected in the free fatty acid fraction (this fraction represents the fatty acid in position 1), although more than 70% of the lipid was hydrolyzed (determined by a densitometric method). Peak number 6 had the same combination as that in peak number 7, whereas 18: 1-20: 5 were considered as contaminants of peak number 5. Nevertheless in this case, 22: 6 was considered to be bound in position 1 in the molecule. Peak number 5 had an almost even amount of fatty acids of 16: 0, 18: 1, 20: 5 and 22: 6. Among these fatty acids, 16: 0 as well as 22: 6 were regarded as contaminants from peak number 6, considering that peak number 6 is larger than peak number 5. It was concluded that peak number 5 was the combination of 18: 1 and 20: 5. Molecular species from other biological sources were also determined in the same manner. The small peaks which have critical pairs were first subjected to Ag^+ -TLC and separated according to their degree of unsaturation. In most of the small peaks in the first half of the HPLC chromatograms of

fish lecithin, only one band appeared on Ag^+ -TLC plate. The complex combinations of molecular species were identified in the following manner. An example of determination of molecular species in the small peaks with critical pairs appeared in the first half of the HPLC chromatogram of chum salmon (captured in fall) is shown in Table 3 (also see Fig. 2-D). Small peak number 1 is obviously a combination of 20: 5 and 16: 1. Small peak number 2 is also a combination of 20: 5 and 16: 1 with 10% unidentified contaminants. The difference in retention time on HPLC between these two peaks was attributed to the differences in binding position of the fatty acid. It was considered that small peak number 1 had 20: 5 in position 1, whereas small peak number 2 had it in position 2 in the molecule, since molecular species which have a highly unsaturated fatty acid in position 2 are likely to elute later than the one which has the same fatty acid in position 1. Small peak number 3 is a combination of 20: 5 in position 1 and 14: 0 in position 2, and

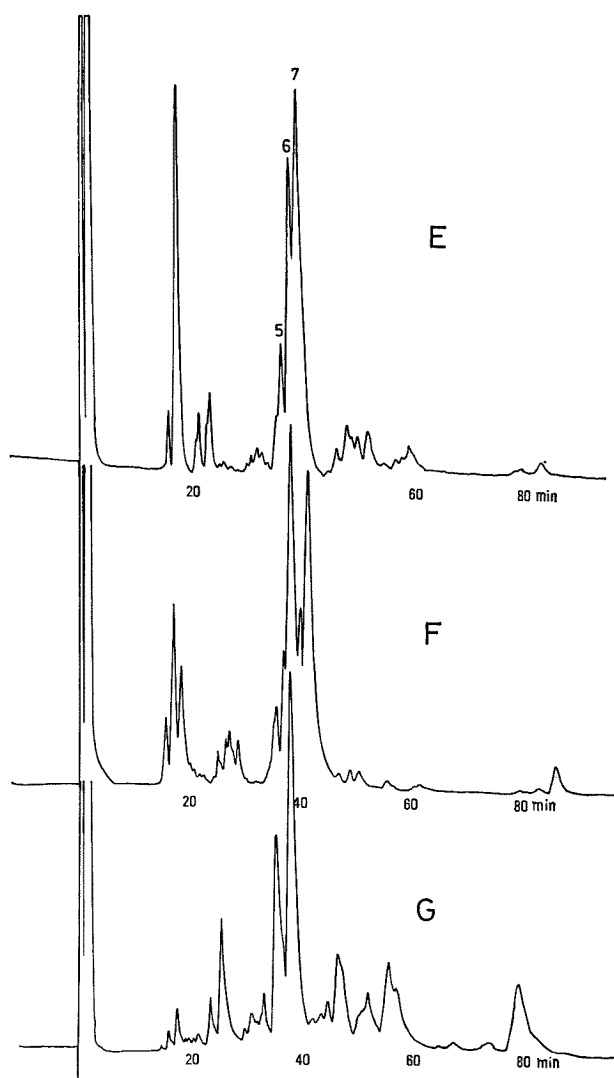


Fig. 3. HPLC chromatograms of big-eyed tuna, Alaska pollack and carp muscle lecithin.
E: Big-eyed tuna. F: Alaska pollack. G: Carp.

also 16:1 in position 1 and 20:5 in position 2 since this peak is composed of two combinations of total acyl carbon numbers of 34 and 36 respectively. However, the latter molecular species, i.e. 16:1-20:5, are considered to be contaminants from the previous peak. From the fatty acid composition and total acyl carbon number, three molecular species were presented in small peak number 4, i.e. combination of 14:0 in position 1 and 20:5 in position 2 for 63.3%; combination of 16:1 in position 1 and 20:5 in position 2 for 5.7%; and combination of 22:6 in position 1 and

16:1 in position 2 for 31.0%. Among these molecular species, 16:1-20:5 can be considered as contaminants from the two previous peaks. In small peak number 5, three molecular species, i.e. 14:0 in position 1 and 20:5 in position 2; 22:6 in position 1 and 14:0 in position 2; and 16:1 in position 1 and 22:6 in position 2 were identified in the same manner as that in small peak number 4. In this case, molecular species of 14:0-20:5 was considered to be a contaminant from the previous peak. In the case of small peak number 6, molecular species of 14:0 in

Table 3. Determination of lecithin molecular species of minor component*¹

Fatty acid	Peak number					
	1	2	3	4	5	6
14:0			21.3	30.6	34.6	44.6
16:1	53.0	47.7	28.3	19.0	19.6	9.7
20:5	47.0	52.3	50.4	36.2	15.6	8.3
22:6				14.2	30.2	37.9
Carbon number* ²						
34			42.6	63.3	32.2	10.8
36	95.0<	90.0	57.4	5.7	41.7	89.2
38		10.0		31.0	26.1	
Carbon number* ²						
34			$\begin{matrix} 20:5 \\ 14:0 \end{matrix}$ 42.6	$\begin{matrix} 14:0 \\ 20:5 \end{matrix}$ 63.3	$\begin{matrix} 14:0 \\ 20:5 \end{matrix}$ 32.2	$\begin{matrix} 14:0 \\ 20:5 \end{matrix}$ 10.8
36	$\begin{matrix} 20:5 \\ 16:1 \end{matrix}$ 95.0<	$\begin{matrix} 16:1 \\ 20:5 \end{matrix}$ 90.0	$\begin{matrix} 16:1 \\ 20:5 \end{matrix}$ 57.4	$\begin{matrix} 16:1 \\ 20:5 \end{matrix}$ 5.7	$\begin{matrix} 22:6 \\ 14:0 \end{matrix}$ 41.7	$\begin{matrix} 14:0 \\ 22:6 \end{matrix}$ 89.2
38				$\begin{matrix} 22:6 \\ 16:1 \end{matrix}$ 31.0	$\begin{matrix} 16:1 \\ 22:6 \end{matrix}$ 26.1	

*¹ % in each peak*² Example of chum salmon (Fall).*² Total acyl carbon number.

position 1 and 20:5 in position 2 were considered to be the contaminants from the previous peak. By analyzing the data in this way, molecular species of all other small peaks were identified.

As illustrated in Fig. 2-D, the HPLC chromatogram of diglyceride acetate of fish muscle lecithin can be divided into four molecular species groups, that is, I: molecular species composed of highly unsaturated fatty acids such as 20:5–22:6, 22:6–22:6 and 20:5–20:5, III: molecular species composed of generally found fatty acids such as 16:0 or 18:1 with combinations of 20:5 or 22:6, that is, 16:0–20:5, 16:0–22:6, 18:1–20:5 and 18:1–22:6, and II and IV: others. In all marine fish examined, groups I and III accounted for more than 70% of the molecular species which were determined in the following manner.

All HPLC chromatograms were traced on section papers, and the area of the identified peaks were measured, though there was some limitation in accuracy caused by the overlapping of the peaks.

The results for fish muscle lecithin are summarized in Table 4. Fish from marine sources contain a significant amount of highly unsaturated fatty acid combinations such as: 22:6–22:6, 22:6–20:5 and 20:5–20:5. All fish lecithin examined contain 16:0–22:6 as the predominant molecular species.

The RRTs of all peaks were determined by dividing the retention time of each peak by the retention time of 16:0–22:6.*¹ In the case of soybean lecithin, 16:0–18:2*² was used as a reference peak and the RRTs were recalculated against 16:0–22:6*¹ by using the RRTs data in egg yolk lecithin. The RRT of each molecular species is summarized in Tables 5–8.

Discussion

We have plotted the RRT of each molecular species semilogarithmically against the partition number (PN) defined as $PN = C^{*3} - 2d^{*4}$ since PN is proportional to the logarithm of RRT.⁽¹⁾ A general expression for this can be written as:

$$PN \propto \log(RRT) \quad (1)$$

After plotting the RRT of each molecular species semilogarithmically, a similar correlation was obtained, that is:

$$PN \propto \log(RRT) + \Delta Q \quad (2)$$

(2) is similar to (1), and if we formulate (2) into an equation

$$PN = P \cdot \log(RRT) + \Delta Q \quad (3)$$

where P becomes the slope of the oblique line, and ΔQ becomes the intersection on the ordinate.

*¹ 16:0 in position 1 and 22:6 in position 2*² 16:0 in position 1 and 18:2 in position 2*³ C: Total acyl carbon number, *⁴ d: Total double bonds.

Table 4. Molecular species of fish muscle lecithin

Chum salmon (Summer)	Chum salmon (Fall)	Big-eyed tuna	Alaska pollack	Carp
$\begin{array}{ c } \hline 16:0 \\ \hline 22:6 \\ \hline \end{array}$ 41.2%	$\begin{array}{ c } \hline 16:0 \\ \hline 22:6 \\ \hline \end{array}$ 18.8%	$\begin{array}{ c } \hline 16:0 \\ \hline 22:6 \\ \hline \end{array}$ 30.3%	$\begin{array}{ c } \hline 16:0 \\ \hline 22:6 \\ \hline \end{array}$ 23.9%	$\begin{array}{ c } \hline 16:0 \\ \hline 22:6 \\ \hline \end{array}$ 24.1%
$\Delta \begin{array}{ c } \hline 16:0 \\ \hline 20:5 \\ \hline \end{array}$ 14.7%	$\begin{array}{ c } \hline 22:6 \\ \hline 22:6 \\ \hline \end{array}$ 10.9%	$\begin{array}{ c } \hline 22:6 \\ \hline 22:6 \\ \hline \end{array}$ 16.5%	$\Delta \begin{array}{ c } \hline 16:0 \\ \hline 20:5 \\ \hline \end{array}$ 24.5%	$\Delta \begin{array}{ c } \hline 16:0 \\ \hline 20:5 \\ \hline \end{array}$ 12.8%
$\begin{array}{ c } \hline 18:1 \\ \hline 22:6 \\ \hline \end{array}$	$\begin{array}{ c } \hline 20:5 \\ \hline 22:6 \\ \hline \end{array}$ 10.4%	$\begin{array}{ c } \hline 22:6 \\ \hline 16:0 \\ \hline \end{array}$ 9.3%	$\begin{array}{ c } \hline 18:1 \\ \hline 22:6 \\ \hline \end{array}$	$\begin{array}{ c } \hline 18:1 \\ \hline 22:6 \\ \hline \end{array}$
$\begin{array}{ c } \hline 22:6 \\ \hline 22:6 \\ \hline \end{array}$ 6.1%	$\begin{array}{ c } \hline 16:0 \\ \hline 20:5 \\ \hline \end{array}$ 18.3%	$\begin{array}{ c } \hline 18:1 \\ \hline 22:6 \\ \hline \end{array}$ 5.4%	$\begin{array}{ c } \hline 22:6 \\ \hline 16:0 \\ \hline \end{array}$ 8.6%	$\begin{array}{ c } \hline 16:0 \\ \hline 18:1 \\ \hline \end{array}$ 9.0%
$\begin{array}{ c } \hline 20:5 \\ \hline 22:6 \\ \hline \end{array}$ 5.1%	$\begin{array}{ c } \hline 18:1 \\ \hline 22:6 \\ \hline \end{array}$	$\Delta \begin{array}{ c } \hline 16:0 \\ \hline 20:5 \\ \hline \end{array}$	$\begin{array}{ c } \hline 20:5 \\ \hline 22:6 \\ \hline \end{array}$ 6.8%	$\begin{array}{ c } \hline 16:0 \\ \hline 20:4 \\ \hline \end{array}$ 8.6%
$\begin{array}{ c } \hline 22:6 \\ \hline 16:0 \\ \hline \end{array}$ 4.6%	$\begin{array}{ c } \hline 22:6 \\ \hline 16:0 \\ \hline \end{array}$ 8.7%	$\begin{array}{ c } \hline 22:5 \\ \hline 22:6 \\ \hline \end{array}$ 4.6%	$\begin{array}{ c } \hline 18:1 \\ \hline 20:5 \\ \hline \end{array}$ 6.3%	$\begin{array}{ c } \hline 16:0 \\ \hline 18:2 \\ \hline \end{array}$
$\begin{array}{ c } \hline 16:0 \\ \hline 18:1 \\ \hline \end{array}$ 4.1%	$\begin{array}{ c } \hline 20:5 \\ \hline 16:0 \\ \hline \end{array}$ 5.5%	$\begin{array}{ c } \hline 20:5 \\ \hline 22:6 \\ \hline \end{array}$ 2.7%	$\begin{array}{ c } \hline 22:6 \\ \hline 22:6 \\ \hline \end{array}$ 3.9%	$\begin{array}{ c } \hline 16:0 \\ \hline 16:1 \\ \hline \end{array}$ 13.0%
$\begin{array}{ c } \hline 14:0 \\ \hline 22:6 \\ \hline \end{array}$ 2.2%	$\begin{array}{ c } \hline 20:5 \\ \hline 20:5 \\ \hline \end{array}$ 3.5%	$\begin{array}{ c } \hline 16:0 \\ \hline 20:4 \\ \hline \end{array}$ 4.2%	$\begin{array}{ c } \hline 16:0 \\ \hline 18:1 \\ \hline \end{array}$ 4.5%	$\begin{array}{ c } \hline 18:0 \\ \hline 22:6 \\ \hline \end{array}$
$\begin{array}{ c } \hline 16:0 \\ \hline 22:5 \\ \hline \end{array}$ 1.7%	$\begin{array}{ c } \hline 14:0 \\ \hline 20:5 \\ \hline \end{array}$ 3.1%	$\begin{array}{ c } \hline 22:5 \\ \hline 16:0 \\ \hline \end{array}$	$\begin{array}{ c } \hline 14:0 \\ \hline 20:1 \\ \hline \end{array}$	$\begin{array}{ c } \hline 18:1 \\ \hline 20:5 \\ \hline \end{array}$ 2.9%
$\begin{array}{ c } \hline 20:5 \\ \hline 18:1 \\ \hline \end{array}$ 1.5%	$\begin{array}{ c } \hline 22:6 \\ \hline 16:1 \\ \hline \end{array}$	$\begin{array}{ c } \hline 16:0 \\ \hline 22:5 \\ \hline \end{array}$ 2.1%	$\Delta \begin{array}{ c } \hline 20:5 \\ \hline 16:0 \\ \hline \end{array}$ 4.3%	$\begin{array}{ c } \hline 16:0 \\ \hline 22:5 \\ \hline \end{array}$ 2.4%
$\begin{array}{ c } \hline 20:5 \\ \hline 20:5 \\ \hline \end{array}$ 1.1%	$\begin{array}{ c } \hline 14:0 \\ \hline 22:6 \\ \hline \end{array}$ 2.9%	$\begin{array}{ c } \hline 20:4 \\ \hline 22:6 \\ \hline \end{array}$ 3.7%	$\begin{array}{ c } \hline 22:6 \\ \hline 18:1 \\ \hline \end{array}$	$\begin{array}{ c } \hline 18:1 \\ \hline 20:4 \\ \hline \end{array}$ 1.9%
$\begin{array}{ c } \hline 20:4 \\ \hline 22:6 \\ \hline \end{array}$	$\begin{array}{ c } \hline 18:1 \\ \hline 20:5 \\ \hline \end{array}$ 2.6%	$\begin{array}{ c } \hline 20:5 \\ \hline 22:5 \\ \hline \end{array}$	$\begin{array}{ c } \hline 14:0 \\ \hline 22:6 \\ \hline \end{array}$ 2.0%	$\begin{array}{ c } \hline 16:1 \\ \hline 22:6 \\ \hline \end{array}$
$\begin{array}{ c } \hline 20:5 \\ \hline 22:5 \\ \hline \end{array}$	$\begin{array}{ c } \hline 16:1 \\ \hline 22:5 \\ \hline \end{array}$	$\begin{array}{ c } \hline 22:6 \\ \hline 18:0 \\ \hline \end{array}$ 1.1%	$\begin{array}{ c } \hline 20:5 \\ \hline 20:5 \\ \hline \end{array}$ 1.9%	$\begin{array}{ c } \hline 18:2 \\ \hline 22:6 \\ \hline \end{array}$ 6.5%
$\begin{array}{ c } \hline 22:5 \\ \hline 22:6 \\ \hline \end{array}$	$\begin{array}{ c } \hline 15:0 \\ \hline 20:5 \\ \hline \end{array}$ 3.8%	$\begin{array}{ c } \hline 18:1 \\ \hline 16:0 \\ \hline \end{array}$ 1.7%	$\begin{array}{ c } \hline 16:1 \\ \hline 22:6 \\ \hline \end{array}$ 2.5%	$\begin{array}{ c } \hline 14:0 \\ \hline 22:6 \\ \hline \end{array}$
$\begin{array}{ c } \hline 16:1 \\ \hline 20:5 \\ \hline \end{array}$ 4.1%	$\begin{array}{ c } \hline 17:1 \\ \hline 22:6 \\ \hline \end{array}$	$\begin{array}{ c } \hline 16:0 \\ \hline 18:1 \\ \hline \end{array}$	$\begin{array}{ c } \hline 18:2 \\ \hline 22:6 \\ \hline \end{array}$	$\begin{array}{ c } \hline 16:1 \\ \hline 20:5 \\ \hline \end{array}$ 2.2%
$\begin{array}{ c } \hline 14:0 \\ \hline 20:5 \\ \hline \end{array}$	$\begin{array}{ c } \hline 20:5 \\ \hline 18:1 \\ \hline \end{array}$	$\begin{array}{ c } \hline 20:4 \\ \hline 16:0 \\ \hline \end{array}$	$\begin{array}{ c } \hline 14:0 \\ \hline 20:5 \\ \hline \end{array}$ 1.3%	$\begin{array}{ c } \hline 18:2 \\ \hline 20:5 \\ \hline \end{array}$
$\begin{array}{ c } \hline 16:1 \\ \hline 22:6 \\ \hline \end{array}$	$\begin{array}{ c } \hline 20:5 \\ \hline 14:0 \\ \hline \end{array}$ 0.9%	$\begin{array}{ c } \hline 17:0 \\ \hline 22:6 \\ \hline \end{array}$ 1.7%	$\begin{array}{ c } \hline 16:1 \\ \hline 20:5 \\ \hline \end{array}$ 1.0%	$\begin{array}{ c } \hline 16:0 \\ \hline 16:2 \\ \hline \end{array}$ 1.9%
$\begin{array}{ c } \hline 22:6 \\ \hline 14:0 \\ \hline \end{array}$	$\begin{array}{ c } \hline 16:1 \\ \hline 22:6 \\ \hline \end{array}$ 0.9%	$\begin{array}{ c } \hline 18:1 \\ \hline 22:5 \\ \hline \end{array}$	$\begin{array}{ c } \hline 18:2 \\ \hline 20:5 \\ \hline \end{array}$	$\begin{array}{ c } \hline 18:1 \\ \hline 22:5 \\ \hline \end{array}$
Others 13.6%	$\begin{array}{ c } \hline 22:6 \\ \hline 14:0 \\ \hline \end{array}$	Others 16.7%	Others 8.5%	Others 14.7%
	Others 9.7%			

Δ Major component in each peak.

This equation (3) suggests that PN is not only a function of RRT alone, but also a function including ΔQ .

Back to the definition of the PN, that is, $PN = C - 2d$, the left member in (3) can be changed as:

$$C - 2d = P \cdot \log(RRT) + \Delta Q \quad (4)$$

So RRT can be expressed as follows:

$$RRT = 10^{\frac{C - 2d - \Delta Q}{P}} \quad (5)$$

If we put ΔQ as $Q_1 + \alpha$, equation (5) may be written as:

$$RRT = 10^{\frac{C - 2d - (Q_1 + \alpha)}{P}} \quad (Q_1 \text{ is a minimum of } \Delta Q) \quad (6)$$

where P and Q are invariables and α is a variable

Table 5. Relation between relative retention time of HPLC and molecular species of soybean lecithin

PN	MS	RT	RRT*
26	$\begin{smallmatrix} 18:2 \\ 18:3 \end{smallmatrix}$	27.6	70.4
28	$\begin{smallmatrix} 18:2 \\ 18:2 \end{smallmatrix}$	37.6	95.8
"	$\begin{smallmatrix} 18:1 \\ 18:3 \end{smallmatrix}$	39.2	99.9
"	$\begin{smallmatrix} 16:0 \\ 18:3 \end{smallmatrix}$	42.0	107.1
30	$\begin{smallmatrix} 18:1 \\ 18:2 \end{smallmatrix}$	51.4	131.0
"	$\begin{smallmatrix} 16:0 \\ 18:2 \end{smallmatrix}$	55.2	140.7
"	$\begin{smallmatrix} 18:0 \\ 18:3 \end{smallmatrix}$	60.0	152.9
31	$\begin{smallmatrix} 17:0 \\ 18:2 \end{smallmatrix}$	62.8	160.1
32	$\begin{smallmatrix} 18:1 \\ 18:1 \end{smallmatrix}$	73.8	188.1
"	$\begin{smallmatrix} 20:1 \\ 18:2 \end{smallmatrix}$	"	"
"	$\begin{smallmatrix} 16:0 \\ 18:1 \end{smallmatrix}$	79.6	202.9
"	$\begin{smallmatrix} 18:0 \\ 18:2 \end{smallmatrix}$	"	"
"	$\begin{smallmatrix} 16:0 \\ 16:0 \end{smallmatrix}$	83.6	213.0
34	$\begin{smallmatrix} 18:0 \\ 18:1 \end{smallmatrix}$	113.2	288.6
"	$\begin{smallmatrix} 18:2 \\ 20:0 \end{smallmatrix}$	"	"

Abbreviations:

PN: Partition number, MS: Molecular species, RRT: Relative retention time.

* $\begin{smallmatrix} 16:0 \\ 22:6 \end{smallmatrix}$ is used as the reference peak.

range of Q_1 . By letting C and d be invariables, RRT becomes a function of α which is:

$$RRT = f(\alpha) \quad (7)$$

Back to equation (4), by putting dQ_1 as $Q_1 + \alpha$, the following equation can be obtained.

$$C - 2d = P \cdot \log(RRT) + Q_1 + \alpha \quad (4')$$

If we let d be invariable and rearrange the expression for C, (4') will become:

$$C = P \cdot \log(RRT) + \alpha + Q_2 \quad (Q_2 = Q_1 + 2d) \quad (8)$$

On the other hand, if we let C be invariable and rearrange the expression for d, (4') will become:

$$d = -\frac{P}{2} \cdot \log(RRT) - \frac{\alpha}{2} + Q_3 \quad (9)$$

$$\left(Q_3 = -\frac{1}{2}(Q_1 - C) \right)$$

These two functions, i.e. (8) and (9) have a deviation which are α for the former and $-\alpha/2$ for the

Table 6. Relation between relative retention time of HPLC and molecular species of egg yolk lecithin

PN	MS	RT	RRT
28	$\begin{smallmatrix} 16:1 \\ 18:2 \end{smallmatrix}$	41.2	100.0
"	$\begin{smallmatrix} 18:2 \\ 18:2 \end{smallmatrix}$	"	"
"	$\begin{smallmatrix} 14:0 \\ 18:2 \end{smallmatrix}$	"	"
26	$\begin{smallmatrix} 16:0 \\ 22:6 \end{smallmatrix}$	"	"
28	$\begin{smallmatrix} 16:0 \\ 18:3 \end{smallmatrix}$	47.0	114.1
"	$\begin{smallmatrix} 18:1 \\ 20:4 \end{smallmatrix}$	"	"
"	$\begin{smallmatrix} 16:0 \\ 20:4 \end{smallmatrix}$	50.4	122.2
30	$\begin{smallmatrix} 16:0 \\ 22:4 \end{smallmatrix}$	54.4	132.0
"	$\begin{smallmatrix} 16:1 \\ 18:1 \end{smallmatrix}$	55.6	134.9
"	$\begin{smallmatrix} 18:1 \\ 18:2 \end{smallmatrix}$	"	"
"	$\begin{smallmatrix} 16:0 \\ 18:2 \end{smallmatrix}$	59.2	143.7
28	$\begin{smallmatrix} 18:0 \\ 22:6 \end{smallmatrix}$	"	"
31	$\begin{smallmatrix} 15:0 \\ 18:1 \end{smallmatrix}$	72.4	175.7
"	$\begin{smallmatrix} 16:0 \\ 17:1 \end{smallmatrix}$	"	"
"	$\begin{smallmatrix} 17:0 \\ 18:2 \end{smallmatrix}$	"	"
30	$\begin{smallmatrix} 18:0 \\ 20:4 \end{smallmatrix}$	"	"
32	$\begin{smallmatrix} 18:1 \\ 18:1 \end{smallmatrix}$	79.0	191.7
"	$\begin{smallmatrix} 18:0 \\ 22:4 \end{smallmatrix}$	"	"
"	$\begin{smallmatrix} 16:0 \\ 18:1 \end{smallmatrix}$	83.6	202.9
"	$\begin{smallmatrix} 18:0 \\ 18:2 \end{smallmatrix}$	"	"
34	$\begin{smallmatrix} 18:0 \\ 18:1 \end{smallmatrix}$	121.8	295.1

Abbreviations:

PN: Partition number, MS: Molecular species, RRT: Relative retention time.

latter. Alfa is considered to be a factor that has a slight effect on RRT, and may be due to the positional isomers such as between 1 and 2 position in the molecule, or by the large differences in number of double bonds between the two fatty acids in the molecule. This leads us to the concept of a matrix since it is convenient to distinguish the positional isomers or the bias in the number of double bonds between the two fatty acids in the molecule. So in order to simplify the equation, we can induce the following equations from (8) and (9) respectively under the conditions of:

Table 7. Relation between relative retention time of HPLC and main molecular species of fish muscle lecithin

Chum salmon (Summer)		Chum salmon (Fall)		Big-eyed tuna		Alaska pollack		Carp	
MS	PN/RRT	MS	PN/RRT	MS	PN/RRT	MS	PN/RRT	MS	PN/RRT
20:5]	20	20:5]	20	20:5]	20	20:5]	20	20:5]	20
20:5]	27.3	20:5]	37.5	22:6]	41.2	20:5]	37.2	22:6]	42.5
20:5]	20	20:5]	20	22:6]	20	20:5]	20	22:6]	20
22:6]	40.7	22:6]	40.4	22:6]	44.6	22:6]	40.4	22:6]	46.2
22:6]	20	22:6]	20	20:4]	22	22:6]	20	16:1]	24
22:6]	44.2	22:6]	44.1	22:6]	55.2	22:6]	44.3	20:5]	62.5
14:0]	24	18:1]	26	22:5]	22	18:1]	26	18:2]	24
22:6]	69.7	20:5]	87.0	20:5]	55.2	20:5]	86.7	20:5]	62.5
16:0]	26	20:5]	26	22:5]	22	20:5]	26	16:1]	24
20:5]	92.2	16:0]	89.5	22:6]	60.0	16:0]	89.8	22:6]	67.7
18:1]	26	16:0]	26	18:1]	26	22:6]	26	18:2]	24
22:6]	92.2	20:5]	91.9	22:6]	93.5	18:1]	89.8	22:6]	67.7
22:6]	26	18:1]	26	16:0]	26	16:0]	26	14:0]	24
16:0]	97.1	22:6]	91.9	20:5]	93.5	20:5]	92.0	22:6]	67.7
16:0]	26	22:6]	26	22:6]	26	18:1]	26	18:1]	26
22:6]	100.0	16:0]	96.7	16:0]	96.8	22:6]	92.0	20:5]	88.3
		16:0]	26	16:0]	26	22:6]	26	16:0]	26
		22:6]	100.0	22:6]	100.0	16:0]	97.1	20:5]	93.7
				16:0]	28	16:0]	26	18:1]	26
				22:5]	134.9	22:6]	100.0	22:6]	93.7
						16:0]	32	16:0]	26
						18:1]	216.6	22:6]	100.0
						14:0]	32	16:0]	28
						20:1]	216.6	20:4]	124.0

Abbreviations:

PN: Partition number, MS: Molecular species, RRT: Relative retention time.

$$C' = \begin{vmatrix} x & d_1 \\ c_2 & d_2 \end{vmatrix} \quad (c_2, d_1 \text{ and } d_2 \text{ are invariables})$$

$$d' = \begin{vmatrix} c_1 & y \\ c_2 & d_2 \end{vmatrix} \quad (c_1, c_2 \text{ and } d_2 \text{ are invariables})$$

$$C' = p_1 \cdot \log(RRT) + q_1 \quad (8')$$

$$d' = p_2 \cdot \log(RRT) + q_2 \quad (9')$$

by analyzing the RRT data of each molecular species.

After plotting the RRT of each molecular species from the source examined against the total acyl carbon number or the number of total double bonds of each molecular species, the RRT plots of molecular species laid almost on a straight line by giving a variable integer x for the carbon number and a variable integer y for the number of double bonds of each fatty acid in the molecular species, when we express the molecular species in matrix relation. The oblique lines were parallel to each other as shown in Fig. 4. By applying these correlations between the RRT and the corresponding molecular species under the condition of matrix for the determination of molecular species, it was

suggested that an unidentified molecular species of lecithins could be identified from RRT on HPLC even if it has the most complicated composition of molecular species such as fish muscle lecithin.

While this manuscript was in preparation, another method for HPLC analysis of phospholipid molecular species was published by PATTON *et al.*¹²⁾ To our surprise, the sequence of each molecular species in elution on HPLC, when the RRTs were plotted against the total acyl carbon number from our biological material, were the same as common molecular species of rat liver which was analyzed by PATTON *et al.*¹²⁾ despite the fact that the analytical conditions were significantly different. This suggests that, although there are differences in retention time or in RRT among the different conditions on HPLC, the sequence in elution of each molecular species might be unchangeable. This leads us to a conclusion that in HPLC, the sequence in elution might be controlled by a fixed correlation, that is matrix relation. By accepting this idea, we can expand this matrix concept to triglycerides. Details will be reported at a later date.

Table 8. Relation between relative retention time of HPLC and molecular species of fish muscle lecithin

Chum salmon (Summer)		Chum salmon (Fall)		Big-eyed tuna		Alaska pollack		Carp	
MS	PN/RRT	MS	PN/RRT	MS	PN/RRT	MS	PN/RRT	MS	PN/RRT
20: 4	22	18: 3	22	22: 6	24	16: 1	24	18: 3	22
22: 6	50.6	22: 6	48.7	16: 1	65.0	20: 5	61.0	22: 6	50.7
22: 5	22	20: 4	22	18: 2	24	18: 2	24	20: 4	22
20: 5	50.6	22: 6	48.7	22: 6	66.9	20: 5	61.0	20: 5	52.3
22: 5	22	22: 5	22	14: 0	24	14: 0	24	22: 5	22
22: 6	52.8	20: 5	48.7	22: 6	71.1	20: 5	64.2	22: 6	54.5
16: 1	24	20: 5	24	22: 4	24	16: 1	24	20: 4	22
20: 5	60.8	16: 1	59.1	22: 6	71.1	22: 6	65.9	22: 6	56.4
14: 0	24	20: 5	24	16: 1	26	18: 2	24	16: 1	26
20: 5	64.8	18: 2	59.1	20: 4	71.1	22: 6	65.9	22: 5	79.4
16: 1	24	16: 1	24	20: 4	24	14: 0	24	18: 2	26
22: 6	65.8	20: 5	60.8	22: 5	74.0	22: 6	69.9	22: 5	79.4
22: 6	24	18: 2	24	22: 6	25	22: 5	28	15: 0	25
14: 0	67.4	20: 5	60.8	17: 1	78.4	16: 0	115.7	22: 6	79.4
15: 0	25	20: 5	24	22: 5	24	16: 0	28	17: 1	25
20: 5	77.8	14: 0	62.2	22: 5	78.4	20: 4	115.7	22: 6	79.4
17: 1	25	14: 0	24	17: 2	25	16: 0	28	16: 1	26
22: 6	82.3	20: 5	64.2	20: 4	78.4	22: 5	120.3	20: 4	82.7
14: 0	26	22: 6	24	17: 2	25	20: 1	28	18: 2	26
22: 5	82.3	16: 1	64.2	22: 5	78.5	20: 5	125.0	20: 4	82.7
15: 0	25	16: 1	24	17: 1	25	18: 0	28	18: 1	28
22: 6	84.4	22: 6	67.1	22: 6	82.7	20: 5	137.5	22: 5	111.7
20: 5	26	22: 6	24	15: 0	25	20: 1	28	16: 0	28
18: 1	87.2	14: 0	67.1	22: 6	84.7	22: 6	137.5	16: 2	111.7
18: 1	26	14: 0	24	20: 5	26	18: 1	30	18: 1	28
20: 5	90.2	22: 6	69.5	18: 1	84.7	16: 1	137.5	20: 4	115.8
20: 5	26	16: 1	26	18: 1	26	18: 1	30	16: 0	28
16: 0	90.2	22: 5	77.8	20: 5	88.2	18: 2	137.5	22: 5	119.3
22: 6	26	15: 0	25	22: 6	26	14: 0	30	16: 0	30
18: 1	90.2	20: 5	77.8	18: 1	91.3	18: 1	150.0	22: 4	133.8
16: 0	28	17: 1	25	20: 5	26	16: 0	30	20: 1	28
22: 5	121.9	22: 6	77.8	16: 0	91.3	16: 1	150.0	22: 6	133.8
16: 0	28	20: 5	26	20: 4	28	16: 0	30	18: 1	30
20: 4	126.6	18: 1	82.2	16: 0	116.7	18: 2	150.0	16: 1	138.4
17: 0	27	22: 5	28	17: 0	27	18: 0	28	18: 1	30
22: 6	126.6	16: 0	116.6	22: 6	116.7	22: 6	150.0	18: 2	138.4
20: 1	28	16: 0	28	18: 1	28	18: 1	32	16: 0	30
20: 5	126.6	22: 5	119.9	22: 5	116.7	16: 0	207.4	18: 2	148.4
20: 1	28	16: 0	28	16: 0	28	20: 1	32	16: 0	30
22: 6	140.6	20: 4	124.6	20: 4	124.6	14: 0	207.4	16: 1	148.4
18: 0	28	20: 1	28	22: 5	28	16: 0	34	18: 0	28
20: 5	140.6	20: 5	124.6	16: 0	130.0	20: 1	301.9	22: 6	148.4
18: 1	30	18: 0	28	22: 6	28	18: 0	34	18: 0	30
16: 1	143.6	20: 5	133.3	18: 0	143.2	18: 1	301.9	20: 4	179.4
18: 0	28	18: 1	30	18: 0	28			18: 1	32
22: 6	152.5	16: 1	137.2	22: 6	148.1			18: 1	195.3
14: 0	30	14: 0	30	18: 1	32			16: 0	32
18: 1	152.5	18: 1	142.9	18: 1	189.6			18: 1	210.1
16: 0	30	16: 0	30	18: 1	32				
16: 1	152.5	16: 1	142.9	16: 0	206.1				
24: 1	32			16: 0	32				
20: 5	265.7			18: 1	215.5				

Abbreviations:

PN: Partition number, MS: Molecular species, RRT: Relative retention time.

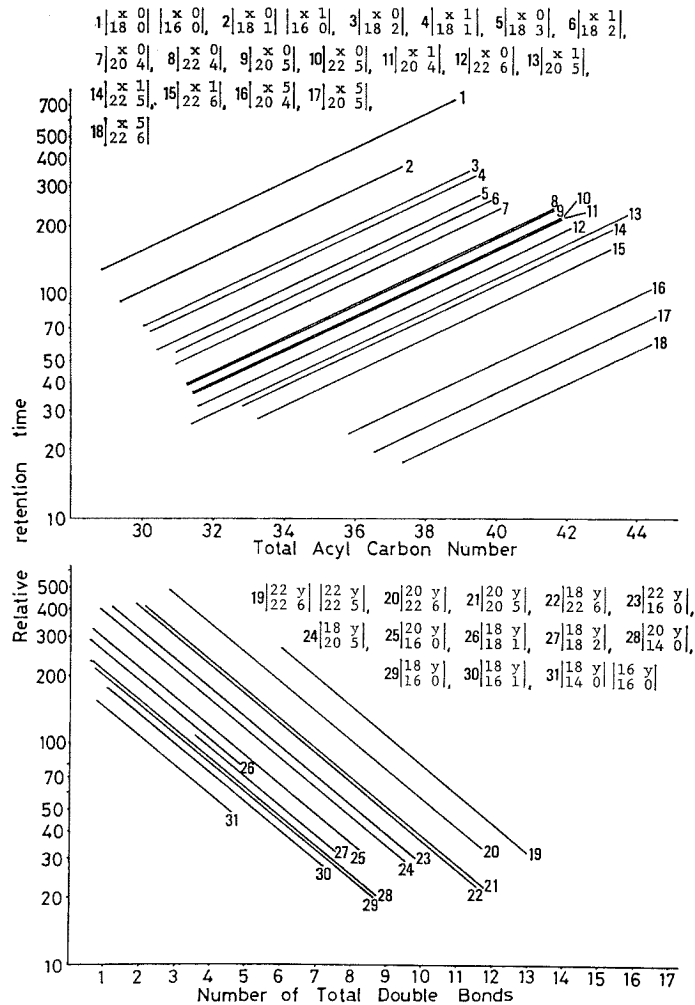


Fig. 4. Relation between relative retention time and total acyl carbon number and relation between relative retention time and total double bonds on HPLC of lecithin.

Acknowledgements

The authors wish to thank J. CHIGIRA of Asahi Chemical Industry Ltd. for the supply of egg yolk lecithin; and W. NAGATA for reading the manuscript.

References

- 1) K. TAKAMA: in "Quality of Fish" (ed. by Japan. Soc. Sci. Fish.), Suisangaku Series No. 4, Koseisha-Koseikaku, Tokyo, 1974, pp. 138-144.
- 2) W. E. M. LANDS and P. HART: *J. Am. Oil Chem. Soc.*, **43**, 290-295 (1966).
- 3) O. RENKONEN: *J. Am. Oil Chem. Soc.*, **42**, 298-304 (1965).
- 4) A. KUKSIS and L. MARAI: *Lipids*, **2**, 217-224 (1967).
- 5) A. F. PREVOT and F. X. MORDRET: *Rev. Fse. Corps Gras*, **23**, 409-423 (1976).
- 6) K. ZAMA, T. MARUYAMA, and K. TAKAHASHI: *Bull. Fac. Fish. Hokkaido Univ.*, **27**, 181-190 (1976).
- 7) K. TAKAHASHI, K. ZAMA, and T. MATSUOKA: *Bull. Fac. Fish. Hokkaido Univ.*, **29**, 378-385 (1978).
- 8) K. TAKAHASHI, F. CABLING, and K. ZAMA: *Bull. Fac. Fish. Hokkaido Univ.*, **29**, 386-391 (1978).
- 9) A. KUKSIS: in "Progress in the Chemistry of Fats

- and Other Lipids" (ed. by R. T. HOLMAN), Vol. 12, Pergamon Press, Oxford, New York, Toronto, 1972, pp. 105-111.
- 10) R. WOOD and F. SNYDER: *Arch. Biochem. Biophys.*, **131**, 478-494 (1969).
- 11) S. WADA, C. KOIZUMI, A. TAKIGUCHI, and J. NONAKA: *Yukagaku*, **27**, 579-584 (1978).
- 12) G. M. PATTON, J. M. FASULO, and S. J. ROBINS: *J. Lipid Res.*, **23**, 190-196 (1982).