

# Assimilation of shrimp farm sediment by *Holothuria scabra*: a coupled fatty acid and stable isotope approach

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**Abstract** – Deposit-feeding sea cucumbers are efficient nutrient recyclers and have the potential to contribute to the limitation of organic matter load in polyculture or integrated aquaculture systems. Assessing how they assimilate organic matter originating from other farmed species is therefore important for the development of such multi-species farming systems. Here, a coupled stable isotope – fatty acid approach was used to characterize the assimilation of organic matter from shrimp (*Penaeus stylirostris*) farming by *Holothuria scabra* in an experimental culture system. *H. scabra* were reared in mesocosms on shrimp farming-originating sediment with and without additional food sources (maize and fish meals). Although fatty acid results did indicate that shrimp-farming sediment was assimilated by holothurids, we found no evidence of maize waste and fish meal contribution to *H. scabra* organic carbon (no effect on  $\delta^{13}\text{C}$ , no accumulation of meal-specific fatty acids). However, a strong effect of fish meal on *H. scabra*  $\delta^{15}\text{N}$  was observed, suggesting that this additional food source could represent an alternative source of nitrogen for holothurids. Finally, this study supports the culture of *H. scabra* as a perspective to reduce sedimentary organic matter excess associated with shrimp farms, and suggest that the addition of selected food sources might contribute to increasing the content in some nitrogen organic compounds in holothurid tissues.

**Keywords:** Fatty acids / stable isotopes / rotational co-culture / shrimp-farming / Holothurid

## 1 Introduction

The combined farming of different species, where each one feeds on wastes from the other one (commonly referred to as polyculture, co-culture, or integrated culture Lutz, 2003; Zamora et al., 2016) aims at increasing economic yields of aquaculture surfaces, reducing ultimate effluents from production units and generating a mass-balanced system mitigating impacts on the adjacent environment (Soto, 2009). Sea cucumbers are considered as ideal models for co-culture systems in association with various taxa (molluscs, crustaceans, fish, seaweed) in both temperate and tropical environments (Zamora et al., 2016). Their high economic value makes them an attractive aquaculture product, with an annual

production evaluated at 166 712t in 2012-2014 (Zhang et al., 2015; Zamora et al., 2016).

Most holothurids in the order Aspidochirota (e.g. *Holothuria scabra*, *Apostichopus japonicus*) are surface or sub-surface deposit-feeders, and can efficiently extract organic matter from coastal sediments (Roberts et al., 2003), which can contribute to mitigate aquaculture environmental impacts through the recycling of organic matter (Zamora et al., 2016). In Réunion Island (SW Indian Ocean), a mixed population of *Holothuria atra* and *Holothuria leucospilota* has been observed to ingest up to  $82 \text{ kg m}^{-2} \text{ y}^{-1}$  of sediment, with an organic matter assimilation efficiency of 10% (Mangion et al., 2004). Individuals of *H. scabra* averaging 1 kg were reported to displace more than  $250 \text{ cm}^3$  of sediments per diurnal cycle (Purcell, 2004). Besides, their burrowing activity limits sediment hypoxia, thereby enhancing organic matter bacterial remineralization (MacTavish et al., 2012). Both *H. scabra* and

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*A. japonicas* have been considered as good candidate to be used in association with other species (i.e. crustaceans, molluscs, fish, seaweeds), in order to limit detrimental effects of organic matter accumulation on the sediment (e.g. Slater and Carton 2007; Watanabe et al., 2012; Neofitou et al., 2019; Zamora et al., 2016). In particular, their ability to selectively extract specific components of sedimentary organic matter, such as bacteria, nitrogen and transparent exopolymer particles (Moriarty 1982; Roberts et al., 2003; Robinson et al., 2016; Sadeghi-Nassaj et al., 2018), may suggest that their use in co-culture systems is relevant. Indeed, their use can contribute to remediate or manage the risks associated with pond sediment degradation in aquaculture ponds, where high densities and/or dystrophy conditions typically result in animal health issues (Lemonnier et al., 2006; Lemonnier, 2007). However, the selective assimilation of specific components of sediment is still poorly documented (but see Moriarty, 1982), especially in the context of high sedimentary organic enrichment (Gao et al., 2011; Slater et al., 2011; Yu et al., 2015; Zamora and Jeffs, 2015), such as for animals reared on aquaculture sediment. In particular, although addition of extra food is generally not required, it has recently been suggested that growth performances of holothurids could be improved through the addition of additional material to the system (Gao et al., 2011; Robinson et al., 2018, 2019).

In this context, trophic biomarkers, such as stable isotopes (SI)  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$ , and fatty acids (FA) are powerful approaches to characterize the diet of marine organisms and assess the pathways of organic matter within food webs. Different food sources usually display different SI compositions, and the SI composition of a consumer reflects its food source, plus a small heavy isotope enrichment, usually  $\approx 1\%$  for carbon and  $\approx 3.4\%$  for nitrogen (Vander Zanden and Rasmussen, 2001). SI have been successfully used to investigate the trophic ecology of sea cucumbers (Slater and Carton, 2010) and their trophic relationships in a polyculture system (Feng et al., 2014). In spite of their usefulness in discriminating food sources contributing to the diet of consumers in low food source diversity systems, SI provide a limited resolution for heterogeneous organic matter pools, such as sediment or suspended particulate organic matter. Due to the higher taxa specificity of FA, their analysis is complementary to SI to investigate the composition of such heterogeneous organic matter pools (Kharamenko et al., 2001). The existence of bacteria, diatom, flagellate and higher plant specific FA makes this approach extremely powerful to disentangle trophic relationships for sediment-associated organisms that might be overlooked otherwise when using bulk SI alone (Kelly and Scheibling, 2012).

The Pacific blue shrimp (*Penaeus stylirostris* Stimpson, 1974) is the second export product of New Caledonia. The island hosts 19 shrimp farms, for an annual production of ca. 2000 tons $\cdot\text{year}^{-1}$  (Lemonnier, 2007). Shrimp farming occurs on ponds varying between 4 and 11 ha, where shrimp densities range from 18 to 35 individuals  $\text{m}^{-2}$  (Della Patrona and Brun, 2008). The accumulation of uneaten food pellets on sediments results in shrimp ponds eutrophication, ultimately favoring the development of pathogens (Lemonnier et al., 2010). The co-culture of shrimps and sea-cucumbers (Purcell et al., 2006; Xu and Zhu, 2002), or their polyculture with jellyfish (Li et al., 2014b, 2014a) has therefore been considered as an alternative

to limited the organic matter excess from shrimp monocultures. Moreover, studies revealed that shrimp wastes could be used to feed sea cucumbers (Chen et al., 2015a, 2015b). These co-culture systems could (1) limit sediment organic enrichment (Slater and Carton, 2007) and (2) develop a new high value exportation pathway. Because simultaneous culture of these two species has been proven deleterious to juvenile sea cucumbers (Bell et al., 2007; Pitt et al., 2004), a rotational culture system, where sea cucumbers are reared in ponds after extraction of shrimps, seems to be the most relevant system.

In this context, the first objective of the present study was to evaluate the assimilation of shrimp farm sediments by *H. scabra*, simulating a context of a *P. stylirostris* – *H. scabra* rotational co-culture system. In a second time, the study tried to assess the relevance of using additional food sources, by characterizing the assimilation of two potential sources (i.e. maize meal and fish meals) in the tissues of the holothurids. Maize meal is a potential low-cost and easily available food supplement which can stimulate the benthic food web through heterotrophic assimilation. Fish meal is more expensive, but also of higher nutritional value. The assimilation of these different food sources by juvenile sea cucumbers was assessed through a combined SI ( $\delta^{13}\text{C}$ ,  $\delta^{15}\text{N}$ ) and FA approach.

## 2 Material and methods

### 2.1 Experimental setup

Experiments were conducted at the experimental station of Saint-Vincent Bay (21°58'S, 165°57'E; New Caledonia) for 7 months, between April and December 2014. Ten mesocosms of 1600 L (1.72  $\text{m}^2$ ) were each filled with 25 cm of sediment from upper 5 cm of a semi-intensive shrimp farm, collected at the end of a shrimp growing period after 3 weeks of dry out, and presenting a content in organic matter of 2.3%, and ammonium concentrations higher than 1000  $\mu\text{mol}\cdot\text{L}^{-1}$ . Mesocosms were filled with seawater on open circuit, with an average daily water renewal of 30%. Juveniles *H. scabra* (initial weight =  $15.2 \pm 0.2$  g) were then placed in 9 tanks at a density of 7 individuals $\cdot\text{m}^{-2}$  (12 individuals per tank). One tank was left without animals as a control for the natural evolution of shrimp farm sediment. The 9 *H. scabra* tanks were then randomly assigned to a treatment (3 replicates / treatment): (1) animals reared on shrimp farm sediment only (unenriched treatment), (2) animals reared on shrimp farm sediment + maize meal (added twice a week, eq. 1.5% of initial biomass $\cdot\text{d}^{-1}$ ) and (3) animals reared on shrimp farm sediment + fish meal (added twice a week, eq. 1.5% of initial biomass $\cdot\text{d}^{-1}$ ). Fish meal was gently spread in the mesocosms to ensure a uniform repartition. Maize meal was hydrated with mesocosm water to make sure it will not float, and then gently spread as for fish meal. Fecal pellets and uneaten food were not removed during the experiment.

### 2.2 Sampling

Because of the limited number of animals available for each treatment, and in order to limit density modifications inside each tank, which could lead to unpredictable density-dependent processes, a limited number of individuals were sampled at each sampling time. Five individuals were

randomly sampled at the beginning of the experiment ( $T_0$ ), before the distribution of animals in the different mesocosms. During experiment, one individuals per tank (3 for each treatment) was sampled after 1 and 3 months ( $T_2$  and  $T_3$ ), and 3 individuals per tank (9 for each treatment) were sampled after 7 months ( $T_f$ ), at the end of the experiment. At each sampling time, sampled individuals were weighed and then dissected in order to collect muscle strips for SI and FA analyses. The sampling of the muscle allowed analysing the same tissues during the all experiment, while gonads were not developed at the beginning of the experiment. Two sub-samples were taken for FA and SI analyses, respectively, and then stored at  $-80^\circ\text{C}$  before further analyses. Every month, the medium weight of animals was assessed for each tank, and mortality was assessed at the end of the experiment through total fishing of the tank.

Surface sediment (upper 5 cm) was also sampled (three 2.6 cm diameter cores pooled for each tank) at  $T_0$  and after 1, 3 and 7 months (end of experiment). Biofilm covering mesocosm walls was also sampled at the end of the experiment by scraping. Fish meal and maize meals distributed to the additional food source treatments throughout the experiment did originate from the same initial stocks, and were therefore sampled at the beginning of the experiment for SI and FA analyses. All these samples were stored at  $-80^\circ\text{C}$  before further analyses.

## 2.3 Laboratory analyses

### 2.3.1 Stable isotope analyses

Muscle samples, biofilm, maize and fish meals were freeze-dried and ground with a mortar and a pestle into a fine and homogeneous powder. About 0.5 g of muscle, 1 mg of sediment, biofilm, and additional food sources were placed in tin capsules. Sediment samples were split into 2 sub-samples for carbon and nitrogen analyses. As carbonates present higher  $\delta^{13}\text{C}$  than organic carbon, one was treated for  $\delta^{13}\text{C}$  analysis, after acidification by 1% HCl solution to remove carbonates, rinsed with distilled water and oven-dried at  $60^\circ\text{C}$  for 24 h (DeNiro and Epstein, 1978). The other subsample for nitrogen isotope analysis was not acidified because acidification results in enrichment in  $\delta^{15}\text{N}$  (Pinnegar and Polunin, 1999). The  $^{13}\text{C}$ : $^{12}\text{C}$  and  $^{15}\text{N}$ : $^{14}\text{N}$  ratios were measured by continuous-flow isotope-ratio mass spectrometry. The spectrometer (Delta V Advantage stable isotope analyzer, Thermo Scientific, Bremen, Germany, with Flash EA-1112 elemental analyzer, Thermo Scientific, Milan, Italy) was operated in dual isotope mode. The analytical precision, estimated from standards analyzed along with the samples, was  $<0.1\%$  for  $\delta^{13}\text{C}$  and  $<0.15\%$  for  $\delta^{15}\text{N}$ . Reference gas and the internal standard used (acetanilide, Thermo Scientific) were calibrated against reference materials (USGS-24, IAEA-CH6, IAEA-600 for carbon; IAEA-N1, N2, -N3, -600 for nitrogen). Data are expressed in the  $\delta$  unit, in permil:  $\delta X = [(R_{\text{sample}}/R_{\text{standard}}) - 1] \cdot 10^3$ . Where  $X = ^{13}\text{C}$  or  $^{15}\text{N}$  and  $R = ^{13}\text{C}/^{12}\text{C}$  or  $^{15}\text{N}/^{14}\text{N}$ , respectively.

### 2.3.2 Fatty acid analyses

A known mass of muscle strips (between 20 and 100 mg of wet weight, according to individual mass) was ground with a Dounce tissue grinder in 6 mL of chloroform:methanol

(2:1, v:v ; HPLC grade, Sigma-Aldrich). Samples were then sonicated 10 min at  $4^\circ\text{C}$  in order to ensure complete lipid extraction. Lipids from sediment, biofilm and additional food sources were also extracted following this method. Lipid extracts were then stored at  $-20^\circ\text{C}$  under nitrogen atmosphere after addition 0.01% by weight of butylated hydroxytoluene (Sigma-Aldrich) as antioxidant, before analysis. After evaporation to dryness under nitrogen and recovering with three washings of 0.5 mL chloroform:methanol (98:2, v:v), an aliquot (1/5) of muscle lipid extract was deposited at the top of a silica gel (silica gel 60, 70–230  $\mu\text{m}$  mesh, Sigma-Aldrich previously heated at  $450^\circ\text{C}$  and deactivated with 6% water by weight) micro-column (40 mm  $\times$  5 mm i.d.). Neutral lipids (NL) were eluted with 10 mL of chloroform:methanol (98:2, v:v) (Marty et al., 1992). NL are FA generally stored in consumer tissues in the same proportions than in their food sources, therefore they mirror the composition of the diet (here sediment, meals, and/or biofilm). In contrast, dietary lipids are assimilated without lipid classes distinction (Arts et al., 2001; Dalsgaard et al., 2003). For this reason, NL were separated from the total lipid pool only for sea cucumber samples, while total FA were analyzed for food sources (sediment, biofilm, meals). Lipids were then transesterified by adding of 800  $\mu\text{L}$  of  $\text{H}_2\text{SO}_4$  (3.4% v:v in methanol, Sigma Aldrich) as a catalyzer, 0.01% and tricosanoic acid (Sigma Aldrich) as an internal standard and heating 10 min at  $100^\circ\text{C}$ . After cooling at room temperature and adding 800  $\mu\text{L}$  of hexane (HPLC grade, Sigma Aldrich), the organic phase-containing FA methyl esters (FAME) was washed three times with 1.5 mL of water-saturated hexane. FA methyl esters were then recovered and analyzed in a Varian CP 8400 gas chromatograph (GC) equipped with a splitless injector and a flame-ionization detector and using hydrogen as mobile phase. FA methyl esters were separated using a polar (ZB-WAX – 30 m  $\times$  0.25 mm i.d.; 0.25  $\mu\text{m}$  thickness, Phenomenex) and a non-polar (ZB-5HT – 30 m  $\times$  0.25 mm i.d.; 0.25  $\mu\text{m}$  thickness, Phenomenex) capillary columns in parallel. Chromatograms were analyzed using the Galaxie software. The internal standard allowed FA quantification. FA were identified by comparing retention times with those of a commercial standard mixture containing 37 FAME (Sigma Aldrich) and other known standard mixtures (Le Grand et al., 2013). FA were designated following the general nomenclature (X:Yn-Z) where X is the number of carbones, Y, the number of double bounds and Z the position of the first double bound counting from the terminal methyl carbon toward the carbonyl carbon. Results were expressed as the proportion (mass percentage) of each FA on total FA of the total lipids (sediment, biofilm and meals) or of the neutral lipids (*H. scabra*).

## 2.4 Data analyses

Due to limited number of replicates for each time / treatment, the effects of treatment at each time on the SI composition, and for each individual NL FA of sea cucumbers were assessed through non parametric Kruskal-Wallis (KW) tests, followed by Conover-Iman multiple comparison tests with Bonferroni's adjustment method.

Only FA representing more than 1% of total FA were taken into account for data analysis. The global FA composition of sea cucumbers and sources was represented



through non-metric multidimensional scalings (nMDS) based on Euclidean distances. The effects of time and treatments on the FA composition of sea cucumbers and sediments were assessed through permutational variance analyses (PERMANOVA). Because the number of replicates of potential food source (sediment, meals and biofilm) samples were lower or equal to 3, it was not possible to compare the effect of treatment at each times by statistical analyses. All statistical analyses and graphics were performed with the free software R (Core Team, 2017), with R Version 3.4.1 (2017 06 30).

## 3 Results

### 3.1 Survival and growth of animals

Survival rates of sea-cucumbers at the end of the experiment ranged between  $86.7 \pm 15.3\%$  for fish meal treatment, to  $90 \pm 17.3\%$  and  $90 \pm 10\%$ , for unenriched and maize meal treatments, respectively. They did not differ significantly among treatments (data not shown). Likewise, individual final weights did not differ among treatments, ranging from  $37.3 \pm 9.1$  g for individuals coming from unenriched treatment, to  $40.6 \pm 13.8$  g for individuals from fish meal treatment, and to  $43.3 \pm 16.3$  g for individuals from maize meal.

### 3.2 Stable isotopes

The SI composition of sediment did not vary significantly among treatments during the experiment (KW tests, for  $\delta^{13}\text{C}$   $H_{25} = 2.39$ ,  $p$ -value = 0.30; and for  $\delta^{15}\text{N}$   $H_{25} = 6.38$ ,  $p$ -value = 0.05). None of the 3 treatments (unenriched, maize meal, fish meal) induced significant variation of sediment  $\delta^{13}\text{C}$  over time, values ranging between  $-19.1\%$  and  $-18.3\%$ , (KW tests; at  $T_0$   $H_{25} = 1.69$   $p$ -value = 0.43; at  $T_3$   $H_{25} = 2.85$ ,  $p$ -value = 0.24; at  $T_f$   $H_{25} = 0.35$ ,  $p$ -value = 0.84), or  $\delta^{15}\text{N}$  over time (KW tests; at  $T_0$   $H_{25} = 3.01$   $p$ -value = 0.22; at  $T_3$   $H_{25} = 3.52$ ,  $p$ -value = 0.17; at  $T_f$   $H_{25} = 5.6$ ,  $p$ -value = 0.06), (Fig. 1). Biofilm ( $n=3$  for each treatment) covering mesocosms walls showed no significant differences of  $\delta^{13}\text{C}$  (KW tests,  $H_{25} = 4.86$ ,  $p$ -value = 0.09) between unenriched, maize and fish meal treatment at  $T_f$ . The post hoc test showed significant difference between  $\delta^{15}\text{N}$  values from fish and unenriched treatments ( $p$ -value < 0.05). Among these 3 treatments, biofilm from unenriched mesocosms displayed the most  $^{13}\text{C}$ -enriched mean values ( $-3.4\%$ ), while fish meal addition resulted in the highest mean  $\delta^{15}\text{N}$  values (8.5‰). The SI composition of maize ( $n=2$ ) and fish ( $n=2$ ) meals differed of 4.3‰ for the  $\delta^{13}\text{C}$  and of 7‰ for  $\delta^{15}\text{N}$  ( $\delta^{13}\text{C} = -12.5\%$  and  $-16.8\%$  and  $\delta^{15}\text{N} = 4.6\%$  and 11.6‰ for maize and fish meals, respectively).

The  $\delta^{15}\text{N}$  of *H. scabra* increased in all treatments during the experiment (Fig. 1). The  $\delta^{15}\text{N}$  was significantly different among treatments only after 7 months (KW test,  $H_{25} = 18.56$ ,  $p$ -value < 0.001). The highest values were found for fish meal treatment (from  $6.0 \pm 0.6\%$  at the beginning to  $11.5 \pm 0.4\%$  at the end of the experiment), significantly different from both other treatments (Conover-Iman multiple comparison tests with Bonferroni's adjustment, both  $p$ -values < 0.05), while muscles of animals from unenriched and maize meal treatments displayed a lower and similar  $^{15}\text{N}$  enrichment

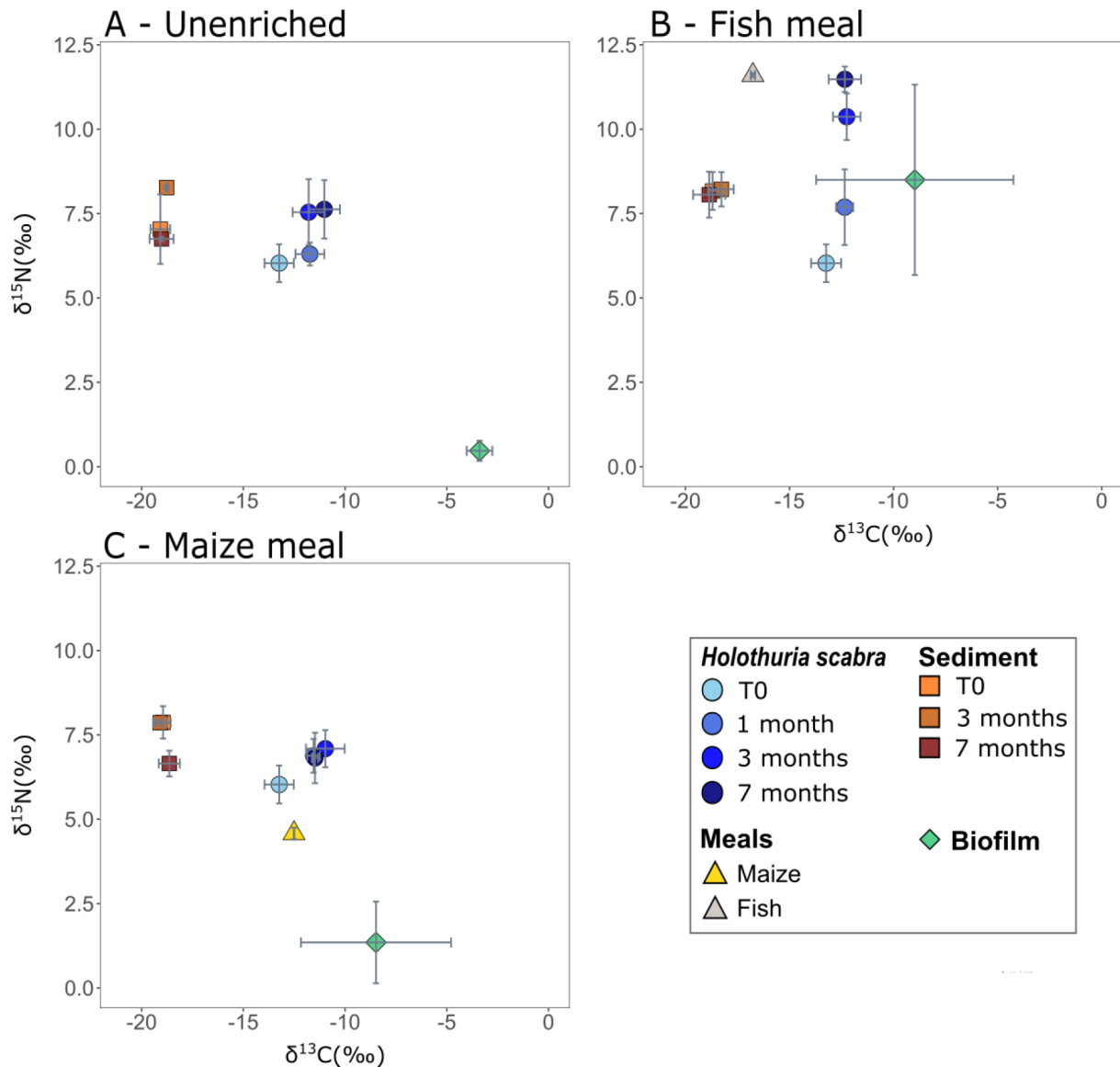
( $7.6 \pm 0.9\%$  and  $6.8 \pm 0.7\%$ , respectively, Conover-Iman multiple comparison tests with Bonferroni's adjustment,  $p$ -value = 0.18). The  $\delta^{13}\text{C}$  also displayed significant variations between treatments at the  $T_f$  (KW test,  $H_{25} = 11.49$ ,  $p$ -value < 0.05). Muscles of animals from unenriched and fish meal treatments displayed significantly different  $\delta^{13}\text{C}$  values (Conover-Iman multiple comparison tests with Bonferroni's adjustment  $p$ -value < 0.001), with muscles from unenriched treatment more  $^{13}\text{C}$  enriched ( $-11 \pm 0.8\%$ ) than muscles from fish meal treatment ( $-12.3 \pm 0.8\%$ ). Muscles of animals from maize meal treatment presented intermediate values ( $-11.5 \pm 0.8\%$ ) showing no significant difference from both others treatments (Conover-Iman multiple comparison tests with Bonferroni's adjustment,  $p$ -value = 0.09 and  $p$ -value = 0.15, when compare to fish and unenriched treatments, respectively).

### 3.3 Fatty acids

The global FA composition of sediments varied significantly among the different treatments (PERMANOVA,  $p$ -value < 0.05,  $df=3$ ,  $R^2=0.19$ ), with all sampling times combined. The FA composition of sediment was dominated by saturated fatty acids (SFA) in the 3 different treatments during all the experiment ( $\Sigma\text{SFA}$  ranging from 41.0% to 47.0%; Tab. 1). All sediment samples but one (maize meal, dominated by 15:0) were dominated by 16:0, and included less than 10% of poly-unsaturated fatty acids (PUFA) (Tab. 1). Even though these differences were supported by slight variations in specific FA, higher levels of 15:0 and 17:1n-8 were found in maize meal sediment, as well as higher levels of branched FA for unenriched and fish meal treatments (Tab. 1). FA whose abundance differed the most between maize and fish meal (18:1n-9, 18:2n6, 20:5n-3 and 22:6n-6, Tab. 2) showed slight variations in sediments from the different treatments (Tab. 1). A higher amount of branched FA was observed in the composition of sediments from unenriched and fish meal treatments at the end of the experiment.

Biofilm contained a higher proportion of PUFA than sediment (between 24.3% and 27.9%, Tab. 2) with 20:5n-3 accounting for 7.6% to 10.5% of total FA and 20:4n-6 reaching  $\pm 4\%$ , while 22:6n-3 was almost absent from biofilms from the 3 different conditions (Tab. 2). SFA were still dominant, accounting for 35.9% to 38.6%, with 16:0 as the principal SFA (Tab. 1). The biofilm was also characterized by high proportions of 16:1n-7 for the three treatment, while 16:1n-9 and 18:1n-9 were found in higher levels in biofilm that developed in tanks supplemented with maize and fish meal (Tab. 2).

Maize and fish meals supplemented to shrimp farm sediment during the experiment displayed highly contrasted FA compositions. Both food sources were dominated by PUFA (between 39.6% and 42.1% of total FA; Tab. 2). Maize meal showed higher levels of monounsaturated FA (MUFA) than fish meals (29.2% and 22.4%, respectively; Tab. 2) and SFA were higher in fish meal than in maize meal (34.9% and 28.4%, respectively; Tab. 2). These two food sources differed markedly in the composition of a few major FA. 18:1n-9 accounted for 26.5% in maize but for only 7.7% in fish meal (Tab. 2). Similarly, 18:2n-6 accounted for 38.7% of total FA in



**Fig. 1.**  $\delta^{13}\text{C}$  vs.  $\delta^{15}\text{N}$  (mean  $\pm$  S.D.) of *Holothuria scabra* and its potential food sources (sediment, biofilm, maize and fish meals) during the feeding experiment. *H. scabra* reared on shrimp-farm sediment only (unenriched treatment) (A). *H. scabra* reared on shrimp-farm sediment and fishmeal (B) and *H. scabra* reared on shrimp-farm sediment and maize meal (C). Fish and maize meal stable isotope compositions did not vary through the experiment.

maize meal but only for 0.9% in fish meal. 20:5n-3 and 22:6n-3 were in much higher proportions in fish meal than in maize meal, accounting for 14.8% and 11.1% of fish meal total FA, respectively, while these FA were virtually absent from maize meal (Tab. 2).

We found no significant interaction between time and treatment in the composition of in *H. scabra* NL (PERMANOVA,  $p$ -value = 0.39,  $df=2$ ,  $R^2=0.04$ ). The global composition of NL varied over time (PERMANOVA,  $p$ -value < 0.001,  $df=2$ ,  $R^2=0.08$ ), and between the different treatments (PERMANOVA,  $p$ -value < 0.05,  $df=4$ ,  $R^2=0.09$ ). For the three treatments and the three sampling times, NL FA in *H. scabra* muscle strips were dominated by SFA (between 31.0% and 45.2%), while MUFA ranged from 21.2% to 31.8% and PUFA from 14.0% to 26.9% (Tab. 3). The content of branched FA ranged from 1.6% to 7.5% (Tab. 3). The main

SFA were 16:0 (between 7.9% and 13.7%) and 18:0 (between 7.3% and 17.7%). The most abundant MUFA was 18:1n-9 (from 3.96% to 7.7%), while PUFA were dominated by 20:4n-6 (from 4.3% to 12.3%) and 20:5n-3 (from 2.9% to 7.9%) (Tab. 3).

The NL FA composition of *H. scabra* in unenriched treatment varied significantly over the experiment (PERMANOVA,  $p$ -value < 0.001,  $df=2$ ,  $R^2=0.34$ ; Fig. 2A), due to an initial strong increase in PUFA (mainly 20:4n-6 and 20:5n-3) from T<sub>0</sub> to after 3 months of experiment, then a slight decrease was observed after the 7 months of experiment (Tab. 3 and Fig. 2A). The NL FA composition of individuals from fish meal treatment was also significantly different over time (PERMANOVA,  $p$ -value < 0.001,  $df=2$ ,  $R^2=0.31$ ; Fig. 2B), as for individuals from maize meal treatment (PERMANOVA,  $p$ -value < 0.05,  $df=2$ ,  $R^2=0.26$ ; Fig. 2C).

**Table 1.** Fatty acid composition (mass % of total FA, mean  $\pm$  S.D.) of sediment from the different treatments at the beginning of the experiment ( $T_0$ ), after 3 months, and after 7 months ( $n=3$  for each time). SFA: saturated FA; MUFA: monounsaturated FA; PUFA: polyunsaturated FA; Branched: FA [15iso+15anteiso+16iso+17iso+17anteiso].

	$T_0$	3 months				7 months			
		Unenriched	Maize meal	Fish meal	Control	Unenriched	Maize meal	Fish meal	Control
$\Sigma$ Branched	6.7 $\pm$ 1.9	10.2 $\pm$ 1.1	6.1 $\pm$ 0.8	8.6 $\pm$ 2.8	12.0	11.8 $\pm$ 0.9	8.2 $\pm$ 2.3	12.4 $\pm$ 2.0	6.7
14:0	5.6 $\pm$ 0.5	4.8 $\pm$ 0.5	4.3 $\pm$ 0.7	5.9 $\pm$ 0.6	5.3	5.0 $\pm$ 0.8	5.3 $\pm$ 1.6	6.5 $\pm$ 1.0	4.9
15:0	8.7 $\pm$ 2.5	9.5 $\pm$ 2.1	15.9 $\pm$ 5.3	6.7 $\pm$ 4.2	5.3	7.2 $\pm$ 3.1	14.0 $\pm$ 7.8	3.8 $\pm$ 1.2	11.3
16:0	22.6 $\pm$ 1.3	16.2 $\pm$ 0.8	14.8 $\pm$ 1.2	19.7 $\pm$ 1.4	17.5	16.9 $\pm$ 1.5	15.8 $\pm$ 3.2	18.2 $\pm$ 0.7	19.7
17:0	2.8 $\pm$ 0.4	2.8 $\pm$ 0.3	2.2 $\pm$ 0.2	1.8 $\pm$ 0.6	2.6	2.8 $\pm$ 0.1	2.4 $\pm$ 0.9	2.6 $\pm$ 0.4	3.3
18:0	2.9 $\pm$ 0.7	2.6 $\pm$ 0.7	3.1 $\pm$ 0.6	4.8 $\pm$ 1.5	4.2	4.6 $\pm$ 1.1	4.9 $\pm$ 3.9	4.6 $\pm$ 0.3	3.3
19:0	0.1 $\pm$ 0.1	0.3 $\pm$ 0.0	0.2 $\pm$ 0.1	0.2 $\pm$ 0.0	0.3	0.3 $\pm$ 0.0	0.2 $\pm$ 0.0	0.3 $\pm$ 0.1	0.5
20:0	0.9 $\pm$ 0.2	1.4 $\pm$ 0.3	1.0 $\pm$ 0.3	0.9 $\pm$ 0.1	1.6	1.6 $\pm$ 0.3	1.0 $\pm$ 0.3	1.4 $\pm$ 0.5	1.3
22:0	1.1 $\pm$ 0.4	1.4 $\pm$ 0.3	1.4 $\pm$ 0.6	0.7 $\pm$ 0.2	1.7	1.6 $\pm$ 0.3	1.3 $\pm$ 0.4	1.2 $\pm$ 0.7	1.1
24:0	2.1 $\pm$ 0.8	3.0 $\pm$ 0.7	2.8 $\pm$ 1.5	0.9 $\pm$ 0.8	3.6	3.6 $\pm$ 0.7	1.9 $\pm$ 0.6	2.5 $\pm$ 1.6	2.0
$\Sigma$ SFA	<b>46.9<math>\pm</math>2.2</b>	<b>41.9<math>\pm</math>1.6</b>	<b>45.8<math>\pm</math>2.5</b>	<b>41.7<math>\pm</math>2.4</b>	<b>42.1</b>	<b>43.6<math>\pm</math>1.4</b>	<b>47.0<math>\pm</math>2.5</b>	<b>41.0<math>\pm</math>1.6</b>	<b>47.4</b>
16:1n-9	1.3 $\pm$ 0.8	2.1 $\pm$ 0.9	0.6 $\pm$ 0.2	1.4 $\pm$ 0.4	1.5	1.9 $\pm$ 0.2	1.4 $\pm$ 0.1	1.9 $\pm$ 0.3	1.1
16:1n-7	11.6 $\pm$ 1.8	7.0 $\pm$ 1.1	6.0 $\pm$ 0.5	7.6 $\pm$ 0.6	6.7	5.2 $\pm$ 0.8	6.0 $\pm$ 2.4	7.3 $\pm$ 1.7	7.3
16:1n-5	1.0 $\pm$ 0.2	1.1 $\pm$ 0.2	0.7 $\pm$ 0.1	0.9 $\pm$ 0.1	1.3	1.2 $\pm$ 0.3	0.9 $\pm$ 0.1	1.0 $\pm$ 0.2	0.8
17:1n-8	4.8 $\pm$ 1.2	5.3 $\pm$ 1.6	9.6 $\pm$ 2.8	3.1 $\pm$ 1.8	2.8	3.6 $\pm$ 1.7	7.1 $\pm$ 4.3	1.8 $\pm$ 0.7	5.0
18:1n-9	2.7 $\pm$ 1.0	3.1 $\pm$ 0.4	2.4 $\pm$ 0.4	4.9 $\pm$ 3.5	3.9	3.8 $\pm$ 1.2	2.7 $\pm$ 0.6	4.4 $\pm$ 0.3	3.3
18:1n-7	3.9 $\pm$ 1.0	4.9 $\pm$ 0.6	4.4 $\pm$ 0.5	6.8 $\pm$ 1.8	5.8	5.1 $\pm$ 0.9	4.7 $\pm$ 1.1	8.2 $\pm$ 1.8	4.3
20:1n-11	0.3 $\pm$ 0.3	0.3 $\pm$ 0.1	0.2 $\pm$ 0.0	0.3 $\pm$ 0.0	0.5	0.4 $\pm$ 0.1	0.2 $\pm$ 0.1	0.7 $\pm$ 0.6	0.4
22:1n-7	0.0 $\pm$ 0.0	0.0 $\pm$ 0.0	0.0 $\pm$ 0.0	0.1 $\pm$ 0.0	0.1	0.0 $\pm$ 0.0	0.1 $\pm$ 0.1	0.0 $\pm$ 0.0	0.0
23:1n-9	0.0 $\pm$ 0.1	0.0 $\pm$ 0.1	0.2 $\pm$ 0.0	0.1 $\pm$ 0.1	0.0	0.0 $\pm$ 0.1	0.1 $\pm$ 0.1	0.0 $\pm$ 0.0	0.1
24:1n-9	0.1 $\pm$ 0.1	0.1 $\pm$ 0.0	0.1 $\pm$ 0.1	0.1 $\pm$ 0.0	0.1	0.1 $\pm$ 0.1	0.0 $\pm$ 0.0	0.1 $\pm$ 0.0	0.1
$\Sigma$ MUFA	<b>25.9<math>\pm</math>3.3</b>	<b>24.2<math>\pm</math>1.4</b>	<b>24.4<math>\pm</math>2.9</b>	<b>25.4<math>\pm</math>3.7</b>	<b>23.0</b>	<b>21.6<math>\pm</math>2.3</b>	<b>23.3<math>\pm</math>0.7</b>	<b>25.7<math>\pm</math>3.6</b>	<b>22.5</b>
18:2n-6	1.1 $\pm$ 0.9	0.9 $\pm$ 0.3	1.1 $\pm$ 0.7	3.0 $\pm$ 3.9	1.1	0.9 $\pm$ 0.2	0.7 $\pm$ 0.2	1.6 $\pm$ 0.8	0.8
18:3n-3	0.7 $\pm$ 0.4	0.8 $\pm$ 0.3	3.4 $\pm$ 5.3	1.1 $\pm$ 0.4	1.3	1.3 $\pm$ 0.3	0.8 $\pm$ 0.2	1.3 $\pm$ 0.3	0.8
18:4n-3	0.6 $\pm$ 0.1	0.5 $\pm$ 0.1	0.5 $\pm$ 0.1	1.1 $\pm$ 0.2	0.4	0.7 $\pm$ 0.2	1.1 $\pm$ 0.5	0.6 $\pm$ 0.3	0.7
20:3n-3	0.3 $\pm$ 0.2	0.4 $\pm$ 0.1	0.3 $\pm$ 0.1	0.4 $\pm$ 0.1	0.7	0.8 $\pm$ 0.5	0.3 $\pm$ 0.1	0.3 $\pm$ 0.3	0.3
20:4n-6	1.1 $\pm$ 0.3	1.2 $\pm$ 0.5	1.3 $\pm$ 0.2	0.8 $\pm$ 0.3	0.3	0.8 $\pm$ 0.2	1.0 $\pm$ 0.5	0.3 $\pm$ 0.1	1.6
20:5n-3	1.3 $\pm$ 0.7	1.3 $\pm$ 0.7	1.5 $\pm$ 0.4	1.7 $\pm$ 1.5	0.2	0.7 $\pm$ 0.1	1.3 $\pm$ 1.2	0.1 $\pm$ 0.1	1.8
22:6n-3	0.1 $\pm$ 0.1	0.1 $\pm$ 0.0	0.0 $\pm$ 0.0	0.5 $\pm$ 0.8	0.2	0.2 $\pm$ 0.0	0.1 $\pm$ 0.0	0.1 $\pm$ 0.1	0.1
$\Sigma$ PUFA	<b>5.6<math>\pm</math>1.4</b>	<b>5.7<math>\pm</math>1.6</b>	<b>8.6<math>\pm</math>6.4</b>	<b>9.2<math>\pm</math>6.0</b>	<b>4.5</b>	<b>5.7<math>\pm</math>0.5</b>	<b>5.8<math>\pm</math>2.2</b>	<b>4.8<math>\pm</math>0.4</b>	<b>6.5</b>
$\Sigma$ Unknown	1.2 $\pm$ 0.3	1.6 $\pm$ 0.3	0.9 $\pm$ 0.2	0.9 $\pm$ 0.3	2	1.9 $\pm$ 0.3	1 $\pm$ 0.2	1.5 $\pm$ 0.8	1.1
$\Sigma$ Other	13.7 $\pm$ 1	16.5 $\pm$ 0.4	14.3 $\pm$ 0.9	14.3 $\pm$ 4.6	16.3	15.3 $\pm$ 1.3	14.7 $\pm$ 0.7	14.5 $\pm$ 0.4	15.9

However, we found a strong overlap in the FA composition of sea cucumbers from the different treatments, owing to a strong inter-tank variability (Fig. 2D). At the end of the experiment, individuals fed maize meal were characterized by high (>5%) content in 16:0, 18:0, 16:1n-9 and 20:3n-3, while individuals fed fish meal displayed high content in branched FA, 16:0 and 18:0 (Tab. 3). Branched FA were found in significant higher amount in *H. scabra* having received additional meals, especially for fish meal treatment, and at intermediate levels in animals fed on maize meal (Tab. 3).

## 4 Discussion

### 4.1 Stable isotope and fatty acid composition of potential food sources

Although maize and fish meal displayed contrasted SI and FA compositions, the addition of these food sources did not result in significant differences in the SI composition of

sediments. A first explanation of this result could be that external inputs are diluted in a larger volume of underlying sediment. This would result in buffering any isotopic variations due to these inputs. However, additional isotopic measurements carried out on surface sediment did not reveal any significant differences compared to those presented, suggesting that these additional food sources did not accumulate on the bottom of the mesocosms (data not shown). It is therefore more likely that these sources are consumed, either by *H. scabra* and/or by micro-organisms (Yokoyama, 2013; Robinson et al., 2018). The consumption of maize/fish meals by *H. scabra* and/or by micro-organisms would release inorganic nutrients that could enhance the growth of fast-growing primary producers forming biofilm, such as filamentous algae or bacteria. This is supported by isotopic differences in biofilms covering the mesocosms' walls, which follow isotopic differences in external sources (i.e.  $^{15}\text{N}$  enriched for fish meal,  $^{13}\text{C}$  enriched for maize meal).

**Table 2.** Fatty acid composition (mass % of total FA, mean  $\pm$  S.D.) of biofilm (after 7 months) ( $n=3$ ), maize and fish meals ( $n=2$ ). SFA: saturated FA; MUFA: monounsaturated FA; PUFA: polyunsaturated FA; Branched: FA [15iso+15anteiso+16iso+17iso+17anteiso].

	Biofilm				Meals	
	Unenriched	Maize meal	Fish meal	Control	Maize	Fish
$\Sigma$ Branched	1.5 $\pm$ 0.3	2.0 $\pm$ 0.7	1.9 $\pm$ 0.4	1.8	0.1	0.7
14:0	5.9 $\pm$ 0.2	9.4 $\pm$ 2.6	5.8 $\pm$ 1.1	6.2	0.3	6.8
15:0	0.4 $\pm$ 0.1	0.8 $\pm$ 0.6	1.2 $\pm$ 0.5	1.6	0.4	0.5
16:0	27.1 $\pm$ 1.6	25.0 $\pm$ 1.3	25.2 $\pm$ 1.5	26.4	17.7	22.0
17:0	0.3 $\pm$ 0.1	0.4 $\pm$ 0.2	0.5 $\pm$ 0.2	0.5	0.3	0.5
18:0	1.8 $\pm$ 1.0	2.0 $\pm$ 0.9	2.1 $\pm$ 0.9	2.5	5.2	4.9
19:0	0.1 $\pm$ 0.0	0.1 $\pm$ 0.0	0.1 $\pm$ 0.0	0.1	0.1	0.1
20:0	0.4 $\pm$ 0.1	0.4 $\pm$ 0.1	0.5 $\pm$ 0.0	0.5	1.2	0.2
22:0	0.3 $\pm$ 0.1	0.5 $\pm$ 0.1	0.5 $\pm$ 0.4	0.2	2.0	0.1
24:0	0.2 $\pm$ 0.3	0.0 $\pm$ 0.0	0.1 $\pm$ 0.0	0.0	1.2	0.3
$\Sigma$ SFA	<b>36.4<math>\pm</math>2.4</b>	<b>38.6<math>\pm</math>2.2</b>	<b>35.9<math>\pm</math>2.7</b>	<b>38.1</b>	<b>28.4</b>	<b>34.9</b>
16:1n-9	0.7 $\pm$ 0.2	5.5 $\pm$ 3.5	4.1 $\pm$ 3.8	8.5	0.2	0.4
16:1n-7	12.6 $\pm$ 4.0	9.7 $\pm$ 1.8	10.2 $\pm$ 8.5	4.2	0.8	7.6
16:1n-5	0.7 $\pm$ 0.1	1.1 $\pm$ 1.3	0.3 $\pm$ 0.0	5.4	0.0	0.2
17:1n-8	0.3 $\pm$ 0.1	0.5 $\pm$ 0.2	0.8 $\pm$ 0.4	0.8	0.0	0.1
18:1n-9	1.6 $\pm$ 0.4	2.7 $\pm$ 2.1	5.9 $\pm$ 2.3	5.4	26.5	7.7
18:1n-7	4.7 $\pm$ 0.8	3.4 $\pm$ 1.4	2.4 $\pm$ 0.6	2.1	0.7	3.7
20:1n-11	0.3 $\pm$ 0.4	0.2 $\pm$ 0.2	0.1 $\pm$ 0.1	0.2	0.3	0.1
22:1n-7	0.0 $\pm$ 0.0	0.1 $\pm$ 0.0	0.0 $\pm$ 0.0	0.0	0.0	0.1
23:1n-9	0.1 $\pm$ 0.1	0.1 $\pm$ 0.0	0.1 $\pm$ 0.0	0.1	0.0	0.0
24:1n-9	1.7 $\pm$ 0.4	1.7 $\pm$ 0.1	0.8 $\pm$ 0.5	1.2	0.0	0.9
$\Sigma$ MUFA	<b>23.0<math>\pm</math>4.4</b>	<b>25.3<math>\pm</math>2.3</b>	<b>25.2<math>\pm</math>6.0</b>	<b>28.5</b>	<b>29.2</b>	<b>22.4</b>
18:2n-6	2.5 $\pm$ 0.6	1.7 $\pm$ 0.5	3.1 $\pm$ 2.2	1.7	38.7	0.9
18:3n-3	3.9 $\pm$ 1.6	1.6 $\pm$ 0.9	4.2 $\pm$ 4.8	3.7	2.2	0.5
18:4n-3	5.9 $\pm$ 1.3	5.1 $\pm$ 1.6	4.4 $\pm$ 3.2	3.3	0.0	2.3
20:3n-3	0.3 $\pm$ 0.1	1.9 $\pm$ 2.5	3.5 $\pm$ 2.8	5.8	0.1	0.1
20:4n-6	4.3 $\pm$ 2.4	3.2 $\pm$ 0.6	3.7 $\pm$ 2.1	1.8	0.1	1.0
20:5n-3	10.5 $\pm$ 2.3	9.9 $\pm$ 3.5	7.6 $\pm$ 0.3	6.3	0.1	14.8
22:6n-3	0.2 $\pm$ 0.2	0.5 $\pm$ 0.1	0.3 $\pm$ 0.1	0.0	0.3	11.1
$\Sigma$ PUFA	<b>27.9<math>\pm</math>1.5</b>	<b>24.3<math>\pm</math>3.1</b>	<b>27.1<math>\pm</math>10.0</b>	<b>22.9</b>	<b>42.1</b>	<b>39.6</b>
$\Sigma$ Unknown	0.2 $\pm$ 0.1	0.4 $\pm$ 0.3	0.5 $\pm$ 0.2	0.1	0.1	0.4
$\Sigma$ Other	11 $\pm$ 1.3	9.5 $\pm$ 2.5	9.5 $\pm$ 1.3	8.5	0.1	1.9

The FA composition of sediment was dominated by SFA, which is common for detritus-dominated sediments (Kelly and Scheibling, 2012). Although the overall FA composition did not vary during the experiment, sediment at the beginning of the experiment was characterized by significant amounts of 16:1n-7, which is generally considered to be abundant in diatoms (Dalsgaard et al., 2003). The amount of this FA decreased during the experiment in the 3 treatments, simultaneously with an increase in branched FA and in 18:1n-7, which have been reported specific to bacteria in tropical environments (Meziane and Tsuchiya, 2000). These results reinforce the previous hypothesis of external food sources being metabolized by bacteria in supplemented treatments. The increase in bacterial FA is also observed in non-enriched treatment, which suggests a natural trend to favor bacterial development of high organic matter shrimp-farm sediments (Burford et al., 1998).

The FA composition of biofilm was also investigated, because *H. scabra* was sometimes observed foraging at the

vicinity of tank walls. At the end of the experiment, the FA composition of biofilm was dominated by 16:1n-7, 18:1n-7 and 20:5n-3, which are biomarkers of diatoms and bacteria (Kharlamenko et al., 2001; Meziane and Tsuchiya, 2000). Higher content in 18:3n-3, 18:4n-3 and 20:3n-3 compared to sediments was also observed, and can be explained by the presence of filamentous Phaeophyta on the walls of mesocosms (Volkman et al., 1989; Kharlamenko et al., 2001).

#### 4.2 Assimilation of shrimp farm sediment by *H. scabra*

The temporal variation of *H. scabra* SI composition for unenriched treatment revealed slight  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  increases during the experiment (+2.2‰ and +1.6‰, respectively). Although at  $T_E$ , SI composition of *H. scabra* seems too  $^{13}\text{C}$  enriched to reveal a diet only based on sedimentary organic matter, this temporal variation could reveal either a diet based on both sedimentary organic matter and biofilm, or dominated

**Table 3.** Fatty acid composition (mass % of total FA, mean  $\pm$  S.D.) of the neutral lipids of *Holothuria scabra* fed the different treatments at the beginning of the experiment ( $T_0$ ) ( $n=5$ ), after 3 months ( $n=3$  for each treatment) and after 7 months ( $n=9$  for each treatment). Different letters indicate significant differences (KW tests followed by Conover-Iman multiple comparison tests with Bonferroni's adjustment method, at significant level  $\alpha < 0.05$ ), indicating variations of FA compositions between treatment at 3 months and 7 months. SFA: saturated FA; MUFA: monounsaturated FA; PUFA: polyunsaturated FA; Branched: FA [15iso+15anteiso+16iso+17iso+17anteiso].

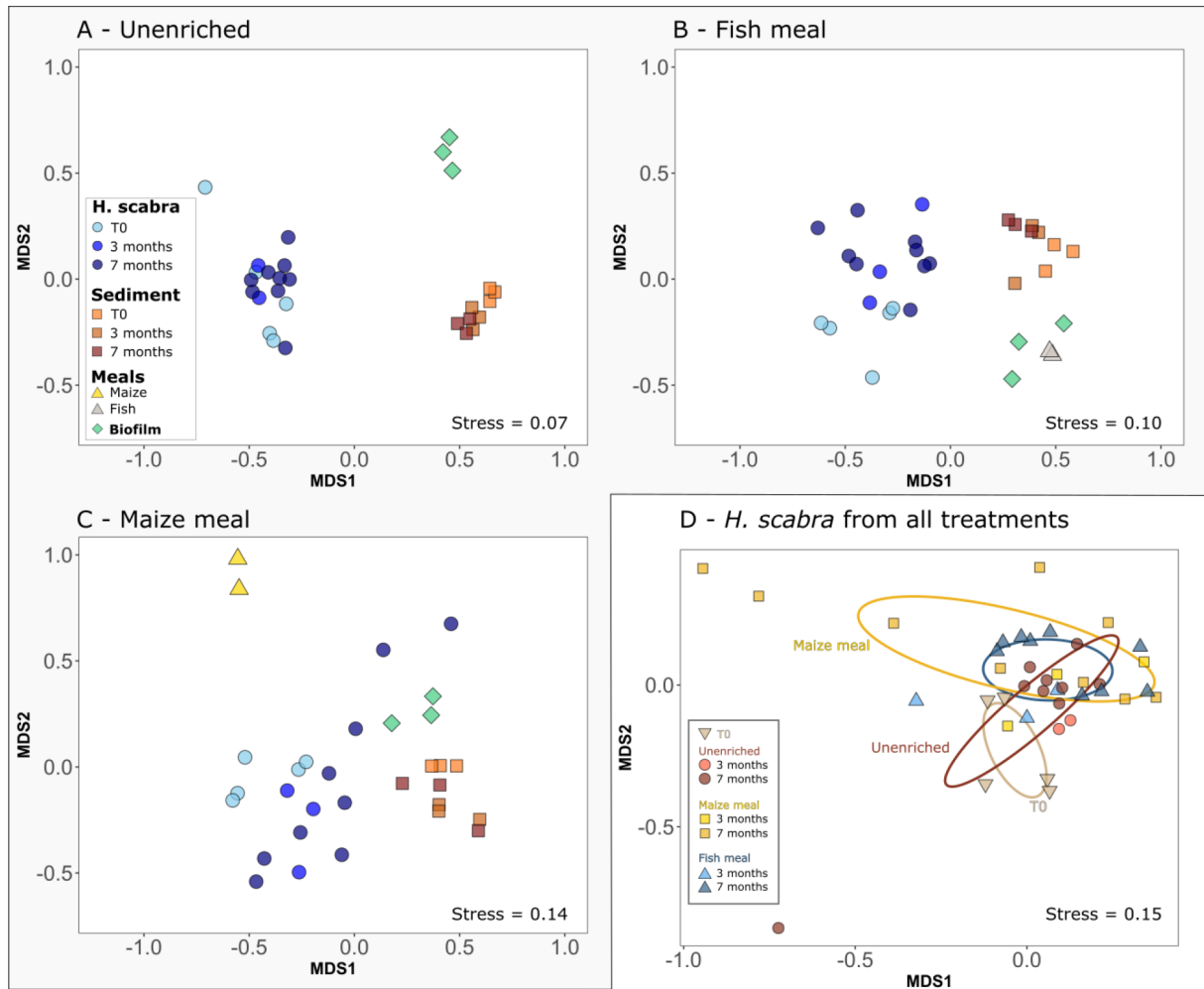
	<i>Holothuria scabra</i>						
	$T_0$	3 months ( $T_3$ )			7 months ( $T_7$ )		
		Unenriched	Maize meal	Fish meal	Unenriched	Maize meal	Fish meal
$\Sigma$ Branched	1.5 $\pm$ 1.2	2.2 $\pm$ 0.4	3.6 $\pm$ 1.4	5.3 $\pm$ 4.9	3.7 $\pm$ 0.8 <sup>a</sup>	4.9 $\pm$ 1.9 <sup>ab</sup>	7.5 $\pm$ 2.5 <sup>b</sup>
14:0	1.8 $\pm$ 0.9	1.7 $\pm$ 0.4	2.4 $\pm$ 0.0	1.8 $\pm$ 0.5	2.6 $\pm$ 0.4	4.0 $\pm$ 2.4	2.6 $\pm$ 0.8
15:0	0.9 $\pm$ 0.7	1.2 $\pm$ 0.4	1.4 $\pm$ 0.6	1.9 $\pm$ 0.8	1.4 $\pm$ 0.2	1.8 $\pm$ 1.0	1.6 $\pm$ 0.6
16:0	13.7 $\pm$ 5.4	10.5 $\pm$ 0.1	7.9 $\pm$ 2.5	10.9 $\pm$ 0.5	11.5 $\pm$ 2.3	12.3 $\pm$ 4.4	11.7 $\pm$ 3.3
17:0	0.7 $\pm$ 0.5	1.1 $\pm$ 0.2	1.3 $\pm$ 0.2	0.8 $\pm$ 0.3	1.3 $\pm$ 0.2	1.2 $\pm$ 0.2	1.8 $\pm$ 0.6
18:0	17.7 $\pm$ 2.7	10.3 $\pm$ 2.5	7.3 $\pm$ 1.2	13.6 $\pm$ 7.5	11.5 $\pm$ 3.2	8.5 $\pm$ 3.5	10.4 $\pm$ 0.8
19:0	1.4 $\pm$ 1.0	1.5 $\pm$ 0.3	1.5 $\pm$ 0.3	1.1 $\pm$ 0.3	1.3 $\pm$ 0.3	1.5 $\pm$ 1.2	1.8 $\pm$ 0.4
20:0	2.3 $\pm$ 0.4	2.1 $\pm$ 0.3	2.5 $\pm$ 0.5	1.6 $\pm$ 0.3	2.7 $\pm$ 0.6	2.1 $\pm$ 1.3	2.5 $\pm$ 0.5
22:0	4.6 $\pm$ 2.1	3.9 $\pm$ 0.2	5.1 $\pm$ 2.7	2.6 $\pm$ 1.7	5.0 $\pm$ 1.2	3.5 $\pm$ 2.1	4.2 $\pm$ 0.6
24:0	1.9 $\pm$ 0.9	1.4 $\pm$ 0.1	1.5 $\pm$ 0.3	1.0 $\pm$ 0.2	1.2 $\pm$ 0.4	0.7 $\pm$ 0.2	1.2 $\pm$ 0.4
$\Sigma$ SFA	<b>45.2 <math>\pm</math> 8.3</b>	<b>33.6 <math>\pm</math> 2.5</b>	<b>31.0 <math>\pm</math> 4.3</b>	<b>35.2 <math>\pm</math> 6.2</b>	<b>38.6 <math>\pm</math> 8.1</b>	<b>35.7 <math>\pm</math> 5.3</b>	<b>37.8 <math>\pm</math> 4.6</b>
16:1n-9	1.8 $\pm$ 1.4	1.8 $\pm$ 0.7	2.3 $\pm$ 1.3	2.6 $\pm$ 1.3	2.4 $\pm$ 1.0	5.4 $\pm$ 6.7	2.1 $\pm$ 2.2
16:1n-7	1.8 $\pm$ 0.6	3.0 $\pm$ 0.3 <sup>ab</sup>	4.2 $\pm$ 0.3 <sup>a</sup>	1.6 $\pm$ 0.4 <sup>b</sup>	3.2 $\pm$ 0.3	2.8 $\pm$ 2.4	2.9 $\pm$ 1.2
16:1n-5	5.6 $\pm$ 3.4	3.8 $\pm$ 2.3	1.1 $\pm$ 1.0	2.0 $\pm$ 1.9	0.3 $\pm$ 0.3	0.4 $\pm$ 0.3	0.2 $\pm$ 0.1
18:1n-9	5.7 $\pm$ 1.6	4.3 $\pm$ 0.2	7.7 $\pm$ 5.7	4.8 $\pm$ 1.3	4.2 $\pm$ 1.7	5.0 $\pm$ 2.1	3.9 $\pm$ 0.7
18:1n-7	1.1 $\pm$ 0.7	2.1 $\pm$ 0.0	4.2 $\pm$ 0.7	2.8 $\pm$ 1.0	2.8 $\pm$ 0.5	2.5 $\pm$ 1.4	3.9 $\pm$ 1.1
20:1n-11	2.1 $\pm$ 0.4	3.5 $\pm$ 1.3	2.4 $\pm$ 0.4	1.9 $\pm$ 1.1	2.6 $\pm$ 0.3	2.8 $\pm$ 2.8	3.2 $\pm$ 2.8
22:1n-7	2.4 $\pm$ 0.9	2.2 $\pm$ 0.2	3.5 $\pm$ 1.4	1.5 $\pm$ 0.9	2.4 $\pm$ 0.2	2.4 $\pm$ 1.7	2.8 $\pm$ 0.6
23:1n-9	0.8 $\pm$ 0.5	1.7 $\pm$ 0.0	1.4 $\pm$ 0.5	1.7 $\pm$ 1.0	2.1 $\pm$ 0.1	2.4 $\pm$ 2.5	3.2 $\pm$ 3.0
24:1n-9	3.2 $\pm$ 1.1	3.1 $\pm$ 0.4	4.7 $\pm$ 2.7	2.9 $\pm$ 1.4	2.9 $\pm$ 0.2	2.7 $\pm$ 2.0	4.0 $\pm$ 0.7
$\Sigma$ MUFA	<b>25.3 <math>\pm</math> 2.8</b>	<b>26.3 <math>\pm</math> 2.2</b>	<b>31.8 <math>\pm</math> 2.0</b>	<b>22.9 <math>\pm</math> 7.3</b>	<b>23.1 <math>\pm</math> 4.2</b>	<b>27.3 <math>\pm</math> 0.6</b>	<b>26.6 <math>\pm</math> 4.1</b>
18:2n-6	4.8 $\pm$ 3.6	4.7 $\pm$ 1.9	1.5 $\pm$ 1.3	2.8 $\pm$ 2.0	1.8 $\pm$ 1.3	1.2 $\pm$ 0.2	0.8 $\pm$ 0.4
18:4n-3	2.0 $\pm$ 2.2	1.8 $\pm$ 0.2	1.6 $\pm$ 1.4	1.4 $\pm$ 0.3	1.4 $\pm$ 0.4	1.1 $\pm$ 0.8	0.8 $\pm$ 0.4
20:3n-3	0.3 $\pm$ 0.4	0.0 $\pm$ 0.0	0.0 $\pm$ 0.0	0.3 $\pm$ 0.2	0.6 $\pm$ 0.4 <sup>b</sup>	8.2 $\pm$ 11.8 <sup>a</sup>	1.7 $\pm$ 1.4 <sup>ab</sup>
20:4n-6	4.3 $\pm$ 0.9	12.3 $\pm$ 2.5	6.5 $\pm$ 1.6	4.4 $\pm$ 2.5	6.4 $\pm$ 1.6	4.3 $\pm$ 3.4	4.4 $\pm$ 2.5
20:5n-3	2.9 $\pm$ 0.5	7.3 $\pm$ 2.9	7.9 $\pm$ 0.9	4.0 $\pm$ 2.2	4.7 $\pm$ 0.8	3.1 $\pm$ 1.7	4.3 $\pm$ 1.8
22:6n-3	0.3 $\pm$ 0.4	0.8 $\pm$ 0.1	1.2 $\pm$ 0.2	1.3 $\pm$ 0.8	1.1 $\pm$ 0.3 <sup>ab</sup>	1.5 $\pm$ 1.5 <sup>a</sup>	1.8 $\pm$ 0.9 <sup>b</sup>
$\Sigma$ PUFA	<b>14.7 <math>\pm</math> 1.9</b>	<b>26.9 <math>\pm</math> 3.8</b>	<b>19.0 <math>\pm</math> 0.6</b>	<b>15.6 <math>\pm</math> 7.2</b>	<b>16.2 <math>\pm</math> 3</b>	<b>19.6 <math>\pm</math> 6.9</b>	<b>14.0 <math>\pm</math> 1.9</b>
$\Sigma$ Unknown	0.7 $\pm$ 0.9	0.5 $\pm$ 0.7	1.2 $\pm$ 0.1	2.3 $\pm$ 1.7	0.7 $\pm$ 0.4	0.8 $\pm$ 0.7	0.6 $\pm$ 0.4
$\Sigma$ Other	12.6 $\pm$ 5.5	10.5 $\pm$ 1.1	13.5 $\pm$ 4.3	18.3 $\pm$ 3.9	17.6 $\pm$ 13.7	11.6 $\pm$ 4.8	13.4 $\pm$ 2.4

by a specific  $^{13}\text{C}$  enriched fraction of sediment. The selective assimilation of a fraction of a composite pool has indeed been reported for various deposit-feeders using SI (Kolasinski et al., 2016). However, the extent of the isotopic lag between sediment and *H. scabra* (8‰) makes the hypothesis of the consumption of a sole fraction of sediment quite unlikely, and strongly suggests that *H. scabra* is also feeding on an alternative source, such as biofilm that may fall down on sediment and thus be incorporated into the pool of sedimentary organic matter.

The NL PUFA/SFA ratio provides information relative to the quality of food assimilated by *H. scabra* (Cripps and Atkinson, 2000). This ratio increased in the first 3 months of growth, before decreasing at the end of the experiment, suggesting a shift in food quality. This shift could be related to a growth stop observed for the same experiment by (Hochard et al., 2016). The growth stopped when the total biomass of

*H. scabra* reached  $220\text{ g m}^{-2}$ , a value similar to those mentioned in other studies (Battaglione et al., 1999; Pitt et al., 2004). This phenomenon was interpreted as a limit carrying capacity of the system for *H. scabra* that was reached, which is supported by the present results. Interestingly, other studies in more controlled environment (filtered seawater, washed sediment) and also testing the food addition were able to reach much higher biomass for the same species ( $600\text{--}900\text{ g m}^{-2}$ ) (Watanabe et al., 2014; Robinson et al., 2015, 2019), showing thus that this limitation can be controlled from initial conditions. The development of the heterotrophic bacterial production through the manipulation of the C:N ration inputs could increase the system carrying capacity (Robinson et al., 2018, 2019). The identification of factors involved in the establishment of the system carrying capacity are still poorly understood and requires further investigation.





**Fig. 2.** Multidimensional scaling based on fatty acid composition (mass %) of *Holothuria scabra* (neutral lipids) and its potential food sources (total lipids); for the unenriched treatment (A), the fish meal treatment (B), and the maize meal treatment (C). (D) Represent neutral lipid fatty acids of *H. scabra* from all treatments and associated ellipses for the different feeding conditions.

The NL FA composition of *H. scabra* muscles in all treatments was dominated by the SFA 16:0 and 18:0, which are generally among the most abundant in any marine consumers worldwide (Kelly and Scheibling, 2012), and not informative in a trophic ecology perspective. The content in PUFA was quite low for all treatments (< 20% for most samples) in comparison to other marine invertebrates, where PUFA often represent up to 50% of NL (e.g. Soudant et al., 1999). The relatively high abundance of 20:5n-3 and 20:4n-6 has been repeatedly reported in sea-cucumbers from various environments (e.g. Kaneniwa et al., 1986; Drazen et al., 2008; Yu et al., 2015). In particular, the content in 20:4n-6 was surprisingly high in *H. scabra* (up to 12%) considering that it was almost absent from all sources, at the exception of the biofilm, where it reached 4% of total FA. This highlights the capacities of *H. scabra* to synthesize *de novo* or to selectively incorporate specific FA. Echinoderms have been shown capable to synthesize several PUFA when they are lacking in their diet, while most other marine animals have very limited PUFA biosynthesis capacities (Kelly and Scheibling, 2012). Finally, branched FA and 18:1n-7 represented non-negligible amounts in muscles of *H. scabra* (their total content reached up to 12%),

which represents an evidence of shrimp farm sediment assimilation by holothurids.

Both tracers support a contribution of biofilm growing on tanks walls to the diet of *H. scabra*. Such conclusion about biofilm contribution should however be only valid for our experimental conclusions. Indeed, because of the relatively small size of the tanks where the experiment took place, the biofilm covering tanks walls did represent an abundant available food source because of high wall surface/sediment area ratio, while its contribution to the diet of holothurids should be reduced in larger ponds, where this ratio is lower. In addition to be a potential food source for animals, some studies shown that the utilization of additional substrates where biofilm can grow contribute to improving water quality, and could also increase the productivity of systems (Milstein, 2005; Khatoun et al., 2007). The accurate evaluation of each food source contribution to the diet of sea cucumbers with SI mixing models requires a reliable evaluation of trophic fractionation, as well as a proper characterization of end-members. The trophic fractionation of organisms during intense growth phases is highly variable (Gorokhova and Hansson, 1999), hence using such a coefficient from the

literature would make no sense for this purpose in this case study. Moreover, our results suggest that only a  $^{13}\text{C}$ -enriched fraction of sedimentary organic matter is assimilated, whose SI characterization is not possible.

### 4.3 Effect of additional food sources on the trophic ecology of *H. scabra*

Sea cucumbers reared on sediment with additional maize meal supplementation displayed a significant increase of their  $\delta^{15}\text{N}$ . At the end of the experiment, their SI composition was in accordance with what could be expected if they were only feeding on maize meal, considering commonly accepted mean isotope fractionation values (e.g. 0.8‰ for  $\Delta\delta^{13}\text{C}$  (Vander Zanden and Rasmussen, 2001), 2.5‰ for  $\Delta\delta^{15}\text{N}$  (Vanderklift and Ponsard, 2003)). However, no significant difference could be observed between individuals for which maize meal was available and those for which this source was absent, suggesting that the similarity between the SI composition of maize meal and *H. scabra* might just as well be a coincidence, and that maize would not be assimilated by *H. scabra*. This hypothesis is strengthened by the analysis of FA composition. The most important maize meal unsaturated FA (18:1n-9 and 18:2n-6) were indeed not present in higher abundances in individuals fed maize meal compared to fish meal and unenriched treatments.

In contrast to other treatments, *H. scabra* subjected to fish meal supplementation displayed a significant marked increase in the  $\delta^{15}\text{N}$ , likely related to the high  $\delta^{15}\text{N}$  of fish meals. This suggests that nitrogen compounds from fish meal are assimilated by *H. scabra*. However, this hypothesis does not fit with  $\delta^{13}\text{C}$  values, for which there was a clear difference (5.3 ‰) between fish meal and *H. scabra*. In fact the  $\delta^{13}\text{C}$  of *H. scabra* from fish meal-supplemented conditions was only significantly different from unenriched but not from maize meal treatment. Moreover, the most abundant FA found in fish meals, especially 20:5n-3 (EPA) and 22:6n-3 (DHA), were not found at higher levels in individuals supplemented with fish meal. Because FA do not contain nitrogen, they only reflect the assimilation of carbon, and do not support a dependence of *H. scabra* towards fish meal. Both SI and FA therefore support the lack of carbon assimilation from fish meal within these mesocosms, although nitrogen seemed to contribute to the growth of holothurids. The differential integration of carbon and nitrogen from this source could arise from a differential digestibility of carbon and nitrogen compounds. Orozco et al. (2014) reported a marked difference in the assimilation of proteins (N dominated compounds) and carbohydrates (C dominated compounds) from shrimp powder by *H. scabra*. The apparent digestibility coefficient was three time higher for proteins (90%) than for carbohydrates (30%). If these inputs of organic matter did enrich the environment (sediment and biofilm), they did not seem to increase the productivity of animals, suggesting limiting mechanisms, which could affect the palatability of food, and/or the alteration of the nutrients transfer via microbial loop. Alternatively, these results could result from an indirect consumption of fish meal in the system as also suggested recently by Robinson et al. (2019). Holothurids are indeed in competition with a diversity of macro and meiofaunal organisms for the assimilation of food,

and can ingest both fresh and recycled organic matter. This illustrates the complexity of the benthic system and show the difficulties emerging from the culture of detritus-feeders.

## 5 Conclusion

This study supports the assimilation of shrimp farm sediment by holothurids, which could therefore represent a solution to pond sediment excessive organic matter enrichment associated with shrimp farming. We also showed that although carbon and nitrogen compounds from maize meal was not assimilated by holothurids, fish meal could represent an indirect source of nitrogen compounds into their tissues. However, discrepancies between the assimilation of nitrogen and carbon from this food source, as well as the growth stop observed during the experiment highlight the need to provide holothurids with a stoichiometrically balanced diet for a more efficient assimilation of sedimentary organic matter. Although our study presents some experimental limitations (in particular linked to density-dependent regulation of growth and feeding) which are inherent, to this kind of approach, we are confident that the patterns observed here concerning nutrient assimilation are quite robust to these biases. Finally, even if the assimilation of sediment was clear, this study could not reveal any significant effect of holothurids on the FA and SI composition of pond sediment. Evaluating the effects of such supplementation on *H. scabra* growth performances and tissue turnover is the next step toward assessing the relevance of this treatment, from both ecological and economic perspectives.

## Conflict of interest

The authors declare no conflict of interest whatsoever concerning the results presented in this manuscript.

## Animal experiments

Animals used in this study were used in experiments in euthanized complying with national rules regarding animal experiments.

## Informed consent

All authors listed here read and agreed with the submitted version of this manuscript.

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