

The fate of dietary lipids in the Arctic ctenophore *Mertensia ovum* (Fabricius 1780)

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Received: 21 December 2006 / Accepted: 1 October 2007 / Published online: 23 October 2007
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Abstract Lipids of the Arctic ctenophore *Mertensia ovum*, collected from Kongsfjorden (Svalbard) in 2001, were analysed to investigate seasonal variability and fate of dietary lipids. Total lipids, lipid classes and fatty acid and alcohol compositions were determined in animals, which were selected according to age-group and season. Changes in lipids of age-group 0 animals were followed during growth from spring to autumn. Total lipids increased from May to September. Lipids as percentage of dry mass were lowest in August indicating their use for reproduction. Higher values occurred in September, which may be due to lipid storage for overwintering. Wax esters were the major lipid class accounting for about 50% of total lipids in age-group 0 animals from July and August. Phospholipids were the second largest lipid fraction with up to 46% in this age-group. The principal fatty acids of *M. ovum* from all age-groups were 22:6(n-3), 20:5(n-3) and 16:0. Wax ester fatty alcohols were dominated by 22:1(n-11) and 20:1(n-9) followed by moderate proportions of 16:0. The unique feature of *M. ovum* lipids was the high amount of free fatty alcohols originating probably from the dietary wax esters. In May, free alcohols exhibited the highest mean proportion with 14.6% in age-group 0 animals. We present the first data describing a detailed free fatty alcohol composition in zooplankton. This composition was very different from the

alcohol composition of *M. ovum* wax esters because of the predominance of the long-chain monounsaturated 22:1 (n-11) alcohol accounting for almost 100% of total free alcohols in some samples. The detailed lipid composition clearly reflected feeding of *M. ovum* on the herbivorous calanoid species, *Calanus glacialis* and *C. finmarchicus*, the abundant members of the zooplankton community in Kongsfjorden. Other copepod species or prey items seem to be less important for *M. ovum*.

Introduction

The true Arctic ctenophore *Mertensia ovum* (Fabricius 1780) accounts for up to 70% of the total abundance of gelatinous zooplankton in Arctic waters (Hop et al. 2002), and its persistent presence and length distribution during most seasons suggest a two to multi-year life-cycle (Percy 1989; Lundberg et al. 2006). It uses its extensive tentacles to entrap and ingest prey and is considered to consume mainly the abundant *Calanus* species, *C. finmarchicus*, *C. glacialis* and *C. hyperboreus* (Swanberg and Båmstedt 1991; Siferd and Conover 1992; Raskoff et al. 2005). It has also been shown that *M. ovum* feeds on smaller copepods (*Pseudocalanus* sp.), copepod nauplii, the pteropod *Limacina helicina* and fish larvae in the pelagic food web (Swanberg and Båmstedt 1991; Granhag et al. 2005). At high abundances ctenophore predation can significantly reduce copepod populations (Swanberg and Båmstedt 1991).

During summer *M. ovum* is the most abundant gelatinous zooplankton species in the Arctic and forms a substantial portion of the zooplankton biomass in Kongsfjorden (Lundberg et al. 2006). The high abundance of zooplankton (Walkusz et al. 2003) offers a variety of prey to the ctenophores. Kongsfjorden harbours a mixture of boreal and

Communicated by M. Wahl.

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Arctic flora and fauna and is influenced by water from the West Spitsbergen Current, which mixes with Arctic-derived water and locally produced fjord water (Walkusz et al. 2003).

The population of *M. ovum* is mainly controlled by the ctenophore *Beroe cucumis* (Falk-Petersen et al. 2002). Ctenophores are often considered to be dead-ends in the pelagic food web, but they may also be preyed upon by higher trophic levels, such as polar cod (*Boreogadus saida*), Atlantic cod (*Gadus morhua*), northern fulmar (*Fulmarus glacialis*) and black legged kittiwake (*Rissa tridactyla*), as well as the medusa *Cyanea capillata* (Lundberg et al. 2006 and citations therein). The food-chain relationships between copepods, *M. ovum* and *B. cucumis* have far reaching implications for the high latitude ecosystem, providing energy for higher trophic levels.

M. ovum is able to store lipids in special storage structures despite its high water content (Larson and Harbison 1989). Its amount of total lipids decreases during summer and reaches highest levels in autumn (Lundberg et al. 2006). The lipid composition of *M. ovum* is strongly influenced by its prey, and the major storage lipids are wax esters, which originate from feeding on lipid-rich zooplankton species (Falk-Petersen et al. 2002). Wax esters are produced in high amounts by calanoid copepods and consist of fatty acid and alcohol moieties (Sargent and Henderson 1986; Kattner et al. 1989). The long-chain monounsaturated fatty acids and alcohols, 20:1(n-9) and 22:1(n-11), can be used as trophic markers because, in the Arctic, they are only synthesised de novo by the herbivorous *Calanus* species (Graeve et al. 1994; Dalsgaard et al. 2003). In the food chain, from copepods via *M. ovum* to *B. cucumis*, both ctenophores contain relatively high amounts of these fatty acids and alcohols. Applying trophic marker lipids enhances our understanding of trophic interactions in polar regions in general and of the linkages between ctenophores and their potential prey in particular (Falk-Petersen et al. 2002; Ju et al. 2004).

The objectives of this study were to examine lipid classes and fatty acids and alcohols of the Arctic ctenophore *M. ovum* and to follow the fate of dietary lipids. The role of free alcohols, which we detected in *M. ovum* and which has been recently reported for the Antarctic ctenophores *Callinaria antarctica* and *B. cucumis* (Phleger et al. 1998; Ju et al. 2004), is still unknown and will be discussed. We also aim to understand the utilisation of specific lipids within different developmental phases and seasons.

Materials and methods

Sampling, dry mass and total lipids

Mertensia ovum was collected in Kongsfjorden, Svalbard (Arctic), during four periods in 2001 (21–24 May, 22–30

July, 5–18 August and 4–30 September) with a WP-3 plankton net onboard RV “Lance”, RV “Oceania” and RV “Haakon Mosby” or with jars mounted on a rod, operated from small boats. Live animals were transported to the laboratory in Ny-Ålesund or on the research vessels, where all individual specimens were measured for length (see details in Lundberg et al. 2006). Specimens without clear signs of prey in their guts were directly transferred into glass vials containing chloroform:methanol (2:1, by vol.) with 0.01% butyl hydroxy toluene (BHT) as antioxidant and immediately frozen at -25°C . All samples consisted of one specimen. In the home laboratory, the solvent was evaporated with nitrogen to dryness. Dried samples were transferred into pre-weighed vials and lyophilised for 48 h (Alfa 1-4, Christ, Germany). Dry mass (DM) was determined gravimetrically. Total lipids were extracted from the freeze-dried samples using dichloromethane:methanol (2:1 by vol.), essentially after Folch et al. (1957), and measured gravimetrically. In some cases, where it was not possible to make reliable gravimetric analyses, only lipids were extracted, and lipid class and fatty acid/alcohol compositions were determined.

Lipid class and fatty acid analysis

Lipid classes were determined by high performance thin layer chromatography (HPTLC) and densitometry essentially after Olsen and Henderson (1989). Briefly, HPTLC silica gel 60 plates (20 × 10 cm, Merck) were spotted with 5 μl of sample extracts and standard solutions using a CAMAG-Linomat 4. The separation of lipid classes was performed in a CAMAG horizontal chamber with hexane:diethylether:acetic acid (80:20:0.2; by vol.). Thereafter the plate was dried in a desiccator under vacuum for 30 min. Lipid classes were visualised by submerging the plate in manganese (II) chloride·(4H₂O), methanol and sulphuric acid reagent in a CAMAG immersion device for 5 s followed by combustion at 120°C for 20 min. The quantification was performed with a TLC Scanner (CAMAG 3) combined with winCATS software (Böer et al. 2005).

For the determination of the fatty acid and alcohol compositions of the major lipid classes (wax esters, free fatty alcohols and phospholipids), lipids were preparatively separated on TLC plates coated with silica gel 60 H. Following separation (see above), lipid bands were visualised with 2',7'-dichlorofluorescein, scraped off and extracted with dichloromethane:methanol (2:1 by vol.) (Albers et al. 1996). Because the amount of lipids of only one specimen was not enough for the separation, the lipid extracts of 2–7 single specimens were combined. In total, eight combined samples were analysed for fatty acids and alcohols of the lipid classes. In few samples the amount of lipids was still insufficient for a reliable analysis of all lipid class compositions so that the number of replicates is different.

The fatty acid and alcohol compositions of the total lipid extracts and the lipid classes were analysed by gas–liquid chromatography according to Kattner and Fricke (1986). Lipids were converted to fatty acid methyl esters (FAME) and alcohols by transesterification in methanol containing 3% concentrated sulphuric acid at 80°C for 4 h. After extraction with hexane, FAME and fatty alcohols were analysed with a Hewlett-Packard 6890 Series gas chromatograph on a DB-FFAP fused silica capillary column (30 m × 0.25 mm inner diameter, 0.25 µm film thickness) using temperature programming (160–240°C at 4°C min⁻¹, hold 15 min). The alcohol composition of the free fatty alcohol fractions was determined without hydrolysis confirming that the alcohols did not originate from wax esters. In addition, free fatty alcohols were analysed following transesterification, which confirms that this fraction did not contain any fatty acids. For recording and integration, Class-VP software (4.3) (Shimadzu, Germany) was used. FAME and alcohols were identified with standard mixtures and, if necessary, additional confirmation was carried out by gas chromatography–mass spectrometry (Kattner et al. 1998).

Age-groups were separated in polymodal length-frequencies using the “mix.dist” package in “R” applying distribution mixture analysis (Lundberg et al. 2006). For all lipid data One-way ANOVA was performed together with multiple range and Kruskal–Wallis tests to identify the specific dependent variables that contributed significantly to seasonal effects. *F*-ratios and the significance at the 5% probability level are presented [ndf = 3, ddf = 11 (lipid classes) and 16 (fatty acids)] in lipids within the group 0 animals. Due to the robustness of the ANOVA we assume that all data are normal distributed with even homogeneity of variance.

Results

Total lipid mass and lipid classes

Length, dry mass, lipid mass and lipid content data for *Mertensia ovum* were combined according to age-groups as defined by Lundberg et al. (2006) and are summarized in

Table 1. Age-group 0 animals from the entire year were analysed. Their mean length increased from 11.6 mm in May to 19.9 mm in September and their mean dry mass from 11.2 to 26.9 mg ind⁻¹. Age-group 1 animals had a length of 20.9 mm and a mean dry mass of 75.7 mg ind⁻¹ in May as well as 38.0 mm and 179 mg ind⁻¹ in August. Group 2 animals were only analysed in May with 32 mm length and a mean dry mass of 77.1 mg ind⁻¹. The mean total lipid mass increased from 0.7 to 2.3 mg ind⁻¹ from May to September in group 0. The highest concentrations were found in group 1 and 2 animals. The lipid content as percentage of dry mass was lowest in animals of group 0 and 1 in August and highest in group 0 animals from September, however, data were not significantly different.

Major lipid classes of *M. ovum* were wax esters and phospholipids (Table 2). Wax esters were the dominant lipids for storage being highly variable within the age-groups and were significantly different within group 0 (*F*-ratio = 4.7; *p* < 0.05). Highest levels (41–57% of total lipid) occurred in group 0 animals in July, August, and September and in group 2 already in May. In contrast, in group 0 animals from May only 17.1% of the total lipids were wax esters, which was significantly different from all other samples (multiple range test). In age-group 1 animals, wax esters accounted for ca. 38% on average. The second most important storage lipids were free fatty alcohols (significantly different within the group 0, *F*-ratio = 8.4; *p* < 0.01). In May, free alcohols exhibited mean proportions of 14.6% (maximum value of 17%) in group 0 animals, being significantly higher than in other age-groups or seasons. In July, August and September proportions decreased to levels between 5.2 and 6.9%. In the other age-groups, mean values were between 2.5 and 7.9%. Triacylglycerols had higher proportions in group 0 animals from July (13.1%) and in age-group 1 from August (10.3%). During the other months, triacylglycerols varied between 1.9 and 5.3%, reflecting their minor importance. Free fatty acids ranged from 1.6 to 6.2% in all samples (Table 2).

Phospholipids, as structural membrane lipids, decreased from 45.6% in age-group 0 animals from May to 18.5% in July and 25.6% in August and increased again to 38.4% in September (significantly different, *F*-ratio = 4.5; *p* < 0.05).

Table 1 *Mertensia ovum* from Kongsfjorden, Svalbard, 2001. Mean and standard deviation of general data [dry mass (DM), total lipid (TL) and percent lipid of DM; (*n*) numbers of replicates (one animal each)]

Age group	0				1		2
	May (6)	July (2)	Aug (3)	Sept (6)	May (2)	Aug (2)	May (2)
Length (mm)	11.6 ± 1.7	16.0 ± 2.2	18.4 ± 1.7	19.9 ± 1.3	20.9 ± 1.2	38.0 ± 1.3	32.0 ± 2.2
DM (mg ind ⁻¹)	11.2 ± 2.5	27.7 ± 30.2	24.6 ± 14.9	26.9 ± 18.7	75.7 ± 41.6	178.9 ± 86.9	77.1 ± 35.0
TL (mg ind ⁻¹)	0.7 ± 0.3	1.7 ± 0.9	1.4 ± 0.8	2.3 ± 1.4	5.2 ± 4.8	6.1 ± 0.6	5.4 ± 1.7
TL (%DM)	6.6 ± 1.9	10.5 ± 8.1	5.5 ± 0.1	10.0 ± 4.4	6.0 ± 3.1	3.8 ± 1.5	7.3 ± 1.2

Table 2 *Mertensia ovum* from Kongsfjorden, Svalbard, 2001. Mean and standard deviation of the lipid class compositions (mass % of total lipids). *PL* Phospholipids, *ST* sterols, *FFAlc* free fatty alcohols, *FFA*

free fatty acids, *TAG* triacylglycerols, *WE* wax esters. (*n*) Numbers of replicates (one animal each)

Age group	0				1		2
	May (5)	July (3)	Aug (4)	Sept (3)	May (10)	Aug (2)	May (3)
PL	45.6 ± 8.2*	18.5 ± 7.2*	25.6 ± 4.8	38.4 ± 21.3	45.8 ± 14.4	24.9 ± 2.8	27.7 ± 0.8
ST	13.7 ± 1.9*	6.1 ± 4.7	5.5 ± 4.1	5.9 ± 4.1	8.0 ± 3.5	10.8 ± 2.5	7.7 ± 3.7
FFAlc	14.6 ± 1.8*	5.2 ± 4.8	6.9 ± 4.0	6.2 ^a	2.5 ± 5.6	7.9 ± 0.5	6.0 ± 5.3
FFA	3.7 ± 2.4*	5.7 ± 6.5	1.6 ± 1.9	5.0 ± 3.6	4.5 ± 2.6	6.2 ± 2.9	3.3 ± 0.6
TAG	5.3 ± 2.0*	13.1 ± 7.5*	3.3 ± 1.1	4.1 ± 5.3	1.9 ± 2.7	10.3 ± 1.9	4.5 ± 1.4
WE	17.1 ± 10.0*	50.1 ± 19.6	57.1 ± 12.1	40.8 ± 26.9	37.0 ± 15.5	39.1 ± 11.7	50.7 ± 10.8

* Significantly different from other seasons

^a FFAlc from September animals is calculated from fatty acid and alcohol composition

In September animals, phospholipids were highly variable. High proportions were also found in age-group 1 in May but decreased in August, whereas phospholipids in group 2 were low in May. The high proportions in May were significantly different from the other samples except group 0 animals from September. The proportions of sterols ranged from 5.5 to 13.7% (Table 2).

Fatty acid and alcohol compositions

The principal fatty acids and alcohols of the total lipid extracts of animals from all age-groups are presented in Table 3 including statistically significant differences for the group 0 animals resulting from One-way ANOVA. Due to the high level of phospholipids in the group 0 animals in May membrane-bound fatty acids [22:6(n-3), 20:5(n-3) and 16:0] were significantly different from other seasons ($p < 0.05$). The highest mean level of the 22:6(n-3) fatty acid was 34.5% in age-group 0 in May. For other periods and age-groups, mean values from 21 to 26% were found. The 20:5(n-3) fatty acid had elevated mean proportions in May for age-groups 0 and 1 (19.7 and 19.6%), compared to other seasons and age-groups with 10–16% on average. The fatty acids, which are important dietary markers, showed some variations during the seasonal development. The 18:4(n-3) fatty acid increased from May to August in group 0 from mean values of 1.8–6.1% with a maximum of 12.1%. The low levels in May were only significantly different from the group 0-values in August. Higher proportions of 18:4(n-3) were also found in group 1 from August. The proportions of 16:1(n-7) were mostly lower than those of 18:4(n-3) except in animals of age-groups 1 and 2 in May. Significant differences within the group 0 animals were found for the proportions of the long-chain monounsaturated fatty acids 20:1(n-9) (F -ratio = 5.1; $p < 0.01$) and 22:1(n-11) (F -ratio = 3.6; $p < 0.05$), which generally increased with the amount of wax esters. The low values in May of group 0

animals were significantly different from the other months. The proportions of all long-chain monounsaturated fatty acid isomers ranged on average from 1.5 to 5.4%.

Fatty alcohols of the total lipid extracts were predominantly 22:1(n-11) followed by 20:1(n-9) as well as moderate to low proportions of 16:0 and 14:0 being significantly different within the group 0 animals (Table 4). The proportion of the 22:1(n-11) alcohol in age-group 0 in May (83.2%) was significantly higher than in the other months. Group 1 animals were also high in the 22:1(n-11) alcohol in May (70.7%). In all other animals, 22:1(n-11) ranged from 54.1 to 65.2% on average. The proportions of the other major alcohol, 20:1(n-9), were lower for the May animals (8.3 and 15.1%) and ranged from 20 to 27.1% in all other specimens. The 16:0 alcohol occurred in proportions of less than 10%, and other alcohols were only minor components. These alcohol compositions of the total lipid extracts have to be interpreted carefully, because they represent a combination of alcohols as moiety of the wax ester molecules and of free fatty alcohols. Below we present the composition of the wax esters and first detailed compositions of free fatty alcohols in marine zooplankton.

For the analysis of fatty acid and alcohol compositions of individual lipid classes several samples have to be combined for a reliable separation by thin layer chromatography. The compositions of the three major lipid classes, wax esters, phospholipids and free fatty alcohols, were analysed (Tables 4, 5). Phospholipids were dominated by three fatty acids: 22:6(n-3) accounting for 28.7%, 20:5(n-3) for 17.5% and 16:0 for 16.9% on average, all with only small variations. The composition of the wax esters of *Mertensia ovum* is shown in comparison with mean wax ester data of the dominant *Calanus* species (Table 4; Albers et al. 1996). The composition was much more variable than that of the phospholipids. Proportions of about 10% each were found for the fatty acids 20:5(n-3), 14:0, 16:1(n-7) and 18:1(n-9); slightly lower proportions occurred for 18:4(n-3), 20:1(n-9)

Table 3 *Mertensia ovum* from Kongsfjorden, Svalbard, 2001. Mean and standard deviation of the fatty acid and alcohol compositions of the different age-group animals (mass %). (n) Numbers of replicates (one animal each)

Age-group	0				1		2	
	Month	May (6)	Jul (3)	Aug (5)	Sept (9)	May (10)	Aug (2)	May (3)
<i>Fatty acids</i>								
14:0		5.1 ± 0.7	6.1 ± 2.7	6.6 ± 1.2	6.6 ± 1.5	7.1 ± 1.3	6.7 ± 0.3	10.5 ± 0.9
15:0		0.4 ± 0.1	0.6 ± 0.2	0.6 ± 0.1	0.6 ± 0.1	0.4 ± 0.2	0.5 ± 0.1	0.6 ± 0.0
16:0		11.8 ± 1.0*	8.1 ± 1.8*	10.4 ± 2.2	13.0 ± 1.8	12.0 ± 1.8	8.6 ± 1.7	11.4 ± 1.2
16:1(n-7)		2.6 ± 0.8	4.0 ± 1.5	3.7 ± 1.0	2.7 ± 1.9	4.4 ± 2.2	3.7 ± 0.7	7.6 ± 1.1
16:1(n-5)		0.2 ± 0.1	0.4 ± 0.2	0.3 ± 0.2	0.3 ± 0.2	0.2 ± 0.1	2.2 ± 2.8	0.4 ± 0.0
16:2(n-4)		0.3 ± 0.1	0.3 ± 0.1	0.4 ± 0.1	0.3 ± 0.1	0.3 ± 0.2	0.4 ± 0.1	0.7 ± 0.1
16:3(n-4)		0.3 ± 0.1	0.5 ± 0.2	0.5 ± 0.3	0.5 ± 0.2	0.7 ± 0.7	0.5 ± 0.0	1.0 ± 0.1
16:4(n-1)		0.2 ± 0.3	0.5 ± 0.2	0.5 ± 0.2	0.4 ± 0.3	0.7 ± 0.6	0.5 ± 0.1	1.4 ± 0.1
18:0		3.7 ± 0.5	4.8 ± 1.2	4.6 ± 1.3	4.8 ± 1.7	3.3 ± 1.1	3.6 ± 0.6	2.4 ± 0.5
18:1(n-9)		3.7 ± 0.3*	7.0 ± 2.2	5.0 ± 0.6	6.5 ± 1.7	3.9 ± 0.6	4.5 ± 0.3	4.0 ± 0.2
18:1(n-7)		1.1 ± 0.1*	0.8 ± 0.1	0.9 ± 0.1	0.8 ± 0.2	1.3 ± 0.3	1.0 ± 0.3	1.3 ± 0.4
18:2(n-6)		1.0 ± 0.4*	6.9 ± 4.4*	2.4 ± 0.6	1.3 ± 0.2	0.7 ± 0.1	1.8 ± 0.2	0.8 ± 0.1
18:3(n-3)		0.6 ± 0.1*	1.0 ± 0.3	0.8 ± 0.1	0.9 ± 0.3	0.4 ± 0.1	0.8 ± 0.2	0.6 ± 0.1
18:4(n-3)		1.8 ± 0.4*	5.0 ± 3.1	6.1 ± 3.5	4.2 ± 2.0	2.4 ± 0.9	5.5 ± 3.2	4.1 ± 0.9
20:0		0.0 ± 0.1	0.7 ± 0.2	0.5 ± 0.2	0.3 ± 0.1	0.1 ± 0.1	0.4 ± 0.1	0.1 ± 0.1
20:1(n-9)		1.8 ± 0.6*	5.4 ± 2.7	4.8 ± 1.3	3.2 ± 1.5	3.2 ± 1.9	5.0 ± 2.3	3.5 ± 2.5
20:1(n-7)		3.0 ± 0.9	1.5 ± 0.9	2.2 ± 0.8	2.5 ± 0.6	4.1 ± 1.5	3.2 ± 1.4	1.9 ± 1.4
20:4(n-6)		0.4 ± 0.2	0.7 ± 0.6	0.5 ± 0.2	0.4 ± 0.1	0.5 ± 0.1	0.8 ± 0.6	0.4 ± 0.1
20:4(n-3)		0.6 ± 0.3	1.1 ± 0.1	1.3 ± 0.1	1.7 ± 0.3	0.7 ± 0.1	1.0 ± 0.2	0.8 ± 0.2
20:5(n-3)		19.7 ± 5.0*	9.5 ± 3.2*	13.5 ± 2.5	14.2 ± 1.7	19.6 ± 2.3	12.4 ± 1.3	15.7 ± 4.2
22:1(n-11)		1.8 ± 0.7*	4.6 ± 1.3	4.3 ± 1.7	3.3 ± 1.7	2.8 ± 2.1	4.2 ± 1.9	5.0 ± 1.0
22:1(n-9)		3.8 ± 2.2	2.4 ± 1.8	3.0 ± 1.1	3.4 ± 2.0	4.4 ± 1.9	3.5 ± 0.3	2.9 ± 1.0
22:5(n-3)		0.0 ± 0.0	0.4 ± 0.4	0.5 ± 0.7	1.7 ± 2.4	0.9 ± 0.6	0.0 ± 0.0	0.3 ± 0.6
22:6(n-3)		34.5 ± 3.9*	25.8 ± 14.5	24.0 ± 3.3	23.8 ± 5.3	24.0 ± 4.3	26.5 ± 4.9	21.2 ± 5.3
<i>Alcohols</i>								
14:0		1.2 ± 0.3*	3.3 ± 1.2	4.0 ± 0.8	3.6 ± 1.5	2.0 ± 0.8	2.9 ± 1.7	2.9 ± 0.6
16:0		3.2 ± 1.5*	7.7 ± 2.9	8.5 ± 1.5	9.9 ± 4.7	5.1 ± 2.5	6.0 ± 2.6	7.8 ± 2.5
16:1(n-7)		0.8 ± 1.3	2.2 ± 0.8	1.6 ± 0.7	1.7 ± 1.2	3.3 ± 2.6	2.1 ± 0.6	5.8 ± 3.0
18:1(n-9)		2.9 ± 1.9	1.7 ± 0.6	1.6 ± 0.3	1.5 ± 0.8	1.5 ± 0.7	1.6 ± 0.3	2.2 ± 0.4
18:1(n-7)		0.5 ± 1.1	1.7 ± 0.6	1.9 ± 0.6	1.4 ± 1.0	2.4 ± 1.7	2.3 ± 0.8	1.4 ± 0.9
20:1(n-9)		8.3 ± 4.3*	27.1 ± 6.4	24.1 ± 6.7	18.1 ± 10.5	15.1 ± 7.7	20.0 ± 6.2	25.9 ± 4.7
22:1(n-11)		83.2 ± 6.9*	56.2 ± 11.6	58.3 ± 8.4	63.8 ± 17.7	70.7 ± 13.6	65.2 ± 8.9	54.1 ± 7.8

* Significantly different from other seasons

and 22:1(n-11). Major alcohols of the wax esters were 22:1(n-11) accounting for 36.9%, 20:1(n-9) for 32.4% and 16:0 for 15.8% on average (Table 4).

The unique feature of the *M. ovum* lipids was the fraction of free fatty alcohols. Data are shown as average of all samples but are also divided into three seasons (Table 5) because they are much more variable than the compositions of the other lipid classes. The predominant component was the 22:1(n-11) alcohol accounting for 93.6% on average in the September animals. In May, the 22:1(n-11) alcohol also exhibited high proportions (75.1%), whereas in July the composition was more similar to that of the wax ester composition, although the 22:1(n-11) alcohol remained the major component (42.8%). The next abundant alcohol was

20:1(n-9). The other typical wax ester alcohols only occurred in higher proportions in the July sample.

Discussion

Lipids of *Mertensia ovum*

Mertensia ovum is able to store lipids in special storage structures (oil sac-like), which are visible in its gelatinous body. These oil sacs are associated with the tentacle bulbs. The lipid proportion of dry mass ranged from 2.7 to 16.3%, which is similar to that reported by Percy and Fife (1981) for *M. ovum* from the Canadian Arctic, but did not reach the

Table 4 *Mertensia ovum* from Kongsfjorden, Svalbard, 2001. Mean and standard deviation of the fatty acid and alcohol compositions of wax esters and phospholipids (mass %). (*n*) Numbers of replicates (several specimens combined for each replicate). Mean data of the dominant *Calanus* species (Albers et al. 1996) are added for comparison

Lipid class	<i>Mertensia ovum</i>		<i>C. glacialis</i>	<i>C. finmarchicus</i>	<i>C. hyperboreus</i>
	Phospholipids (7)	Wax esters (4)	Wax esters	Wax esters	Wax esters
<i>Fatty acid</i>					
14:0	5.4 ± 1.6	10.1 ± 1.4	13.1	26.3	6.4
15:0	0.3 ± 0.3	0.3 ± 0.4	0.3	0.7	–
16:0	16.9 ± 3.2	6.5 ± 1.0	6.1	9.8	5.8
16:1(n-7)	1.6 ± 0.5	10.0 ± 5.5	32.9	6.7	11.7
16:1(n-5)	0.1 ± 0.1	0.3 ± 0.4	0.3	0.9	0.7
16:2(n-4)	0.1 ± 0.1	1.0 ± 0.3	1.2	0.6	1.0
16:3(n-4)	0.2 ± 0.1	1.3 ± 1.0	–	0.9	0.5
16:4(n-1)	0.0 ± 0.0	2.2 ± 1.9	–	0.5	2.0
18:0	6.0 ± 1.5	3.6 ± 3.8	–	0.9	0.6
18:1(n-9)	4.0 ± 1.7	9.9 ± 4.9	5.5	5.3	5.8
18:1(n-7)	1.2 ± 0.4	0.9 ± 0.2	1.1	0.3	1.6
18:2(n-6)	1.3 ± 0.2	0.9 ± 0.7	1.0	1.2	3.6
18:3(n-3)	0.7 ± 0.1	1.1 ± 0.4	0.3	1.5	1.6
18:4(n-3)	2.3 ± 0.7	7.9 ± 0.3	0.5	13.7	6.2
20:0	0.1 ± 0.1	1.6 ± 1.3	0.1	–	0.8
20:1(n-9)	1.6 ± 0.4	8.6 ± 3.4	23.0	7.8	19.0
20:1(n-7)	2.5 ± 1.2	1.2 ± 0.1	1.0	0.9	1.5
20:4(n-6)	0.5 ± 0.3	0.2 ± 0.2	–	0.4	–
20:4(n-3)	1.0 ± 0.3	1.3 ± 0.5	–	0.5	0.7
20:5(n-3)	17.5 ± 3.0	11.9 ± 3.0	2.7	11.4	7.0
22:1(n-11)	1.4 ± 0.5	7.4 ± 1.8	8.3	7.0	17.3
22:1(n-9)	3.2 ± 2.0	1.1 ± 0.4	2.0	0.2	3.2
22:1(n-7)	0.9 ± 1.9	0.6 ± 0.4	–	–	–
22:5(n-3)	0.6 ± 0.3	1.6 ± 0.6	–	0.2	0.5
22:6(n-3)	28.7 ± 2.8	2.9 ± 0.7	0.8	2.2	2.4
<i>Alcohols</i>					
14:0		5.4 ± 1.0	2.1	3.9	4.4
16:0		15.8 ± 5.6	9.3	14.6	11.2
16:1(n-7)		4.9 ± 2.6	5.3	3.4	1.6
18:1(n-9)		2.5 ± 0.3	–	–	–
18:1(n-7)		2.2 ± 0.1	–	–	–
20:1(n-9)		32.4 ± 3.8	58.4	39.3	27.8
22:1(n-11)		36.9 ± 4.0	25.0	38.8	55.0

high percentage of up to 27.7% found in the smallest stages in Kongsfjorden by Lundberg et al. (2006). The lipid content was higher than in most other ctenophores, especially those from Antarctic regions, which exhibit only about 3% lipid of DM (Larson and Harbison 1989). The moderate lipid stores of the Arctic *M. ovum* may enhance their ability to survive longer periods of food scarcity (Percy 1988) and may also serve as buoyancy aid although ctenophores usually have a body composition, which makes them essentially neutral in buoyancy (Clarke and Peck 1990).

M. ovum is able to store lipids feeding on lipid-rich prey, especially *Calanus* species, which has been observed to be important prey in Kongsfjorden. *M. ovum* may even control

a copepod population (Swanberg and Båmstedt 1991). Its major storage lipids were wax esters, which are normally utilized for metabolic requirements. It has also been suggested that these lipid deposits might be a possibility to sequester excess of lipids from the diet (Clarke and Peck 1990).

For the determination of feeding preferences, fatty acid and alcohol compositions can be used (Graeve et al. 1994; Dalsgaard et al. 2003). The high proportions of 16:1(n-7) and 18:4(n-3) in the *M. ovum* wax esters suggest that *Calanus* species are the major prey of *M. ovum* as also reported by Falk-Petersen et al. (2002). The predominant 20:1(n-9) and 22:1(n-11) alcohols in the wax esters of *M. ovum* are also comparable to the major alcohols in *Calanus* wax esters

Table 5 *Mertensia ovum* from Kongsfjorden, Svalbard, 2001. Mean and standard deviation of the free fatty alcohol compositions (mass %). (n) Numbers of replicates (several specimens combined for each replicate)

Month	May (2)	Jul (1)	Sept (2)	All samples (8)
14:0	0.6 ± 0.9	13.0	0.8 ± 0.9	2.0 ± 4.5
16:0	2.3 ± 3.2	14.4	1.2 ± 0.4	2.8 ± 4.9
16:1(n-7)	1.7 ± 2.4	–	0.4 ± 0.4	0.5 ± 1.2
18:1(n-9)	0.5 ± 0.7	–	0.2 ± 0.2	0.3 ± 0.4
18:1(n-7)	0.5 ± 0.7	–	0.0 ± 0.0	0.1 ± 0.3
20:1(n-9)	19.4 ± 3.5	29.7	3.9 ± 3.6	15.3 ± 16.0
22:1(n-11)	75.1 ± 4.4	42.8	93.6 ± 5.1	79.0 ± 21.0

(Table 4). Herbivorous calanoid copepods are the only species in the Arctic to biosynthesise large amounts of wax esters with long-chain monounsaturated alcohols esterified with fatty acids of dietary origin or with fatty acids synthesised de novo (Sargent and Henderson 1986; Kattner and Graeve 1991; Scott et al. 2002). The compositions of copepod wax esters show species-specific differences. The ratio of both alcohols (20:1 and 22:1) in females and stages IV and V of *C. hyperboreus* is about 2, whereas the ratio is ca. 1 in *C. glacialis* and *C. finmarchicus* (Kattner et al. 1989; Scott et al. 2002). Both alcohols were almost equally distributed in the separately analysed wax esters of *M. ovum*, which reflects that it feeds preferentially on the abundant *C. glacialis* and *C. finmarchicus* but probably less on other zooplankton, like the less abundant *C. hyperboreus*. The calanoid copepods are certainly of differing importance during the different seasons as, for example, *C. finmarchicus* was the most abundant species in Kongsfjorden in autumn 1997 (Scott et al. 2000).

The second important neutral lipids were free fatty alcohols exhibiting a maximum value of 17%. The occurrence of free fatty alcohols is very unusual in marine organisms and has only been reported for the Antarctic ctenophores *Beröe cucumis* and *Callinara antarctica* (Phleger et al. 1998; Ju et al. 2004). This class of lipids may also be important in other marine zooplankton and animals, but may have been overlooked in other studies. In Table 5, we present the first data on free fatty alcohol compositions in marine zooplankton. This composition was very different from the alcohol composition of *M. ovum* wax esters, as it was predominated by the long-chain monounsaturated 22:1(n-11) alcohol accounting in maximum for almost 100% of total free alcohols in some samples. Free fatty alcohols certainly originate from the hydrolysis of dietary wax esters, which are ingested by *M. ovum*. The 22:1(n-11) alcohol seems to be selectively retained and might result from a certain order in the catabolism of free fatty acids and alcohols due to different enzymatic activities. The conversion of alcohols to fatty acids, which is necessary for the catabolism (β -oxidation), could be a slow process originating in an excess of free alcohols. In addition, the enzyme

might be more active on short-chain alcohols and thus, the conversion of the 22:1(n-11) alcohol might be slow or even impossible. If the 22:1(n-11) alcohol cannot be catabolised, then it has to be egested to get rid of indigestible lipids or excess of dietary lipids as hypothesised by Clarke and Peck (1990). However, it is also possible that this alcohol is actively retained because of its high energetic value (Albers et al. 1996). Proportions of 93.6% found in autumn indicate that the 22:1(n-11) alcohol might serve as an energy source for overwintering. However, these hypotheses are speculative because pathways for conversion and catabolism of fatty alcohols in ctenophores are still unknown.

The other but less important neutral lipids were triacylglycerols. Comparable low levels between 1 and 6% of total lipids are also found in ctenophores from Arctic and Antarctic regions (Nelson et al. 2000; Falk-Petersen et al. 2002). We assume that *M. ovum* and probably ctenophores in general do not convert dietary lipids or other dietary items to triacylglycerols for energy storage, which is consistent with the low levels of total lipids and the direct utilisation of prey for growth and reproduction.

Phospholipids, as structural lipids of membranes, are a major lipid class in *M. ovum* because of the generally low proportions of storage lipids. The typical phospholipid fatty acids, 16:0, 20:5(n-3) and 22:6(n-3), are less variable and showed only small variations within the different age-groups. Similarly high proportions of these fatty acids in phospholipids are also found in Arctic copepods (Albers et al. 1996; Scott et al. 2002) and most other zooplankton species (Lee et al. 2006). The dominance of the three fatty acids in marine phospholipids seems to be typical for most marine organisms. The polyunsaturated fatty acids are essential for animals, and the original source is phytoplankton. They are selectively incorporated into membranes but less frequently into storage lipids (Albers et al. 1996). Tracing of the carbon uptake with ^{14}C labelled algae showed a considerable uptake of carbon into the polar lipid fraction (Graeve 1992). The high amounts of polyunsaturated fatty acids seem to be responsible for membrane fluidity, however, Hazel (1995) concluded that the diversity of membrane adaptations to temperature is unlikely to be captured by lipid-based adjustments alone. The high proportions of 20:5(n-3) and 22:6(n-3) fatty acids in phospholipids of tropical zooplankton (G. Kattner et al., unpublished) further challenges the cold adaptation.

Seasonal lipid dynamics

The population structure of *Mertensia ovum* was applied from the study of Lundberg et al. (2006) as revealed from the distribution mixture analysis. Animals of three different size groups, found in Kongsfjorden in 2001, were analysed corresponding to age-group 0 throughout the year, and in

addition, to age-group 1 in May and August and to group 2 in May.

In spring, the age-group 0 animals, probably originating from spring spawning (Lundberg et al. 2006), contained small amounts of lipids, which consisted primarily of phospholipids and moderate amounts of wax esters as storage lipids. This suggests that they invest most of their prey lipids into growth. The proportions of wax esters were highly variable reflecting the different feeding conditions of the individual specimens. The greatest proportions of free fatty alcohols occurred in these young stages, which might be due to a limited ability to cope with the alcohol moiety of the wax esters. In general, the amounts of lipids as well as the lipid class and fatty acid composition in *M. ovum* are widely dependent on the catching success. The successful ingestion of single lipid-rich copepodids and adults will considerably determine the lipids of *M. ovum*. The young age-group 0 animals might only be able to catch nauplii and young copepodite stages but not the large wax ester-rich *Calanus* stages. Enhanced amounts of wax esters in *Calanus* species occur firstly in copepodid II (Kattner and Krause 1987; Wold et al. 2007). Nauplii probably do not accumulate lipids, but the lipid composition of nauplii is still unknown. In May 2001, a variety of prey was highly abundant in Kongsfjorden consisting mainly of *Calanus* and smaller copepod species (Basedow et al. 2004). Younger stages of *C. glacialis* and *C. hyperboreus* were probably more important as prey for the *M. ovum* than of *C. finmarchicus* in spring since *C. glacialis* can reproduce at low rates without food, and *C. hyperboreus* reproduces prior to feeding, whereas *C. finmarchicus* is dependent on food availability (Lee et al. 2006). The spring phytoplankton bloom in Kongsfjorden usually starts in early May (Hop et al. 2002).

In July, the amount of lipids in age-group 0 animals increased, being dominated by wax esters, which indicate summer growth and good condition of the population. During the summer season copepods are highly abundant, dominated by late copepodite stages CIV–CV and adults of *C. glacialis* (Kwasniewski et al. 2003). *M. ovum* is able to considerably diminish the copepod biomass and may even control a copepod population (up to 9% in the Barents Sea; Swanberg and Båmstedt 1991). Some specimens in Kongsfjorden had visible prey in their guts, mostly composed of *Calanus*, krill and unidentified matter (Lundberg et al. 2006). Fatty acids and alcohols as trophic markers in *M. ovum* clearly show the ingestion of wax ester-rich *Calanus* lipids with high proportions of the long-chain monounsaturated moieties. The composition of the free fatty alcohols was more variable than during the other seasons. We assume that if ingestion of wax ester-rich copepods is high, the free alcohol fraction in *M. ovum* reflects more the wax ester alcohol composition of the copepods. This confirms the hypothesis that there is a certain order in utilisation of fatty alcohols.

In August, the lipids in age-group 0 animals decreased, probably due to intensified reproduction (Lundberg et al. 2006), but lipids were still dominated by wax esters. The increase in the trophic marker fatty acid 18:4(n-3) clearly shows the changes in the lipid composition of the prey, changing from a spring to a more summer/autumn-like signature in both phyto- and zooplankton (Kattner et al. 1989; Leu et al. 2006). This is probably accompanied by a change in copepod composition from the dominance of *Calanus glacialis* to *C. finmarchicus*, associated with advection of transformed Atlantic water masses into the fjord (Basedow et al. 2004; Willis et al. 2006). *C. glacialis* is always rich in the 16:1(n-7) fatty acid (a marker for feeding on diatoms) but have never been reported to contain enhanced amounts of 18:4(n-3), a flagellate marker (e.g. Tande and Henderson 1988; Kattner et al. 1989; Scott et al. 2002; Wold et al. 2007). In contrast, both fatty acids can reach high amounts in *C. finmarchicus* and *C. hyperboreus*. The fatty acid 16:1(n-7) is generally enriched in copepods feeding in the spring bloom period, whereas 18:4(n-3) increases during summer and autumn feeding (Kattner et al. 1989). Very high proportions of 18:4(n-3) were also determined in the wax esters of *M. ovum* and *Beröe cucumis* during August/September 1997 in Kongsfjorden, but a study by Falk-Petersen et al. (2002) still assigned *C. glacialis* as the major prey. Towards autumn, *M. ovum* became lipid-richer probably replenishing lipids for overwintering. The highest proportions of the free fatty alcohol 22:1(n-11), accounting in maximum for almost 100% of the total free alcohols, occurred during this season. This indicates that this energy-rich component might be retained for winter survival instead of being egested.

Age-group 1 animals in spring, which have overwintered, still are of the same size as group 0 animals from the preceding autumn and probably do not grow owing to low prey abundance during winter (Lundberg et al. 2006). Roughly half of the lipid stores had been utilised during winter. Wax esters were the major storage lipid, but free fatty alcohols were strongly reduced. We found no clear indications of lipid accumulation in spring. Prey might be either utilised for growth or for early reproduction. The fatty acid composition also gives no hint regarding any special kind of prey or lipid accumulation. Until August, animals of group 1 grew fast but did not accumulate lipids. The lowest lipid content as percentage DM was found for these animals, which probably have invested most of their prey lipids into reproduction rather than into storage. They are probably not able to overwinter again although animals may be assigned to age-group 2 occurring the following spring. These animals had similar amounts of lipids as animals of age-group 1 and were not significantly different in their lipid compositions. Our data give no clue as to whether *M. ovum* has a life cycle of more than 2 years.

Acknowledgments We are grateful to the captain and crew of “Oceania”, RV Haakon Mosby and “Lance” and the staff of Kings Bay AS in Ny-Ålesund for their professional support during field experiments. We would like to thank Charlotte Gannefors, Anette Wold and Marthi Wolff for their important assistance. This work was supported by the Personnel Exchange Programme between the Research Council of Norway and Deutscher Akademischer Austauschdienst (DAAD).

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