

## Short term summer to autumn variability of dissolved lipid classes in the Ligurian sea (NW Mediterranean)

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**Abstract.** Changes in concentration and composition of *l*atrosca*n*-measured dissolved lipids were examined at a daily to month scale, in relation to the hydrological and biological context at a central site of the Ligurian sea, NW Mediterranean during the PECHÉ-DYNAPROC 2 experiment (14 September to 17 October 2004). Lipid concentrations (excluding hydrocarbons) (TLd-HC) and TLd-HC to DOC ratios in the 0–1000 m water column, varied from 5.3 to 48.5  $\mu\text{g l}^{-1}$  and 0.01 to 0.09, respectively. The highest TLd-HC concentration values were found in the 0–50 m surface layer, coinciding with phytoplankton biomass. Significant correlations ( $p < 0.01$ ,  $n = 87$ ) between glycolipids from chloroplast membranes, namely the monogalactosyldiacylglycerols, a major component of dissolved lipids (25.1  $\pm$  10.8% of TLd-HC,  $n = 166$ ), and various phytoplankton pigments (chlorophyll *cs*-170, violaxanthin, diadinoxanthin, zeaxanthin, and lutein), suggested that picoeucaryote phytoplankton were a major source of dissolved lipids. Lipid metabolites (free fatty acids, alcohols, diacylglycerols and monoacylglycerols), an other important component of TLd-HC (37.6  $\pm$  11.1%,  $n = 166$ ), showed a greater degree of degradation of lipids in this transitional period than previously observed earlier in the year. Zooplankton wax and steryl ester biomarkers (WSE) and triacylglycerols showed a distinct periodicity in the mesopelagic layer throughout the

period investigated. Concentrations of WSE (5.5–13.6  $\mu\text{g l}^{-1}$ ) increased in the 0–150 m surface layer, mid-way through the cruise (4–6 October), before the winter mixing. WSE were observed later and deeper in the mesopelagic layer (6–11 October), accompanied by rebounds in hydrocarbons (6–8 October) and phospholipid concentrations (12 October) in the 400–1000 m depth layer. Zooplankton migration and/or fecal pellet egestion, followed by DOM release from POM, were likely responsible for the appearance of these lipid signatures in the mesopelagic layer. Because we observed these signatures during low wind period only (<15 knots: 28 September–12 October), it may indicate that this organic matter transfer to depth was related to undisturbed trophic web in the water column above. The low salinity water lenses that appeared twice during the cruise in the 40–80 m surface layer had little effect on dissolved lipid concentrations. Lower concentrations in phosphoglycerides and hydrocarbons (HC) than in nearby sea water suggested different microbial assemblages and different level of HC contamination in this low salinity water.

### 1 Introduction

Marine dissolved organic matter (DOM) is an essential reservoir for the carbon cycle on Earth (Hansell and Carlson, 2002). In the ocean upper layer, photosynthesis yields organic material from inorganic carbon. Upon cell death, the



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photosynthetically produced substances are released to the medium as dissolved compounds or particulate detritus that may be further transformed into DOM. Thus, primary production is one of the most important source of DOM even though DOM can be produced at each step of the food web. Proteins, carbohydrates and lipids are the major constituents of the fraction of DOM characterized at molecular level. Considerable attention has been given to this easily identifiable fraction because it is biologically labile, it contains bioavailable moieties of biogeochemical interest like phosphorus and nitrogen and is a major substrate for marine bacteria. In addition, they may be used as biogeochemical markers to track the source, fate and transport of freshly biosynthesized DOM from the euphotic layer through the mesopelagic domain, under environmental forcings. Hence, despite the low percentage of these compounds (less than 11%) in the DOM surface pool (Benner, 2002), understanding variations in concentrations, resistance to remineralization and transformation to more complex and refractory substances in the water column, is essential to better depict the process involved in the sequestration of photosynthetically-produced organic carbon. Although DOM complex molecular assemblages would be better candidate for further characterisation (Mopper et al. 2007), the distribution of peptides, carbohydrates and lipids has not yet been clearly documented in the marine environment mainly due to limitations in methodologies.

Lipids represent from a few percent to 40% of the easily identifiable carbon fraction of DOM. They are membrane compounds, reserve products or metabolites from anabolism or catabolism in plankton cells (Parrish, 1988). Qualitative changes in their composition can be studied through the variability of selected lipid classes (Parrish, 1988), namely the cell membranes phospholipids and sterols from living organisms, the chloroplast membrane lipids (pigments and glycolipids) from autotrophic cells and phytodetritus, the reserve lipids from zooplankton (wax esters and triacylglycerols) and phytoplankton (triacylglycerols) and the degradation metabolites (free fatty acids, alcohols, di- and monoacylglycerols) of these source lipids, indicating the degradation status of the organic material (Goutx et al., 2000, 2003, 2005). They are released from living organisms through exudation, cell lysis or grazing, fueling the dissolved pool, and are further used by heterotrophic bacteria and/or exported to depth. The constitutive fatty acids of various lipids have been extensively used as biogeochemical markers of sources and diagenetic process of organic matter in suspended, sinking particles, and sediments (Saliot et al., 1991; Wakeham et al., 1995). Comparatively, dissolved lipid classes concentrations in the marine environment are poorly documented.

Dissolved lipids are operationally defined by filters that separate suspended particles from material including dissolved lipids, colloids, viruses and bacteria (Liu et al., 1998). Intact dissolved lipid classes have been studied in the extracellular medium of microalgae cultures to evaluate their pro-

duction of food web substrates (Kattner et al., 1983; Parrish and Wangersky, 1987; Lombardi and Wangersky, 1991), to characterize extracellular substances that may be responsible for the toxicity of phytoplankton blooms (Parrish et al., 1994) and to identify tensio active substances in bacterial culture growing on hydrophobic substrates (Goutx et al., 1987, 1990). Recently, the need to find new biomarkers in the polar lipid fatty acids (glycolipids, phospholipids), and to understand the potential role of phospholipids in phosphorus cycling, has revived research on intact polar lipids and revealed the lack of data, on phospholipids in particular, and other intact dissolved lipids, in the marine environment (Suzumura, 2005). Comprehensive studies of the distribution and dynamics of dissolved lipid classes have mainly focussed on the transport of lipid DOM by rivers, estuaries and urban discharge at land/ocean interfaces (Parrish, 1987; Leveau et al., 1990; Jaffé et al., 1995; Derieux et al., 1998; Mannino and Harvey, 1999; Rütters et al., 2002). Only a few studies have reported on the distribution of dissolved lipid classes in the open sea (Parrish et al., 1988; Goutx et al., 1990; Gérin and Goutx, 1994; Liu et al., 1998).

Analytical limitations due to low lipid class concentrations in the open ocean are a major obstacle to the study of dissolved lipids in marine systems. The traditional methods of thin-layer or column chromatography, require a significant amount of material, which can be problematic in field studies. The TLC/FID technique involving an "Iatroscan" apparatus allows detection of low concentrations of lipids without prior splitting of the extract, and opens very interesting prospects by removing a major obstacle to the study of neutral and polar lipid classes in the marine environment. This analytical technique involves a qualitative separation of the lipid extract by thin-layer chromatography on Chromarods coupled to a quantification of separated compounds by flame ionization detection. Developments in the Iatroscan lipid analytical protocol have been proposed by Delmas et al. (1984), Gérin and Goutx (1993), Parrish et al. (1996) and Striby et al. (1999). As this methodology requires a small extract for lipid class separation compared to current methodologies, it requires less seawater and therefore seemed to be a priori a technique of choice enabling the exhaustive sampling required for describing short temporal or spatial scale variability of chemical organic species in marine ecosystems.

The project PECHE (Production and Export of Carbon: control by Heterotrophic organisms at short temporal scale) and its main operation, the oceanographic campaign DYNAPROC 2 (DYNAmics of rapid PROCesses), was designed by Valérie Andersen with the specific goal of assessing variability in the pelagic food web and its effect on biogeochemistry of the mesopelagic zone, targeting an oligotrophic system, the NW Mediterranean Sea at the end of the summer 2004. In the present study, using TLC/FID, we examined the short term changes in abundance and composition of neutral and polar dissolved lipid classes in the 0–1000 m water column. By processing a large set of ~90 samples at daily and

monthly time scales at a fixed station in the open Ligurian sea, our overall objective was to better understand relationships between physical and biological forcings, and the short time variability of the organic chemical content in the water column, during the summer oligotrophy to autumn transitional period, in Northwestern Mediterranean.

## 2 Material and methods

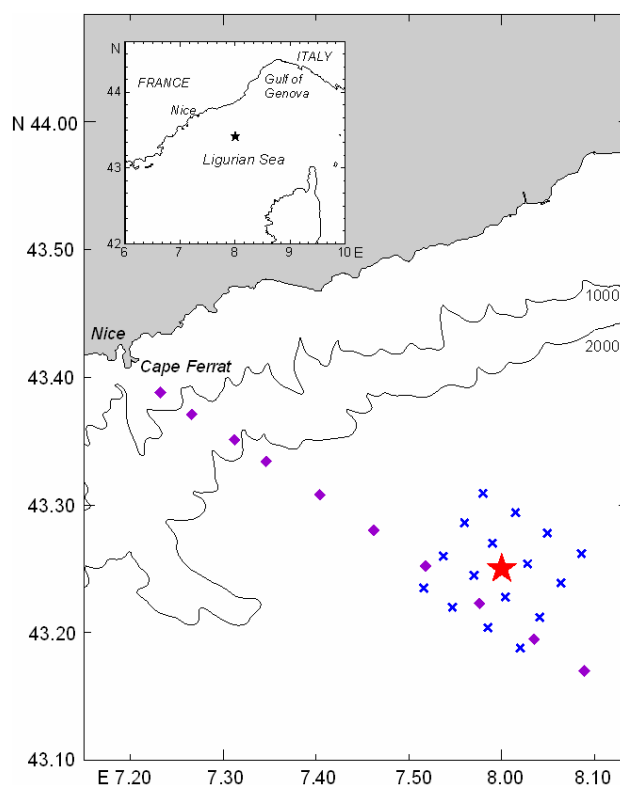
### 2.1 Study area and sample collection

Samples were collected at a Time Series Station (TSS, 43°25'N, 8°E; ca. 48 km offshore) in the Ligurian sea (North-Western Mediterranean) during the 2004 DYNAPROC 2 cruise (14 September to 17 October) on board the O/V *Thalassa* (Fig. 1). The dynamics of the biological system, in particular the phyto-, microzoo- and macrozooplankton biomasses and bacterioplankton, were studied in the 0–1000 m water column during four cycles of 5 days each (Andersen et al., 2009), two cycles during Leg 1 (C1 and C2; 17–29 September) and two cycles during Leg 2 (C3 and C4; 3–16 October). Sampling frequency was 3 h or 12 h depending on the parameter. For dissolved lipids, a minimum of two 0–1000 m CTD profiles were sampled at the beginning of each cycle (two profiles at C1, C2 and C4, and five profiles at C3) at 12 h interval (one around midnight and one around midday). Additional samples for lipid analysis were taken on two 0–150 m surface CTD/rosette profiles at C1, and at the sub-surface and chlorophyll maximum depths during the whole cruise, in order to obtain a continuous pattern of dissolved lipids in the surface layer (see the Supplement Table for the complete list of samples <http://www.biogeosciences.net/6/1229/2009/bg-6-1229-2009-supplement.pdf>).

### 2.2 Dissolved lipid analysis

Water was transferred from the Niskin bottles with a Teflon tube into 5 litre glass bottles previously washed with 2 M HCl and rinsed with MilliQ water. Four liters were gently filtered (pressure < 15 mm Hg) onto pre-combusted (6 h at 450°C) 47 mm glass fiber filters (GF/F) using all-glass cylinders and filtration kits to separate suspended particles from dissolved organic material. All glassware was previously washed with RBS detergent, 2 M HCl, then rinsed with MilliQ water and combusted for 6 h at 450°C or rinsed with acetone.

After adding the internal standard (I.S.) (100 µl of an hexadecanone solution at 0.2 mg ml<sup>-1</sup> in dichloromethane), dissolved fractions (2 l) were liquid-liquid extracted on board using 4 times 100 ml dichloromethane (Rathburn Chemical Ltd, HPLC grade) (2 times at pH < 2). After phase separation, the organic phases were combined, the volume reduced with a rotary evaporator and stored at -18°C under nitrogen atmosphere until analysis. The recovery of dissolved lipids was 70% ± 13 (n=167) based on the ketone internal standard. In order to examine the possibility of interference between



**Fig. 1.** Location of the time series station (★) sampled during the Dynaproc2 cruise, and grid of stations that enabled to characterize the physics and hydrology of the area.

I.S. and natural ketones that might affect our extraction rates, a few additional samples were extracted without addition of the I.S. during extraction. In the particular case of these dissolved lipid extracts, in our analytical conditions, material at retention time similar to that of the I.S. remained below the detection level.

Dissolved lipid classes were separated on Chromarods by thin layer chromatography and quantified with flame ionisation detection (TLC/FID) in an Iatrosan apparatus model MK-6 s (hydrogen flow, 160 ml min<sup>-1</sup>; air flow, 2000 mL min<sup>-1</sup>) coupled to a compatible PC equipped with an i-Chromstar 6.1 integration system (SCPA-Bremen, Germany). Samples (usually 4–6 µl of a 30 µl solution of the lipid extract in dichloromethane) were spotted using a 2 µl Hamilton syringe onto Chromarods-SIII previously calibrated with standard compounds (Sigma Chemical Ltd, GC grade). Analyses were run in triplicate.

The separation scheme involved five elution steps in solvents of increasing polarity: hexane+diethyl ether+formic acid, acetone, chloroform+methanol and chloroform+methanol+ammonium hydroxide according to a modified procedure of Goutx et al. (1990) described in Striby et al. (1999). The use of these elution systems separated neutral lipid classes (hydrocarbons, sterol esters co-eluting

with wax esters, ketone as internal standard, triacylglycerols, free fatty acids, alcohols, sterols and diglycerides), chloroplast lipids (pigments, glycolipids) and monoglycerides, and non-nitrogen containing phospholipids (diphosphatidylglycerides co-eluting with phosphatidylglycerides) from nitrogen containing phospholipids (phosphatidylethanolamine and phosphatidylcholine). The relative standard deviation of replicate samples ( $n=3$ ) for Iatroscan TLC-FID analysis was below 12%. Detection limit was  $0.02 \mu\text{g L}^{-1}$ . Total dissolved lipids concentration (TLD-HC) was calculated by the summing of all lipid classes except hydrocarbons. As the biogenic origin of hydrocarbons could not be ascertained here, this lipid class is not included in TLD-HC that are thus referred to as biogenic lipids. Hydrocarbon dynamics were then treated separately. Molarity was calculated on the basis of the molecular weight of the reference standard for each lipid compound class.

### 2.3 Dissolved organic carbon

The samples were filtered through 2 precombusted (24 h,  $450^\circ$ ) glass fiber filters (Whatman GF/F, 25 mm), collected into precombusted glass tubes (closed with a screw cap and a Teflon liner), acidified with Orthophosphoric acid ( $\text{H}_3\text{PO}_4$ ) and immediately analysed on board on a Shimadzu TOC-V analyser. Dissolved organic carbon (DOC) was analyzed by high temperature catalytic oxidation (HTCO) (Cauwet, 1994, Cauwet 1999). Typical analytical precision is  $\pm 0.1$ – $0.5$  (SD) or  $0.2$ – $1\%$  (CV). Deep Sargasso Sea reference water ( $47 \mu\text{M}$ ,  $\pm 0.5$  SE, <http://www.rsmas.miami.edu/groups/biogeochem/CRM.html>) was injected every 10 to 12 samples to insure stable operating conditions.

### 2.4 Ancillary parameters

Meteorological (wind, rain), hydrological (temperature, salinity, density), biogeochemical (nutrients, oxygen, colored dissolved organic matter and pigments) and biological (bacteria, flagellates and ciliates, microphytoplankton, pico and nanophytoplankton, meso- and macrozooplankton) parameters were measured during this multidisciplinary cruise. CTD profiles were performed by using a Seabird SBE 25 probe measuring temperature, salinity, pressure, fluorescence,  $\text{O}_2$  and irradiance. Composition of phytoplankton was determined on the basis of a fine HPLC resolution of chlorophyll and carotenoid pigments, which for example included all chlorophylls, i.e. chl-*a* and divinyl chl-*a*, chl-*b* and divinyl chl-*b*, and chl-*c* including c1, c2 and cs-170 (Marty et al., 2008). Colored dissolved organic matter (CDOM) substances were extracted from 500 ml sea water samples by adsorption on C18 microcolumns. CDOM was eluted from the microcolumn with methanol and quantified by light-absorbance at 412 nm. Data are available on the web site, in previously published papers (Marty et al., 2008) and in companion papers in the present BGS volume (Ghiglione et al.,

2008; Lasternas et al., 2008; Mevel et al., 2008; Mousseau et al., 2009; Raybaud et al., 2008, 2009)

### 2.5 Statistics

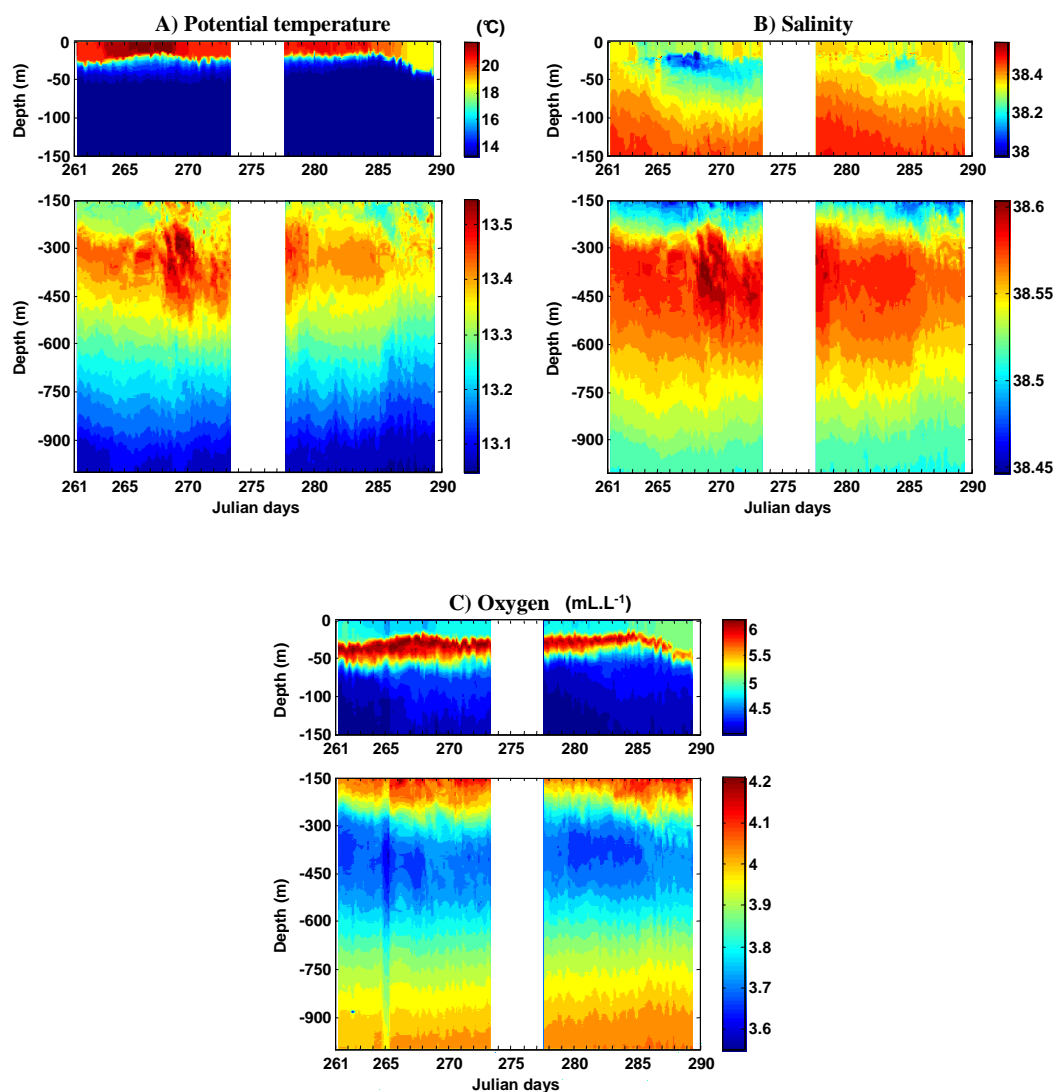
Correlation analyses between lipid classes and ancillary parameters were conducted using the statistics package of Microsoft Excel (Microsoft Office Professional Edition 2003). Pearson's correlation coefficients were determined for pairwise comparisons of total lipids, individual lipid classes and other environmental parameters (depth, salinity, temperature, oxygen, nutrients, DOC, particulate carbon, CDOM, Chl-*a*, phytoplankton pigments) with a significance level of  $P < 0.05$  assessed by Student's two-tailed *t*-test being considered significant. Values of  $P < 0.001$  were considered highly significant. The significance of mean comparison was assessed by ANOVA using the StatView 5 software.

## 3 Results

### 3.1 General hydrological conditions

The main hydro-biological characteristics during the cruise are presented in Fig. 2. Briefly, the DYNAPROC 2 cruise was conducted during the seasonal transitional period between summer and fall, a period marked by a decline in air temperature and the collapse of the summer water column stratification due to successive meteorologically forced mixing events. The cruise extended for two legs, 14–29 September (Leg1) and 2–17 October (Leg2). Concentrations of nutrients were low throughout the cruise (Marty et al., 2008). During Leg 1, two wind events from the NE occurred (17 and 25 September). At the end of the cruise, a succession of wind gusts ( $>20$  Knots) from opposite directions: SW, NE and SW occurred over periods of 24 h, 60 h and longer until the end of the cruise from 10, 11 and 16 October, respectively. Temperature profiles showed a highly stratified water column from the beginning of the cruise to 10 October, being partially disrupted at the end of the cruise (Fig. 2a). The thermocline was strongly marked, with a mixed-layer temperature higher than  $20^\circ\text{C}$  ( $22^\circ\text{C}$  during weak wind periods). This thermocline was located at approximately 25 m depth throughout the cruise. At the end of the cruise, the thermocline deepened to 40 m depth after the last two successive strong wind events accompanied by a decrease of air temperature and a cooling of the mixed water layer, which suggested the beginning of fall mixing.

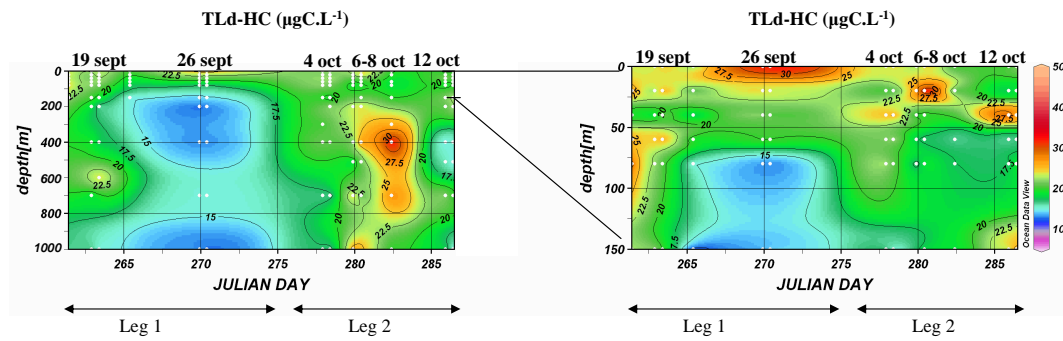
Two intrusions of low salinity water masses (LSW) ( $<38.3$  psu) occurred below the thermocline (between 40 and 80 m) (Fig. 2b). The first intrusion (LSW-1) lasted 5 days from Julian day (JD) 265 (21–30 September) and the second intrusion (LSW-2) lasted for 4 days from JD283 (9–12 October). The coastal LSW did not affect the depths exceeding 300 m. In 2004, the Levantine water was much more noticeable at the DYFAMED site than in previous years



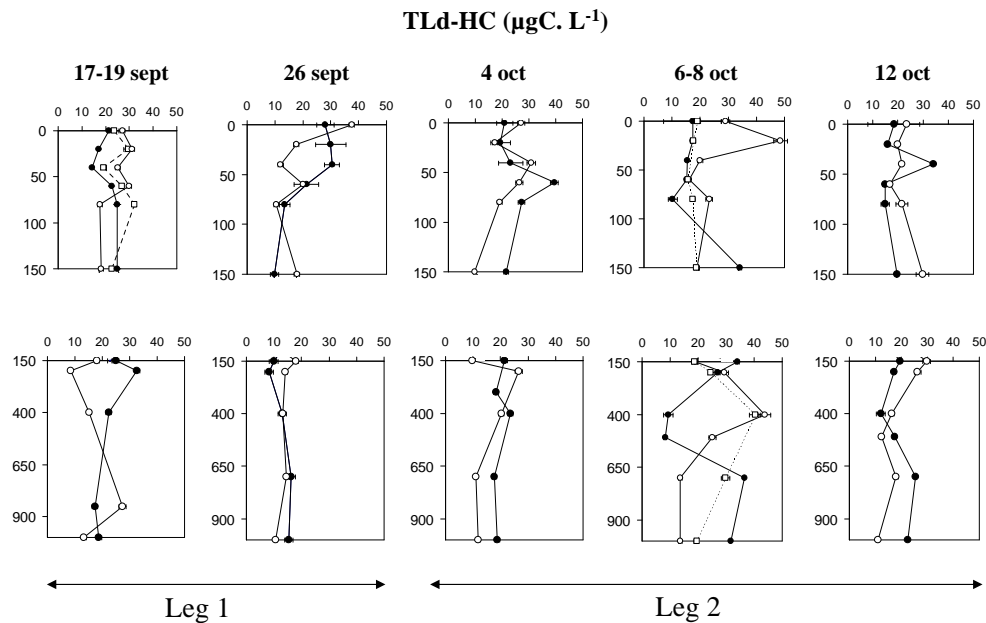
**Fig. 2.** Potential temperature ( $^{\circ}\text{C}$ ), salinity (p.s.u.) and oxygen ( $\text{mL L}^{-1}$ ) at the Dynaproc 2 time series station ( $43^{\circ}25\text{ N}$ ,  $8^{\circ}\text{ E}$ ) during the late summer to autumn transitional period.

(1970–1995) and remained at levels between 250 to 400 m depth in contrast to the frontal zone where it is found between 350 and over 600 m depth, taking salinity  $>38.54$  and temperature  $\theta >13.4^{\circ}\text{C}$  as a criterion for Levantine water at depth (below the temperature minimum of the surface layer). In 2004, the minimum of oxygen that characterized the Levantine water was  $\text{O}_2 <3.8\text{ mL L}^{-1}$  (Fig. 2c). Variations in the thickness of the Levantine water during the DYNAPROC2 cruise remained within the bounds of small-scale spatial variability in the center of the Ligurian Sea. At the beginning of the cruise, two deep-chlorophyll maxima (DCM, 50–60 m and 90 m depth) were detected, displaying a phytoplankton biomass exceptionally high for this time of the year (integrated Chl-*a* concentration of 35–40  $\text{mg m}^{-2}$ ). After two days at the fixed station, only one

DCM (19 September, JD263), was observed at 40–50 m depth with a 0–150 m integrated value Chl-*a* concentration of 20–25  $\text{mg m}^{-2}$  (Marty et al., 2008). The integrated primary production was 200  $\text{mg C m}^{-2}\text{ d}^{-1}$  on average, varying from 400  $\text{mg C m}^{-2}\text{ d}^{-1}$  at the beginning of the cruise (thus 50% higher than usually observed), down to 100  $\text{mg C m}^{-2}\text{ d}^{-1}$  at the end of the cruise. The overall characteristics of the site show that the DYNAPROC2 cruise covered a period of the transitional regime between late summer and autumn (Andersen et al., 2009), but with hydrological and nutrient resources (N/P) similar to summer oligotrophic conditions (Marty et al., 2008).



**Fig. 3.** Concentrations of total dissolved lipids (hydrocarbons not included) ( $\mu\text{gC.L}^{-1}$ ) in the 1000 m (left) and in the surface 0–150 m (right) water column. Extrapolation ODV X and Y scale length 150.

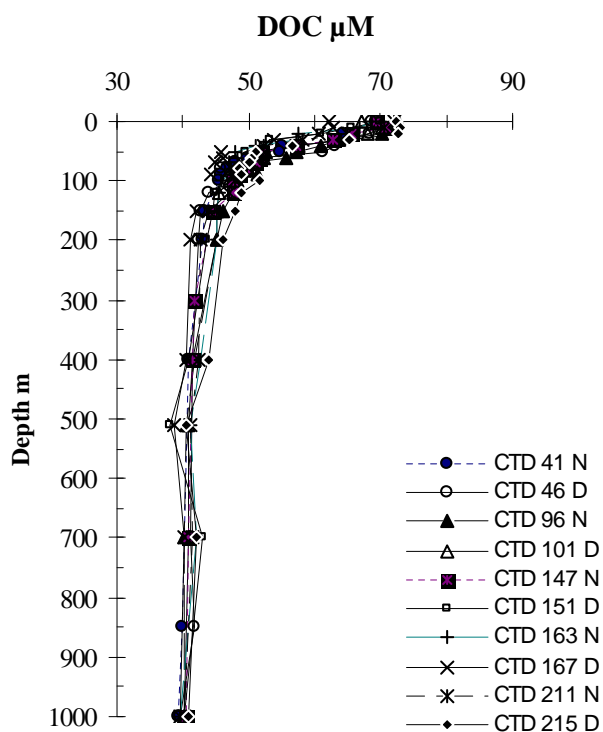


**Fig. 4.** Profiles of total dissolved lipids (hydrocarbons not included) ( $\mu\text{gC.L}^{-1}$ ) in the 0–1000 m of the water column. Days (open symbols) and nights (filled symbols). Two additional day profiles (17 September and 8 October) are in dotted line.

### 3.2 TLd-HC, TLd-HC to DOC ratios

Lipid concentrations, hydrocarbons not included (TLd-HC), varied from  $5.3$  to  $48.5 \mu\text{gC.L}^{-1}$  ( $0.4$  to  $4 \mu\text{M}$ ) (See the supplemental data table <http://www.biogeosciences.net/6/1229/2009/bg-6-1229-2009-supplement.pdf>). The highest values were found in the upper 50 m of the water column (with exception of the data collected 19 September and 4 October) (Fig. 3). The concentration gradient with depth was more pronounced during Leg1, with TLd-HC values  $\leq 18 \mu\text{gC.L}^{-1}$  below 100 m, than during Leg 2 when TLd-HC concentrations below the surface layer were up to  $40 \mu\text{gC.L}^{-1}$  (Fig. 3). An increase of concentrations in the mesopelagic layer occurred twice during the cruise, shortly at the beginning of Leg 1 ( $27.4 \mu\text{gC.L}^{-1}$ , 19 September) at 600 m depth, and sev-

eral times during Leg 2 (up to  $43.9 \mu\text{gC.L}^{-1}$ , between 6 and 12 October) in the 400–800 m layer. There was no clear day/night periodicity pattern in the TLd-HC profiles throughout the period investigated (Fig. 4): in the 0–150 m surface layer, peaks were observed at night in 3 over the 5 day/night profiles (26 September, 4 and 12 October). During leg 2 below 500 m, TLd-HC concentrations were higher during the night than during the day (Fig. 4). DOC varied by a factor of 2 from surface to depth in the range  $38.6$ – $73.8 \mu\text{M}$ . Concentrations ( $70 \pm 2.5 \mu\text{M}$ ) were maximal in the 0–10 m subsurface waters, then decreased following a moderate gradient down to  $50.7 \pm 2.1 \mu\text{M}$  in the 60–70 m water layer and were almost homogeneous down to a minimum value at 1000 m depth ( $39.3 \pm 0.6 \mu\text{M}$ ). There was no apparent difference between days and nights (Fig. 5). The contribution of TLd-HC

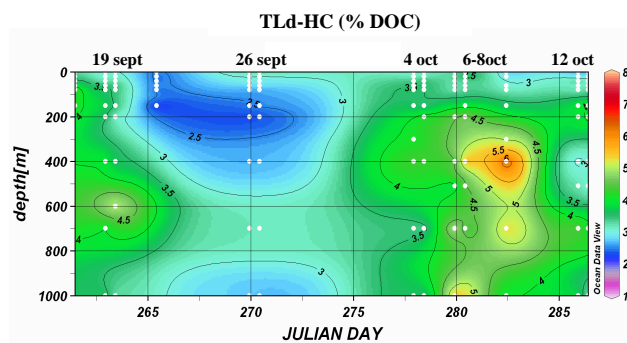


**Fig. 5.** Dissolved organic carbon concentrations ( $\mu\text{M}$ ). N=night, D=day.

to DOC varied in the range 0.9–9.1% (Fig. 6). The highest values ( $\geq 7\%$ ) were observed within the 400–800 m depth layer, during the Leg 2 on 6 October during day and night, and 8 October during the day only, a pattern due to the increase of lipid concentrations while DOC concentrations remained quite stable. An increase in the percentage of TLd-HC in DOC ( $>5\%$ ) was occasionally observed on a day profile, during Leg 1 (19 September at 700 m, Julian day 263). In order to examine the lipid pattern in DOC at a seasonal scale, the average concentrations of DOC and total lipids were computed in the 0–150 m surface and the 150–1000 m mesopelagic layers during Leg 1 and 2 (Table 1). Overall, there was an increase in the contribution of dissolved lipids to DOC in the mesopelagic layer from an average  $3.3 \pm 1.1\%$  in September (Leg 1) to  $4.3 \pm 1.8\%$  in October (Leg 2) (ANOVA,  $p < 0.05$ ,  $n = 31$ ), whereas the TLd-HC/DOC ratio in the 0–150 m surface layer was similar ( $2.9 \pm 1.1\%$  and  $3.1 \pm 1.1\%$ , during Leg 1 and Leg 2, respectively). There were no obvious differences in the daily variations of TLd-HC contribution to DOC (data not shown).

### 3.3 Lipid classes

Lipid class concentrations are presented in supplemental Table 1 <http://www.biogeosciences.net/6/1229/2009/bg-6-1229-2009-supplement.pdf>. The relative contribution of phospholipids, chloroplast lipids (glycolipids and

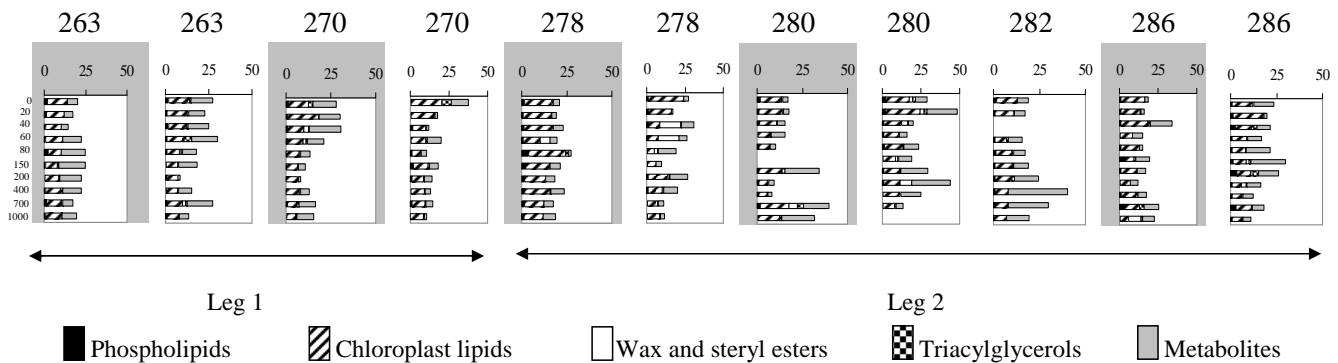


**Fig. 6.** Relative percentage of total dissolved lipids (hydrocarbons not included) in DOC. Extrapolation ODV X et Y scale length 150.

pigments), triacylglycerols (TG), wax and steryl esters (WSE) and metabolites (free fatty acids, alcohols, di- and monoacylglycerols) to the lipid pool showed a quite stable pattern during the sampling period (Fig. 7) with the exception of an increase of WSE and TG contribution to TLd-HC, from  $5.1$  ( $n=1$ ) to  $15.1 \pm 11.7\%$  ( $n=8$ ) and from  $4.1 \pm 1.8$  ( $n=12$ ) to  $4.0 \pm 3.0\%$  ( $n=19$ ) (data computed from supplemental Table 1 <http://www.biogeosciences.net/6/1229/2009/bg-6-1229-2009-supplement.pdf>), respectively, beyond 200 m depth in the mesopelagic layer between Leg 1 and Leg 2. Table 2 provides the average concentration and percentage of lipid classes in TLd and TLd-HC during the whole cruise (all depths). Except for hydrocarbons, the glycolipids (monogalactosyl diacylglycerols) (25.09%) were the major lipid class, followed by pigments (13.58%), free aliphatic alcohols (12.34%), the monoacylglycerols (11.84%) and free fatty acids (11.38%), and finally the phospholipids (PL). Thus the so-called chloroplast lipids, which include glycolipids and pigments, varied between  $2.1 \mu\text{gCL}^{-1}$  and  $21.9 \mu\text{gCL}^{-1}$  (see supplemental Table 1 <http://www.biogeosciences.net/6/1229/2009/bg-6-1229-2009-supplement.pdf>) and dominated the lipid pool ( $38.7 \pm 8.5\%$  on average,  $n=166$ ) (Table 2), which suggested a phytoplankton source for DOM. The highest values were measured in the 0–80 m upper layer, at the beginning of Leg 2. High values (ca.  $10\text{--}14 \mu\text{gCL}^{-1}$ ) were also found at depth in the 400–800 m layer (6 and 8 October) (see supplemental Table 1 <http://www.biogeosciences.net/6/1229/2009/bg-6-1229-2009-supplement.pdf>). The other most important class of compounds were the lipid metabolites (FFA, ALC, MG and DG) ( $37.6 \pm 11.1\%$  on average,  $n=166$ ) (Table 2 and supplemental Table 1 <http://www.biogeosciences.net/6/1229/2009/bg-6-1229-2009-supplement.pdf>), indicating hydrolysis of lipid DOM. The average concentration of each lipid class per depth (data not shown) showed that free fatty acids had a maximal average concentration ( $5.2 \pm 0.7 \mu\text{gCL}^{-1}$ ) at 400 m depth, at the bottom of the Levantine water body. However, we did not sample properly the Levantine waters (only one sample was taken at 400 m

**Table 1.** Average concentrations ( $\pm$ standard deviation) of dissolved organic carbon (DOC) and dissolved total lipids (TLd-HC) (hydrocarbons not included) and percentage of TLd-HC in DOC, in the water column (0–1000 m), the surface (0–150 m) and the mesopelagic layers (150–1000 m) during the two legs of the cruise, September (Leg 1) and October (Leg 2).

Dissolved Matter			DOC $\mu\text{M}$	TLd-HC $\mu\text{g C l}^{-1}$	TLd-HC/DOC %
Water column	0–1000 m	September	53.6 $\pm 11.1$ ( $n=62$ )	18.4 $\pm 7.24$ ( $n=72$ )	3.0 $\pm 1.2$ ( $n=62$ )
		October	52.4 $\pm 11.3$ ( $n=91$ )	21.2 $\pm 8.42$ ( $n=95$ )	3.5 $\pm 1.5$ ( $n=90$ )
Surface	0–150 m	September	57.9 $\pm 9.75$ ( $n=46$ )	19.6 $\pm 7.3$ ( $n=46$ )	2.9 $\pm 1.1$ ( $n=46$ )
		October	58.0 $\pm 10.1$ ( $n=63$ )	21.0 $\pm 7.9$ ( $n=64$ )	3.1 $\pm 1.1$ ( $n=63$ )
Mesopelagic	150–1000 m	September	41.3 $\pm 1.9$ ( $n=16$ )	16.3 $\pm 6.5$ ( $n=16$ )	3.3 $\pm 1.3$ ( $n=16$ )
		October	41.8 $\pm 1.8$ ( $n=32$ )	21.5 $\pm 8.9$ ( $n=31$ )	4.3 $\pm 1.8$ ( $n=31$ )



**Fig. 7.** Time series of TLd-HC ( $\mu\text{g L}^{-1}$ ) and dissolved lipid class distribution (% of TLd-HC) during night (■) and days (□) sampling. Julian days are above the graph.

depth in this water mass) and further characterisation of the lipid distribution in Levantine waters may not be performed straightforward. Other intact lipids (phospholipids, triacylglycerols, wax and steryl-esters) were present in lower proportions (on average  $9.5 \pm 6.0\%$ ). Phospholipids (PL) were the least abundant lipid class in the concentration range  $0\text{--}3.7 \mu\text{gCL}^{-1}$  (mean  $1.0 \pm 0.6 \mu\text{gCL}^{-1}$ ,  $n=166$ ). A marked increase in PL concentrations was noticeable during leg2 at 150–200 m and 700 m depth (mean  $3.1 \pm 0.3 \mu\text{gCL}^{-1}$ ,  $n=4$ ) (Fig. 8, 12 October, upper panel). Finally, the hydrocarbon pool varied in the range  $0.9\text{--}21.9 \mu\text{gCL}^{-1}$  (on average  $6.3 \pm 1.8 \mu\text{gCL}^{-1}$ ), which represented  $24.7 \pm 3.2\%$  of the TLd (Table 2). An increase of concentrations at 400–800 m depth occurred in 6 and 8 October (Fig. 8, lower panel).

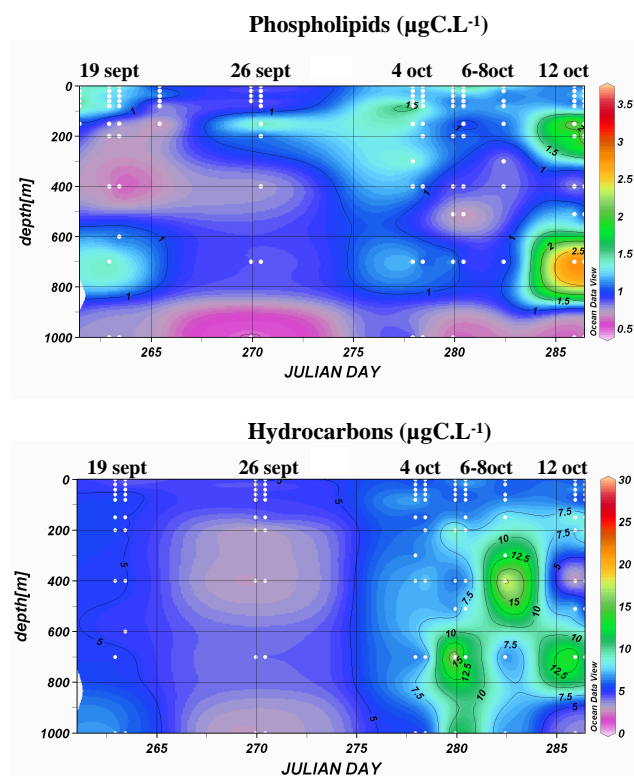
The storage lipids, triacylglycerols (TG) and wax and steryl esters (WSE), varied in the range  $0\text{--}6.0$  and  $0\text{--}13.6 \mu\text{gCL}^{-1}$ , respectively. They were minor compounds not always present in the total lipids. Triacylglycerols are re-

serve lipids in microalgae and zooplankton (Parrish, 1988). Wax and steryl esters cannot be easily separated by Iatroscan TLC/FID and there is no reports on wax versus steryl esters in dissolved lipids by using this technic. Steryl esters usually represent a small contribution to particulate WSE (usually less than 8%) (Wakeham et al., 1980; Hudson et al., 2001). However, it is not always the case as shown by the high concentration measured in mesopelagic sediment trap in the Atlantic ocean (Wakeham et al., 1980). Ciliated protozoa also, are able to synthesize quantity of steryl esters (Kaneshiro, 1987). Whatever their relative proportions, wax and steryl esters are related to the presence of zooplankton (Sargent et al., 1976; Sargent et al., 1977; Sargent, 1978; Cavaletto et al., 1989; Kattner, 1989). In our study, the pattern of variation of TG and WSE over time and depth was well described by comparing the 0–150 m and 150–1000 m average concentrations (Fig. 9). During Leg 1, TG night average concentration values were always lower at surface (0–150 m) than in

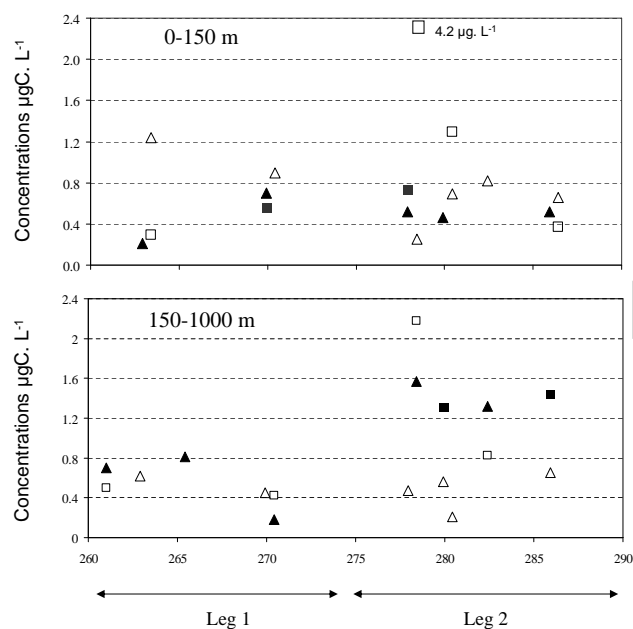


**Table 2.** Mean dissolved lipid class concentrations ( $n=167$ ) in the water column at the Dynaproc 2 central station during the summer to autumn transition period (14 September–17 October). Abbreviations used for lipid classes. Contribution of each lipid class to total dissolved lipids including HC (TLd) and contribution to total dissolved lipids (not including HC).

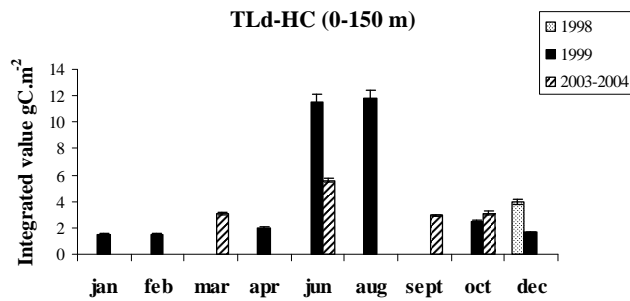
Lipid class	Abbreviation	Concentration		Contribution to TLd		Contribution to TLd-HC	
		$\mu\text{g L}^{-1}$	$\pm\text{SD}$	%	$\pm\text{SD}$	%	$\pm\text{SD}$
Hydrocarbons	HC	6.33	1.81	24.69	3.25		
Wax and steryl esters	WE	0.53	1.08	2.10	1.51	2.32	1.61
Triglycerides	TG	0.68	0.31	2.60	0.84	2.94	1.14
Free fatty acids	FFA	2.62	2.00	10.10	3.65	11.38	4.62
Free aliphatic alcohols	ALC	2.84	0.89	11.10	2.42	12.34	2.57
1.3 Diglycerides	1.3DG	0.08	0.34	0.32	0.39	0.37	0.49
Free sterols	ST	0.01	0.08	0.04	0.10	0.05	0.11
1.2 Diglycerides	1.2DG	0.01	0.06	0.04	0.06	0.05	0.08
Chlorophyll pigments	PIG	3.13	1.08	12.17	2.09	13.58	2.45
Monoglycerides	MG	2.73	1.36	10.64	2.85	11.84	3.37
Monogalactosyldiglycerides	MGDG	5.78	0.94	22.38	4.60	25.09	6.11
Phosphoglycerides	PG	0.86	0.23	3.31	0.80	3.71	1.00
Phosphatidylethanolamines	PE	0.11	0.05	0.41	0.11	0.46	0.12
Phosphatidylcholines	PC	0.01	0.02	0.04	0.03	0.04	0.03



**Fig. 8.** Distribution of specific lipid markers ( $\mu\text{gC L}^{-1}$ ). Dissolved phospholipids (upper panel), and hydrocarbons (lower panel). Extrapolation ODV X and Y scale length 150.



**Fig. 9.** Triacylglycerols (triangle) and wax esters (square) average concentrations ( $\mu\text{gC L}^{-1}$ ) based on integrated concentrations in the 0–150 m surface (upper panel) and mesopelagic 150–1000 m (lower panel) layers. Day (open symbols) and night (filled symbols).



**Fig. 10.** Composite of total lipid concentrations measured in the framework of the Dyfamed time series observations (1998-1999) and in the framework of the PECHE project (March and June, 2003).

the mesopelagic layer, if we exclude night value observed at julian day 278. An opposite trend was observed for the TG day values, they were always higher at surface than in the mesopelagic layer. WSE occasionally appeared in the lipid pool during Leg 1 (Fig. 9). Like TG, WSE night values were lower at surface than in the mesopelagic layer but the day pattern was less obvious.

### 3.4 Relationships between total lipids, individual lipid classes, and hydrological and biogeochemical parameters

Correlations between bulk organic compounds (DOC, particulate carbon, CDOM), TLd and TLd-HC, individual lipid classes, and environmental parameters (salinity, temperature, nutrients, Chl-*a*, phytoplankton pigments) were examined using Pearson's product-moment correlation coefficient. Correlations were computed on two matrices, a 0–150 m matrix including all parameters cited above, and a 0–1000 m matrix not including pigments because they were analyzed in the surface water column only. In the 0–1000 m water column, significant correlations were found between total dissolved lipids (both TLd and TLd-HC) and concentrations of hydrocarbons, triacylglycerols, chloroplast lipids and metabolites (free fatty acids and alcohols) ( $p < 0.01$ ,  $n = 167$ ) whereas the minor lipid classes (wax esters and phospholipids) were not correlated with total lipids (data not shown).

As chloroplast lipids (CL) were the major components of the lipid pool, the correlation between TLd-HC and CL suggests that phytoplankton were one major source of lipids in the 0–1000 m water column. In the 0–150 m surface water column, the monogalactosylglycerides (MGDG), which are constituents of CL in microalgae, were the only class of lipids that was significantly correlated with several individual pigments (chlorophyll *cs*-170 (0.54,  $p < 0.01$ ), violaxanthin (0.43,  $p < 0.01$ ), diadinoxanthin (0.47,  $p < 0.01$ ), zeaxanthin (0.52,  $p < 0.01$ ), and lutein (0.49,  $p < 0.01$ )) which all together support the idea of a picoplanktonic sources of

MGDG in the dissolved lipid pool (Table 3). Such correlations between one component of the dissolved lipid fraction and pigments that are located inside the cells, suggest that a direct transfer of MGDG from the picoplankton intracellular lipid pool to dissolved lipids possibly occurred. MGDG was also correlated with DOC and particulate carbon (PC) (0.57 and 0.45,  $p < 0.01$ ) (Table 3). This suggests that both PC and DOC contained a similar picoplankton material as TLd and/or that a PC to DOC transfer through picoplankton lysis and grazing is possible. In contrast, MGDG and coloured dissolved organic matter (CDOM) were not related to each other. CDOM, unlike MGDG, was not related to DOC, but was correlated with particulate carbon ( $p < 0.05$ ,  $n = 81$ ), total fluorescence and almost all individual pigments including the five picoplankton pigments cited above (Table 3) indicating different CDOM sources from that of DOC.

## 4 Discussion

Our study provides the first high frequency time series measurements of total lipid concentrations and lipid class composition in DOM from the surface to the mesopelagic layer in the open Ligurian sea, which represents a reference for the late summer to autumn period in North Western Mediterranean. The amount of lipids that we measured in 2004 were in the lower range of values previously reported for marine coastal inlets and open sea areas ( $11\text{--}580 \mu\text{g L}^{-1}$ ) (Table 4).

### 4.1 Seasonal variations of the total lipid pool

Previous measurements of dissolved lipid concentrations (unpublished data; Ghiglione et al., 2007; Bourguet et al., 2009) at the Dyfamed TSS (Marty et al., 2002) during the 1998–1999 years, and during two additional PECHE short-cruises conducted in March and June 2003 (Andersen et al., 2009), enable to extend the temporal scale of the average lipid class concentrations measured in the 0–150 m euphotic layer during the Dynaproc2 cruise to an annual time series (Fig. 10): dissolved lipid concentrations were generally low in winter, maximal during summer oligotrophic period (June–August 1999 and June 2003) and returned to low values in autumn. Interestingly, this pattern suggests that lipid accumulations that occurred during summer oligotrophy (Van Wambeke et al., 2001; Bourguet et al., 2009) did not persist at the surface until the winter mixing. During late summer and autumn, lipids may be either remineralized, transformed through biotic and abiotic process into another type of material which is not captured by our analytical methodology, or exported to depth by other means than the mixing of water masses, namely through aggregation/dissolution and migration of zooplankton.



to bacterial attack. Degradation rates have been reported by several authors (Harvey et al., 1995, 1997; Goutx et al., 2003, 2007; Caradec et al., 2004; Moriceau et al., 2008). Goutx et al. (2003) and Moriceau et al. (2008) reported that as soon as mineral dissolution started, MGDG turned over more rapidly than the other AMPL lipids, i.e. the monoglycerides and pigments. Hence, due to differences in residence times, lipid classes resulting from diagenetic processes may have masked the origin of dissolved lipids.

In our study, the AMPL peak was fractionated into its various components (Striby et al., 1999), which may be the cause of the significant relation observed between a dissolved lipid class, the glycolipids (i.e. MGDG), and specific picoplankton pigments. This relation points out the picoeucaryotes as a major source of dissolved lipids during the period investigated.

One other reason for the lack of significant relationships between Chl-*a* and dissolved lipids in our study and in previous works, may be that Chl-*a* and AMPL that are enriched in large microphytoplankton cells, in particular diatoms, are preferentially exported to depth by aggregate formation through coagulation, and/or meso- and macrozooplankton grazing. Hence, large cells are less susceptible to breakdown through lysis and grazing in surface layers, having less potential for fueling the surface dissolved pool with intracellular Chl-*a* and chloroplast lipids than small picoplankton cells do. The small picoplankton which is rich in xanthopigments, is most likely to enter the microbial loop, which would favour the export of its intracellular compounds to the dissolved pool through sloppy feeding.

A striking feature of the relationship between phytoplankton lipid biomarkers and organic matter descriptors was the significant correlations between particulate carbon (PC), DOC, glycolipids (MGDG) and picoplankton pigments on the one hand, and PC, CDOM, Chl-*a*, and most pigments on the other hand (cf. Results Sect. 3.3 and Table 3). This pattern suggested differences in sources and formation processes of bulk DOC and CDOM during the summer to autumn transitional period. The dynamics of DOC concentrations would mostly be controlled by release of matter from picoplankton that dominated the phytoplankton biomass (Marty et al., 2008) at this time of the year, probably through lysis, grazing and sloppy feeding. CDOM would rather be formed through intramolecular reactions of a larger range of organic biomolecules from the decaying pico- and microphytoplankton, under a combination of biotic and abiotic factors including light radiation (Yentsch and Reichert, 1961; Sieburth and Jensen, 1969).

Below 60 m, the decrease of lipid concentrations suggested that stratification limited the transfer of phytoplankton lipids (and by extrapolation, of other phytoplankton material) from the productive layer to the mesopelagic layer. These conditions may result in a carbon limitation for the microbial population below this depth, which is consistent with

observations made by Van Wambeke et al. (2009) during the Dynaproc2 cruise.

### 4.3 Degradation status of the dissolved lipid pool

An other important component of the dissolved lipid pool was the lipid metabolites (FFA, ALC, MG and DG). Release of metabolites occurs through the enzymatic cleavage of esters bounds of source glyceride lipids (Goutx et al., 2003). When the readily available molecules derived from phytoplankton start to be depleted, bacteria enzymatic activities cleave biopolymers into monomers (Hoppe, 1993). During the Dynaproc2 experiment, bacterial production was significantly correlated with the various organic matter descriptors (particulate carbon, DOC, MGDG and CDOM) (Mével et al., 2008; Ghiglione et al., 2008), which suggests that bacteria most probably were taking benefit from a wide range of biopolymers during this transitional period, because of varying growth limiting factors (Van Wambeke et al., 2009). Enzyme activity and bacterial production are not always tightly coupled, and accumulation of monomers (i.e. metabolites) is a function of this coupling/decoupling (Van Wambeke et al., 2001; Mullholand et al., 2007; Bourguet et al., 2009). Small lipid metabolites may be less reactive to bacteria than more complex and fresh biopolymers (Loh et al., 2004). In addition, lipid metabolite accumulation may reflect the presence of zooplankton digestive enzymes (Weeks et al. 1993). Finally, the accumulation of acyl lipid metabolites in the system may result from the maturation and shift of communities from autotrophic to heterotrophic as already reported in the Ligurian sea (Van Wambeke et al., 2001). This view is supported by the high contribution of metabolites to total dissolved lipids during late summer to autumn 2004 (on average  $37.6 \pm 11.1\%$ ,  $n=166$ ), compared to spring and summer 2003 period ( $23.9 \pm 16.9\%$  ( $n=12$ ) and  $28.1 \pm 6.0\%$  ( $n=16$ ), respectively) (Bourguet et al. 2009). At the end of the stratified period, monoacylglycerols constituted the largest pool of metabolites, which may be related to the low reactivity of these compounds to bacterial lipase attack (Gurr and James, 1980).

### 4.4 Inputs of dissolved lipids to depth

Few studies have considered the dynamics of dissolved lipids beyond the euphotic layer. In our study, lipid concentrations exhibited a remarkable stability with a maximum of 40% variability in concentrations from surface to 1000 m depth throughout the period investigated. Such stability characterized several other biological and chemical parameters (Ghiglione et al., 2008; Lasternas et al., 2008; Marty et al., 2008; Mevel et al., 2008; Raybaud et al., 2008, 2009) that were studied during the Dynaproc cruise, probably due to the exceptional hydrological calm encountered during this late summer to autumn transitional period.

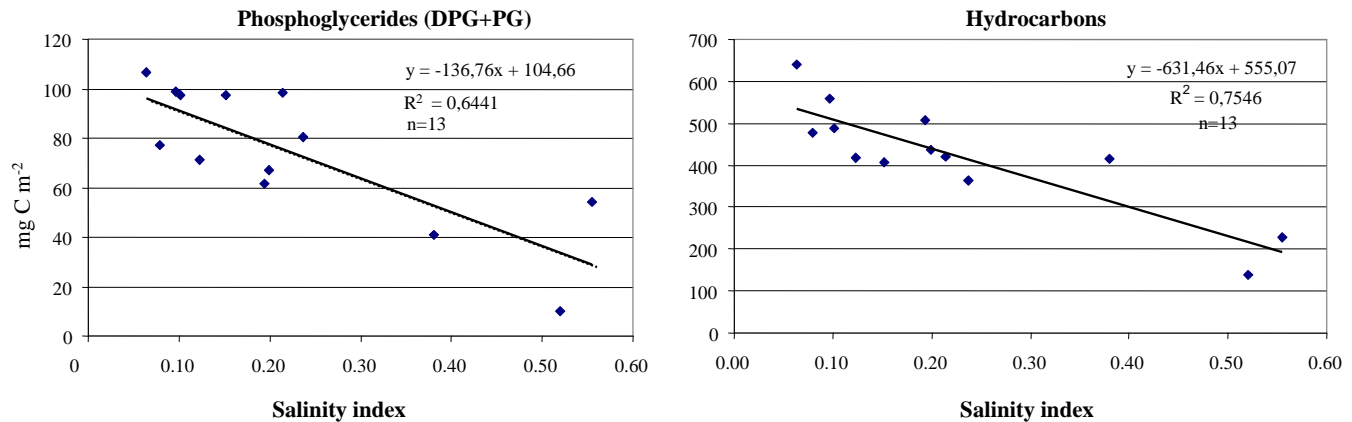
In such hydrological conditions with limited physical and hydrological forcings, the impact of biology on dissolved lipid distribution likely prevailed. The larger variations of lipid concentrations in the 0–150 m surface layer compared to the 150–1000 m layer (cf. Table 1), were due to the high phytoplankton production and bacterial activity that characterized the euphotic layer during the sampling period, as reported by Marty et al. (2008) and Mevel et al. (2008), respectively, and that presumably released and consumed lipids at various rates. Below 150 m depth during leg 2, peaks of dissolved lipid and individual biomarker concentrations (cf. Figs. 2 and 8) reflected an input of lipids from surface to depth that was not related to the DOC export through the deepening of the mixed layer that we observed the last days of the cruise. It suggests that the mechanism responsible for lipid transport to the mesopelagic layer was different from that transporting the other DOC components. Because of the presence of zooplankton biomarkers in the 0–150 m layer in the early Leg 2 and later in the 400–700 m deep lipid peaks, one potential carrier may be the zooplankton through its daily migration (i.e., Raybaud et al., 2008), and production of fecal pellets, either in the productive layer or in the mesopelagic layer, followed by sedimentation and/or dissolution. Indeed, there were a clear day/night periodicity in the mesopelagic distributions of dissolved TG and WSE throughout the period investigated. The double signature of TG and WSE in sinking particles has already been related to fecal pellets of carnivorous euphausiids during Dynaproc 1 experiment (Goutx et al., 2000). Moreover, carnivorous zooplankton were identified as a major trophic element during the leg 2 (Mousseau et al., 2009).

The export of hydrophobic contaminants like petroleum hydrocarbons has been previously related to the vertical transport of lipid material in sedimenting particles (Andersson et al., 1993). As inferred during periods of increasing productivity, the nature of sinking particles may also impact the flux of these hydrocarbons. Bouloubassi et al. (2006) showed that the flux of fecal pellets was able to carry a significant fraction of petroleum hydrocarbons to depth. During the Dynaproc 2 cruise, the high proportion observed at 700 m during leg 2 may be related to egestion of fecal pellets from zooplankton, followed by disaggregation/dissolution of the pellets within the mesopelagic layer, a scenario supported by the presence of zooplankton biomarkers at depth during Leg 2. At the surface, the origin of hydrocarbons may probably result from a background contamination of the lipid pool by petroleum hydrocarbons, as revealed by very high proportion ( $24.7\% \pm 3.2$ ) in the dissolved lipids (Goutx, 1988).

Phospholipid signatures were noticeable later on during leg 2 (12 October) in the mesopelagic zone (Fig. 8). This change in phospholipid concentrations at 700 m depth between 6–8 October and 12 October amounted  $1.5\text{--}2 \mu\text{g L}^{-1}$ . Marine bacteria contain a range of  $\sim 2\text{--}7\%$  lipids in total carbon, free or bound to proteins and polysaccharides, with most lipids in the form of membrane phospholipids ( $88 \pm 5\%$  TL)

(Goutx et al., 1990; Goutx, unpublished). Because most bacteria may pass through the  $0.7 \mu\text{m}$  membrane, operationally defined for isolation of suspended particles from dissolved matter, bacteria may account for a substantial fraction of this deep “phospholipid” pool, as suggested by Wakeham et al. (2003). However, between 6–8 October and 12 October, the total bacterial biomass did not increase at this depth (Mevel et al., 2008). Increases were observed in free or non-particle associated bacterial production ( $\times 5$ ) and abundance (above  $9 \times 10^5 \text{ cells L}^{-1}$ ) (Mevel et al., 2008) and ecto-enzyme activities (F. Van Wambeke, personal communication, 2008). Using the conversion factor of Findlay et al. (1989) for naturally occurring mixed populations ( $3.4 \times 10^9$  cells or 192 mg of C per 100 nmol of phospholipid), we evaluated the phospholipid increase as equivalent to  $10^7\text{--}10^8 \text{ cells L}^{-1}$  bacteria, which is far above estimated bacterial concentrations, and does not support the idea that these phospholipids would represent only a bacterial biomass increase. Phospholipids may rather indicate the presence of cell remains either derived from the flux of sinking particles described above and/or from egestion of fecal pellets by migrator zooplankton in the mesopelagic layer. It is most probable that this input of fresh biopolymers at these depth occurred through particle dissolution and bacterial enzyme attack, stimulating the production of free living bacteria and promoting a mesopelagic microbial food web, which limited the bacterial abundance according to Tanaka and Rassoulzadegan (2004). We sampled sea water along 6 profiles for POC measurement in the 0–1000 m water column during the cruise. Among the four Leg 2 profiles, three profiles (11–13 October) clearly exhibited deep POC accumulation at 700 m depth ( $\times 1.5$  to  $2.5$  folds the concentration in the 500 m depth above) (data not shown).

Finally, the above observations combine to show that an egestion of fecal pellets at the surface or in the mesopelagic layers by zooplankton and/or migrators, and their subsequent disintegration/dissolution through bacterial activity, sloppy feeding and grazing in the mesopelagic layer is a reasonable assumption that would explain the presence of lipid signatures in the deep water layers. This scenario is consistent with previous observations that slowly sinking particles derived from trophic chains based on small particles would rather dissolve within the mesopelagic layer at Dyfamed (Goutx et al., 2007). During Dynaproc 2, fecal pellets coming from the upper layers probably were emitted during the low wind period (25 September to 8 October), when high POC fluxes were recorded at 200 m depth (Marty et al., 2009), which is consistent with a slow sinking velocity for these aggregates. These observations also support the conclusion of Bendtsen et al. (2002) about the Greenland Sea, that deep-sea labile DOC may only be explained by a DOC released from the sinking flux of POC. Dissolution of POM would lead to formation of colloids known to preferentially shelter phospholipids (Liu et al., 1998).



**Fig. 11.** Relationships between lipid biomarkers and salinity index. Lipid concentrations and Salinity index are integrated value between surface and 70 m depth.

#### 4.5 Relationship between lipid classes and low salinity index

In order to compare chemical and biological parameters with low salinity events, an index in the range 0–1 of the presence of lowered salinity or diluted waters in the upper water column was calculated, based on average salinity  $S$  in the 40–70 m water layer, following the formula:  $\text{Index}(\text{sta}) = (S_{\text{max}} - S(\text{sta})) / (S_{\text{max}} - S_{\text{min}})$ ;  $S(\text{sta})$  is the average salinity in the 40–70 m layer,  $S_{\text{min}}$  is that of the water taken as reference for low salinity water (coastal waters in our case),  $S_{\text{max}}$  is that of the salinity waters without anomaly.

Average lipid class concentrations were calculated similarly in the 40–80 m (we had no sample at 70 m depth). Regression between the low salinity index and lipid classes was significant (Fig. 11) showing that the low salinity water (LSW) had lower concentrations of phosphoglycerides and hydrocarbons than the nearby sea water (HSW). Phosphoglycerides are among the major phospholipids in cell membranes of prokaryotes and eukaryotes. Recent studies have attempted to determine their relative distribution among bacteria, pico- and microphytoplankton (Van Mooy et al., 2006). The decrease of PG concentration in the LSW reflected the change in microbial assemblages in LSW compared to HSW, reported by Mevel et al. (2008) and Lasternas et al. (2008). Dissolution properties of hydrophobic compounds may account for the difference in hydrocarbons (HC) concentrations. It has been long established that organic compounds are generally less soluble in aqueous salt solutions, such as seawater, than in pure water (reduced by 1.36), a phenomenon termed “the salting-out effect” (Xie et al., 1997). Studies examining the influence of salinity on hydrocarbons and polycyclic aromatic hydrocarbon (PAH) solubility in seawater have yielded similar conclusions. For 12 aromatic hydrocarbons, Sutton and Calder (1975) and McAuliffe (1987) found that the mean reduction in solu-

bility at 25°C in seawater was about 68%, relative to fresh water. In our study, solubility was certainly not involved in the distribution of hydrocarbons (HC) since there was an opposite relationship between HC dissolution and salinity. Interactions between aromatic hydrocarbons and particles can be affected by salinity. Unfortunately particle size distribution was not investigated during the Dynaproc2 cruise. In our study, the higher concentrations of hydrocarbons in the high salinity water was surprising. Indeed, a higher contamination would be expected in waters coming from the coast, rather than from offshore sea water, due to ship concentration and/or resuspension of HC rich sediments in coastal zones. This finding suggests that chronic petroleum hydrocarbon discharge are important in NW Mediterranean and/or that other sources like atmospheric inputs may be significant. Finally, the time series of total dissolved hydrocarbon concentrations constitutes a unique data set for these compounds in offshore Mediterranean waters that may be of further use in environmental studies.

#### 5 Conclusions

This study contributes to advances in DOM characterization by documenting the concentration and composition of dissolved lipids along the 0–1000 m surface and mesopelagic water layers in the central ligurian sea, NW Mediterranean. We conclude that different processes were involved in variability of dissolved lipids at short temporal and spatial scale throughout the water column. Changes in concentration and composition of dissolved lipid classes varied along the water column but significant variations also occurred during the summer to autumn transition. Values were high in the surface layer in relation to phytoplankton biomass where picoeukaryotes appeared to be the major source of dissolved lipids. In parallel, the variations of the system over time during the seasonal transition marked by nutrient limitations, corresponded

to lipid degradation and metabolite accumulation that were higher than previously observed earlier in the year. During the transitional period, the stratification of the system was a factor limiting the distribution of fresh dissolved matter to depth. Under these conditions, the lipid biomarker pattern in the mesopelagic layer suggested strongly that the mesopelagic community was benefitting from DOM release from sinking POM due to zooplankton trophic behaviour, before the winter convection.

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