Trophic niche of the invasive gregarious species *Crepidula fornicata*, in relation to ontogenic changes

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Abstract

Crepidula fornicata is a common and widespread invasive gregarious species along the European coast. Among its life-history traits, well documented ontogenic changes in behavior (i.e., motile male to sessile female) suggest a potential shift in feeding strategy across its life stages. Considering the ecological significance of this species in colonized areas, understanding how conspecifics share the trophic resource is crucial. Using fatty acids (FA) and stable isotopes (SI) as complementary trophic markers, we conducted a field survey between late winter and spring to investigate the trophic niche of three ontogenic stages of *C. fornicata* that bear different sexual (male/female) and motility (motile/sessile) traits. Potential trophic sources were characterized by their pigment, FA and SI compositions and showed well discriminated compositions over the studied period. We showed that the biofilm covering *C. fornicata* shells harbored a higher biomass of primary producers (i.e., chlorophytes and diatoms) than the surrounding sediment. Over the studied period, we observed a covariation between the three ontogenic stages for both FA and SI compositions which suggest that the trophic niche of *C. fornicata* does not change significantly across its benthic life. During periods of low food availability, slipper limpets displayed an opportunistic suspension-feeding behaviour, relying on both fresh and detrital organic matter, likely coming from superficial sedimentary organic matter. However, during high food availability (i.e., spring phytoplankton bloom), all ontogenic stages largely benefited from this fresh supply of organic matter (pelagic diatoms in this case). The three ontogenic stages showed consistent differences in FA composition, and to a lesser extent in SI composition. These differences persist over time, as they originate from ontogenic physiological changes (differential growth rates, metabolic rate or gametogenesis) rather than diet discrepancies. This study revealed that multiple trophic markers allow high complementary to characterize organic matter as well as food partitioning between conspecific organisms.

Key words: *Crepidula fornicata*, trophic niche, ontogenic shift, fatty acids, stable isotopes, pigments, Bay of Brest

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1. Introduction

The slipper limpet *Crepidula fornicata* is a non-indigenous and invasive gastropod originating from the East coast of the US (Blanchard, 1997). This species extensively colonized shallow soft bottom habitats of European coasts, from Norway to the Mediterranean Sea (Blanchard, 1997). Because of its introduction in many parts of the world and its potential cascading effect on food web functioning (Arbach Leloup et al., 2008; Chauvaud et al., 2000; Cugier et al., 2010), several studies have closely investigated its diet and inferred potential trophic overlap with co-occurring benthic species (Blanchard et al., 2008; P Decottignies et al., 2007; Priscilla Decottignies et al., 2007; Lefebvre et al., 2009; Riera, 2007; Riera et al., 2002). *C. fornicata* is overall considered has an opportunistic suspension-feeder, able to feed on a large array of trophic sources (e.g., phytoplankton, microphytobenthos, macroalgae, bacteria), depending on their availability. Based on stable isotope ratios, it has been hypothesised a potentially large contribution of microphytobenthos, and more specifically benthic diatoms, in the diet of *C. fornicata* (P Decottignies et al., 2007; Guérin, 2004; Lefebvre et al., 2009; Riera, 2007). However, the unexpected presence of inorganic carbonates in *C. fornicata* soft tissues have led to overestimate $\delta^{13}\text{C}$ ratios in the consumer and then to overestimate the trophic role of microphytobenthos (Androuin et al., 2019).

C. fornicata is a hermaphroditic gregarious species, which begins its benthic life as a low-motile male and become a sessile female. Sessile adults form stacks of several non-moving individuals, while juveniles and small males (~10 mm) are motile (Coe, 1936). Adult females are suspension-feeders, but contrary to bivalves, they lack labial palp and showed no anatomical or functional potential for qualitative selection (Beninger et al., 2007). They form a food cord in a groove at the distal end of their gill filaments and ultimately catch this cord with their radula before ingesting it (Shumway et al., 2014). For the related species *Crepidula fecunda*, which exhibits ontogenic behavior changes comparable to those of *C. fornicata*, it has been demonstrated that newly settled individuals first adopt a grazing feeding mechanism and gradually shift to a suspension-feeding behaviour once their gill are fully developed (Montiel et al., 2005). Young individuals are able to use both feeding mechanisms (i.e., grazing and suspension-feeding) during the motile phase of their life cycle (size < 28 mm), whereas females are exclusive suspension-feeders (Chaparro et al., 2002; Navarro and Chaparro, 2002). Such observations have also been suggested for *C. fornicata* but without further behavioral evidence nor quantitative measurements (Breton and Huriez, 2010; Yee and Padilla, 2015). Since *C. fornicata* often occurs in large densities (up to 2000 ind. m²) on the seafloor with all ontogenic stages grouped in stacks (Guérin, 2004; Martin et al., 2006), one can expect strong intraspecific interactions for food. These interactions could be either facilitative or competitive depending on ontogenic feeding ecology. While purely suspension-feeding slipper limpets should compete for food among ontogenic stages, recent works suggested that younger individuals may be facilitated by adults, both via a higher substrate availability (de Montaudouin and Accolla, 2018) and through the grazing of microphytobenthic microalgae colonizing adult shells (Androuin et al., 2018). Given that *C. fornicata* often proliferates on muddy and turbid habitats with high suspended inorganic load, grazing behavior of motile males could also prevent the overloading of their digestive tract with inert matter of low nutritional quality (Navarro and Chaparro, 2002).

Different trophic markers have long been used to investigate the trophic niche of marine benthic invertebrates (e.g., Blanchet-Aurigny et al., 2015; Cresson et al., 2016; Dubois and Colombo, 2014) and to describe the origin of assimilated particulate organic matter (hereafter OM) (Ke et al., 2017; Lavaud et al., 2018; Liénart et al., 2017). As mentioned earlier, carbon and nitrogen stable isotopes (SI) are broadly used used to infer trophic niche of consumers (Fry and Sherr, 1984; Layman et al., 2012). Classically, nitrogen isotope ratio informs about the trophic position of a species and carbon isotope ratio reflects the origin of assimilated food sources (e.g., continental vs. oceanic). In coastal ecosystems, the diet of most of benthic primary consumers is composed of a mixture of OM from various origins (phytoplankton, macroalgae, continental detritus, zooplankton, etc) which are often difficult to disentangle with isotopes of only two elements, namely carbon and nitrogen. This diversity of food sources implies that complementary trophic markers are relevant to complement SI intel (Majdi et al., 2018). For instance, pigment analyses have been widely used to study community composition of microscopic primary producers in the water column or in the sediment, since some pigments are specific of clades of algae (Brotas and Plante-Cuny, 2003; Roy et al., 2011). To a lesser extent, fatty acid compositions can be also specific of group of organisms, such as diatoms, bacteria, copepods or vascular plants, (Dalsgaard et al., 2003; Kelly and Scheibling, 2012). Recently, the combined use of SI, FA and pigments improved our understanding of trophic pathways from the sources of particulate OM to benthic primary consumers (Lavaud et al., 2018; Majdi et al., 2018).

In this study, we investigated the trophic niche of *C. fornicata* and quantified intra-specific diet shift associated with ontogenic behavior changes (i.e., motile male to sessile female). For this purpose, we conducted a field survey and characterized potential OM sources by their SI, FA and pigments compositions and inferred their assimilation in *C. fornicata* tissues using both SI and FA trophic markers. Based on previous experimental study in stimulated microphytobenthic biofilm in *C. fornicata* beds (Androuin et al., 2018), we expect ontogenic trophic shift to happen within stacks, with a higher contribution of biofilm to motile males than to sessile males and females.

2. Materials and methods

2.1. Sampling strategy

The bay of Brest (Brittany, France) is a 180 km² semi-enclosed marine ecosystem. The sampling site is located near the Elorn estuary (48°23'N, 4°23', average depth: 10 m) in a dense *C. fornicata* beds (~2000 ind. m⁻²) (Guérin, 2004). Potential OM sources and *C. fornicata* individuals were collected by SCUBA divers at five sampling dates (S1 = 26th February, S2 = 21th March, S3 = 28th March, S4 = 12th April and S5 = 14th June) around mid and flood tide to ensure homogeneous mixing between estuarine and oceanic water. The late winter - spring period was chosen to encompass a period with potentially contrasted OM sources availability (e.g., spring blooms) (Figure 1).

Figure 1: Sampling dates (S1 to S5) at the study site superimposed with weekly chlorophyll *a* concentration at the entrance of the bay of Brest in 2018 (data from the French Coastal Monitoring Network SOMLIT; <http://somlit.epoc.u-bordeaux1.fr/fr/>).

Suspended particulate organic matter (SPOM) was sampled using two 8-liters Niskin bottles at 50 cm above the sediment-water interface, immediately filtered on board onto a 200 µm nylon mesh to remove large zooplankton and particles. In the laboratory, between 1 and 1.5 L was filtered on pre-combusted (450°C for 5 hours) GF/F filters (0.7 µm). Three replicates for each of the three analyses (SI, FA and pigment) were obtained. Superficial sedimentary organic matter (SSOM) was sampled from three cores of 15 cm diameter and 15 cm depth. In the laboratory, the sediment-water interface was re-suspended by flushing seawater with a 30 ml syringe following a standardized process: 60 ml of SSOM was pre-filtered on a 200 µm nylon mesh to being consistent with SPOM samples and filtered on pre-combusted (450°C during 5 hours) GF/F filters (0.7 µm). Three replicates for each of the three analyses were obtained. Biofilm from one *C. fornicata* stacks was scrapped off using a toothbrush and suspended in 600 ml of filtered seawater (0.7 µm). 200 ml of the suspended solution was filtered on pre-combusted (450°C for 5 hours) on GF/F filters (0.7 µm). Three replicates for each of the three analyses were obtained. Filters for FA analysis were put in glass tubes containing 6 ml of chloroform-methanol (2:1, v:v) solution and stored at -80°C before analysis, whereas filters for pigment and SI analysis were immediately stored at -80°C.

Females of *C. fornicata* were sampled at the bottom of the stacks (mean shell length 33 ± 6 mm), attached to a dead *C. fornicata* shell. Sessile and motile males were sampled if they had a penis and a mean

shell length of 20 ± 8 mm and 10 ± 1 mm, respectively. We used the digestive gland as a relevant trophic integrator tissue because it has a higher turnover rate than muscle tissue and is an energy storage organ enriched in lipids (McCutchan et al., 2003; Vander Zanden et al., 2015). However, since digestive gland and gonad are fused in a single organ in *C. fornicata*, we analysed both tissues together for sessile males and females. Because gonads are comparatively small in motile males, the whole body was used to ensure sufficient lipid concentration. At each date and for each ontogenic stage, both stable isotope (SI) and fatty acid (FA) analyses of *C. fornicata* were performed on subsamples originating from the same tissue sample.

2.2. Pigment analysis

The photosynthetic communities of SSOM, biofilm and SPOM have been analyzed by the quantification of pigments by High Performance Liquid Chromatography (HPLC) according to Brotas and Plante-Cuny (2003). Filters were crushed and extracted in 3 ml of 95 % cold buffered methanol (2 % ammonium acetate) for 20 min at -20°C in the dark. Samples were centrifugated for 3 minutes at 3000 g after the extraction period. Extracts were then filtered with Whatman membrane filters (0.2 mm) immediately before HPLC analysis. Pigment extracts were analysed using an Agilent 1260 Infinity HPLC composed of a quaternary pump (VL 400 bar), a UV-VIS photodiode array detector (DAD 1260 VL, 190–950 nm), and a 100 μl sample manual injection loop (overfilled with 250 μl). Chromatographic separation was carried out using a C18 column for reverse phase chromatography (Supelcosil, 25 cm long, 4.6 mm inner diameter). The solvents used were A: 0.5 M ammonium acetate in methanol and water (85:15, v:v), B: acetonitrile and water (90:10, v:v), and C: 100 % ethyl acetate. The solvent gradient followed the Brotas and Plante-Cuny method (2003), with a flow rate of 0.5 mL min^{-1} . Identification and calibration of the HPLC peaks were performed with chlorophyll *a*, β -carotene, chlorophyll *c*2, diatoxanthin, diadinoxanthin and fucoxanthin standards. All peaks detected were identified by their absorption spectra and relative retention times using the Open Lab CDS software (ChemStation Edition for LC/MS Systems, Agilent Technologies). Quantification was performed by repeated injections of standards over a range of dilutions to establish a standard curve of concentrations. Pigment percentages were expressed relatively to the surface/volume sampled ($\mu\text{g.cm}^{-2}$ for biofilm and SSOM, and $\mu\text{g L}^{-1}$ for SPOM). We measured the mean surface of three stacks of *C. fornicata* to standardize surfaces.

2.3. Stable isotope analysis

$\delta^{15}\text{N}$ and $\delta^{13}\text{C}$ analyses were carried out independently for both OM sources and *C. fornicata* tissues. For OM sources, filters were freeze-dried and split in two equal parts. Half of the filter was scrapped off and weighed in tin capsules for $\delta^{15}\text{N}$ analysis. The second half was decarbonated using acid-flume (10 N hydrochloric acid solution) for 7 hours (Lorrain et al., 2003), dried at 40°C for 12 h, scrapped off and weighed in tin capsules for $\delta^{13}\text{C}$ analysis. *C. fornicata* samples were freeze-dried and ground into homogenous powder using a mortar and pestle. Approximately 400 μg of powder was weighed in tin capsules for $\delta^{15}\text{N}$ analysis. Because both lipids content and inorganic carbonates can influence $\delta^{13}\text{C}$ (Androuin et al., 2019; McCutchan et al., 2003), approximately 400 μg of powder was added to 1 ml of cyclohexane in Eppendorf tubes. Tubes were vortexed and centrifuged at 3000 g during 5 min. The supernatant was discarded, and the tubes dried at 40°C during 12 h. If the supernatant remained coloured, the sample was re-processed. Lipid-free tissues were then weighed in silver capsules and in-cup decarbonated using 1N HCl. Each capsule was visually checked, dried at 40°C during 1 h, and sealed. Samples were analysed for $\delta^{15}\text{N}$ and $\delta^{13}\text{C}$ by continuous flow on a Thermo Scientific Flash EA 2000 elemental analyser coupled to a Delta V Plus mass spectrometer at the Pôle de Spectrométrie Océan (PSO, Plouzané, France). Results are expressed in standard δ notation based on international standards (Vienna Pee Dee Belemnite for $\delta^{13}\text{C}$ and atmospheric nitrogen for $\delta^{15}\text{N}$) following the equation:

$$\delta = [(R_{\text{sample}}/R_{\text{standard}}) - 1] \times 10^3 \text{ (in ‰)}, \text{ where } R \text{ is } ^{13}\text{C}/^{12}\text{C} \text{ or } ^{15}\text{N}/^{14}\text{N}.$$

International isotopic standards of known $\delta^{15}\text{N}$ and $\delta^{13}\text{C}$ values were used: IAEA-600 Caffeine, IAEA-CH-6 Sucrose, IAEA-N-1 and IAEA-N-2 Ammonium Sulphate. The analytical precision was estimated using the standard deviation of an internal standard (Thermo Acetanilide, $n = 8$), as $\pm 0.11 \text{ ‰}$ and $\pm 0.07 \text{ ‰}$ for $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values, respectively.

2.4. Fatty acids analysis

Freeze-dried powder of *C. fornicata* was subsampled for FA analyses: between 2 and 20 mg - depending on ontogenic stages - were immediately put in glass tubes (previously heated for 6 h at 450°C) containing 6 mL of a chloroform/methanol mixture (2:1, v:v), and extracted with a Dounce homogenizer. OM sources and *C. fornicata* samples were sonicated during 10 min and kept at -20°C until further analysis. The total lipid fractions were analyzed in OM sources, whereas only the neutral lipids were analyzed in *C. fornicata* samples. The detailed analysis method for separation and methylation is detailed in Le Grand et al. (2014). Fatty acid methyl esters (FAME) were analyzed in a Varian CP 8400 gas chromatograph (GC) equipped with a split/splitless injector and a flame-ionization detector (FID). FAMES were identified using two different capillary columns (ZBWAX 30 m × 0.25 mm i.d., 0.25 µm thickness, Phenomenex®; and ZB-5HT 30 m × 0.25 mm i.d., 0.25 µm thickness, Phenomenex®) by means of a standard 37 component FAME mix (Sigma Aldrich®) and other known standard mixtures. FAs were expressed as the molar percentage of the total FA content.

2.5. Statistical analyses

Pigment and FA compositions of OM sources, and FA compositions of ontogenic stages of *C. fornicata* were represented using a non-metric multidimensional Scaling (n-MDS). Homogeneity of the data was tested using permutational analyses of multivariate dispersion (PERMDISP) (Anderson, 2001). Statistical analyses on OM sources (pigments and FA) and *C. fornicata* (FA) were conducted using a non-parametric distanced-based permutation multivariate analysis of variance (PERMANOVA) based on a Bray-Curtis distance. Analyses were performed using two variables: OM sources or ontogenic stages (3 levels' factors) and sampling dates (5 levels' factor). Each date was considered independent due to the relative high turnover rate of both microorganisms found in the OM sources as well as the cells in the sampled tissues in *C. fornicata*. Following significant PERMANOVA results, *post hoc* tests were carried out using multiple pairwise comparisons with Bonferroni correction to identify differences among factors (Martinez Arbizu, 2017). However, the number of samples at each sampling date ($3 < n < 5$) was not sufficient to allow significant differences among the two factor levels in interaction, because of lack of statistical power when using Bonferroni correction in too many multiple comparisons. Therefore, *post hoc* comparisons of interaction term were not investigated. Finally, a SIMPER analysis was used to identify the FA explaining most of the dissimilarities between sampling dates and OM sources/ontogenic stages of *C. fornicata*.

Temporal variations and differences in SI ratios and pigment ratios/FA markers between OM sources/ontogenic stages of *C. fornicata* and sampling dates were assessed using two-way factorial analyses of variance (ANOVA). When significant, *post hoc* multiple comparisons were carried out using Tukey HSD. Normality and homogeneity of residuals were graphically assessed. Because concentrations in SPOM were not comparable with biofilm and SSOM (surface vs. volume), only SSOM and biofilm concentrations were compared together using the same procedure. Statistical analyses were performed in R version 3.3.0 (R Core Team, 2016) using packages 'vegan', 'plyr', 'FactoMiner', and 'ggplot'.

3. Results

3.1. Organic matter sources

3.1.1. Pigments and fatty acids compositions

Overall, OM sources were well discriminated by their pigment compositions (Figure 2a), and by their FA compositions (Figure 2b). Pigment compositions significantly differed between OM sources and sampling dates ($p < 0.001$ in both cases) and the interaction between the two factors was significant ($p < 0.001$). SIMPER analysis revealed that 80 % of the variability was explained by 7 pigments (Table S1). The biofilm was characterized by higher percentages of chlorophyll b and neoxanthin, together with one unknown pigment. SSOM was characterized by pheophytin *a*, pheophorbide *a*, and to a lesser extent lutein, whereas fucoxanthin and alloxanthin mainly discriminated SPOM. Temporal variations were mainly driven by a constant increase in both pheophorbide *a* and pheophytin *a* percentages for all OM sources. Fucoxanthin also showed an increase over time, except for SPOM and SSOM at S5 (Table S1).

Similarly, FA compositions also significantly differed between OM sources and sampling dates ($p < 0.001$ in both cases), and the two factors showed a significant interaction ($p < 0.001$). According to the SIMPER analysis, SSOM was characterized by higher percentages of 22:0, 16:1n-7, 18:1n-7 and a lower percentage of 16:0 (Table S2). The FA 18:0 and 20:4n-6 mostly discriminated biofilm whereas SPOM had higher percentages of 14:0 and 22:2n-6, and a lower percentage of 20:5n-3. In terms of temporal variations, both biofilm and SPOM showed similar decrease in saturated FA (i.e., 16:0 and 18:0) and increase in 16:1n-7 and 20:5n-3, especially between S4 and S5 (Table S2). SSOM exhibited less variable FA composition over time.

Figure 2: n-MDS based on the total pigment (a) and fatty acid (b) compositions of organic matter sources (Biofilm, suspended particulate organic matter (SPOM), superficial sedimentary organic matter (SSOM)). S1 to S5 correspond to the sampling dates (S1 = 26th February, S2 = 21st March, S3 = 28th March, S4 = 12th April and S5 = 14th June).

Total FA concentration did not show significant temporal variations for any OM sources (Figure 3a), with the biofilm always exhibiting higher concentration of total FA than SSOM ($p < 0.05$ at each sampling date). Chlorophyll (chl) *a* concentration increased over time in PPOM up to sampling date S4 ($1.8 \pm 0.2 \mu\text{g L}^{-1}$) ($p < 0.05$) as well as in biofilm (reaching $3.9 \pm 1 \text{ mg m}^{-2}$ at S5) even if differences were not significant due to high between-samples variability (Figure 3b). Chl *a* concentration in SSOM remained constant and was lower than in biofilm for each date ($p < 0.05$ in all cases). Fucoxanthin concentration increased over time for both SPOM and SSOM ($p < 0.05$ and $p < 0.05$, respectively), followed by a decrease in S5 (Figure 3c). Fucoxanthin concentration is the highest in biofilm, which showed a similar increasing trend that is not statistically supported because of high variability between samples. The chl *b*: chl *a* ratio was 3 to 7-fold higher for biofilm than for SSOM ($p < 0.001$) and SPOM ($p < 0.001$) (Figure 3d). Fucoxanthin: chl *a* (Figure 3e) and chl *c*: chl *a* (Figure 3f) ratios did not show clear temporal patterns for any OM sources and. However, they were 5 to 50-fold higher in SSOM than in SPOM ($p < 0.001$) and biofilm ($p < 0.001$) over the studied period, respectively.

Figure 3: Total fatty acids (a), chlorophyll a (b) and fucoxanthin (c) concentrations, and ratios between chlorophyll b (d), fucoxanthin (e), and chlorophyll c (f) over chlorophyll a (mean $\pm$ SD, $n \geq 2$) of organic matter sources (Biofilm, suspended particulate organic matter (SPOM), superficial sedimentary organic matter (SSOM)). S1 to S5 correspond to the sampling dates (S1 = 26th February, S2 = 21st March, S3 = 28th March, S4 = 12th April and S5 = 14th June).

3.1.2. Stable isotopes composition

The three OM sources were well discriminated by their $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values over the studied period. Their $\delta^{13}\text{C}$ signal varied significantly according to both OM sources and sampling dates, and the interaction between the two factors was significant ($p < 0.001$, $p < 0.01$ and $p < 0.001$, respectively). PPOM was always depleted in ^{13}C compared to RPOM ($p < 0.001$ at each date) and biofilm ($p < 0.01$ at each date) (Figure 4a, Table S3). Significant temporal $\delta^{13}\text{C}$ variations were only observed in biofilm, with higher values at S5 than at S2 ($p < 0.001$) or S4 ($p < 0.001$). Biofilm was significantly enriched in ^{15}N compared to both PPOM ($p < 0.001$) and RPOM ($p < 0.001$) (Figure 4b, Table S3). There was no interaction between OM sources and sampling dates ($p = 0.26$).

Figure 4: $\delta^{15}\text{N}$ (a) and $\delta^{13}\text{C}$ (b) isotopic compositions (mean $\pm$ SD, $n = 3$), and overall isotopic biplot (c) obtained of organic matter sources (Biofilm, suspended particulate organic matter (SPOM), superficial sedimentary organic matter (SSOM)). S1 to S5 correspond to the sampling dates (S1 = 26th February, S2 = 21st March, S3 = 28th March, S4 = 12th April and S5 = 14th June).

3.2. *Crepidula fornicata*

3.2.1. Fatty acids composition

FA composition of *C. fornicata* significantly differed between ontogenic stages and sampling dates ($p < 0.001$ in both cases) and the interaction between the two factors was significant ($p < 0.001$). The analysis of multivariate dispersion was also significant ($p < 0.05$), indicating that multivariate dispersion was not homogeneous. This was clearly illustrated by the n-MDS (when comparing the convex hull areas) where motile males showed much higher variation than sessile females and sessile males exhibited an intermediate level of variation (Figure 5). Pairwise SIMPER analyses between ontogenic stages revealed that sessile males were mainly characterized by saturated FA 16:0 and 18:0, especially at the two first sampling dates (Table S4). Sessile females differed from both motile and sessile males by higher percentages of C₂₀ FA such as 20:5n-3, 22:6n-3 and 20:1n-11, but also higher percentages of odd branched FA as iso17:0 (Table S4). Sessile males showed an overall comparable FA composition than sessile females but exhibited higher variability between sampling dates, as shown by the n-MDS. The SIMPER analyses performed between dates revealed that FA that most contributed to the observed temporal changes were the FA 16:0, 18:0 and 22:6n-3 decreasing over time, and the FA 20:5n-3 and 16:1n-7 increasing over time, all accounting for approximately 45 % of the dissimilarity over the 5 sampling dates (Table S4).

Figure 5: n-MDS based on the total fatty acid compositions of ontogenic stages of *Crepidula fornicata* (motile males, sessile males, sessile females). S1 to S5 correspond to the sampling dates (S1 = 26th February, S2 = 21st March, S3 = 28th March, S4 = 12th April and S5 = 14th June).

All FA percentages and ratios changed significantly between sampling dates ($p < 0.001$ in all cases) and between ontogenic stages ($p < 0.05$ in all cases, except for 16:1n-7 and 18:4n-3). The interaction terms were always significant ($p \leq 0.05$). Overall, temporal variations in FA composition were the highest in motile males, the lowest for females and intermediate for sessile males.

PUFA/SFA ratio increased over time for both motile and sessile males but remained constant in females (Figure 6a). There was no significant difference between ontogenic stages at sampling date S5. The relative abundance of branched FA was quite variable but significantly higher in sessile females than in motile males ($p > 0.05$) (Figure 6b). The highest values in branched FA was recorded in sessile females at S1 (11.2 ± 4.3) and the lowest in sessile males at S5 (3.8 ± 1). The ratio between 20:5n-3 and 22:6n-3 exhibited temporal variations for all ontogenic stages (Figure 6c), with a strong increase in S5 ($p < 0.001$ in all cases) where values ranged from 2.5 ± 0.5 in sessile females to 4 ± 0.6 in motile males. The FA 16:1n-7 followed the same trend as the 20:5n-3 / 22:6n-3 ratio with a more gradual increase over time for all ontogenic stages (Figure 6d). The highest values (from 3.6 ± 0.9 in sessile females to 6.4 ± 1.9 in motile males) were also recorded at S5. The FA 18:4n-3 also showed strong temporal variations in all ontogenic stages (Figure 6e), with an increase up to S4 followed by a decrease at S5. Finally, the n-3/n-6 ratio showed no temporal variation for sessile females but increased significantly over time up to S4 in both sessile ($p < 0.01$ in all cases) and motile males ($p < 0.05$ in all cases) (Figure 6f).

Figure 6: Relative abundance of fatty acids (FA) in the three ontogenic stages of *Crepidula fornicata* (motile males, sessile males, sessile females) (mean $\pm$ SD, $n = 5$): (a) Polyunsaturated FA / Saturated FA ratio, (b) Branched FA, (c) 20:5n-3 / 22:6n-3 ratio, (d) 16:1n-7, (e) 18:4n-3 and (f) n-3/n-6 ratio. S1 to S5 correspond to the sampling dates (S1 = 26th February, S2 = 21st March, S3 = 28th March, S4 = 12th April and S5 = 14th June).

3.2.2. Stable isotopes composition

Overall, the three ontogenic stages exhibited similar isotopic patterns over time (Figures 7a and 7b, respectively). No interactions were found between ontogenic stage and sampling date, both for carbon ($p = 0.6$) and nitrogen ($p = 0.27$). Motile males were significantly depleted in ^{13}C compared to sessile females ($p < 0.05$) and significantly depleted in ^{15}N compared to both sessile females ($p < 0.001$) and males ($p < 0.05$). Significant temporal variations were observed for $\delta^{13}\text{C}$ ($p < 0.001$) with a marked ^{13}C enrichment at S5 compared to all other sampling dates ($p < 0.05$ in all cases), up to 2 ‰ when compared to S4. A significant temporal decrease in $\delta^{15}\text{N}$ was found ($p < 0.001$) for all ontogenic stages.

Figure 7: $\delta^{15}\text{N}$ (a) and $\delta^{13}\text{C}$ (b) isotopic compositions (mean $\pm$ SD, $n = 5$), and corresponding isotopic biplot (c) for three ontogenic stages of *Crepidula fornicata* (motile males, sessile males, sessile females). S1 to S5 correspond to the sampling dates (S1 = 26th February, S2 = 21st March, S3 = 28th March, S4 = 12th April and S5 = 14th June).

4. Discussion

C. fornicata is a widespread invasive gregarious gastropod. The aim of this study was to investigate the trophic niche of this species and quantified intra-specific diet shift associated with ontogenic behavior changes (i.e., motile male to sessile female). A multi trophic markers assessment was conducted over spring to characterized potential OM sources and inferred their assimilation in *C. fornicata* tissues.

4.1. Composition and availability of potential food sources

Suspended particulate (SPOM) and superficial sedimentary (SSOM) organic matter as well as biofilm associated with *C. fornicata* shells were well discriminated by their pigment, fatty acid and stable isotope compositions, for each sampling date. When compared to marine POM (-22.3 ± 1.2 ‰), as measured over the same period close to the mouth of the bay of Brest (data from the French Coastal Monitoring Network SOMLIT; <http://somlit.epoc.u-bordeaux1.fr/fr/>) and terrestrial POM (-27.5 ± 1.2 ‰), as measured in terrestrial inputs nearby our study site by Mortillaro et al. (2014), the $\delta^{13}\text{C}$ obtained for SPOM (-25.4 ± 0.7 ‰) suggests a significant terrestrial influence at our study site, as mentioned by Marchais et al. (2013). SPOM over our studied period showed comparable chlorophyll *a* biomass ($0.4 - 1.8 \mu\text{g L}^{-1}$) than currently encountered in other areas of the bay of Brest (ca. $0.3 - 5 \mu\text{g L}^{-1}$, Chatterjee et al. 2013). High percentage of fucoxanthin and alloxanthin suggested the presence of Bacillariophyta (i.e., diatoms) and Cryptophyta in the water column, respectively (Brotas and Plante-Cuny, 2003; Roy et al., 2011).

Using the SOMLIT weekly monitoring of chlorophyll *a* and marine POM $\delta^{13}\text{C}$, a typical ^{13}C enrichment (~ 3 ‰) was noticeable during a phytoplankton bloom that occurred between May 25th and June 1st in the bay. This bloom was mainly composed by three diatoms: *Cerataulina pelagica* ($1.1 \cdot 10^6$ cells L^{-1}), *Leptocylindricus danicus* ($4.7 \cdot 10^4$ cells L^{-1}) and *Rhizosolenia imbricata* ($1.2 \cdot 10^4$ cells L^{-1}) (data extracted from the REPHY network, IFREMER). While this bloom was not sampled in the PPOM sampling set, the ^{13}C enrichment was evidenced in the individuals of *C. fornicata* collected at S5 (14th June).

Among the three sources of organic matter, SSOM showed the most homogeneous isotopic composition over time, which is unexpected as SSOM is often considered as a mixture of pelagic and benthic OM sources and consequently highly variable according to OM sources proportions and isotopic compositions. SSOM appeared here as a complex mixture of low and high quality OM (Lefebvre et al., 2009; Rigolet et al., 2014). On one hand, it was characterized by i) pheophorbide *a* and pheophytin *a*, which are degradation products of chlorophyllide *a* and chlorophyll *a*, respectively (Brotas and Plante-Cuny, 1998; Cartaxana et al., 2003), ii) odd branched FA (such as ant15:0) and the 18:1n-7 indicating the presence of

bacteria (Hubas et al., 2017; Jaschinski et al., 2011; Meziane et al., 1997) and iii) the long chain saturated fatty acids 22:0 which confirms the presence of a refractory terrestrial contribution in this environment (Canuel, 2001). On the other hand, SSOM exhibited the highest PUFA/SFA ratio most of the time (Table S1), suggesting higher quality/labability compared to SPOM and biofilm (Connelly et al., 2015, 2016; Parrish et al., 2005). This was confirmed by high Fucoxanthin / Chl *a* and Chl *c* / Chl *a* ratios which indicated a higher contribution of diatoms in SSOM than in SPOM and biofilm (Brotas and Plante-Cuny, 2003). Interestingly, fucoxanthin concentrations measured both in SSOM and biofilm were comparable to those observed on intertidal mud flats (Barranguet et al., 1997; Cartaxana et al., 2006). This blue-light absorbing pigment, and the very low light irradiance of our study site (0.01 %, Figure S1) strongly suggested that low light acclimated diatoms contributed to the SSOM at our study site (McGee et al., 2008).

Biofilm scrapped on shells of *C. fornicata* showed higher chlorophyll *a* concentration than the surrounding sediment, suggesting higher biomass of primary producers on shells (Androuin et al., 2018). The high percentages of chl *b* and neoxanthin, as well as the Chl *b* / Chl *a* ratio in the biofilm also suggest that chlorophytes were abundant on shells (Brotas and Plante-Cuny, 2003). While these results were not supported by FA (e.g., 18:3n-3 or 18:4n-3 characterizing chlorophytes, Fleurence et al. 1994), mollusk shells are currently inhabited by microchlorophytes or macrochlorophytes propagules (Barillé et al., 2017; Mineur et al., 2007). FA characterizing the biofilm were 18:0 and 20:4n-6. While 18:0 is an ubiquitous FA in marine environment (Kelly and Scheibling, 2012), the 20:4n-6 can be found in large proportion in red algae but also in brown ones (Fleurence et al., 1994; Kelly and Scheibling, 2012). It is worth noting that *C. fornicata* shells were partly covered with crustose red algae in our study site (pers. obs.). The high concentration of fucoxanthin also suggests the presence of high biomass of diatoms on these shells, as already mentioned by Ní Longphuirt et al. (2007). For instance, the biomass in biofilm was ~10 times higher than in the surrounding SSOM. The fact that biofilm was ¹⁵N-enriched by ~2.5 ‰ compared to SSOM may indicate that primary producers that compose this biofilm could use dissolved nitrogen derived from *C. fornicata*'s excretion products, which is a well-known process in benthic coastal ecosystems (Arzul, 2001; Prins et al., 1998; Ragueneau et al., 2002). While the excretion product (i.e., ammonium) is expected to be ¹⁵N-depleted relative to *C. fornicata* tissues (DeNiro and Epstein, 1981), it could still be ¹⁵N-enriched compared to the available dissolved nitrogen in the environment (Cifuentes et al., 1989; Raimonet et al., 2013; Wainright and Fry, 1994). This could indicate that nutrients are not a limiting factor for primary producers inhabiting *C. fornicata* shells, making them available for benthic consumers throughout the year provided that sufficient light reach the sea floor. The fact that chlorophyll *a* concentration decreased in SPOM after the spring bloom, but not in the biofilm, strongly support this hypothesis.

4.2. Trophic niche of *C. fornicata*, in relation to ontogenic changes

Trophic markers suggested an overall similar trophic niche of *Crepidula fornicata* across ontogenic stages, as shown by the covariation in both SI and FA compositions over the studied period. The slipper limpet is an opportunistic suspension-feeder that exploits both pelagic and benthic particulate OM in varying proportions according to the season and sources availability. However, differences in FA composition, and to a lesser extent in SI composition, were noticeable between ontogenic stages. These differences at each sampling date likely reflect ontogenic physiological changes link to growth rate and energetic demand rather than profound changes in diet.

Overall, SI ratios showed that all ontogenic stages had similar isotopic niches, although the niche of sessile females does not fully overlap with those of motile males. According to the respective SI ratios of potential food sources (Biofilm, SPOM, SSOM and marine POM) and those of *C. fornicata* tissues, and considering classical diet to consumer trophic enrichment factor (~ 0.75 - 1 ‰ for carbon and ~2.5 - 2.74 ‰ for nitrogen, Caut et al., 2009; McCutchan et al., 2003), it is likely that *C. fornicata* relies either on SSOM or marine POM depending of the season and food availability in the water column. During lower food availability period (i.e., end of winter and early spring), it is difficult to disentangle SSOM from marine POM since their SI signals do not differ. However marine POM was sampled close to the Bay entrance with higher oceanic influence, whereas SPOM was sampled just above the *C. fornicata* beds around mid and flood tide to ensure mixing with oceanic water. Therefore, SPOM was readily more available than marine POM for the slipper limpet. It is then reasonable to assume that SI composition of *C. fornicata* refers to SSOM rather than marine POM, which can be exploited through regular resuspension events linked to tidal currents (Beudin, 2014). After the spring phytoplankton bloom that occurred in the bay of Brest at the end of May, producing

a ^{13}C enrichment in the water column ($\sim 3\text{‰}$), a similar ^{13}C enrichment was found for all stages of *C. fornicata* ($\sim 2\text{‰}$). These results clearly showed that adults as well as young individuals of *C. fornicata* benefited from the spring bloom. However, minor but consistent isotopic differences were also found between ontogenic stages at each sampling date, which cannot necessarily be attributed to an ontogenic diet shift. Indeed, inferring diet shift using SI ratios may be hampered by the effects of physiological changes occurring during ontogeny such as gonadal maturation, metabolic rate or differential tissue growth between youngs and adults. Such physiological processes can modify the isotope signal of tissues even without significant change of diet, as evidenced in several species (Blanchet-Aurigny et al., 2012; Hentschel, 1998; Rossi et al., 2004). The isotopic dynamic also depends on tissue turnover rate (McCutchan et al., 2003; Vander Zanden et al., 2015). Lefebvre and Dubois (2016) analysed trophic enrichment factor and turnover rate in several marine benthic invertebrates, including *C. fornicata*. They showed a clear negative relationship between growth and enrichment factor values: when the body mass is increasing rapidly and the individual growing fast (high turnover), enrichment factors are expected to be low, and conversely. Young motile males of *Crepidula fornicata* are growing more rapidly than larger sessile males or even larger females (Hoagland, 1978; Walne, 1956), and consequently have a higher growth rate and a higher metabolic rate (Bayne and Newell, 1983). Enrichment factors are then expected to be much smaller for motile males, than sessile males or females, for which energy allocation is mainly directed to gamete production rather than other tissue growth. So, a similar diet in all ontogenic stages could very likely lead to differences in SI ratios such as those presented in Figure 7c. Even if motile males were analysed *in toto* (including muscle tissue with longer turnover rate), we believe that this does not biased interpretations of between-stages differences over the studied period. Analysing only tissues with fast turnover rate (as we did for sessile individuals) would have increased these between-stages differences.

Neutral lipids represent essential energy reserves for sustaining early life stages of marine molluscs and play a key role in their settlement, habitat selectivity and recruitment (Barbier et al., 2017; Pernet and Tremblay, 2004; Tremblay et al., 2007). Contrary to polar lipids, mostly involved in membrane regulation, FA incorporated in the neutral fraction are largely unaltered and reflect the diet in a more straightforward manner than polar lipids (Dalsgaard et al., 2003). Therefore, extracting FA from this specific class of lipids from a tissue with a rapid turnover (i.e., digestive gland) should allow assessing rapid changes in the diet (McCutchan et al., 2003). As suggested above by SI data, FA profiles showed a clear temporal variation in food sources utilization for the three ontogenic stages. This temporal pattern resulted from three distinct groups of sampling dates (Figure 5). The two first ones (S1 and S2) likely corresponded to a period that integrated the trophic signal of winter season's food sources, whereas the last one (S5) clearly corresponded to the assimilation of the spring phytoplankton bloom. The two-intermediate sampling dates (S3 and S4) corresponded to the transition with an increase in food availability. As shown earlier with SI, all ontogenic stages of *C. fornicata* may have probably exploited SSOM before the spring bloom when food in the water column is less available. In this pool of OM, FA revealed that slipper limpets likely fed on benthic diatoms (as suggested by 16:1n-7 and 20:5n-3; Dunstan et al. 1992; Napolitano et al. 1997; Passarelli et al. 2012), dinoflagellates (22:6n-3; Zhukova and Aizdacher 1995; Lavaud et al. 2018) and bacteria (Branched FA and 18:1n-7; Perry et al. 1979; Zhukova et al. 1992; Haack et al. 1994), which is in agreement with previous interpretations done in other comparable coastal bays (Dubois et al., 2014; Leroy et al., 2013). In sediment, bacteria are often associated with detritus and are therefore not considered as a high-quality food source (Dalsgaard et al., 2003). This is confirmed by the PUFA/SFA ratio, a biomarker of fresh vs. detritic OM (Connelly et al., 2015, 2016; Parrish et al., 2005), which was lower in the slipper limpet (~ 1.5 , our study) than in other suspension-feeding species of the bay of Brest, such as *Pecten maximus* (~ 2.8 , Lavaud et al. 2018) or *Ophiotrix fragilis* (~ 2 , Blanchet-Aurigny et al. 2015). The fact that *C. fornicata* lacks pre-ingestive mechanisms for particle selection likely explains their opportunistic trophic behaviour based on both fresh and detritic organic matter (Beninger et al., 2007). After spring bloom, the percentages of diatom's markers 16:1n-7 and 20:5n-3 drastically increased in *C. fornicata*'s tissue, as well as the 20:5n-3/22:6n-3 ratio, confirming that all ontogenic stages of *C. fornicata* benefit from this food supply from the water column (Budge and Parrish, 1998; Lavaud et al., 2018).

Besides, the FA 18:4n-3 showed an increasing contribution over time for ontogenic stages. According to the literature, this FA may originate from different primary producers such as dinoflagellates (Budge and Parrish, 1998) or green macroalgae (Fleurence et al., 1994; Kelly and Scheibling, 2012). Considering the absence or low temporal variation observed for others dinoflagellate biomarkers (peridinin pigment and 22:6n-3 FA) in the OM sources and the frequent seasonal accumulation of green macroalgae

near our study site (Study Centre for Algal Promotion, <http://www.ceva.fr>; Ragueneau et al. 2018), we can expect a seasonal trophic role of these green macroalgae for *C. fornicata* at our study site, probably in the form of detrital particles. The FA 20:1n-11 was found in relative high abundance in *C. fornicata* (4-6 %). Although it is known to be a biomarker of copepod in tropical estuaries (Bachok et al., 2003), the ¹⁵N-enrichment of *C. fornicata* compared to SPOM was not high enough to suggest a significant contribution of zooplankton in its diet (Kopp et al., 2015). Comparable percentages of 20:1n-11 were reported for *Ophiotrix fragilis* in the bay of Brest (Blanchet-Aurigny et al., 2015), evidencing that this FA was not a good biomarker of zooplankton for the species considered.

Contrary to SI, FA compositions of the three ontogenic stages of *C. fornicata* showed low overlap, especially between motile and sessile limpets. Considering that lipids and fatty acids profiles are age- and sex-specific (Correia et al., 2003; Pernet et al., 2012), some changes in FA compositions are then likely to originate from physiological changes between ontogenic stages. During winter period, motile males were characterized by higher proportions of SFA such as 16:0 (25-34 %) and 18:0 (27-36 %). These FA are very common in marine organisms and do not necessarily reflect a specific diet (Dalsgaard et al., 2003; Kelly and Scheibling, 2012). Moreover, despite the dietary interest of short-chain SFA, where energy is more efficiently released via beta-oxidation than for PUFA (Langdon and Waldock, 1981), temperature may also influence the process of their utilization (Pernet et al., 2007). For example, it has been experimentally demonstrated that cold-acclimated oysters (5-7°C) have a clear preference for PUFA (n-3) over SFA (16:0) as fuel for energy compared to 'temperate' oysters (Chu and Greaves, 1991). The lower utilization of SFA in cold-acclimated oysters has been attributed to the fact that SFA are not in the liquid phase under cold temperature, thus making them less accessible for catabolic processes. In *Crepidula fornicata*, young individuals have proportionally less energy reserves than adults (Guérin, 2004) and are more subjected to low temperature effects due to a higher surface-to-volume ratio (Diederich et al., 2015). Hence, the lower utilization of SFA could explain their higher SFA percentages during winter period in the bay of Brest, where temperature fall down to 7°C (Figure S2). Moreover, the weight-specific metabolic rate, which is higher in smaller organisms (Bayne and Newell, 1983; Bougrier et al., 1995), could be exacerbated in *C. fornicata* because motile males are more active than adults through their motility (Coe, 1936; Hoagland, 1978; Walne, 1956). Together, these results explain the higher variability in their FA compositions, among individuals but also between sampling dates. They also suggest that young motile individuals of *C. fornicata*, having less energetic storage in winter while having more energetic needs, are probably in poor energetic condition during this period.

As we measured neutral FA in digestive gland and gonad simultaneously (because the digestive gland cannot be isolated from the gonad), the level of lipid storage and FA composition may also depend on their sexual development stage. Indeed, females allocate more energy than males in the reproduction due to maternal gametogenesis (Deslous-Paoli and Héral, 1986; Leroy et al., 2013). As an illustration of FA composition changes, the n-3/n-6 ratio increased over time for both sessile and motile males whereas it remains unchanged in females. This may be linked to a preferential allocation of n-3 to early embryos, which showed an increase over the reproductive period of the slipper limpet (Leroy et al., 2013).

Finally, we found a surprisingly high amount of non-methylene interrupted (NMI) FA in neutral lipids (5-10 % on average in sessile stages) while these FA are preferentially incorporated in polar lipids (e.g., 8 % of NMI FA reported in polar lipids of *C. fornicata* by Dagorn et al. 2014). NMI FA are specific PUFA *de novo* synthesized by marine molluscs (Barnathan, 2009; Zhukova, 1991). Although their biological role and function are not well understood, the NMI FA have an unusual unsaturation pattern that confers to cell membranes a higher resistance to oxidative processes and microbial lipases than for the common PUFA. The content of NMI FA may thus represent a biochemical adapting feature of benthic organisms to their specific habitat (Barnathan, 2009). Because *C. fornicata* showed ontogenic histological changes in its foot (the largest part of the body) when becoming completely sessile (Androuin et al., 2019; Chaparro et al., 1998), high amount of such FA may act as a protection against potential negative effect of the sessility.

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898 **7. Supplementary materials**

899

900 **Table S1: Pigment (% , mean $\pm$ SD, n $\geq$ 2) composition of organic matter sources (Biofilm, suspended particulate organic matter (SPOM), superficial**
 901 **sedimentary organic matter (SSOM)) over the sampling survey. UK: Unknown pigments.**

Pigments	S1 = 26th February			S2 = 21th March			S3 = 28th March			S4 = 12th April			S5 = 14th June		
	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM
Alloxanthin	4 $\pm$ 2.2	11.7 $\pm$ 1.3	3.9 $\pm$ 0.1	1.2 $\pm$ 0.8	7.1 $\pm$ 1.2	3 $\pm$ 0.8	1.7 $\pm$ 1	9.3 $\pm$ 0.3	3.3 $\pm$ 0.3	2 $\pm$ 0.2	5.1 $\pm$ 0.8	4.1 $\pm$ 0.3	1.7 $\pm$ 0.6	15.1 $\pm$ 1.3	2.1 $\pm$ 0.4
β caroten	1.8 $\pm$ 0.6	0 $\pm$ 0	0.7 $\pm$ 0.2	0.2 $\pm$ 0.1	0.3 $\pm$ 0.2	0 $\pm$ 0	0.4 $\pm$ 0.1	0.6 $\pm$ 0.1	0 $\pm$ 0	1.5 $\pm$ 0.9	3.2 $\pm$ 3.4	0.1 $\pm$ 0.1	1.5 $\pm$ 0.5	2.8 $\pm$ 0.5	1.9 $\pm$ 1.6
Chlorophyll <i>a</i>	4.7 $\pm$ 0.6	7.7 $\pm$ 1.4	0.8 $\pm$ 0.2	4.1 $\pm$ 0.4	6.1 $\pm$ 0.7	0.3 $\pm$ 0.1	4.1 $\pm$ 0.4	7.2 $\pm$ 0.1	0.4 $\pm$ 0.1	4.7 $\pm$ 0.6	6.3 $\pm$ 0.6	0.3 $\pm$ 0.1	5 $\pm$ 0.4	4.7 $\pm$ 0.1	0.7 $\pm$ 0.1
Chlorophyll <i>b</i>	19.8 $\pm$ 1.7	7.8 $\pm$ 0.7	0.6 $\pm$ 0.1	25.8 $\pm$ 2.8	4.8 $\pm$ 0.5	0.3 $\pm$ 0.1	21.7 $\pm$ 0.1	7.1 $\pm$ 0.1	0.3 $\pm$ 0.1	19.6 $\pm$ 2.9	4.3 $\pm$ 0.6	0.2 $\pm$ 0.1	16 $\pm$ 0.6	5.1 $\pm$ 0	0.3 $\pm$ 0.2
Chlorophyllide	0.3 $\pm$ 0.1	0 $\pm$ 0	0.1 $\pm$ 0	0.3 $\pm$ 0.3	0.1 $\pm$ 0.1	0.1 $\pm$ 0	0.4 $\pm$ 0.1	0.2 $\pm$ 0	0.1 $\pm$ 0	0.5 $\pm$ 0.4	0.3 $\pm$ 0.1	0.1 $\pm$ 0	0.4 $\pm$ 0.1	0.2 $\pm$ 0	0 $\pm$ 0
Diadinoxanthin	0.8 $\pm$ 0.1	2.7 $\pm$ 0.4	1.1 $\pm$ 0.1	0.6 $\pm$ 0.1	3.4 $\pm$ 0.2	1 $\pm$ 0.1	0.7 $\pm$ 0.1	3.4 $\pm$ 0	0.9 $\pm$ 0.3	0.8 $\pm$ 0.1	2.1 $\pm$ 0.1	1.2 $\pm$ 0	1 $\pm$ 0	1.5 $\pm$ 0	1 $\pm$ 0.2
Xanthophyll	1.1 $\pm$ 0	2.5 $\pm$ 0.1	3.5 $\pm$ 0.2	0 $\pm$ 0	1.8 $\pm$ 1.6	2.9 $\pm$ 0.7	0.6 $\pm$ 0.1	2 $\pm$ 0.1	3.3 $\pm$ 0.3	0.7 $\pm$ 0.2	2.1 $\pm$ 0.2	3.1 $\pm$ 0.1	0.6 $\pm$ 0.1	3 $\pm$ 0.2	2.1 $\pm$ 0.5
Fucoxanthin-like	1.1 $\pm$ 0.1	2.7 $\pm$ 0.2	4.2 $\pm$ 0.4	1.2 $\pm$ 0.1	3.5 $\pm$ 0.1	5.6 $\pm$ 0.8	1.4 $\pm$ 0.4	4 $\pm$ 0.3	5.1 $\pm$ 0.2	2.8 $\pm$ 1.9	3 $\pm$ 0.1	5.7 $\pm$ 0.6	1.7 $\pm$ 0.2	3.1 $\pm$ 0.1	4.5 $\pm$ 0.5
Fucoxanthin	12.5 $\pm$ 2.8	22.5 $\pm$ 1.8	5.8 $\pm$ 0.9	10.8 $\pm$ 2.4	21.5 $\pm$ 1.4	5 $\pm$ 1.2	12.2 $\pm$ 3.4	19.2 $\pm$ 0.1	6.1 $\pm$ 1.3	12.3 $\pm$ 1.3	22.1 $\pm$ 0.9	6.8 $\pm$ 0.3	16.1 $\pm$ 2.3	11.3 $\pm$ 0.1	4.3 $\pm$ 1.1
Pheophorbide	14.8 $\pm$ 1.2	20 $\pm$ 2.7	32.8 $\pm$ 1.7	6.7 $\pm$ 2.4	23.7 $\pm$ 3	25.3 $\pm$ 1.6	8.8 $\pm$ 1.2	15.8 $\pm$ 0.5	26.9 $\pm$ 1.4	13.2 $\pm$ 3.4	22.3 $\pm$ 1.7	24.3 $\pm$ 0.6	12 $\pm$ 1.1	18.8 $\pm$ 2.8	20.8 $\pm$ 1.1
Lutein	3 $\pm$ 0.1	8.5 $\pm$ 0.8	13.9 $\pm$ 1.6	1.5 $\pm$ 0.5	7.5 $\pm$ 0.7	11.8 $\pm$ 2.3	2.2 $\pm$ 0.3	2.9 $\pm$ 0.4	10.6 $\pm$ 1	1.2 $\pm$ 0.3	4.3 $\pm$ 0.6	8.3 $\pm$ 1.6	1 $\pm$ 0.1	2 $\pm$ 0	3.8 $\pm$ 1.5
Neoxanthin	7.7 $\pm$ 1.2	1.7 $\pm$ 0.2	0 $\pm$ 0	14.8 $\pm$ 0.7	2.7 $\pm$ 0.4	0 $\pm$ 0	12.9 $\pm$ 0.5	5.1 $\pm$ 0.2	0 $\pm$ 0	6.1 $\pm$ 4.3	2.1 $\pm$ 0.1	2.9 $\pm$ 0.1	7.2 $\pm$ 0.9	2 $\pm$ 0.2	0.7 $\pm$ 1.2
Pheophytin	4.1 $\pm$ 3.3	0 $\pm$ 0	23.2 $\pm$ 1.1	1 $\pm$ 0.7	0.8 $\pm$ 1.4	32.9 $\pm$ 2.4	2.9 $\pm$ 0.6	1.3 $\pm$ 0.1	31.1 $\pm$ 1.8	7.9 $\pm$ 1.4	9.5 $\pm$ 5.8	30.1 $\pm$ 1.9	15 $\pm$ 1.5	17.9 $\pm$ 1.8	51.4 $\pm$ 2.6
Chlorophyll <i>c</i>	1.5 $\pm$ 0.5	4.8 $\pm$ 0.5	3.7 $\pm$ 0.2	0 $\pm$ 0	12.5 $\pm$ 1.5	6.4 $\pm$ 0.2	1.3 $\pm$ 0.2	15.2 $\pm$ 0.1	7.2 $\pm$ 0.7	1.3 $\pm$ 0.3	6.6 $\pm$ 1	8 $\pm$ 1.4	1.9 $\pm$ 0.1	4.1 $\pm$ 0.3	3.9 $\pm$ 0.4
UK 3	1.1 $\pm$ 0.3	0.9 $\pm$ 0	0.8 $\pm$ 0.1	1.6 $\pm$ 0.2	0.9 $\pm$ 0.4	0.7 $\pm$ 0.1	1.1 $\pm$ 0	1.6 $\pm$ 0.1	0.6 $\pm$ 0	1.1 $\pm$ 0.3	0.7 $\pm$ 0.4	1 $\pm$ 0.2	0.7 $\pm$ 0	0.7 $\pm$ 0.2	0.4 $\pm$ 0.1
UK 6	17.3 $\pm$ 1.3	0 $\pm$ 0	0 $\pm$ 0	26.7 $\pm$ 4.5	0 $\pm$ 0	0 $\pm$ 0	23 $\pm$ 1.3	0 $\pm$ 0	0 $\pm$ 0	20.4 $\pm$ 3.8	0 $\pm$ 0	0 $\pm$ 0	14.5 $\pm$ 0.9	0 $\pm$ 0	0 $\pm$ 0
UK 7	1.2 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0	1.7 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0	1.1 $\pm$ 0.3	0 $\pm$ 0	0 $\pm$ 0	1.6 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0	0.9 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0
UK 8	0.6 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0	1.1 $\pm$ 0.6	0 $\pm$ 0	0 $\pm$ 0	1 $\pm$ 0.8	0 $\pm$ 0	0 $\pm$ 0	0.7 $\pm$ 0.6	0 $\pm$ 0	0 $\pm$ 0	1.3 $\pm$ 0.5	0 $\pm$ 0	0 $\pm$ 0
UK 9	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0.8 $\pm$ 0.3	0 $\pm$ 0	0 $\pm$ 0	0.8 $\pm$ 0	0.2 $\pm$ 0.2
Violaxanthin	0.6 $\pm$ 0.2	1.3 $\pm$ 0.3	0 $\pm$ 0	0 $\pm$ 0	0.7 $\pm$ 0.7	0 $\pm$ 0	0.9 $\pm$ 0.1	3.3 $\pm$ 0.4	0 $\pm$ 0	0.3 $\pm$ 0.5	1.8 $\pm$ 1.7	0 $\pm$ 0	0.2 $\pm$ 0.4	4.5 $\pm$ 0.1	0 $\pm$ 0
Zeaxanthin	2 $\pm$ 0.1	5.2 $\pm$ 0.2	5 $\pm$ 0.5	0.8 $\pm$ 0.4	2.5 $\pm$ 0.4	4.8 $\pm$ 1.1	1.6 $\pm$ 0	1.7 $\pm$ 0.1	4.2 $\pm$ 0.3	1.1 $\pm$ 0	3.2 $\pm$ 0.3	3.7 $\pm$ 0.6	0.9 $\pm$ 0.1	2.3 $\pm$ 0.1	1.9 $\pm$ 0.6
Chl <i>c</i> / Chl <i>a</i> ratio	0.3 $\pm$ 0.1	0.6 $\pm$ 0.2	4.5 $\pm$ 0.7	0 $\pm$ 0	2.1 $\pm$ 0.5	27.3 $\pm$ 17.1	0.3 $\pm$ 0.1	2.1 $\pm$ 0	17.1 $\pm$ 1.2	0.3 $\pm$ 0.1	1.1 $\pm$ 0.1	28 $\pm$ 14.9	0.4 $\pm$ 0.1	0.9 $\pm$ 0	5.8 $\pm$ 1.1
Chl <i>b</i> / Chl <i>a</i> ratio	4.2 $\pm$ 0.2	1 $\pm$ 0.1	0.7 $\pm$ 0.1	6.3 $\pm$ 0.7	0.8 $\pm$ 0	1.2 $\pm$ 0.5	5.3 $\pm$ 0.5	1 $\pm$ 0	0.7 $\pm$ 0.2	4.2 $\pm$ 0.6	0.7 $\pm$ 0	0.6 $\pm$ 0	3.2 $\pm$ 0.4	1.1 $\pm$ 0	0.5 $\pm$ 0.3
Fuco / Chl <i>a</i> ratio	2.6 $\pm$ 0.3	3 $\pm$ 0.8	7 $\pm$ 1.7	2.6 $\pm$ 0.6	3.6 $\pm$ 0.6	19.3 $\pm$ 6.7	2.9 $\pm$ 0.6	2.7 $\pm$ 0	14.5 $\pm$ 3.2	2.7 $\pm$ 0.5	3.5 $\pm$ 0.4	23 $\pm$ 9.8	3.2 $\pm$ 0.2	2.4 $\pm$ 0.1	6.3 $\pm$ 1.4

902 **Table S2: Fatty acid (FA) (% , mean $\pm$ SD, n = 3) composition of organic matter sources (Biofilm, suspended particulate organic matter (SPOM), superficial**
903 **sedimentary organic matter (SSOM)) over the sampling survey. Only FA accounting for more than 0.5 % of total FA in at least one sample was shown.**
904 **BFA: Branched FA; SFA: saturated FA; MUFA: monounsaturated FA; PUFA: polyunsaturated FA; UK FA: Unknown FA; EPA: 20:5n-3; DHA: 22:6n-**
905 **3.**

Fatty acids	S1 = 26 th February			S2 = 21 th March			S3 = 28 th March			S4 = 12 th April			S5 = 14 th June		
	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM
TMTD	0.1 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0	0.1 $\pm$ 0.1	0.4 $\pm$ 0	0.6 $\pm$ 0	0.1 $\pm$ 0.1	0 $\pm$ 0	0.2 $\pm$ 0.2	0 $\pm$ 0	0 $\pm$ 0	0.3 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0
15:0iso	1.8 $\pm$ 0.7	1 $\pm$ 0.2	2.7 $\pm$ 0.4	0.6 $\pm$ 0.5	1 $\pm$ 0.1	1.9 $\pm$ 0.8	1.3 $\pm$ 0.9	1.1 $\pm$ 0.1	2 $\pm$ 0.1	1.2 $\pm$ 0.4	1.3 $\pm$ 0.2	2.5 $\pm$ 0.2	1.4 $\pm$ 0.3	1 $\pm$ 0.2	2 $\pm$ 0.2
15:0ant	0.6 $\pm$ 0.1	1.2 $\pm$ 0.1	3.7 $\pm$ 0.7	0.3 $\pm$ 0.1	0.9 $\pm$ 0.1	2.9 $\pm$ 0.6	0.8 $\pm$ 0.2	1 $\pm$ 0.1	3 $\pm$ 0.4	0.5 $\pm$ 0.2	1.1 $\pm$ 0.2	3.2 $\pm$ 0.3	0.2 $\pm$ 0.2	0.4 $\pm$ 0.3	2.3 $\pm$ 0.8
16:0iso	0.8 $\pm$ 0.2	0.9 $\pm$ 0.2	2.3 $\pm$ 0.4	0.2 $\pm$ 0	0.5 $\pm$ 0	0.9 $\pm$ 0.1	0.6 $\pm$ 0.2	0.5 $\pm$ 0	0.8 $\pm$ 0.2	0.5 $\pm$ 0.2	0.5 $\pm$ 0.1	0.9 $\pm$ 0.1	0.6 $\pm$ 0.1	0.2 $\pm$ 0.1	0.5 $\pm$ 0.4
17:0iso	0.9 $\pm$ 0.3	1.7 $\pm$ 0.2	1.4 $\pm$ 0.2	0.1 $\pm$ 0.1	0.3 $\pm$ 0.1	1.2 $\pm$ 0.1	0.5 $\pm$ 0.4	2.8 $\pm$ 0.1	1.2 $\pm$ 0.1	0.8 $\pm$ 0.5	2.7 $\pm$ 0.2	1.6 $\pm$ 0.1	1.8 $\pm$ 0.6	0.8 $\pm$ 0.5	0.7 $\pm$ 0.6
17:0ant	2.5 $\pm$ 2.4	0.3 $\pm$ 0.3	0.9 $\pm$ 0.1	0.9 $\pm$ 0.6	1.2 $\pm$ 0	1.1 $\pm$ 0.1	2.6 $\pm$ 0.8	0 $\pm$ 0	0.6 $\pm$ 0	1.6 $\pm$ 0.5	0 $\pm$ 0	0.2 $\pm$ 0.2	1.2 $\pm$ 0.2	0.6 $\pm$ 0.1	0.9 $\pm$ 0.8
18:0iso	0.1 $\pm$ 0.1	1.2 $\pm$ 0.9	1.1 $\pm$ 0.2	0.1 $\pm$ 0.1	0.4 $\pm$ 0	0.6 $\pm$ 0.1	1.1 $\pm$ 0.1	0.9 $\pm$ 0.2	0.4 $\pm$ 0.1	0.4 $\pm$ 0.3	0.9 $\pm$ 0.3	0.6 $\pm$ 0.1	1.7 $\pm$ 0.4	1.2 $\pm$ 0	1.7 $\pm$ 0.4
Σ BFA	6.8 $\pm$ 2.4	6.3 $\pm$ 0.3	12.1 $\pm$ 1.4	2.2 $\pm$ 0.2	4.4 $\pm$ 0.1	8.5 $\pm$ 1.6	6.8 $\pm$ 0.7	6.2 $\pm$ 0.4	8 $\pm$ 0.7	5 $\pm$ 1.3	6.6 $\pm$ 0.9	9 $\pm$ 0.4	7 $\pm$ 0.6	4.2 $\pm$ 0.3	8.1 $\pm$ 1.5
14:0	3.1 $\pm$ 0.3	7.3 $\pm$ 1.3	4.9 $\pm$ 0.8	1.5 $\pm$ 0.4	5.2 $\pm$ 0.5	3.5 $\pm$ 0	4.2 $\pm$ 0.6	12 $\pm$ 0.8	4.1 $\pm$ 0.2	6.1 $\pm$ 2.1	8.6 $\pm$ 1	4.1 $\pm$ 0.3	4.9 $\pm$ 0.2	5.4 $\pm$ 1.1	5.3 $\pm$ 0.6
15:0	0.9 $\pm$ 0.3	2.6 $\pm$ 0.4	1.5 $\pm$ 0.2	0.2 $\pm$ 0.1	1.9 $\pm$ 0.1	1.2 $\pm$ 0.1	0.9 $\pm$ 0.2	1.7 $\pm$ 0.1	1 $\pm$ 0.1	0.8 $\pm$ 0.1	1.4 $\pm$ 0.2	1.2 $\pm$ 0.1	0.7 $\pm$ 0.1	0.8 $\pm$ 0.2	0.7 $\pm$ 0.6
16:0	21.5 $\pm$ 3.4	33 $\pm$ 4.9	22.4 $\pm$ 3.1	38.6 $\pm$ 0.7	26.3 $\pm$ 2.3	17.6 $\pm$ 1.6	29.5 $\pm$ 4.3	32.7 $\pm$ 1.9	14.8 $\pm$ 1.4	30.5 $\pm$ 4.4	34.1 $\pm$ 2.3	18.2 $\pm$ 0.2	19.9 $\pm$ 4.6	25 $\pm$ 4.7	19.3 $\pm$ 4.3
17:0	1.6 $\pm$ 0.7	1.6 $\pm$ 0.3	1.3 $\pm$ 0.2	0.8 $\pm$ 0.8	0.7 $\pm$ 0	0.9 $\pm$ 0.1	0.8 $\pm$ 0.2	0.9 $\pm$ 0.1	0.8 $\pm$ 0.1	0.7 $\pm$ 0.1	0.9 $\pm$ 0.1	1 $\pm$ 0.1	1.6 $\pm$ 0.4	0.2 $\pm$ 0.2	1.8 $\pm$ 0.5
18:0	16.5 $\pm$ 1.3	11.8 $\pm$ 1.7	9.6 $\pm$ 1.3	42.3 $\pm$ 2.6	13.3 $\pm$ 1.4	10.9 $\pm$ 2.5	21.5 $\pm$ 4.8	9.2 $\pm$ 0.7	6.8 $\pm$ 1	21.9 $\pm$ 3.8	14.9 $\pm$ 5.3	7.6 $\pm$ 0.3	6.7 $\pm$ 2.5	6.5 $\pm$ 1.3	8 $\pm$ 2.1
20:0	0.6 $\pm$ 0.1	1.7 $\pm$ 0.1	1.9 $\pm$ 1.7	0.6 $\pm$ 0	1.3 $\pm$ 0	2.8 $\pm$ 0.3	0.8 $\pm$ 0.2	0.9 $\pm$ 0	2.6 $\pm$ 0.3	0.9 $\pm$ 0.1	0.4 $\pm$ 0.7	3 $\pm$ 0.2	1.1 $\pm$ 0.7	0.9 $\pm$ 0.2	2.2 $\pm$ 0.3
22:0	1 $\pm$ 0.1	1.4 $\pm$ 1.2	3.2 $\pm$ 3	0.3 $\pm$ 0.2	1.6 $\pm$ 0	4.4 $\pm$ 0.7	1.1 $\pm$ 0.3	0.7 $\pm$ 0.6	4.5 $\pm$ 0.5	1.4 $\pm$ 0.4	1.4 $\pm$ 0.3	5.4 $\pm$ 0.3	1.2 $\pm$ 0.9	0.7 $\pm$ 0.1	3.5 $\pm$ 0.8
24:0	0.1 $\pm$ 0	0 $\pm$ 0	0.5 $\pm$ 0.4	0 $\pm$ 0	0 $\pm$ 0	0.1 $\pm$ 0.3	0.7 $\pm$ 0.3	4.4 $\pm$ 2	6.1 $\pm$ 5.2	0.1 $\pm$ 0.1	0.1 $\pm$ 0.2	0.3 $\pm$ 0.3	1.9 $\pm$ 1.5	1.1 $\pm$ 0.2	4.9 $\pm$ 1.2
Σ SFA	45.3 $\pm$ 4.7	59.4 $\pm$ 7.6	45.4 $\pm$ 3.3	84.2 $\pm$ 2.1	50.3 $\pm$ 3.8	41.4 $\pm$ 3.2	59.6 $\pm$ 9.9	62.4 $\pm$ 1.1	40.7 $\pm$ 7.3	62.3 $\pm$ 7.4	61.8 $\pm$ 5.9	40.7 $\pm$ 0.5	38.1 $\pm$ 10	40.5 $\pm$ 7.4	45.7 $\pm$ 8.3
14:1n-5	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0.1 $\pm$ 0.1	0 $\pm$ 0	0.5 $\pm$ 0.1	0.2 $\pm$ 0.2	0 $\pm$ 0	0 $\pm$ 0	0.2 $\pm$ 0.3	0.1 $\pm$ 0.1	0 $\pm$ 0
16:1n-11	0.3 $\pm$ 0.3	0 $\pm$ 0	0 $\pm$ 0	0.1 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0.1 $\pm$ 0.1	0 $\pm$ 0	0.6 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0
16:1n-9	0.9 $\pm$ 0.1	1.5 $\pm$ 0.9	0.3 $\pm$ 0.5	0.2 $\pm$ 0.2	2 $\pm$ 0.2	1.2 $\pm$ 0.2	0.6 $\pm$ 0.3	0.4 $\pm$ 0.4	0.7 $\pm$ 0.2	0.9 $\pm$ 0.2	0.5 $\pm$ 0.1	0.9 $\pm$ 0.1	0.5 $\pm$ 0.4	0.8 $\pm$ 0.1	0.5 $\pm$ 0.4
16:1n-7	3.8 $\pm$ 0.1	1.9 $\pm$ 1.5	5.6 $\pm$ 3	0.8 $\pm$ 0.4	4.5 $\pm$ 0.5	8.8 $\pm$ 0.1	2.7 $\pm$ 1.8	0.9 $\pm$ 0.2	7.3 $\pm$ 2.4	2.5 $\pm$ 1.1	1.6 $\pm$ 0.7	0 $\pm$ 0	7.9 $\pm$ 0.6	3.4 $\pm$ 0.8	9 $\pm$ 1.9

16:1n-5	0.5 ± 0.1	0.8 ± 0.1	0.4 ± 0.4	0.1 ± 0	0.6 ± 0.1	0.7 ± 0.1	0.4 ± 0	0.5 ± 0.4	0.8 ± 0.2	0.3 ± 0.1	0.4 ± 0.1	1.2 ± 0.3	0.5 ± 0.4	0.1 ± 0.2	0.8 ± 0.7
17:1n-7	0.1 ± 0.2	0.4 ± 0.1	0.8 ± 0.3	0.2 ± 0.1	0.4 ± 0.1	0.6 ± 0.1	0.6 ± 0.3	0.8 ± 0.1	0.3 ± 0.2	0.7 ± 0.3	0.8 ± 0.1	0.7 ± 0.2	0.4 ± 0.3	1.2 ± 0.2	0.1 ± 0.3
18:1n-11	0.5 ± 0.3	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0.1 ± 0.1	0 ± 0	0.3 ± 0	0 ± 0	0 ± 0	0 ± 0
18:1n-9	1.6 ± 0.7	0.3 ± 0.5	1.3 ± 2.3	1 ± 0.7	2.5 ± 0.7	2.9 ± 0.2	0.9 ± 0.8	0 ± 0	1.9 ± 1	2.2 ± 0.9	0.2 ± 0.3	3.1 ± 1.1	1.8 ± 0.3	5.8 ± 1.4	2.3 ± 0.2
18:1n-7	2 ± 0.2	0 ± 0	0 ± 0	1.1 ± 0.3	2.7 ± 0.5	5.8 ± 0.3	2.1 ± 1.3	0.5 ± 0.5	4.2 ± 1.5	2 ± 0.8	0.6 ± 0.7	6.9 ± 1.4	4.3 ± 0.6	3.7 ± 1	4.6 ± 1.1
18:1n-5	0.6 ± 0.1	0.1 ± 0.2	0 ± 0	0.1 ± 0.1	0.1 ± 0.2	0 ± 0	0.4 ± 0	0 ± 0	0.3 ± 0.2	0.3 ± 0.2	0 ± 0	0.3 ± 0.3	0.6 ± 0.3	0 ± 0	0 ± 0
20:1n-11	1.2 ± 0.4	1.1 ± 1.5	0.3 ± 0.5	0.2 ± 0.1	3.6 ± 0.6	0.5 ± 0.4	0.5 ± 0.4	1.2 ± 1	0.6 ± 0.6	0.6 ± 0.3	0.4 ± 0.7	0.8 ± 0	0.5 ± 0.4	3.6 ± 0.8	0.1 ± 0.3
20:1n-9	0.3 ± 0	0.3 ± 0.5	0.3 ± 0.4	0.1 ± 0	0 ± 0	0.3 ± 0.3	0.3 ± 0.2	0 ± 0	0.4 ± 0.8	0.3 ± 0	0.5 ± 0.4	0.1 ± 0.2	0.3 ± 0.3	0 ± 0	0 ± 0
20:1n-7	0.6 ± 0.5	0 ± 0	0 ± 0	0.1 ± 0	0 ± 0	0 ± 0	0.2 ± 0.1	0 ± 0	0 ± 0	0.2 ± 0.1	0 ± 0	0.1 ± 0.2	0.3 ± 0.3	0 ± 0	0 ± 0
22:1n-9	0.2 ± 0.1	0 ± 0	0.3 ± 0.6	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0
Σ MUFA	12.7 ± 1.3	6.6 ± 4.2	9.2 ± 6.4	3.9 ± 0.5	16.5 ± 2.7	20.7 ± 0.6	8.7 ± 4.8	4.3 ± 2.4	17 ± 6.6	10.4 ± 2.5	5.2 ± 1.9	15.1 ± 3.8	17.3 ± 3.4	18.6 ± 4.3	17.5 ± 4.7
16:2n-4	0.2 ± 0.2	0 ± 0	0.2 ± 0.4	0.2 ± 0.2	0.1 ± 0.1	0.7 ± 0.1	2.1 ± 0.7	0 ± 0	0.4 ± 0.5	0.9 ± 1.1	0.3 ± 0.3	0.2 ± 0.2	0.9 ± 0.2	1.5 ± 0.3	0.8 ± 0.7
16:3n-4	0.6 ± 0.1	0.3 ± 0.5	0.7 ± 0.6	0.1 ± 0.1	1.2 ± 0.1	1.3 ± 0.1	0 ± 0	0 ± 0	0 ± 0	0.5 ± 0.1	0 ± 0	1 ± 0.9	0.5 ± 0.4	0.3 ± 0.3	0.5 ± 0.4
16:3n-6	0.4 ± 0.2	0.2 ± 0.3	0 ± 0	0.1 ± 0.1	0 ± 0	0.3 ± 0.3	0.4 ± 0.3	0 ± 0	0.2 ± 0.4	0.3 ± 0.1	0.1 ± 0.2	1.1 ± 0.2	0.3 ± 0.3	0 ± 0	0 ± 0
16:3n-3	0.2 ± 0.1	0 ± 0	0.2 ± 0.3	0.1 ± 0.1	1 ± 0.1	1.2 ± 0.3	0.6 ± 0.5	0 ± 0	1.2 ± 0.9	0.4 ± 0.1	0.3 ± 0.3	0.9 ± 0.2	0.1 ± 0.1	0.1 ± 0.2	0.2 ± 0.3
16:4n-3	0.3 ± 0.1	1.3 ± 1.1	1.2 ± 0.1	0.2 ± 0.2	1.3 ± 0.1	0.7 ± 0.1	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0
18:2n-6	4.2 ± 2.1	0.9 ± 0.8	0.7 ± 1	1.9 ± 0.4	1.2 ± 0.2	1.1 ± 0.2	3 ± 1.8	0.9 ± 0.1	1.2 ± 0.3	2.3 ± 0.9	0.9 ± 0.2	1.4 ± 0.3	3.1 ± 0.6	2.5 ± 0.5	1.3 ± 0.4
18:2n-4	0.2 ± 0.1	0.2 ± 0.4	0 ± 0	0 ± 0.1	0.2 ± 0.2	0.4 ± 0.1	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0
18:3n-6	0.2 ± 0	0.1 ± 0.3	0.4 ± 0.7	0.1 ± 0	0.3 ± 0	0 ± 0	0.3 ± 0	0 ± 0	0.2 ± 0.2	0.2 ± 0.2	0.1 ± 0.2	0.4 ± 0.1	0.2 ± 0.1	0.2 ± 0.3	0 ± 0
18:3n-4	0.6 ± 0.1	1.4 ± 0.5	5.9 ± 1.7	0.1 ± 0	1.2 ± 0.4	1.6 ± 0.1	0.4 ± 0	0.8 ± 0.3	2.5 ± 1.7	0.2 ± 0	0.7 ± 0.3	0.7 ± 0.1	0 ± 0.1	0.2 ± 0.1	3.2 ± 0.4
18:3n-3	0.7 ± 0.3	0.6 ± 0.7	0.3 ± 0.5	0.2 ± 0.1	1.3 ± 0.2	0.7 ± 0	0.5 ± 0.2	0.2 ± 0.3	0.4 ± 0.4	0.3 ± 0.2	0.4 ± 0.4	0.7 ± 0.1	0.5 ± 0.4	2.8 ± 0.8	0.3 ± 0.5
18:4n-3	1.1 ± 0.9	2.7 ± 0.3	1.2 ± 0.2	0.2 ± 0.1	3.8 ± 0.5	1.6 ± 0.2	1 ± 0.4	2.5 ± 0.5	2.2 ± 0.6	0.8 ± 0.3	1.9 ± 0.6	2.5 ± 0.5	1.1 ± 0.4	4.3 ± 1.2	0.9 ± 0.9
20:2n-6	0.6 ± 0.2	1.3 ± 0.2	0.9 ± 0.2	0.1 ± 0	0.1 ± 0.2	0.2 ± 0.3	0.3 ± 0	0 ± 0	0.4 ± 0.7	0.4 ± 0.4	1.1 ± 0.2	1.3 ± 0.1	0.5 ± 0.4	1.6 ± 0.5	1.3 ± 0.6
20:3n-6	0.2 ± 0	0.5 ± 0	0 ± 0	0.1 ± 0	0.5 ± 0	0.5 ± 0.1	0.2 ± 0.1	0 ± 0	0.4 ± 0.1	0.3 ± 0.2	0.3 ± 0.6	0.4 ± 0	0.2 ± 0.2	0 ± 0	0 ± 0
20:3n-3	0.2 ± 0	0.5 ± 0.9	0 ± 0	0.1 ± 0	0 ± 0	0.3 ± 0.2	0 ± 0	0 ± 0	0 ± 0	0.2 ± 0.1	0 ± 0	0.4 ± 0	1.7 ± 1.5	0.4 ± 0.2	0.2 ± 0.3
20:4n-6	6.3 ± 0.9	0 ± 0	0.7 ± 0.6	1.6 ± 0.2	0.2 ± 0.2	1.5 ± 0.1	2.8 ± 1.7	0.2 ± 0.3	1.2 ± 0.4	2 ± 0.7	0.1 ± 0.2	2.2 ± 0.4	5.9 ± 1.6	0.2 ± 0.2	1.8 ± 0.7
20:4n-3	0.4 ± 0.2	1.6 ± 0.2	1.1 ± 0.2	0.1 ± 0	0.7 ± 0	0.7 ± 0.1	0.8 ± 0.3	2.3 ± 0.5	0.6 ± 0.1	1.3 ± 0.5	2.2 ± 0.9	0.5 ± 0.4	0.2 ± 0.2	1.7 ± 1	0 ± 0
20:5n-3	8.2 ± 2.1	2.2 ± 1.2	4.6 ± 1	1.4 ± 0.7	4.5 ± 0.6	6.3 ± 1.2	3.4 ± 2.5	0.8 ± 0.3	3 ± 3	2.2 ± 0.6	1.7 ± 1	9.7 ± 2.3	11.2 ± 4.4	6.7 ± 1.8	8.5 ± 3.1
21:5n-3	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	3 ± 5.2	2.3 ± 3.4	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0

22:2n-6	0.3 ± 0.1	0.4 ± 0	0.9 ± 0.1	0.3 ± 0	0.9 ± 0	1.3 ± 0.2	2.1 ± 1.5	7.3 ± 1	2.8 ± 2	1.5 ± 0.8	9.1 ± 3.2	3 ± 2.5	3 ± 2.2	2.7 ± 1.2	2.9 ± 1.6
22:4n-6	0.6 ± 0.5	0.5 ± 0.1	0 ± 0	0.1 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0
22:5n-6	0.7 ± 0.1	2.1 ± 1.3	1.7 ± 0.5	0.1 ± 0	0.4 ± 0	0.6 ± 0.1	1.9 ± 0.2	4.4 ± 0.5	2.5 ± 0.9	1.6 ± 0.6	2.3 ± 0.8	0.9 ± 0.4	1 ± 1	1.7 ± 0.7	0.2 ± 0.3
22:5n-3	0.7 ± 0.8	0 ± 0	0 ± 0	0.1 ± 0	0.1 ± 0.2	0 ± 0	0.7 ± 0.1	2.1 ± 0.6	2.1 ± 1.7	0.6 ± 0.5	2.2 ± 0.5	2.3 ± 0.7	0.5 ± 0.3	1.3 ± 1.2	0.6 ± 0.6
22:6n-3	4.3 ± 2.8	0.9 ± 1.1	0.7 ± 0.7	0.5 ± 0.2	2.9 ± 0.4	1.9 ± 0.2	1.7 ± 0.3	4 ± 0.5	4.5 ± 0.8	1.5 ± 2.1	0 ± 0	3.6 ± 1.7	3.7 ± 1.7	6 ± 1.8	2.9 ± 1.1
Σ PUFA	31.2 ± 6.5	17.6 ± 4.7	21.5 ± 3.2	7.8 ± 1.6	22 ± 1.8	22.9 ± 1.8	22.3 ± 6.1	25.6 ± 1.6	28.7 ± 2.3	19.8 ± 4.7	23.9 ± 4.9	33.3 ± 3.1	34.5 ± 7	33.9 ± 4.9	25.6 ± 5.6
A	0.5 ± 0.4	0 ± 0	0 ± 0	0.1 ± 0	0.1 ± 0.1	0 ± 0	0.2 ± 0	0 ± 0	0 ± 0	0.3 ± 0	0 ± 0	0 ± 0	1.2 ± 1.4	0 ± 0	0 ± 0
M	0.8 ± 0.2	0.4 ± 0	0.2 ± 0.4	0.2 ± 0.1	0.4 ± 0	0 ± 0	0.5 ± 0.2	0 ± 0	0 ± 0	0.4 ± 0.2	0 ± 0	0 ± 0	0.4 ± 0.3	0 ± 0	0 ± 0
N	0.4 ± 0.1	0.5 ± 0	1.4 ± 1.8	0.2 ± 0.1	0.5 ± 0	0.6 ± 0.1	0.5 ± 0.2	0 ± 0	0.1 ± 0.2	0.4 ± 0	0.7 ± 0.6	0.4 ± 0	0 ± 0	0 ± 0	0 ± 0
O	0.8 ± 0	5.2 ± 2.1	5.5 ± 3.9	0.8 ± 0.6	3.1 ± 0.6	0.5 ± 0.5	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	1 ± 0.6	2.2 ± 1.5	2.7 ± 2.1
P	1.4 ± 0	4 ± 0.6	4.9 ± 1.1	0.5 ± 0.2	1.9 ± 0.1	3.9 ± 0.5	1.1 ± 0.5	0.9 ± 0.8	4.8 ± 1.3	1.4 ± 1.5	1.2 ± 0.3	0.6 ± 0.6	0.4 ± 0.3	0.3 ± 0.3	0.5 ± 0.5
W	0 ± 0	0 ± 0	0 ± 0	0.1 ± 0	0.6 ± 0	0.8 ± 0.1	0.1 ± 0.1	0.6 ± 0	0.5 ± 0	0.2 ± 0	0.7 ± 0.6	0.6 ± 0.1	0.2 ± 0.2	0.2 ± 0.2	0 ± 0
Σ UK FA	4 ± 0.7	10.1 ± 1.5	11.9 ± 5.2	1.7 ± 1.1	6.5 ± 0.6	5.8 ± 1.1	2.4 ± 0.6	1.5 ± 0.8	5.4 ± 1.3	2.6 ± 1.4	2.6 ± 1.3	1.6 ± 0.6	3.2 ± 1.2	2.7 ± 1.5	3.2 ± 1.9
PUFA/SFA	0.7 ± 0.2	0.3 ± 0.1	0.5 ± 0.1	0.1 ± 0	0.4 ± 0.1	0.6 ± 0.1	0.4 ± 0.2	0.4 ± 0	0.7 ± 0.2	0.3 ± 0.1	0.4 ± 0.1	0.8 ± 0.1	1 ± 0.4	0.9 ± 0.3	0.6 ± 0.2
EPA/DHA	2.3 ± 1	NA	NA	2.8 ± 0.5	1.6 ± 0	3.4 ± 0.4	1.9 ± 1.3	0.2 ± 0.1	0.8 ± 0.8	NA	NA	3.1 ± 1.6	3.1 ± 0.6	1.1 ± 0	2.9 ± 0.1
Σ n-3/ Σ n-6	1.2 ± 0.6	1.7 ± 0.4	1.8 ± 0.5	0.7 ± 0.3	4.4 ± 0	2.5 ± 0.4	0.8 ± 0.2	0.9 ± 0.1	2.1 ± 1.1	1.2 ± 0.4	0.6 ± 0.1	1.9 ± 0.4	1.3 ± 0.3	2.7 ± 0.7	2 ± 1.1

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907

908 **Table S3: Stable isotope (mean $\pm$ SD, n = 3) composition of organic matter sources (Biofilm, suspended particulate organic matter (SPOM), superficial**
 909 **sedimentary organic matter (SSOM)) over the sampling survey.**

	S1 = 26 th February			S2 = 21 th March			S3 = 28 th March			S4 = 12 th April			S5 = 14 th June		
	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM	Biofilm	SPOM	SSOM
$\delta^{13}\text{C}$	-22.6 $\pm$ 0.1	-25.2 $\pm$ 0.2	-23 $\pm$ 0.2	-23.4 $\pm$ 0.5	-25.3 $\pm$ 0.1	-22.7 $\pm$ 0.3	-22.6 $\pm$ 0.4	-25.3 $\pm$ 0.1	-22.6 $\pm$ 0.1	-23.4 $\pm$ 1	-24.6 $\pm$ 0.4	-22.5 $\pm$ 0.1	-21.8 $\pm$ 0.2	-25.1 $\pm$ 0.3	-22.5 $\pm$ 0.1
$\delta^{15}\text{N}$	9.4 $\pm$ 0.6	7.3 $\pm$ 0	7 $\pm$ 0.7	10.3 $\pm$ 0.5	6.7 $\pm$ 0.3	7.1 $\pm$ 0.1	9.2 $\pm$ 0.7	6.6 $\pm$ 0.1	7 $\pm$ 0.1	9.2 $\pm$ 0.7	6.9 $\pm$ 0.9	7.1 $\pm$ 0.1	9.4 $\pm$ 0.7	6.4 $\pm$ 0.5	6.9 $\pm$ 0.2
C/N	3.9 $\pm$ 0.3	6.6 $\pm$ 1.1	5.1 $\pm$ 1.3	5.8 $\pm$ 3	5.9 $\pm$ 0.6	5.3 $\pm$ 0.6	5.3 $\pm$ 0.4	6 $\pm$ 0.5	5.5 $\pm$ 0.6	5 $\pm$ 0.5	5.6 $\pm$ 0.9	5.4 $\pm$ 0.5	3.6 $\pm$ 0.6	3.9 $\pm$ 0.4	5.9 $\pm$ 0.8

910

911 **Table S4: Fatty acid (FA) (% , mean $\pm$ SD, n = 5) composition of the three ontogenic stages of *Crepidula fornicata* (motile males, sessile males, sessile**
912 **females) over the sampling survey. Only FA accounting for more than 0.5 % of total FA in at least one sample was shown. BFA: Branched FA; SFA:**
913 **saturated FA; MUFA: monounsaturated FA; PUFA: polyunsaturated FA; NMI FA: non-methyl-interrupted FA; DMA: Dimethyl acetals FA; EPA:**
914 **20:5n-3; DHA: 22:6n-3.**

Fatty acids	S1 = 26 th February			S2 = 21 th March			S3 = 28 th March			S4 = 12 th April			S5 = 14 th June		
	Sessile female	Sessile male	Motile male	Sessile female	Sessile male	Motile male	Sessile female	Sessile male	Motile male	Sessile female	Sessile male	Motile male	Sessile female	Sessile male	Motile male
TMTD	1.3 $\pm$ 0.3	2.1 $\pm$ 1.1	0.8 $\pm$ 0.6	1.3 $\pm$ 0.2	0 $\pm$ 0.1	0.5 $\pm$ 0.4	1.3 $\pm$ 0.7	2.5 $\pm$ 0.4	0.9 $\pm$ 1.2	2.1 $\pm$ 1	2.9 $\pm$ 0.8	2.5 $\pm$ 1	2.1 $\pm$ 1.7	2 $\pm$ 0.7	1 $\pm$ 0.6
15:0iso	0.2 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0.1 $\pm$ 0.1	0.1 $\pm$ 0.1	0.1 $\pm$ 0.2	0.1 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0	0.1 $\pm$ 0.1	0.1 $\pm$ 0.1	0.3 $\pm$ 0.4	0.1 $\pm$ 0.1	0.5 $\pm$ 0.3	1 $\pm$ 0.2
15:0ant	0.1 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0	0.1 $\pm$ 0.1	0 $\pm$ 0	0.3 $\pm$ 0.5	0 $\pm$ 0	0.4 $\pm$ 0.3	0.7 $\pm$ 0.1
16:0iso	1.3 $\pm$ 0.5	1.1 $\pm$ 0.3	1.1 $\pm$ 0.1	0.7 $\pm$ 0.1	0.9 $\pm$ 0.2	0.1 $\pm$ 0.2	0.9 $\pm$ 0.1	1 $\pm$ 0.1	0.6 $\pm$ 1	0.8 $\pm$ 0.2	0.5 $\pm$ 0.1	0.6 $\pm$ 0.4	0.7 $\pm$ 0.2	0.3 $\pm$ 0	0.4 $\pm$ 0.2
17:0iso	5.4 $\pm$ 1.8	4.1 $\pm$ 1	3.4 $\pm$ 0.5	3.1 $\pm$ 1.8	1.7 $\pm$ 2.4	1.1 $\pm$ 1	3.5 $\pm$ 2	2.8 $\pm$ 1.4	2.5 $\pm$ 1.1	3.8 $\pm$ 1	2.4 $\pm$ 0.1	3 $\pm$ 0.3	3 $\pm$ 1	1.3 $\pm$ 0.2	1.4 $\pm$ 0.3
17:0ant	3.6 $\pm$ 1.6	3.2 $\pm$ 0.7	2.4 $\pm$ 0.6	2.1 $\pm$ 0.5	3.5 $\pm$ 1	1.5 $\pm$ 0.3	2.8 $\pm$ 0.7	3.1 $\pm$ 0.4	3 $\pm$ 1	2.4 $\pm$ 0.7	1.8 $\pm$ 0.2	2.5 $\pm$ 0.3	2.4 $\pm$ 0.9	1.2 $\pm$ 0.2	1.5 $\pm$ 0.3
18:0iso	0.7 $\pm$ 0.4	0.6 $\pm$ 0.4	0.3 $\pm$ 0.4	0.3 $\pm$ 0.3	0.2 $\pm$ 0.3	0 $\pm$ 0	0.5 $\pm$ 0.5	0.9 $\pm$ 0.2	0.2 $\pm$ 0.4	0.4 $\pm$ 0.6	0.7 $\pm$ 0.1	2.1 $\pm$ 2.9	0.5 $\pm$ 0.4	0.1 $\pm$ 0.1	0 $\pm$ 0
Σ BFA	11.2 $\pm$ 4.3	8.9 $\pm$ 2.1	7.1 $\pm$ 1.3	6.3 $\pm$ 2.3	6.4 $\pm$ 2.6	2.8 $\pm$ 1	7.8 $\pm$ 2.6	7.8 $\pm$ 1.8	6.3 $\pm$ 2.2	7.5 $\pm$ 2.2	5.6 $\pm$ 0.3	8.9 $\pm$ 4	6.6 $\pm$ 2.6	3.8 $\pm$ 1	5 $\pm$ 0.9
14:0	1.5 $\pm$ 0.3	1.3 $\pm$ 0.7	1.4 $\pm$ 0.4	1.3 $\pm$ 0.2	3 $\pm$ 0.9	1.6 $\pm$ 0.1	1.3 $\pm$ 0.4	1.4 $\pm$ 0.3	0.8 $\pm$ 0.8	2 $\pm$ 0.5	2.3 $\pm$ 0.4	1.9 $\pm$ 0.7	2.4 $\pm$ 0.6	2.7 $\pm$ 0.6	2.2 $\pm$ 0.2
15:0	0.6 $\pm$ 0.1	0.6 $\pm$ 0.2	0.8 $\pm$ 0.2	0.4 $\pm$ 0	0.5 $\pm$ 0.1	0.6 $\pm$ 0.4	0.5 $\pm$ 0.1	0.4 $\pm$ 0	0.1 $\pm$ 0.3	0.5 $\pm$ 0.1	0.5 $\pm$ 0.1	0.6 $\pm$ 0	0.4 $\pm$ 0	0.7 $\pm$ 0.4	1.5 $\pm$ 0.3
16:0	11.9 $\pm$ 0.8	16.7 $\pm$ 3.1	25.2 $\pm$ 3.5	11.7 $\pm$ 1	11.6 $\pm$ 3.3	33.7 $\pm$ 1.7	10.9 $\pm$ 1	13 $\pm$ 0.9	12.3 $\pm$ 2.3	12 $\pm$ 0.4	12.6 $\pm$ 3.1	14.2 $\pm$ 2.1	11.2 $\pm$ 0.9	9.7 $\pm$ 1.4	7.6 $\pm$ 0.9
17:0	1.5 $\pm$ 0.1	1.2 $\pm$ 0.3	1 $\pm$ 0.1	1.4 $\pm$ 0.1	1.2 $\pm$ 0.1	0.7 $\pm$ 0.4	1 $\pm$ 0.5	0.8 $\pm$ 0.1	0.4 $\pm$ 0.5	0.9 $\pm$ 0.5	0.8 $\pm$ 0.1	1 $\pm$ 0.1	1.1 $\pm$ 0.2	0.7 $\pm$ 0.1	0.7 $\pm$ 0.1
18:0	6.3 $\pm$ 0.9	17.7 $\pm$ 7.3	27.5 $\pm$ 4.2	6.8 $\pm$ 0.7	9.2 $\pm$ 3.9	35.6 $\pm$ 0.9	6.6 $\pm$ 0.9	10.4 $\pm$ 1.3	14.8 $\pm$ 4.5	6.9 $\pm$ 0.4	9.1 $\pm$ 5.6	14.3 $\pm$ 3.1	6.9 $\pm$ 1.2	5.6 $\pm$ 1.1	7.5 $\pm$ 1.1
20:0	0.5 $\pm$ 0.1	0.7 $\pm$ 0.2	0.5 $\pm$ 0.3	0.5 $\pm$ 0.3	0.2 $\pm$ 0.2	0.6 $\pm$ 0.3	0.2 $\pm$ 0	0.2 $\pm$ 0.3	0 $\pm$ 0	0.2 $\pm$ 0	0.2 $\pm$ 0.1	0.1 $\pm$ 0.2	0.4 $\pm$ 0.1	0.2 $\pm$ 0.1	0.3 $\pm$ 0
22:0	0.5 $\pm$ 0.1	0.3 $\pm$ 0.2	0.1 $\pm$ 0.2	0.4 $\pm$ 0.1	0.2 $\pm$ 0	0 $\pm$ 0	0.4 $\pm$ 0.1	0.1 $\pm$ 0.1	0 $\pm$ 0	0.3 $\pm$ 0.1	0.2 $\pm$ 0.1	0 $\pm$ 0	0.3 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0
24:0	0.4 $\pm$ 0.1	0.6 $\pm$ 0.4	0.6 $\pm$ 0.4	0.2 $\pm$ 0	0.2 $\pm$ 0.1	0.7 $\pm$ 0.2	0.3 $\pm$ 0.2	0.1 $\pm$ 0.1	1.6 $\pm$ 1.8	0.4 $\pm$ 0.1	0.3 $\pm$ 0.1	1.2 $\pm$ 0.5	0.3 $\pm$ 0.1	0.1 $\pm$ 0.1	0 $\pm$ 0
Σ SFA	23.2 $\pm$ 1.5	39.2 $\pm$ 9.6	57 $\pm$ 8.2	22.9 $\pm$ 1.5	26 $\pm$ 7.1	73.5 $\pm$ 2.4	21.4 $\pm$ 1.1	26.4 $\pm$ 1.7	30.1 $\pm$ 5.9	23.3 $\pm$ 0.9	25.8 $\pm$ 8.6	33.3 $\pm$ 4.6	23.1 $\pm$ 1.7	19.9 $\pm$ 1.2	19.7 $\pm$ 1.6
14:1n-5	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0.1 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0.1 $\pm$ 0.1	0.2 $\pm$ 0.5	0.1 $\pm$ 0.1	0 $\pm$ 0	0 $\pm$ 0
16:1n-11	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0 $\pm$ 0	0.4 $\pm$ 0.1	0.7 $\pm$ 0.4	0 $\pm$ 0	0.1 $\pm$ 0.1	0.2 $\pm$ 0.1	1.1 $\pm$ 1	0 $\pm$ 0	0.1 $\pm$ 0.1	0.2 $\pm$ 0.1
16:1n-9	0.7 $\pm$ 0.2	0.5 $\pm$ 0.3	0.3 $\pm$ 0.3	0.7 $\pm$ 0.2	0.1 $\pm$ 0.2	0.1 $\pm$ 0.2	0.5 $\pm$ 0.1	0.7 $\pm$ 0.5	0.4 $\pm$ 1	0.5 $\pm$ 0	0.6 $\pm$ 0.1	0.7 $\pm$ 0.3	0.3 $\pm$ 0	0.5 $\pm$ 0.4	4.1 $\pm$ 2.9
16:1n-7	1.4 $\pm$ 0.6	0.9 $\pm$ 0.2	0.8 $\pm$ 0.3	1.7 $\pm$ 0.2	1.3 $\pm$ 0.4	0.3 $\pm$ 0.4	1.5 $\pm$ 0.3	1.2 $\pm$ 0.4	2.6 $\pm$ 1.3	2.3 $\pm$ 0.6	2.9 $\pm$ 0.8	2.1 $\pm$ 0.3	3.6 $\pm$ 0.9	5.8 $\pm$ 1.9	6.4 $\pm$ 1.9
16:1n-5	0.4 $\pm$ 0.4	0 $\pm$ 0	0 $\pm$ 0	0.3 $\pm$ 0.1	0.1 $\pm$ 0.1	0 $\pm$ 0	0.3 $\pm$ 0.1	0.8 $\pm$ 0.4	1.4 $\pm$ 1.4	0.5 $\pm$ 0.2	0.4 $\pm$ 0.2	1.8 $\pm$ 1.2	0.9 $\pm$ 0.4	3.4 $\pm$ 2.1	2.2 $\pm$ 4.3

17:1n-7	0.1±0.1	0±0	0±0	0.1±0.1	0±0	0±0	0.4±0.5	0.1±0.3	0±0	0.3±0.4	0±0.1	0.2±0.3	0±0	0±0	0±0
18:1n-11	0.5±0.2	0.4±0.1	0.1±0.3	0.5±0.1	0.4±0.1	0±0	0.5±0.2	0.9±0.2	1.6±1	0.8±0.2	0.5±0.1	1.4±1	0.7±0.2	2.1±1.4	6±1
18:1n-9	0±0.1	0±0	5.1±2.3	2.1±0.3	2.1±0.2	4±0.8	2±0.2	2.3±0.1	1.6±1	2±0.2	2±0.1	1.8±0.4	1.5±0.2	1.5±0.2	1.2±0.2
18:1n-7	2.6±0.2	1.8±0.3	1.4±0.2	2.8±0.3	2.1±0.3	1±0.1	2.5±0.2	1.7±0.3	0.7±1.1	2.3±0.3	2.3±0.5	2±0.9	3.3±0.4	4.4±1	2.6±0.8
18:1n-5	0.4±0	0.2±0.2	0.1±0.2	0.4±0	0.4±0	0.2±0.4	0.4±0	0.2±0.2	0.3±0.4	0.3±0	0.3±0.1	0.5±0.3	0.5±0.1	0.6±0.1	0.6±0.1
20:1n-11	5.8±0.5	5.3±0.7	3.2±0.7	6.1±0.8	6.4±0.9	1.9±0.5	5.6±0.6	4±0.9	2.3±1.5	5.2±0.9	4.1±1	2.3±0.6	5.9±0.6	4.3±0.8	2.5±0.3
20:1n-9	0.9±0.1	1±0.2	0.5±0.4	0.8±0	1.2±0.1	0.1±0.2	0.8±0.2	1±0.2	0.4±0.5	0.8±0.1	0.9±0.1	0.7±0.1	0.6±0.2	0.7±0.1	0.4±0.1
20:1n-7	4.7±0.5	3.9±0.6	2.4±0.8	4.6±0.2	5.2±0.9	1.3±0.2	4.6±0.4	3.5±0.8	5.9±4.3	4.2±0.3	4.2±1.1	4±1	5.5±0.9	5.5±0.7	5±0.8
Σ MUFA	17.8±1.1	14±2.1	13.9±2.5	20.4±0.6	19.4±1.6	8.8±0.7	19.9±1	17.2±1.5	17.2±1.5	19.9±0.9	18.8±2.8	18.9±2.8	22.9±1.1	28.7±1.4	31.3±0.5
16:2n-7	0±0	0±0	0±0	0.7±1.5	2.6±2.5	0.7±1	0.8±1.7	0±0	0±0	0±0	0±0	0±0	0±0	0±0	0±0
16:2n-4	0.4±0.5	0.1±0.1	0±0	0.2±0	0.5±0.4	0.2±0.5	0.7±0.3	1±0.9	0.2±0.4	0.4±0.2	0.7±0.3	0.1±0.3	0.5±0.2	0.7±0.4	0.6±0.2
16:3n-4	0.1±0.1	0±0	0±0	0.1±0.1	0±0.1	0±0	0±0.1	0.1±0.1	0±0	0.1±0.2	0.3±0.2	0.3±0.3	0.2±0.1	0.3±0.1	0.1±0.1
16:3n-6	0.4±0.3	0.6±0.2	0.2±0.3	0.5±0.1	0.1±0.2	0±0	0.1±0.2	0.1±0.3	0±0	0.5±0.2	0.2±0.1	0±0	0.5±0.2	0±0	0±0
16:3n-3	0.1±0.1	0±0	0±0	0.1±0.1	0±0.1	0±0	0.1±0.1	0.2±0.2	0.5±0.7	0.3±0.1	0.2±0.1	1.8±2.7	1.6±0.9	0.6±0.2	0.8±0.5
16:4n-3	0.3±0.2	0.2±0.2	0±0	0.3±0.1	0.6±0.5	0.1±0.2	0±0	0±0	0±0	0±0	0±0	0±0	0.2±0.1	0.4±0.1	0.6±0.1
18:2n-6	1.3±0.2	1.4±0.3	1.6±0.3	1.2±0.2	0.9±0.2	0.7±0.4	1.1±0.1	1.5±0.2	0.9±0.6	1.3±0.1	1.4±0.2	1.6±0.5	0.9±0.1	3±1.3	6.3±1.4
18:2n-4	0.4±0.1	0.2±0.2	0±0	0.5±0.1	0.3±0.1	0±0	0.5±0	0.4±0	0±0	0.4±0.1	0.3±0.1	0±0	0.8±0.2	0.7±0.2	0.4±0.3
18:3n-6	0.1±0	0±0	0±0	0.1±0	0.1±0.2	0±0	0.2±0	0±0.1	0±0	0.1±0.1	0.1±0.1	0±0	0.3±0.2	0.2±0.1	0.2±0.3
18:3n-4	1±0.5	2.7±1.2	3.1±1.7	0.4±0.3	0.4±0.5	0±0	1.1±0.3	3.5±1.4	1.3±1.3	1±0.4	0.6±0.3	2.4±0.8	0.3±0.1	0.3±0	0±0
18:3n-3	0.9±0.2	0.2±0.2	0±0	0.9±0.2	1±0.8	1.7±1	0.9±0.1	1.2±0.3	2.1±3.3	1.1±0.2	1.6±0.4	0.7±0.4	0.5±0.1	0.9±0.2	0.5±0.2
18:4n-3	0.7±0.6	0±0	0.6±0.6	1.8±0.4	1.4±0.5	2.3±0.2	1.7±0.3	3±0.6	2.9±0.7	3.1±1	4.4±0.8	3.2±0.8	2.7±0.5	2±0.3	1.5±0.2
18:4n-1	0.1±0	0±0	0±0	0.1±0.1	0.1±0.1	0±0	0±0.1	0±0	0.4±1	0±0	0±0	0±0	0.1±0.1	0.1±0.1	0±0
20:2n-6	1.2±0.1	2.2±0.5	1.1±0.2	1.2±0.2	2.1±0.6	0.9±0.2	1.1±0.2	1.1±0.2	0.4±0.5	1±0.2	1±0.1	0.5±0.4	0.9±0.1	0.6±0.1	0.4±0
20:3n-6	0.1±0	0±0	0±0	0.1±0.1	0±0.1	0±0	0.2±0.1	0±0	0.3±0.6	0.1±0.1	0±0.1	0.1±0.3	0.2±0.1	0.1±0.1	0.1±0.1
20:3n-3	0.4±0.1	0.1±0.1	0±0	0.4±0.1	0.2±0.1	0±0	0.4±0.1	0.3±0.3	0±0	0.4±0.1	0.4±0.1	0±0	0.3±0.1	0.2±0	0.1±0.1
20:4n-6	2.7±0.3	3.5±0.5	2.3±0.6	2.7±0.4	3.5±0.3	1.2±0.1	2.6±0.6	2.6±0.5	0.8±1.2	2.5±0.4	1.7±0.4	1.4±0.4	1.7±0.2	1.2±0.1	0.7±0.1
20:4n-3	0.6±0.1	0.3±0.2	0.3±0.3	0.6±0.2	0.4±0.1	0±0	0.5±0.1	0.6±0.1	0.5±0.8	0.6±0.1	0.8±0.2	0.3±0.3	0.4±0.1	0.5±0.1	0.4±0.1
20:5n-3	10.5±2.6	5.7±1.2	3.3±0.7	12.1±1.3	8.3±1.3	2.5±0.5	11.1±1.8	6.9±0.9	3.4±1.7	11.4±1.8	11.1±2.6	5.8±1.9	14.6±2.4	16.3±3.4	9.4±2.1
21:5n-3	0.4±0	0.3±0.2	0.1±0.3	0.5±0.1	0.6±0.1	0±0	1.5±0.6	0.7±0.2	0±0	0.5±0.1	0.8±0.2	0.1±0.2	0.9±0.2	0.8±0.1	0.5±0.1

22:2n-6	0.2 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0.4 ± 0.1	0.2 ± 0.3	0 ± 0	0 ± 0	0.1 ± 0.1	0.1 ± 0.1	0 ± 0	0.3 ± 0.1	0.1 ± 0	0.4 ± 0.2
22:4n-6	0.6 ± 0.2	0.4 ± 0.1	0.1 ± 0.2	0.5 ± 0.3	0.4 ± 0.1	0 ± 0	0.6 ± 0.1	0.2 ± 0.2	0 ± 0	0.6 ± 0.2	0.2 ± 0.1	0.4 ± 0.4	0.3 ± 0.1	0.2 ± 0	0 ± 0.1
22:5n-6	0.4 ± 0.1	0.3 ± 0.2	0.1 ± 0.2	0.5 ± 0.1	0.4 ± 0.1	0 ± 0	0.4 ± 0.1	0.4 ± 0.4	0.5 ± 1	0.4 ± 0.1	0.4 ± 0.1	0.3 ± 0.3	0.2 ± 0.1	0.2 ± 0	0 ± 0.1
22:5n-3	2 ± 0.5	1.2 ± 0.3	0.5 ± 0.5	2.4 ± 0.3	1.2 ± 0.2	0.4 ± 0.9	2.2 ± 1.3	1.4 ± 0.4	2.1 ± 2.2	2.2 ± 0.5	2.8 ± 2.2	2.8 ± 1.8	1.6 ± 0.3	1.4 ± 0.1	0.9 ± 0.1
22:6n-3	9.7 ± 2.1	4.9 ± 0.9	2.6 ± 0.5	11.1 ± 1.4	6 ± 0.6	1.9 ± 0.5	10.3 ± 1.6	7 ± 0.7	4.5 ± 2.4	9.4 ± 2	7.8 ± 1.5	5.4 ± 1.5	6.1 ± 1.4	4.3 ± 0.7	2.3 ± 0.3
Σ PUFA	34.7 ± 5.1	24.5 ± 4.5	15.7 ± 2.5	38.8 ± 4.6	31.1 ± 2.5	13 ± 1.8	38.4 ± 3.5	32.4 ± 2.4	21.1 ± 10.3	37.9 ± 3.4	37.3 ± 3.7	29 ± 4.2	36.5 ± 2.9	38 ± 2.3	33.8 ± 2.6
20:2i	1.1 ± 0.3	1.1 ± 0.2	0.7 ± 0.5	0.9 ± 0.2	1.6 ± 0.6	0 ± 0	0.9 ± 0.2	0.8 ± 0.3	1.4 ± 2.8	0.9 ± 0.2	1 ± 0.3	0.1 ± 0.3	1.1 ± 0.1	1.3 ± 0.5	0.8 ± 0.2
20:2j	0.5 ± 0.1	0.4 ± 0.1	0.2 ± 0.2	0.4 ± 0	0.6 ± 0.1	0 ± 0	0.4 ± 0.1	0.3 ± 0.3	0.2 ± 0.5	0.3 ± 0.1	0.4 ± 0.1	0 ± 0	0.4 ± 0.1	0.4 ± 0.1	0.3 ± 0.1
22:2i	1.9 ± 0.3	2.6 ± 0.5	1.3 ± 0.7	1.6 ± 0.5	3.8 ± 1.5	0.1 ± 0.2	1.7 ± 0.4	1.9 ± 0.6	0.3 ± 0.6	1.5 ± 0.3	1.7 ± 0.8	0.8 ± 0.3	1 ± 0.1	0.7 ± 0.2	0.3 ± 0.2
22:2j	5.4 ± 0.8	6 ± 0.6	3.2 ± 2	5.1 ± 1.1	8.7 ± 3.2	1.4 ± 0.3	4.2 ± 0.7	4.4 ± 1.4	1.8 ± 1.7	4.6 ± 0.8	4.5 ± 1.9	2 ± 0.7	5.3 ± 1	3.6 ± 0.6	1.7 ± 0.9
22:3i	0.5 ± 0.2	1.1 ± 0.3	0.1 ± 0.2	0.5 ± 0.2	1.4 ± 0.6	0 ± 0	0.5 ± 0.2	0 ± 0	0 ± 0	0.5 ± 0.1	0.6 ± 0.3	0 ± 0	0 ± 0	0 ± 0	0 ± 0
Σ NMI FA	9.4 ± 1.4	11.3 ± 1.2	5.4 ± 3.3	8.4 ± 1.8	16 ± 5.6	1.5 ± 0.3	7.8 ± 1.3	7.3 ± 2.4	3.7 ± 3.4	7.8 ± 1.4	8.2 ± 3.1	3 ± 1.1	7.9 ± 1.2	6 ± 0.5	3 ± 1
16:0DMA	0.1 ± 0	0.3 ± 0.2	0 ± 0	0 ± 0	0.1 ± 0.1	0 ± 0	0.2 ± 0.2	0.4 ± 0.2	0.4 ± 0.5	0.5 ± 0.2	0.5 ± 0.2	1.8 ± 2.1	0.5 ± 0.4	3.2 ± 2.3	7.6 ± 1.9
18:0DMA	1 ± 0.5	0 ± 0	0 ± 0	1.2 ± 0.4	0.1 ± 0.1	0 ± 0	1.7 ± 1	4 ± 1.9	13.5 ± 5.8	0.8 ± 0.4	0.8 ± 0.6	2.7 ± 2	0.5 ± 0.1	1.5 ± 0.6	4.9 ± 1.1
20:1n-7DMA	0.6 ± 0.3	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	1.2 ± 0.5	1.8 ± 1	7.1 ± 3	0.2 ± 0.1	0.2 ± 0.2	1.5 ± 0.8	0 ± 0	0.1 ± 0.1	1.2 ± 0.2
Σ DMA FA	1.7 ± 0.7	0.3 ± 0.2	0 ± 0	1.3 ± 0.5	0.2 ± 0.1	0 ± 0	3.1 ± 1.4	6.2 ± 3	20.9 ± 8.9	1.5 ± 0.7	1.4 ± 1	6 ± 1.8	1.1 ± 0.5	4.9 ± 2.9	13.7 ± 3
A	0.3 ± 0.1	0.2 ± 0.2	0.1 ± 0.2	0.2 ± 0	0.4 ± 0.1	0 ± 0	0.2 ± 0	0.3 ± 0.4	0 ± 0	0.2 ± 0.1	0.3 ± 0.1	0.1 ± 0.2	0.1 ± 0.1	0 ± 0	0 ± 0
B	0.2 ± 0	0 ± 0	0 ± 0	0.2 ± 0	0.1 ± 0.1	0 ± 0	0.1 ± 0.1	0 ± 0	0.2 ± 0.3	0 ± 0.1	0.1 ± 0.1	0.2 ± 0.4	0.1 ± 0	0 ± 0	0 ± 0
W	0.3 ± 0.1	0.6 ± 0.1	0.1 ± 0.3	0.3 ± 0.1	0.7 ± 0.2	0 ± 0	0.2 ± 0.2	0.4 ± 0.2	0 ± 0	0.3 ± 0.1	0.3 ± 0.1	0.1 ± 0.2	0.2 ± 0	0.2 ± 0	0.1 ± 0.1
Σ UK FA	0.8 ± 0.1	0.8 ± 0.2	0.2 ± 0.4	0.6 ± 0.1	1.1 ± 0.3	0 ± 0	0.5 ± 0.2	0.7 ± 0.6	0.2 ± 0.3	0.5 ± 0.1	0.6 ± 0.1	0.4 ± 0.4	0.3 ± 0.1	0.2 ± 0.1	0.1 ± 0.1
PUFA/SFA	1.5 ± 0.3	0.7 ± 0.2	0.3 ± 0.1	1.7 ± 0.3	1.3 ± 0.3	0.2 ± 0	1.8 ± 0.2	1.2 ± 0.1	0.8 ± 0.5	1.6 ± 0.2	1.6 ± 0.6	0.9 ± 0.2	1.6 ± 0.2	1.9 ± 0.2	1.7 ± 0.2
EPA/DHA	1.1 ± 0.1	1.2 ± 0.2	1.3 ± 0.1	1.1 ± 0.1	1.4 ± 0.2	1.3 ± 0.1	1.1 ± 0.1	1 ± 0.2	0.8 ± 0.3	1.2 ± 0.2	1.4 ± 0.1	1.1 ± 0.2	2.5 ± 0.5	3.8 ± 0.3	4 ± 0.6
Σ n-3/Σ n-6	3.7 ± 1	1.5 ± 0.1	1.4 ± 0.4	4.5 ± 0.6	2.6 ± 0.5	2.8 ± 0.7	4.5 ± 0.5	3.6 ± 0.3	Inf ± NA	4.6 ± 0.9	5.9 ± 0.4	5.5 ± 2.8	5.6 ± 1.1	5.3 ± 1.9	2.2 ± 0.5

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917 **Table S5: Stable isotope (mean $\pm$ SD, n = 5) composition of the three ontogenic stages of *Crepidula fornicata* (motile males, sessile males, sessile females)**
918 **over the sampling survey.**

	S1 = 26 th February			S2 = 21 th March			S3 = 28 th March			S4 = 12 th April			S5 = 14 th June		
	Sessile female	Sessile male	Motile male	Sessile female	Sessile male	Motile male	Sessile female	Sessile male	Motile male	Sessile female	Sessile male	Motile male	Sessile female	Sessile male	Motile male
$\delta^{13}\text{C}$	-19.5 $\pm$ 0.3	-19.4 $\pm$ 0.4	-19.4 $\pm$ 0.5	-19.8 $\pm$ 0.3	-19.7 $\pm$ 0.3	-20.1 $\pm$ 0.5	-19.2 $\pm$ 0.6	-19.6 $\pm$ 0.4	-19.9 $\pm$ 0.3	-20.1 $\pm$ 0.4	-20.3 $\pm$ 0.8	-20.6 $\pm$ 0.5	-18.1 $\pm$ 0.3	-18.5 $\pm$ 0.1	-18.5 $\pm$ 0.3
$\delta^{15}\text{N}$	9.3 $\pm$ 0.4	8.9 $\pm$ 0.8	8.9 $\pm$ 0.3	9.6 $\pm$ 0.5	8.9 $\pm$ 0.5	8.4 $\pm$ 1.1	9.7 $\pm$ 0.3	9.4 $\pm$ 0.6	8.5 $\pm$ 0.4	8.4 $\pm$ 0.7	8.4 $\pm$ 0.6	8.3 $\pm$ 0.5	8.6 $\pm$ 0.6	8.3 $\pm$ 0.5	7.5 $\pm$ 0.3
C/N	4 $\pm$ 0.3	3.8 $\pm$ 0.8	4 $\pm$ 0.2	3.2 $\pm$ 0.8	3.8 $\pm$ 0.6	3.5 $\pm$ 0.7	3.6 $\pm$ 0.4	3.3 $\pm$ 0.4	3.8 $\pm$ 0.2	3.6 $\pm$ 0.4	4.3 $\pm$ 0.2	3.8 $\pm$ 0.2	3.5 $\pm$ 0.6	4.1 $\pm$ 1.2	3.4 $\pm$ 0.3

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Figure S1: Photosynthetic available radiations (PAR) at the surface and at the bottom of our study site.

Figure S2: Mean daily temperature ($\pm$ SD) at the bottom in our study site