



Opportunistic feeding on various organic food sources by the cold-water coral *Lophelia pertusa*

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Abstract. The ability of the cold-water coral *Lophelia pertusa* to exploit different food sources was investigated under standardized conditions in a flume. The tested food sources, dissolved organic matter (DOM, added as dissolved free amino acids), bacteria, algae, and zooplankton (*Artemia*) were deliberately enriched in ¹³C and ¹⁵N. The incorporation of ¹³C and ¹⁵N was traced into bulk tissue, fatty acids, hydrolysable amino acids, and the skeleton (¹³C only) of *L. pertusa*. Incorporation rates of carbon (ranging from 0.8–2.4 μg C g⁻¹ DW d⁻¹) and nitrogen (0.2–0.8 μg N g⁻¹ DW d⁻¹) into coral tissue did not differ significantly among food sources indicating an opportunistic feeding strategy. Although total food assimilation was comparable among sources, subsequent food processing was dependent on the type of food source ingested and recovery of assimilated C in tissue compounds ranged from 17 % (algae) to 35 % (*Artemia*). De novo synthesis of individual fatty acids by *L. pertusa* occurred in all treatments as indicated by the ¹³C enrichment of individual phospholipid-derived fatty acids (PLFAs) in the coral that were absent in the added food sources. This indicates that the coral might be less dependent on its diet as a source of specific fatty acids than expected, with direct consequences for the interpretation of in situ observations on coral nutrition based on lipid profiles.

1 Introduction

Cold-water corals (CWCs) form reef structures in the cold and deep oceanic waters around the world (Roberts et al., 2009a; Davies and Guinotte, 2011). These reefs form

hotspots of biodiversity (Roberts et al., 2006) and are important in carbon cycling along continental margins (van Oevelen et al., 2009). The high metabolic demand of CWC communities implies high food processing rates and indeed a close relationship between food availability and the occurrence of CWC reefs has been reported in many studies (Roberts et al., 2006; Thiem et al., 2006).

In the North Atlantic Ocean, CWC reef communities are primarily formed by the scleractinian coral *Lophelia pertusa* (Roberts et al., 2006). Recent in situ investigations using stable isotope and fatty acid analyses point to a close coupling of *L. pertusa* with pelagic resources such as zooplankton and phytodetritus (Spiro et al., 2000; Duineveld et al., 2004; Kiriakoulakis et al., 2005). While tidal pumping (Davies et al., 2009) and internal waves (Frederiksen et al., 1992; Duineveld et al., 2004) deliver a diverse range of particles to the coral, ranging from fresh to resuspended material, the vertical migration (daily or seasonal) of zooplankton can also contribute to the linkage between surface-water production and CWC nutrition (Hind et al., 2000; Valle-Levinson et al., 2004; Dodds et al., 2009).

Although organic food sources of various size, type and quality reach the CWC reefs, very little is currently known about their importance to the metabolism of CWCs or the biogeochemical processing that occurs following nutritional uptake. Analysis of natural stable isotope signatures in tissues allow insight into the coral nutrition as described above (Duineveld et al., 2004; Kiriakoulakis et al., 2005; Dodds et al., 2009; Roberts et al., 2009b; van Oevelen et al., 2009), but to enhance the resolution of bulk tissue isotope data, fatty acids are often used as a biomarkers in these studies

(Kiriakoulakis et al., 2005; Dodds et al., 2009; Duineveld et al., 2012). However, the processing and production of fatty acids by CWCs has not yet been studied. This limits their interpretation since the use of fatty acids as biomarker for a specific food source critically depends on the assumption that these markers cannot be synthesized by the consumer itself (Kelly and Scheibling, 2012).

Feeding studies of *L. pertusa* so far have focused on uptake rates of *Artemia salina* (Purser et al., 2010; Tsounis et al., 2010) in the laboratory, but other food sources and particle sizes below 100 μm have not been considered yet. The nutritional importance of a food source does not only depend on its availability, but also on uptake and physiological processing by *L. pertusa*. In addition to particulates, dissolved resources may also contribute to coral dietary requirements. For tropical corals, it is known that dissolved organic matter (DOM) can be an important food source even under low ambient DOM concentrations (Hoegh-Guldberg and Williamson, 1999; Grover et al., 2006, 2008). Recently, Naumann et al. (2011) reported DOM uptake by the CWC *Desmophyllum dianthus*; thus, DOM may be an additional resource for *L. pertusa* that has not yet been accounted for.

In addition to a demand of organic resources for energy and tissue growth, *L. pertusa* also needs an inorganic carbon source to sustain calcification. Two carbon sources are possibly involved in the calcification process in corals: dissolved inorganic carbon (DIC) from the surrounding seawater or metabolically generated CO_2 . While calcification in tropical zooxanthellate scleractinian corals is mainly (70–75 %) based on metabolic CO_2 (Furla et al., 2000), the opposite seems to be true for the azooxanthellate octocoral *Leptogorgia virgulata* (Lucas and Knapp, 1997). Based on stable isotope data ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$), Adkins et al. (2003) suggested that C for calcification in azooxanthellate scleractinian CWCs is primarily derived externally rather than metabolically. Direct measurements distinguishing between the different calcification pathways in CWCs are however not available.

In this study we used standardized food quantities in a laboratory setting to trace the incorporation of stable isotope labelled food sources (^{13}C and ^{15}N) to measure food uptake rates and potential preferences of the CWC *L. pertusa*. Different food sources were selected to cover different particle sizes and nutritional values (C:N ratio), comprising dissolved organic matter, bacteria, algae and zooplankton (*Artemia*). Following food uptake we traced ^{13}C and ^{15}N into two of the most important biochemical components, hydrolysable amino acids (HAAs) and lipids represented by the sum of total fatty acids (TFAs) to follow metabolic processing. Within the lipid pool we especially followed food-derived C further into phospholipid-derived fatty acids (PLFAs) to have a more detailed look into the synthesis of structural and therefore functional fatty acids. Additionally, we traced food-derived C into the coral skeleton to investigate the metabolic contribution to coral calcification.

2 Methods

2.1 Sampling location and maintenance

All corals used in the experiment were collected at the Tisler Reef, one of the largest and shallowest inshore reefs known so far. It is located at a depth of 70 to 155 m in the Skagerrak, on a sill forming the submarine border between Norway and Sweden. The current velocity over the reef varies from 0 to 50 cm s^{-1} throughout the year while the flow direction fluctuates irregularly between NW and SE (Lavaleye et al., 2009; Wagner et al., 2011). The amount and quality of particulate organic carbon (POC) entering the reef depend on the location within the reef, and concentrations vary between 43 to 106 $\mu\text{g POC L}^{-1}$ (Wagner et al., 2011). Temperature at the reef site typically fluctuates between 6 and 9 $^{\circ}\text{C}$ throughout the year (Lavaleye et al., 2009; Wagner et al., 2011).

Corals were collected from a depth of 117 m (N58 $^{\circ}$ 59'800" E10 $^{\circ}$ 58'045") using an ROV (Remotely Operated Vehicle, Sperre Subfighter 7500 DC). Within a few hours after sampling, corals were transported in cooling boxes filled with cold seawater (7–8 $^{\circ}\text{C}$, salinity 31) to the laboratory at the Sven Lovén Centre for Marine Sciences, Tjärnö, Sweden. Until the start of the experiment (3 months), corals were kept in a dark thermo-stated room at 7 $^{\circ}\text{C}$ with a flow through of sand-filtered bottom water (particle size 1–2 mm) from 45 m depth in the adjacent Koster fjord (7–8 $^{\circ}\text{C}$, salinity 31). During this time the corals were fed with larvae (nauplii) of the Brine Shrimp *Artemia spp.* every 3 to 4 days. The Sven Lovén Centre has kept *L. pertusa* alive and growing under these conditions for a number of years.

Coral samples used in this experiment were clipped to approximately the same size with 4.5 ± 1.9 g dry weight (DW) and 8 ± 4 polyps per fragment (average \pm SD) about a week before the experiments started.

2.2 Preparation of particulate labelled substrates

CWCs are thought to thrive mainly on particulate organic matter (Duineveld et al., 2012). To test for assimilation of different food particle sizes, we chose bacteria to represent picoplankton and microalgae to represent nanoplankton. *Artemia* nauplii were chosen to represent mesozooplankton because they can be cultured in high densities, are the essential food source for successfully keeping *L. pertusa* in the laboratory and have been used in earlier CWC feeding studies (Purser et al., 2010; Tsounis et al., 2010; Naumann et al., 2011). While DOM was added in the form of a commercially available algal-derived mixture of dissolved free amino acids (Cambridge Isotopes, U ^{13}C 97–99 %, U ^{15}N 97–99 %, CNLM-452-0.5), the labelled POM food sources were prepared by culturing the respective organisms in the presence of ^{13}C and ^{15}N labelled substrates. A natural community of bacteria was derived from a few mL of natural seawater, obtained from the Oosterschelde estuary (salinity 30)

in the SW of the Netherlands. The water sample was added to 1 L culture medium (M63) containing 0.02 mol L^{-1} glucose (10 atom % ^{13}C , Cambridge Isotopes) and 0.01 mol L^{-1} ammonium chloride (10 atom % ^{15}N , Cambridge Isotopes). After 3 days of culturing in the dark, bacteria were concentrated by centrifugation ($14\,500 \text{ g}$) and rinsed with $0.2 \mu\text{m}$ filtered seawater to remove residual labelled substrates. Bacteria in the concentrate were kept frozen until use in the experiment.

The diatom *Skeletonema costatum* was cultured axenically in 4 L F/2 culture medium containing 0.8 mmol L^{-1} NaNO_3 (10 atom % ^{15}N , Cambridge Isotopes) and 2 mmol L^{-1} NaHCO_3 (20 atom % ^{13}C , Cambridge Isotopes, 99 % ^{13}C). After 3 weeks of culturing in 12 h light 12 h dark cycles (at a cell density of around $3\text{--}4 \times 10^6 \text{ cells mL}^{-1}$), algae were concentrated by centrifugation at 450 g , rinsed three times with $0.2 \mu\text{m}$ filtered seawater to remove residual labelled substrates and kept frozen until use in the experiment.

For culturing ^{13}C and ^{15}N enriched *Artemia* nauplii, 6 times 0.1 g *Artemia* cysts (Sera) were incubated in 5 L incubation chambers filled with $0.2 \mu\text{m}$ filtered seawater under natural light conditions and light aeration. After the larvae had developed to the state that they take up particulate food (1 to 2 days after eclosion of larvae), they were fed every second day with around 7 to 10 mg C derived from ^{13}C and ^{15}N enriched pre-cultured algae (cultured as described above, 4 atom % ^{13}C , 10 atom % ^{15}N). The uptake of algae by *Artemia* was visually confirmed under the microscope. After seven days of feeding, the larvae were concentrated by filtration, rinsed with filtered seawater, counted under the binocular and stored frozen. Within the filtrate, different early larvae stages could be identified.

To standardize the amount of carbon added to the incubations, all substrates were measured for carbon and nitrogen content. ^{13}C and ^{15}N enrichment and the fatty acid composition (PLFAs) of organic food sources were also measured to trace and calculate coral food uptake (see below for methodological description).

2.3 Experimental setup and procedure

During the incubations, corals were placed in recirculation flumes (60 L) in a thermo-stated room at 7°C (Fig. 1). Water circulation was maintained by a motor-driven propeller situated in the returning pipe (for more details see Purser et al., 2010). Prior to the experiment, the flume was filled with $0.2 \mu\text{m}$ filtered seawater from 45 m depth out of the Koster fjord (salinity 33.7°C , pH 7.9 on NBS scale) and the motor was set to ensure a flow speed of 7 cm s^{-1} , which is within the natural range found at the Tisler Reef (Lavaleye et al., 2009; Wagner et al., 2011).

Three coral fragments were randomly selected and placed in the test section of each flume (Fig. 1). The three pieces were gently inserted into a 1 cm elastic silicone tube on an acrylic plate that could be attached to the flume base (Purser

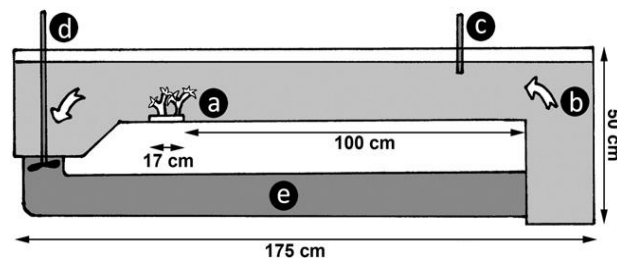


Fig. 1. Scale diagram of recirculation Plexiglas flume setup adopted from Purser et al. (2010). (a) test section with coral branches in plastic mount (depth 8 cm), (b) direction of circulation, (c) food delivery point, (d) motor, (e) opaque plastic return pipe.

et al., 2010). Corals were left in the flume for 12 h to acclimatize to the conditions. After acclimation, 10 mg C of the respective food source per treatment was gently pipetted into the water column of each flume (final concentration $170 \mu\text{g C L}^{-1}$). Visual observations confirmed that the circulating water kept the particulate food in suspension. Each flume contained three coral pieces and each food source was replicated twice. As a control treatment, corals were incubated for the same time without any food addition. After an incubation time of 4 days in darkness, coral samples were frozen at -20°C , freeze-dried and stored frozen for further analysis.

2.4 Sample treatment and analyses

After freeze-drying, coral samples were weighed and homogenized by grinding with a ball mill for 20 s (MM 2000, Retsch, Haan, Germany). A subsample of around 30 mg of ground material (organic and inorganic fraction) was transferred to pre-combusted silver boats and analyzed for isotopic ratio and C/N content using a Thermo Electron FlashEA 1112 analyzer (EA) coupled to a Delta V isotope ratio mass spectrometer (IRMS). Another subsample was transferred to a silver boat, acidified stepwise with drops of increasing concentrated HCl until the inorganic C fraction was removed (no bubbling after acid addition). The remaining material was analyzed on the EA-IRMS for isotopic ratio and organic C content. Incorporation of C in the inorganic skeleton was determined by subtraction of the organic carbon fraction (tissue + organic matrix) from the total carbon pool.

Total fatty acids (TFAs) were extracted with an adjusted Blight Dyer method. A part of the total fatty acid extract was further eluted over a silica column (Merck Kieselgel 60) to isolate the phospholipid-derived fatty acids (PLFAs) (Boschker et al., 1999). The TFA and PLFA extracts were then further derivatized by mild alkaline transmethylation to obtain fatty acid methyl esters (FAME). Preparation of methyl esters was carried out following the method of Boschker et al. (1999). For extraction, 0.7 g DW of coral

samples, $\sim 100 \mu\text{L}$ of DOM and smaller particulate food sources, and 100 *Artemia* (equivalent to $500 \mu\text{g C}$ and $50 \mu\text{g C}$ respectively) were used. Concentration and carbon isotopic composition of individual TFAs and PLFAs were measured on a gas-chromatograph combustion-interface isotope-ratio mass spectrometer (GC-c-IRMS) according to Boschker et al. (1999).

Hydrolysable amino acids (HAAs) were extracted and analyzed following Veuger et al. (2005). Ground coral samples were first treated with 12 mol L^{-1} HCl to dissolve the skeleton by repeated addition of HCl drops to avoid loss of sample through bubbling. The remaining material was then hydrolyzed in 6 mol L^{-1} HCl at 110°C for 20 h and purified by cation-exchange chromatography (Dowex 50WX8 resin). The hydrolysable amino acids were derivatized with isopropanol and pentafluoropropionic anhydride and analyzed by GC-c-IRMS for concentrations and ^{13}C and ^{15}N content.

Stable isotope data are expressed in delta notation as $\delta X\text{‰} = (R_{\text{sample}}/R_{\text{ref}} - 1) \cdot 1000$, in which X represents C or N, R_{sample} is the heavy/light isotope ratio in the sample (e.g., $^{13}\text{C}/^{12}\text{C}$) and R_{ref} is the heavy/light isotope ratio of the reference material ($R_{\text{ref}} = 0.01118$ for C and $R_{\text{ref}} = 0.00368$ for N). The atomic % of the heavy isotope in a sample (e.g., $^{13}\text{C}/(^{13}\text{C} + ^{12}\text{C})$) was calculated as $F = R_{\text{sample}}/(R_{\text{sample}} + 1)$. The excess (above background) atom% is the difference between the F in an experimental sample and the atom% in a background (untreated) coral sample: $E = F_{\text{sample}} - F_{\text{background}}$, so that zero values of E imply no uptake of the isotopically labelled food source and positive values indicates food uptake. To correct for differences in isotope enrichment among the food sources, the excess incorporation was divided by the atom% of each specific food source, e.g., 0.09 for bacteria C and 0.1 for bacteria N. To arrive at total elemental uptake, E was multiplied with C or N content of 1 g DW sample ($\mu\text{g C g}^{-1}$ DW sample, $\mu\text{g N g}^{-1}$ DW sample). All results are reported as average \pm SD derived from all coral pieces per treatment ($n = 6$).

2.5 Statistical analyses

To be able to perform statistical analyses of the obtained data, we treated pseudo-replicates as true replicates if no significant different between flumes were found by using Kruskal–Wallis Test ($p \geq 0.05$). However, all statistical values still to be considered with care because of limited replication. The potential influence of food sources on C and N uptake in bulk tissue and specific components was investigated using Kruskal–Wallis Test since a normal distribution for some data could not be achieved by data transformation. However, in cases where all requirements for an ANOVA were met, the results did not differ from the ones obtained by Kruskal–Wallis Tests. Therefore we decided to use Kruskal–Wallis Tests for all factors. Differences among treatments were then further investigated using a Wilcoxon rank-sum test for pair-

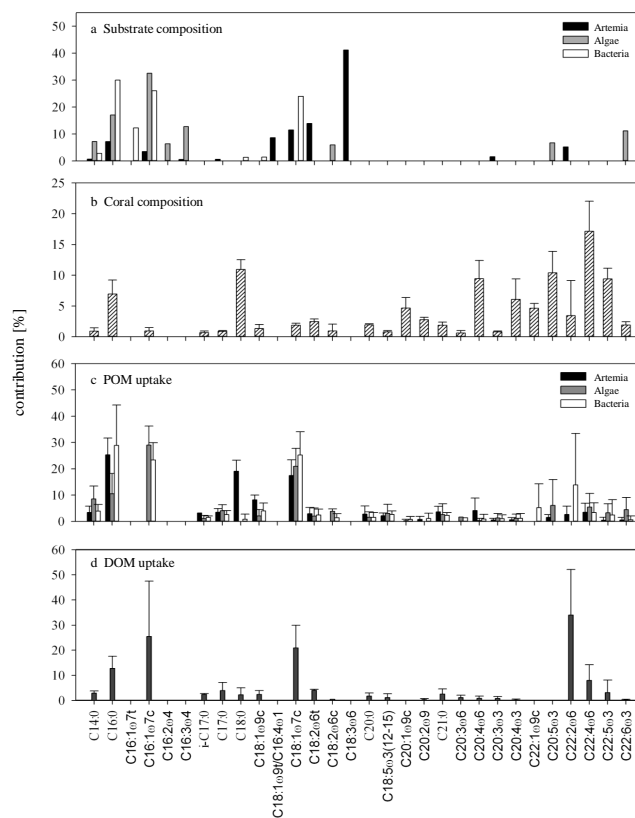


Fig. 2. PLFA profiles of (a) the particulate organic food sources *Artemia*, algae and bacteria (% contribution to total concentration), (b) *L. pertusa*, (c) POM-derived C incorporation into coral PLFAs by *L. pertusa* and (d) DOM transformation into coral PLFAs by *L. pertusa* (% contribution to total uptake). The bars in each figure represent average \pm SD.

wise comparison with an adjustment of p values by the Bonferroni method.

3 Results

3.1 Biogeochemical characteristics of particulate food sources

All food sources were significantly isotopically enriched above background and differed considerably in food quality represented by C/N ratios and PLFA contents as well as compositions in case of particulate organic food sources (Table 1, Fig. 2a). Accordingly, the bacteria-derived PLFA pool was dominated by C16:1 ω 7c/t, C16:0 and C18:1 ω 7c and the algal-derived PLFA pool by C16:0, C16:1 ω 7c, followed by C16:3 ω 4, C20:5 ω 3 and C22:6 ω 3 (Fig. 2a). *Artemia*-derived PLFAs mainly comprised C18:3 ω 6, followed by C16:0, C18:1 ω 7c, C18:1 ω 9t/c and C18:2 ω 6t/c (Fig. 2a).

Table 1. Characteristics of food sources used during the experiment.

Food source	¹³ C (at%)	¹⁵ N (at%)	Molar C/N ratio	PLFA content (% C)
Amino acids (DOM)	99	99	5.0	0.00
Bacteria (BAC)	9	10	3.6	0.31
Algae (ALG)	3	10	9.8	0.03
Zooplankton (ART)	3	2	4.2	0.32

3.2 Biogeochemical characteristics of *L. pertusa*

Corals used in this experiment had a total C content of $128 \pm 5 \text{ mg C g}^{-1}$ DW coral. This total C was partitioned into an organic tissue fraction of 13 % and inorganic skeleton fraction of 87 %. The organic C in the tissue fraction could be further partitioned into 24 % of THAAs, 7 % TFAs and 0.04 % PLFAs (Table 2). Dominant PLFAs in the coral tissue were C16:0, C18:0, C20:4 ω 3, C20:4 ω 6, C20:5 ω 3, C22:4 ω 6 and C22:5 ω 3 (all > 5 %), followed by C20:1 ω 9c and C22:1 ω 9c (~ 5 %, Fig. 2b). No significant difference in coral C content in tissue, TFAs or PLFAs was detected between the coral pieces in the different food treatments (Kruskal–Wallis $p > 0.05$ for all comparisons). Organic N content of corals used in this experiment was $3 \pm 1 \text{ mg N g}^{-1}$ DW coral, with 12 % of the N present in the THAA fraction (Table 2). No significant difference in coral N content in tissue and THAAs was detected between the coral pieces in the different food treatments (Kruskal–Wallis $p > 0.05$).

3.3 Total C and N assimilation

The assimilation of C from the food sources into coral tissue was not significantly different among treatments (Wilcoxon $p > 0.05$ for all comparisons, Fig. 3a). *Artemia*-derived C was assimilated at a rate of $1884 \pm 1067 \text{ ng C g}^{-1}$ DW coral d^{-1} , algal-derived C with $1520 \pm 498 \text{ ng C g}^{-1}$ DW coral d^{-1} , bacterial-derived C with $750 \pm 458 \text{ ng C g}^{-1}$ DW coral d^{-1} and DOM with $2393 \pm 1221 \text{ ng C g}^{-1}$ DW coral d^{-1} . Also N was assimilated comparably among different food sources by *L. pertusa* (Wilcoxon $p > 0.05$ for all comparisons) with *Artemia*-derived N assimilated with $797 \pm 399 \text{ ng N g}^{-1}$ DW coral d^{-1} , algal-derived N with $247 \pm 174 \text{ ng N g}^{-1}$ DW coral d^{-1} bacterial-derived N with $399 \pm 200 \text{ ng N g}^{-1}$ DW coral d^{-1} and DOM with $797 \pm 258 \text{ ng N g}^{-1}$ DW coral d^{-1} (Fig. 3a).

With the exception of corals fed with DOM (Kruskal–Wallis $p = 0.006$, Fig. 3a), C assimilation did not differ significantly from N assimilation among corals fed with particulate sources (Kruskal–Wallis $p > 0.05$), regardless of different N additions per treatment due to fixed C additions and variable C/N ratios of food sources. This points to a higher retention of nitrogen during assimilation and metabolic processing (Fig. 3e).

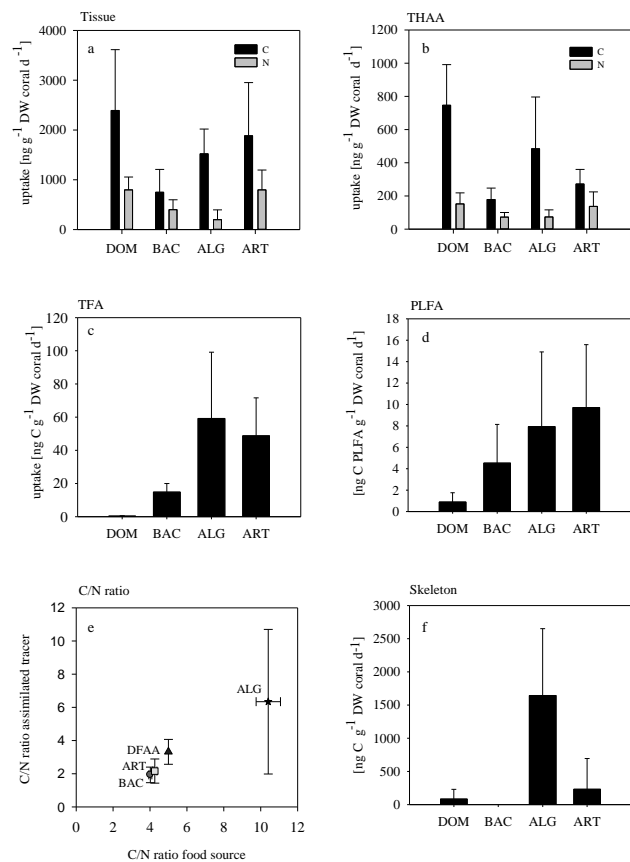


Fig. 3. (a) C and N uptake in coral tissue (note: different C/N ratios of sources), (b) C and N uptake in THAAs of coral samples (note: different C/N ratios of sources), (c) C uptake in TFAs of coral samples, (d) C uptake in PLFAs of coral samples, (e) C/N ratio of food provided and of assimilation, (f) C uptake in coral skeleton. The bars in each figure represent average \pm SD.

3.4 Food processing tracer incorporation in amino acids

Between 14 to 32 % of the total assimilated carbon was incorporated into the total hydrolysable amino acid pool (THAA) of *L. pertusa*. DOM-derived C was assimilated into THAAs at a significantly higher rate ($746 \pm 244 \text{ ng C g}^{-1}$ DW coral d^{-1}) than bacteria and *Artemia*-derived C ($178 \pm 69 \text{ ng C g}^{-1}$ DW coral d^{-1}

Table 2. Biogeochemical composition of *L. pertusa*: Concentration of total carbon (Total C), organic carbon (Org C), total nitrogen (Total N), total hydrolysable amino acids (THAAs), total fatty acids (TFAs) and phospholipid-derived fatty acids (PLFAs) in coral samples. Values are presented in (mg C g^{-1} DW coral \pm SD) and (mg N g^{-1} DW coral \pm SD), respectively.

Total C	Org C	Total N	THAAs	TFAs	PLFAs
128.05 \pm 4.95	1.60 \pm 4.81		3.61 \pm 0.55 (C)	1.02 \pm 0.03	0.06 \pm 0.02
		2.79 \pm 0.95 (N)	0.34 \pm 0.05 (N)		

and $272 \pm 88 \text{ ng C g}^{-1}$ DW coral d^{-1} , respectively, Wilcoxon $p_{\text{DOM-BAC/ART}} = 0.03$). The incorporation of algal-derived C ($484 \pm 311 \text{ ng C g}^{-1}$ DW coral d^{-1}) did not differ significantly from that of the other food sources (Wilcoxon $p > 0.05$ for all comparisons).

The incorporation of N into THAAs represented up to 30% of the total N assimilated by the coral and did not differ significantly between food treatments (Kruskal–Wallis $p = 0.2$) with $152 \pm 67 \text{ ng N g}^{-1}$ DW coral d^{-1} for DOM, $137 \pm 87 \text{ ng N g}^{-1}$ DW coral d^{-1} for *Artemia*, $73 \pm 43 \text{ ng N g}^{-1}$ DW coral d^{-1} for algae and $72 \pm 27 \text{ ng N g}^{-1}$ DW coral d^{-1} for bacteria (Fig. 3b).

3.5 C incorporation into fatty acids

Of the total C assimilated up to 4% was traced in the total fatty acid pool (TFA) of *L. pertusa*. The ingested food source hereby significantly influenced the amount of C incorporated into TFAs (Kruskal–Wallis $p = 0.0003$, Fig. 3c). DOM-derived C incorporation was significantly lower than that of the particulate sources ($0.3 \pm 0.2 \text{ ng C g}^{-1}$ DW coral d^{-1} , Wilcoxon $p_{\text{DOM-ART/ALG/BAC}} = 0.03$). *Artemia*- and algal-derived C (Wilcoxon $p > 0.05$) were incorporated in TFAs at comparable rates, 49 ± 23 and $59 \pm 40 \text{ ng C g}^{-1}$ DW coral d^{-1} , respectively, but bacteria-derived C was incorporated at a significantly lower rate of $15 \pm 5 \text{ ng C g}^{-1}$ DW coral d^{-1} (Wilcoxon $p_{\text{BAC-ART}} = 0.03$, Wilcoxon $p_{\text{BAC-ALG}} = 0.05$, Fig. 3c).

The incorporation into the phospholipid-derived fatty acid (PLFA) C-pool accounted for 0.6% of total assimilated C. Like for TFAs, C incorporated into PLFAs was also significantly different between different food sources (Kruskal–Wallis $p = 0.002$). Again, DOM-derived C was incorporated at a significant lower rate ($1 \pm 1 \text{ ng C g}^{-1}$ DW coral d^{-1}) than any of the particulate sources (Wilcoxon $p_{\text{DOM-ART/ALG}} = 0.03$, Wilcoxon $p_{\text{DOM-BAC}} = 0.05$, Fig. 3d). The particulate sources however did not differ significantly in their incorporation (Wilcoxon $p > 0.05$ for all comparisons). On average *Artemia*-derived C was incorporated with $10 \pm 6 \text{ ng C g}^{-1}$ DW coral d^{-1} , algal-derived C with $8 \pm 7 \text{ ng C g}^{-1}$ DW coral d^{-1} and bacteria-derived C with $5 \pm 4 \text{ ng C g}^{-1}$ DW coral d^{-1} .

The incorporation of C into PLFAs in corals fed with DOM was solely caused by de novo synthesis, since the DOM source did not contain any PLFAs. The PLFAs C16:0, C16:1 ω 7 and C18:1 ω 7c showed highest C incorporation

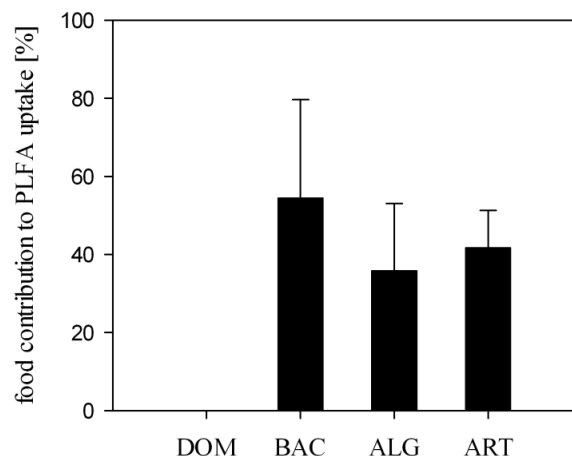


Fig. 4. Contribution of food-derived PLFAs to total PLFA synthesis in corals during the experiment. Based on the assumption that food-derived PLFAs are directly incorporated into animal PLFAs, the food contribution was calculated by summing the uptake in food characteristic PLFAs by *L. pertusa* and dividing it by its total PLFA uptake. The bars in the figure represent average \pm SD.

but also long chain PLFAs like C22:2 ω 6 and C22:4 ω 6 incorporated tracer C in DOM fed corals (Fig. 2d). For particulate sources over 40–60% of the assimilated PLFA-C by the corals was incorporated in PLFAs characterizing the respective food source (Fig. 4). PLFAs not present in the diet but with substantial tracer incorporation ($> 4\%$ contribution to tracer uptake) were C18:0, C18:1 ω 9c and C20:4 ω 6 in *Artemia* fed corals, C18:1 ω 7c and C22:4 ω 6 in algae fed corals and C22:2 ω 6 and C22:1 ω 9c in bacteria fed corals (Fig. 2c).

3.6 Carbon incorporation into coral skeleton

Incorporation of metabolic C derived from the processing of organic food sources into the inorganic carbonate skeleton was highly variable among coral samples (Fig. 3f), partly because only 1–2 coral pieces out of 6 showed measurable incorporation. Incorporation into the coral skeleton was highest in the algal treatment ($1.6 \pm 1.0 \text{ ug C g}^{-1}$ DW coral d^{-1}), followed by the *Artemia* ($0.2 \pm 0.5 \text{ ug C g}^{-1}$ DW coral d^{-1}) and DOM ($0.08 \pm 0.15 \text{ ug C g}^{-1}$ DW coral d^{-1}) treatment. Coral pieces fed with bacteria did not incorporate tracer C in their skeleton.

4 Discussion

4.1 Biochemical characteristics of *L. pertusa*

The overall contribution of amino acids and fatty acids to C tissue is in agreement with observations on other organisms, including tropical corals, in which proteins form the largest fraction before sugars and lipids (Szmant-Froelich and Pilson, 1980; Achituv et al., 1994).

The concentration of total fatty acids (20 mg g⁻¹ DW tissue with tissue DW contributing 5 % to total DW, as observed in this study), however, was below the range of 55–124 mg g⁻¹ DW tissue reported by Dodds et al. (2009) for *L. pertusa* from Rockall Bank, Mingulay Reef and New England Seamounts. Local differences can be responsible for this discrepancy as Larsson et al. (2013a) reported storage fatty acids concentrations of 15 to 19 mg g⁻¹ DW tissue from *L. pertusa* collected at the Tisler Reef but also the maintenance in aquaria (3 month in this study) might have altered the lipid content (Larsson et al., 2013b).

4.2 Food assimilation and source preferences

In line with aquaria studies and in situ observations (Buhl-Mortensen, 2001; Freiwald et al., 2002; Tsounis et al., 2010) our study confirms that *L. pertusa* can utilize various particulate resources from a broad range of sizes including bacteria, algae and zooplankton. The assimilation of DOM by *L. pertusa* is in accordance with the observation that *Desmophyllum dianthus* took up DOM (~ 6 µg C g⁻¹ DW d⁻¹) during core incubations at a 10–100x lower DOC concentration (Naumann et al., 2011). The comparable assimilation rates among resources hereby suggest that *L. pertusa* feeds rather unselectively at equivalent concentrations. This, together with the indication that coral food uptake is primarily driven by external factors such as food availability and current velocity (Purser et al., 2010), suggests that *L. pertusa* is an opportunistic feeder that utilizes resources depending on availability.

Rates of C assimilation into the tissue/biomass of *L. pertusa* in this study (~ 2 µg POC g⁻¹ DW coral d⁻¹) are lower than the *Artemia* capture rates measured by Purser et al. (2010) under comparable flow and food conditions (324 µg POC g⁻¹ DW coral d⁻¹). These measurements however are difficult to compare, because capture rates may overestimate actual ingestion if not every prey item is successfully transferred to the gut (Purser et al., 2010) and tissue assimilation only represents a fraction of the total uptake as respiration and mucus excretion are ignored. Especially the latter can be a significant component of the energy budget of *L. pertusa* (Wild et al., 2008; Maier et al., 2011).

4.3 Food composition governs transfer into amino acids and lipids

Although the food sources were unselectively assimilated, there were clear differences in the metabolic processing. The most pronounced difference was observed between DOM- and POM-derived C incorporation. While POM-derived C was incorporated at a higher rate than DOM-derived C in fatty acids, DOM-derived C was incorporated at higher rates in amino acids than POM-derived C, excluding algal-derived C. This difference was most likely caused by the differences in composition between the sources. The DOM consisted solely of dissolved free amino acids, which can be directly incorporated into coral tissue amino acids, whereas fatty acids had to be produced de novo using amino acids as C-substrate. The POM sources contained among others amino acids and fatty acids. The comparatively high POM incorporation into coral fatty acids most likely results from their availability in the source and their (direct) assimilation by the coral as illustrated by the effective incorporation of dietary PLFAs into coral PLFAs (Fig. 4).

Our results further indicate that not only the quantity (concentration) of amino acids/fatty acids, present in a food source, but also their quality (composition) might affect food source processing. This is especially illustrated by the assimilation of algae in comparison to other POM sources. Algal-derived C was incorporated into coral PLFAs at a comparable rate to other POM sources, despite containing 10 times less PLFA-derived C (Table 1). Additionally, algal-derived N incorporation into coral tissue and amino acids did not significantly differ from the incorporation of other POM sources despite a higher C/N ratio (i.e., a lower N concentration in the algal source). This suggests that while total assimilation can be comparable among sources, their nutritional importance in sustaining tissue components can still differ.

4.4 Assimilation and synthesis of PLFAs

Fatty acids are often used as biomarkers to infer the diet of animals (Braeckman et al., 2012; Kelly and Scheibling, 2012). This approach relies on the assumption that the relative composition of the PLFA pool in the animal reflects that of its food source(s) and hence that there is no alteration/modification or production of PLFAs occurring during the transfer from food source to animal. However this approach is only valid if the respective animal cannot synthesize the biomarker.

Overall, PLFA assimilation profiles (Fig. 2c, d) followed the biomarker concept and more closely reflected the dietary PLFA profiles (Fig. 2a) than that of the coral (Fig. 2a) with highest tracer recovery in PLFAs < C20 : 0 chain length. However, a detailed profile comparison revealed that direct assimilation of dietary PLFAs from particulate sources could only explain 40 to 60 % of the total C incorporation into the coral PLFAs. The remainder of the incorporation was due to

de novo synthesis or alteration of dietary PLFAs. This capability of *L. pertusa* of de novo synthesis was also evident from the assimilation of DOM into coral PLFAs because this resource solely contained amino acids and no PLFAs. The assumption of the biomarker concept “you are what you eat” therefore might not be completely true for *L. pertusa*.

Additionally, the de novo synthesis of FAs by *L. pertusa* includes also FAs which have previously been used as biomarkers. For example, C20:5 ω 3 and C22:6 ω 3 (biomarkers for diatoms and dinoflagellates, respectively, Harwood and Russell, 1984) were synthesized by *L. pertusa* from *Artemia*. Furthermore C22:1 ω 9c, a zooplankton biomarker (Sargent and Falkpetersen, 1988), was synthesized by *L. pertusa* when fed bacteria (detected in one of the three samples). Although the variation in de novo synthesis is high, it may complicate the interpretation of fatty acid profiles from field collected specimens, because presumed “unique” fatty acids may have been (partly) synthesized by *L. pertusa*. Our study thus adds to the growing evidence that care should be taken using fatty acids as dietary tracers in benthic food webs (Kelly and Scheibling, 2012; McLeod et al., 2013).

4.5 Metabolic versus external C incorporation into coral skeleton

Inorganic carbon to sustain calcification can either originate from an external (dissolved inorganic C) or from an internal pool (metabolic-derived C). In this study we directly measured the transfer of metabolic-derived C transfer into the coral skeleton, which ranged from 0.1–1.6 $\mu\text{g C day}^{-1} \text{g}^{-1}$ DW coral depending on the food source. This rate is considerably lower than separate, but comparable, incubations with ^{13}C -bicarbonate that showed a direct external inorganic C uptake of $46 \pm 25 \mu\text{g C g}^{-1}$ DW coral d^{-1} (Mueller et al. unpubl. data). Maier et al. (2009) measured a total calcification rate, i.e., sum of internal and external usage, of 23–78 $\mu\text{g C day}^{-1} \text{g}^{-1}$ DW coral with ^{45}Ca on freshly collected corals. These results indicate that metabolic-derived C only plays a minor role as C source for calcification, which confirms the suggestion by Adkins et al. (2003) based on isotopic data analysis in the coral skeleton ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$), that calcification of *L. pertusa* mainly relies on an external DIC source which decreases the effect of isotopic fractionation during the calcification process. This makes *L. pertusa* skeletons an excellent study object for climatic reconstructions (Adkins et al., 2003).

5 Conclusions and implications

In this study we investigated the capability of *L. pertusa* to take up different food sources ranging from dissolved organic matter, bacteria and algae to zooplankton. The comparable assimilation rates of the different food sources hereby indi-

cated that *L. pertusa* is an opportunistic feeder that is able to exploit a wide variety of food sources even including DOM.

Our main focus was to investigate the uptake of food sources across a broad size spectrum, but the implications of these findings for field conditions depend partly on how representative the experimental food sources are for natural conditions. Zooplankton can be an important food item for CWCs (Freiwald, 2002; Kiriakoulakis et al., 2005; Dodds et al., 2009; Duineveld et al., 2012), but it is logistically challenging to use local and living copepod species in laboratory studies, especially when these need to be enriched with stable isotopes. Therefore, we decided to use laboratory-reared *Artemia* as substitute for natural zooplankton, like most other feeding studies (Purser et al., 2010; Tsounis et al., 2010; Larsson et al., 2013b). Although *Artemia* may differ from copepods in their biochemical composition (Evjemo and Olsen, 1996; Helland et al., 2003), they are in the same size range as natural copepods. The microalgae selected for experimentation, *Skeletonema costatum*, contributes to phytoplankton blooms in the Skagerrak (Lange et al., 1992) and, together with dissolved organic matter, may reach the *Lophelia* reefs during downwelling events (Wagner et al., 2011). The cultured bacteria used in the experiment likely differed in composition from that in nature, but it gives first-order information about how *L. pertusa* deals with the natural picoplankton fraction comprising a broad variety of microbes (bacteria, eukarya and archaea, Jensen et al., 2012). We are therefore confident that our experimental food sources are good representatives for natural food items and are able to illustrate the opportunistic feeding strategy of the coral.

The ability to utilize DOM hereby underlines the nutritional flexibility of *L. pertusa* and might be especially relevant for natural reefs like on Rockall bank, where POM concentrations can be very low during several months (Duineveld et al., 2007). The assimilation of external DOM and the re-assimilation of DOM from coral mucus release, responsible for high labile DOM concentrations above CWC reefs (van Duyl et al., 2008; Wild et al., 2009) might help the coral to withstand several months without POM supply.

Next to the uptake of different sized food source, we also explored for the first time the processing of these sources by *L. pertusa*. The observed differences in food processing hereby suggest that the nutritional value of a food source is at least partly determined by its composition (quantity and quality of fatty acids and amino acids). Our findings further indicate that phytoplankton is a valuable resource for *L. pertusa* due to the efficient transformation into coral fatty acids. This might be especially relevant in locations where downwelling events (Tisler Reef, Mingulay Reef) supply the reefs with a high availability of phytoplankton (Duineveld et al., 2004; Davies et al., 2009; Wagner et al., 2011). Furthermore, the high flow velocities characterizing many CWC reef locations (Messing, 1990; Thiem et al., 2006; White et al., 2007) might also favor the uptake of smaller particles such as algae, since particle retention is negatively affected by particle

size especially at higher flow velocities (Shimeta and Koehl, 1997).

Additionally, we found de novo synthesis within the fatty acid metabolism, indicating that corals do not only rely on dietary fatty acids to obtain certain fatty acids. This especially concerns bacteria, since they are often considered as low quality food based on their lack of long-chain fatty acids (Phillips, 1984; Leroy et al., 2012). However, given the ability of the coral to synthesized long-chain fatty acids such as C22:2 ω 6 and C22:1 ω 9c from bacteria, our results suggest that bacteria can be a valuable addition to coral nutrition. This might be especially relevant since bacteria occur in high abundance around cold-water coral reefs and food uptake in *L. pertusa* is positively correlated with prey abundance (Purser et al., 2010). Additionally, bacteria are constantly available to the coral since they are fertilized by the coral itself via mucus production (Wild et al., 2009; Maier et al., 2011), while POM availability can vary spatially and temporally (Duineveld et al., 2007).

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