# The Fatty Acids of Antarctic Phytoplankton and Euphausiids. Fatty Acid Exchange among Trophic Levels of the Ross Sea

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#### Abstract

The fatty acids of 3 samples of Euphausia superba, 7 samples of E. crystallorophias, and 12 samples of phytoplankton collected in the Ross Sea, Antarctica, during Eltanin Cruise 51 were examined. The fatty acid profiles of the samples of E. superba resembled each other closely. The fatty acid profiles of the E. crystallorophias samples were also similar to each other but different quantitatively from those of E. superba. Phytoplankton fatty acid patterns varied with the geographical location and species composition of the samples. The fatty acids of euphausiids were compared to those of the phytoplankton from the corresponding locations. Rather similar fatty acid patterns in phytoplankton and E. superba corroborate the herbivorous nature of this euphausiid. On the other hand, phytoplankton and E. crystallorophias showed quite different fatty acid patterns. The differences were mostly due to the presence of waxes among the lipids of E. crystallorophias. It is not clear whether these waxes are of dietary origin or are synthesized endogenously.

### Introduction

The study of krill has become intensive in recent times, perhaps as a result of its potential importance as food. A variety of organisms is usually included under that generic name, but in the Southern Oceans the name Euphausia superba has been considered almost a synonym for krill. However, due to a quite defined geographical distribution, there are certain areas in which E. superba is replaced by other members of the same genus. For example, E. superba is very seldom found in shallow waters close to ice, such as the Ross Ice Shelf, but the smaller E. crystallorophias predominates in such areas (Marr, 1962; Mauchline and Fisher, 1969).

The chemical composition of Euphausia superba is reasonably well known. Its fatty acids, in particular, have been the subject of several studies in the past few years (Nonaka and Koizumi, 1964; Tsuyuki et al., 1964a, b; Hansen, 1969; Pierce et al., 1969; Hansen and Meiklen, 1970; Sidhu et al., 1970; Van der Veen et al., 1971). On the other hand, very little is known about the lipids of E

crystallorophias, except that they increase in the late austral summer and decrease gradually during the winter (Littlepage, 1964). The present report describes studies on the fatty acids of E. superba and E. crystallorophias from various locations in the Ross Sea. The fatty acid patterns of the two euphausiids are compared to each other as well as to the fatty acids of phytoplankton from corresponding locations. An attempt is made to determine the flow of fatty acids through trophic levels.

#### Materials and Methods

Phytoplankton samples were collected with a 35  $\mu$ net, by vertical hauls to a depth of 200 m. After microscopic examination, samples containing less than 80% phytoplankton were discarded. Microzooplankton constituted the major contaminant. Euphausiids were collected with a 1 m mid-water trawl, at depths varying between 0 and 300 m. Shortly after being sorted by hand the samples were extracted for lipids with a chloroform:methanol (2:1, v/v) mixture (Folch et  $\alpha l$ ., 1957). Quantities of 5 to 10 mg total lipids were converted into fatty acid methyl esters by saponification, followed by reflux with methanol in the presence of boron trifluoride (American Oil Chemists' Society, 1970). The fatty acid methyl esters were studied by gas-liquid chromatography on a 6' 1/8" column of siliconized polyethylene glycol succinate (DGSS-X, Applied Science Co., State College, Pennsylvania) 10% w/w on Chromosorb (Johns-Manville, Denver, Colorado) at 170°C. A dual-flame model GC-5 Beckman gas chromatograph (Fullerton, California) was used connected to an Infotronics (Columbia Scientific Industries, Austin, Texas) digital integrator. Results are expressed as weight percent. Fatty acid methyl esters were identified by co-chromatography with known standards and by plotting relative retention times versus chain length before, and in most cases, after hydrogenation.

Results and Discussion

Fatty Acids of Euphausia superba

Samples of Euphausia superba were collected from Stations 8 9 and 11 of Eltania Cruico 51 (Fig. 1)



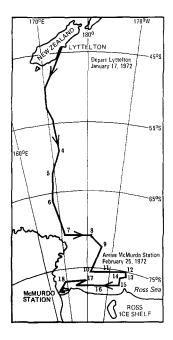


Fig. 1. Ross Sea, Antarctica: track of Eltanin Cruise 51, showing stations

These stations are located in an area in which two currents of water mix (Marr, 1962), thus providing the turbulent environment that *E. superba* seems to prefer for a habitat (Ivanov, 1970).

There is a remarkable similarity in the fatty acid compositions of the samples collected from the three stations (Table 1). In addition, these patterns resemble quite closely those reported by Hansen and Meiklen (1970) and by Sidhu et al. (1970). The present results, however, differ somewhat from those of Nonaka and Koizumi (1964) and of Van der Veen et al. (1971). Of all samples of Euphausia superba studied so far, only those of the present study were extracted immediately after capture. The others were frozen, transported and then extracted. This might explain some of the differences.

In two groups of krill (Stations 8 and 11), the hepatopancreas and stomach were excised and their fatty acids were studied separately from those of the whole animal. In one case (Station 11), the fatty acids of the remaining carcass were also studied. Since the stomachs were empty in all cases, the values in Table 1 under the heading hepatopancreas and stomach correspond essentially to hepatopancreas lipids only. The remarkable similarity among organ, remaining carcass, and whole

Table 1. Euphausia superba. Fatty acids (as weight per cent of total acids)

Fatty acid <sup>a</sup>	Station 8		Station 9	Station 11					
	Whole krill	HP+Sb	Whole krill	Whole krill	HP+S	Remaining carcass			
14:0	14.9	10.7	12.9	14.3	12.9	13.5			
16:0	21.2	21.2	20.9	24.7	22.3	23.4			
18:0	0.7	1.2	0.9	1.4	1.3	1.4			
16:1(n-7)	9.0	6.7	10.7	8.9	8.2	8.0			
18:1(n-9)	18.2	17.1	22.8	21.7	21.8	21.5			
20:1(n-9)	0.6	0.9	1.1	0.9	1.2	1.1			
18:2(n-3)	2.6	2.5	2.7	2.0	2.1	1.9			
18:3(n-3)	1.1	1.2	1.4	1.0	1.0	1.1			
18:4(n-3)	2.2	1.9	2.6	3.3	3.6	3.8			
20:5(n-3)	16.0	22.2	11.8	11.4	13.9	11.6			
22:6(n-3)	8.6	9.4	8.3	7.3	8.1	9.4			
Minor fatty acids <sup>c</sup>	4.9	5.0	3.9	3.1	3.6	3.3			

The number preceding the colon gives the number of carbon atoms in the chain, the number following the colon the number of double bonds; (n-x): number of carbons in the chain minus number of carbons between the methylend and the nearest double bond.

<sup>&</sup>lt;sup>C</sup>Only those fatty acids present at a level of 1% or more are included.



<sup>&</sup>lt;sup>b</sup>Hepatopancreas plus stomach.

body composition suggest that there is very little differentiation of organ lipids in *Euphausia superba*.

### Fatty Acids of Euphausia crystallorophias

In contrast to Euphausia superba, which prefers turbulent waters, E. crystallorophias is usually found in shallow waters in the proximity of the Continental Shelf (Marr, 1962; Mauchline and Fisher, 1969). We know from Littlepage (1964) that E. crystallorophias' lipids decrease in amount at the end of the austral winter and rise in late summer. My own studies (Bottino, in press) show that 20 to 40% of the lipids of E. crystallorophias are waxes, the rest being mostly complex lipids and small amounts of neutral lipids.

The Euphausia crystallorophias collected from Stations 11, and 13 through 17 during Eltanin Cruise 51 show closely similar fatty acid patterns (Table 2). Comparison of the fatty acids of the two euphausiids show (Table 3) that E. crystallorophias contains about twice as much oleic acid (average 44% of total fatty acids) as E. superba (average 21%). Most of this oleic acid comes from the waxes, since about 83% of the wax fatty acids is oleic acid (Bottino, in press). The levels of highly unsaturated fatty acids (HUFA), mostly 20:5(n-3) and 22:6(n-3) are quite similar in both euphausiids.

In conclusion, comparison of the fatty acids of both euphausids shows that Euphausia crystalloro-phias lipids are more unsaturated than E. superba

lipids on account of the larger amount of oleic acid in the former. This different degree of lipid unsaturation might be related to the different environment in which the two euphausiids live. Whereas *E. crystallorophias* dwells near the ice all year around, *E. superba* is probably in contact with the ice only during the winter months (Mackintosh, 1970).

### Phytoplankton Fatty Acids

Phytoplankton samples from Stations 8, 9, 11, 13-15, and 18 were studied. According to microscopic and macroscopic observations 1, the nature of the phytoplankton population changes with the stations, and this is reflected in quantitative variations in the fatty acid patterns (Table 3). Qualitatively, however, the fatty acid patterns showed some common characteristics: (1) The presence in most samples of significant amounts (up to 15%) of C8 to C<sub>13</sub> fatty acids with both even and odd carbon chains (Table 3); most of these acids are not detected or are present at much lower levels in euphausiid lipids (Tables 1 and 2). (2) All phytoplankton samples contained HUFA, mainly 20:5(n-3) and 22:6(n-3) in levels ranging from less than 1% to about 23% (Table 3). This suggests that the

Table 2. Euphausia crystallorophias. Fatty acids (as weight per cent of total acids)

Fatty acid	Station 11	Station	13	Station 14	Station 15	Station 16	Station 17	
		Adults	Juvenile					
14:0	2.3	2.2	2.4	2.6	2.6	2.2	2.6	
16:0	15.3	12.1	13.3	17.2	15.5	13.1	16.0	
18:0	0.3	0.4	0.6	0.3	0.5	0.5	0.8	
16:1(n-7)	8.6	6.5	7.9	7.9	8.4	9.3	6.5	
18:1(n-9)	39.6	49.8	48.8	47.8	49.0	47.6	40.7	
18:2(n-3)	1.7	2.4	2.2	2.1	2.3	2.3	1.6	
18:3(n-3)	1.0	1.0	1.0	1.0	0.8	0.7	0.8	
18:4(n-3)	0.7	1.8	1.5	1.2	1.3	0.8	1.0	
20:4(n-6)	0.9	-	-	0.7	0.4	-	1.0	
20:5(n-3)	18.2	14.1	12.5	12.3	12.8	14.9	16.5	
22:6(n-3)	9.9	7.2	7.7	6.0	5.3	5.6	11.1	
Minor fatty acids <sup>a</sup>	1.5	2.5	2.1	0.9	1.1	3.0	1.4	

Except for stearic acid (18:0) only those fatty acids present at a level of 1% or more



<sup>&</sup>lt;sup>1</sup>Dr. S. El-Sayed provided qualitative and semiquantitative microscopic data on the composition of the phytoplankton samples.

Bottino: Fatty Acids of Antarctic Phytoplankton and Euphausiids

Table 3. Fatty acids of Antarctic phytoplankton and euphausiids (as weight per cent of total acids)

Fatty acid	Phyto	plankt	on at	Station	ısa								Euphausia superba (average of 3 stations)	Euphausia crystallorophias (average of 7 stations)
	8	9a	9b	11a	116	13	14	15a	15b	15c	18a	18b		
8:0	_	_	-	0.4	_	1.2	1.0	_	-	0.7	_	_	_	
9:0	0.2	0.3	2.9	0.9	0.2	1.4	2.5	1.5	0.1	1.7	-	-	-	0.2
10:0	-	1.5	-	1.7		0.9	4.0	2.4	0.1	2.2	_	-	_	0.2
11:0	-	_	-	0.7	1.3	1.2	1.9	1.1	trace	0.3	-	-	_	
12:0	1.3	2.3	2.3	1.9	0.4	2.0	2.2	2.5	0.4	0.8	0.2	-	0.2	0.1
13:0	-	1.1	-	-	-	0.4	3.0	0.8	-	_	-	-	_	0.1
14:0	9.7	22.5	11.5	15.9	22.9	25.5	20.7	17.4	19.3	5.1	13.0	9.5	14.0	2.4
15:0	1.9	0.3	1.6	2.6	1.7	2.6	-	3.4	0.7	1.1	0.3	_	0.4	0.1
16:0	20.4	20.1	19.9	16.0	18.8	21.7	18.5	18.7	17.2	16.1	17.3	14.9	22.3	14.6
18:0	7.0	2.1	3.3	2.0	2.0	2.6	2.2	1.9	1.8	5.2	1.7	1.5	1.0	0.5
10:1(n-?)	0.6	2.2	2.5	_	0.8	-	-	-	-	-	0.2	-	-	_
11:1(n-?)	-		-	0.7	-	1.9	2.0	1.3	0.1	0.9		-	-	_
12:1(n-?)	-	2.8	1.5	1.6	2.4	0.7	1.2	1.6	0.2	0.9	1.5	0.3	_	-
13:1(n-?)	-	0.5	-	1.0	-	0.6	1.8	1.4	0.1	1.1	_	-	-	-
15:1(n-?)	0.5	0.7	0.6	1.3	0.6	1.2	1.0	2.7	0.4	0.7	1.8	_	trace	_
16:1(n-?)	12.4	8.3	7.5	6.2	5.7	5.3	3.4	7.8	3.6	3.2	13.1	10.3	9.5	7.9
17:1(n-?)	1.6	0.3	0.9	0.6	0.3	0.2	-	0.3	0.3	trace	1.4	0.9	0.5	0.3
18:1(n-9)	12.1	16.0	15.6	16.2	18.3	16.3	24.8	11.4	20.2	12.5	17.3	18.7	20.8	46.2
20:1(n-9)	0.4	-	0.8	trace	-	1.7	-	0.1	trace	-	0.3	0.3	0.9	0.2
18:2(n-6)	2.1	0.1	0.3	0.2	0.1	_	-	-	0.1	_	4.1	3.8	0.2	-

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18:2(n-3)	3.7	3.3	2.4	3.1	3.3	3.0	2.7	2.1	7.1	12.0	1.2	1.7	2.4	2.1	
22:2(n-6)	~	-	-	-	-	0.8	0.9	1.6	2.0		_	-	-	-	
22:2(n-3)	-	0.6	0.7	1.4	1.4	_	-		-	_	-	-	-	_	
18:3(n-6)	0.3	0.3	0.3	0.2	0.2	-	-	-	0.2	0.3	0.2	0.3	0.2	0.1	
18:3(n-3)	0.9	0.7	0.6	0.7	0.7	0.7	0.3	0.2	0.1	0.2	0.2	0.3	1.2	0.9	
20:3(n-6)	0.4	0.2	_	trace	0.9	trace		0.3	2.6	0.1	-	0.1	-	-	
20:3(n-3)	0.2	0.2	0.3	_	-	-	-	trace	0.9	1.0	-	0.2	0.5	0.3	
16:4(n-1)	-	_	0.5	-	-	-	-	-	-	6.3	-	_	-	-	
18:4(n-3)	2.0	3.1	3.5	5.2	6.0	3.0	2.7	3.2	6.2	0.9	2.2	2.5	2.7	1.2	
20:4(n-6)	-	-	-	0.4	-	-	-	-	_	4.7	-	-	0.4	0.4	
20:4(n-3)	0.2	-	0.3	0.2	-	-	-	0.1	trace	_	-	0.2	0.4	0.1	
22:4(n-6)	-	-	-	-	-	-	-	-	-	trace	_	_	0.2	_	
22:4(n-3)	1.3		trace	_	-	-	-	trace	-	trace	-	_	-	-	
20:5(n-3)	11.4	4.8	9.2	7.0	6.4	1.7	2.1	5.3	6.0	2.1	18.4	23.4	13.1	14.4	
22:5(n-6)	1.1	-	-	_	-	-	-	-	-	_	-	-	=	_	
22:5(n-3)	0.3	0.3	-	-	-	-	-	0.1	-	2.1	-	_	0.2	_	
22:6(n-3)	6.1	4.9	7.9	8.4	5.5	0.9	0.8	7.1	7.8	16.5	5.5	11.0	8.1	7.5	
Minor fatty acids <sup>b</sup>	3.3	1.5	4.3	4.6	1.0	3.2	0.6	4.0	2.8	1.8	0.5	0.9	0.8	0.4	

<sup>&</sup>lt;sup>a</sup>Microscopic examination indicated that the following genera predominated in each station. Station 9: Eudorina, Pandorina; Station 10: Thalassiosira, Fragilaria, Nitzchia, Corethron, Silicoflagellates; Station 11: Corethron, Fragilaria, Chaetoceros, Silicoflagellates; Station 12: Corethron, Fragilaria, Nitzchia, Tintinnids; Station 13: Complex mixture; Station 14: Phaeocystis; Station 15: Phaeocystis, Chaetoceros, Nitzchia, Thalassiosira, Fragilaria; Stations 17 and 18: Fragilaria, Nitzchia, Coscinodiscus, Dinoflagellates, Tintinnids.

Only those fatty acids present at a level of 1% or more are included. See footnotes to Tables 1 and 2 for further explanation.

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