

1 **Deconvolving feeding niches and strategies of abyssal holothurians from their stable**
2 **isotope, amino acid, and fatty acid composition**

3
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15
16 **Abstract**

17 Holothurians are the dominant megabenthic deposit feeders in the Peru Basin (South-East
18 Pacific) and feed to various degrees of selectively on the heterogenous pool of sedimentary
19 detritus, but diet preferences for most holothurian species are unknown. This study reconstructs
20 the diets of 13 holothurian species of the orders Elaspodida, Holothuriida, and Synallactida,
21 from bulk stable isotope analyses ($\delta^{13}\text{C}$, $\delta^{15}\text{N}$) of holothurian body walls and guts, gut contents,
22 and feces that were combined with compound-specific stable isotope analyses of amino acids,
23 phospholipid-derived fatty acids, and neutral lipid-derived fatty acids in the body wall. Fatty acid
24 concentrations showed high levels of storage lipids, an likely adaption to limited food supply to
25 abyssal plains. Amino acid $\delta^{15}\text{N}$ isotope values allowed estimating trophic levels of holothurian
26 species and calculating heterotrophic re-synthesis of amino acids. Fatty acids served as trophic
27 markers for feeding on diatom- and dinoflagellate derived phytodetritus, bacteria, Foraminifera,
28 and detritus containing the PUFA C22:1 ω 9-*cis*. Several holothurian species seemed to be
29 secondary consumers of detritus, while bacteria in their guts were primary consumers of this
30 detritus. A Sørensen–Dice coefficient based cluster analysis using data of trophic levels, levels of
31 heterotrophic re-synthesis of amino acids, feeding selectivity, and food sources/ diet suggested
32 three trophic groups, characterized by different trophic levels. We show that this multi-biomarker
33 driven approach allows to deconvolve trophic niches and feeding selectivity in one of the most
34 challenging environments on earth and to identify dependence of deep-sea species to organic
35 matter inputs that vary with season and/or climate.

36
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38
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47

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50

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54

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57

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60 revising the manuscript to its final version which was approved by all.

61 Introduction

62 Holothurians are one of the most abundant epifauna in the deep sea (Billett et al. 2001; Ruhl
63 2007; Alt et al. 2013; Stratmann et al. 2018) and they can be suspension and deposit feeders
64 (Massin 1982). On soft sediment, deposit feeding holothurians either dig into the sediment as
65 funnel-feeder or conveyor belt-feeder or scavenge the surface sediment as rake feeders (Massin
66 1982). In this way, they take up particulate organic matter that is deposited on or buried in the
67 sediment (Roberts et al. 2000). Holothurians selectively feed on the organic sources in the
68 sediment. The analysis of gut contents from holothurians collected at the Porcupine Abyssal
69 Plain (PAP, NE Atlantic) showed that e.g. *Amperima rosea* Perrier, 1886, *Peniagone diaphana*
70 Théel, 1882, and *Oneirophanta mutabilis mutabilis* Théel, 1879, feed selectively on fresh
71 phytodetritus (FitzGeorge-Balfour et al. 2010). However, when fresh phytodetritus is scarce, *O.*
72 *mutabilis mutabilis* feeds on more refractory detritus material (FitzGeorge-Balfour et al. 2010)
73 which is primarily consumed by the microbial community in its gut (Romero-Romero et al.
74 2021). Other species have a less selective feeding behavior, e.g. *Psychropotes longicauda* Théel,
75 1882, *Molpadiodemas villosus* Théel, 1886, and *Molpadia blakei* Théel, 1886, (FitzGeorge-
76 Balfour et al. 2010). Though it seems that feeding selectivity and diet preferences of holothurians
77 are well known, this is actually true for very few species. For most abyssal holothurians, in fact,
78 these information are very rudimentary (e.g. Billett 1991; Roberts et al. 2000)).

79
80 Whereas holothurians alter the chemical composition of detritus in the sediment, this detritus
81 composition also affects the species composition of holothurians (Wigham et al. 2003;
82 FitzGeorge-Balfour et al. 2010). At PAP, especially *A. rosea*, *P. diaphana* and *O. mutabilis*
83 *mutabilis* had a high concentration of carotenoids in their ovaries which are important for the
84 reproductive success of the species (Tsushima 2007; Svensson and Wong 2011). Therefore,
85 (Wigham et al. 2003) suggested that higher concentrations of carotenoids in the gonads of *A.*
86 *rosea* as compared to other holothurians might give this species a reproductive advantage which
87 could explain the so-called ‘Amperima’ event. During this event, the density of *A. rosea*
88 increased by three orders of magnitude due to large-scale recruitment events that followed
89 changes in the organic carbon flux to the abyssal plain, even though the total megafauna biomass
90 did not change significantly (Billett et al. 2010).

91
92 Amino acids, the building stones of proteins, are required to produce enzymes, structural tissue
93 of fauna, and cell walls of bacteria (Phillips 1984; Libes 2009). Half of the 20 most common
94 amino acids in faunal proteins can be synthesized by the organism itself (Phillips 1984), whereas
95 the other half has to be taken up with the diet and are therefore called ‘essential’ amino acids
96 (EAA) (Phillips 1984). Amino acids include ‘source amino acids’ (i.e., glycine, serine,
97 phenylalanine, tyrosine, lysine), which preserve their $\delta^{15}\text{N}$ values along the trophic chain because
98 no new bonds are formed to the N atom nor are bonds cleaved (Chikaraishi et al. 2009). Other
99 amino acids are ‘metabolic amino acids’ (i.e., threonine) and ‘trophic amino acids’ (i.e.,
100 asparagine, glutamine, alanine, isoleucine, leucine, valine, proline). The $\delta^{15}\text{N}$ values of ‘trophic
101 amino acids’ become enriched during metabolic transamination when nitrogen bonds are cleaved
102 (Chikaraishi et al. 2009). The larger the difference between the ‘source amino acids’ and the
103 ‘trophic amino acids’, the higher is the trophic level of an organism, so the ratio of the $\delta^{15}\text{N}$
104 values of glutamic acid and phenylalanine has been used to estimate the trophic level of an
105 organism following (Chikaraishi et al. 2009).

106

107 Fatty acids, the main components of lipids, serve as energy source, are involved in the
108 transduction of signals, in gene expression, and are components of membranes (Burdge and
109 Calder 2014). They contain neutral lipid-derived fatty acids (NLFAs) and phospholipid-derived
110 fatty acids (PLFAs) (Dalsgaard et al. 2003). NLFAs are required to build wax esters and the
111 storage lipids triacylglycerols, whereas PLFAs are necessary to build structural phospholipids of
112 cell membranes (Dalsgaard et al. 2003). Fatty acids may be unsaturated or saturated and
113 generally a higher number of unsaturated bonds implies that the fatty acid is more labile than a
114 fatty acid with fewer unsaturated bonds (Pond et al. 1997). ‘Essential’ fatty acids have to be
115 taken up with the diet because they can generally only be synthesized *de novo* by primary
116 producers (Dalsgaard et al. 2003), except for a few hydrothermal vent shrimp species and worms
117 that are also able to synthesize them (Pond et al. 1997, 2002). Since several fatty acids are
118 transferred conservatively (i.e., untransformed) from primary producers and primary consumers
119 to higher trophic levels, they may serve as trophic markers and inform about diets (Dalsgaard et
120 al. 2003).

121
122 To decipher feeding types and diet preferences of holothurians from the Peru Basin, compound-
123 specific stable isotope analyses of bulk tissue, gut content, and feces were combined with
124 compound-specific stable isotope analysis of amino acids and fatty acids. We addressed the
125 following research questions: (1) Do the holothurian species have different trophic levels? (2)
126 Can specific feeding strategies and diet preferences identified for the different species?

127

128 **Material and methods**

129 **Sampling of holothurians**

130 Holothurians of the putative species Elpidiidae gen sp. Théel, 1882 (n = 1), *Amperima* sp.
131 Pawson, 1965 (n = 4), *Benthodytes* sp. Théel, 1882 (n = 2), *Benthodytes typica* Théel, 1882
132 (n = 1), *Galatheathuria* sp. Hansen & Madsen, 1956 (n = 1), *Oneirophantha* sp. Théel, 1879
133 (n = 1), *Psychronaetes hansenii* Pawson, 1983 (n = 1), *P. longicauda* (n = 1), *Psychropotes*
134 *semperiana* Théel, 1882 (n = 1), *Synallactes* sp. Ludwig, 1894 (morphotype “pink”; n = 1), and
135 Synallactidae gen sp. Ludwig, 1894 (n = 2) were collected opportunistically with the ROV with
136 the ROV suction sampler in the Peru Basin (Table 1). As a result, sampling of several species
137 was not balanced, but due to logistical constraints it was often limited to n = 1 or n = 2. Aboard
138 RV *Sonne*, the length, height, and width of each holothurian specimen was measured and the
139 specimens were dissected to separate the gut and its content from the remaining tissue. All
140 samples were shock-frozen in liquid nitrogen and stored frozen at -20°C.

141

142 **Table 1.** Details of sampling location and collected holothurian specimens from RV *Sonne*
143 research cruise SO242-2.

Date	Latitude (N)	Longitude (E)	Depth (m)	Putative species
05.09.2015	-7.074	-88.451	4137.0	<i>Amperima</i> sp. (n = 1), <i>Benthodytes</i> sp. (n = 1)
05.09.2015	-7.074	-88.451	4137.5	<i>Mesothuria</i> sp. (n = 1), <i>Amperima</i> sp. (n = 2),
05.09.2015	-7.074	-88.451	4136.4	<i>Oneirophanta</i> sp. (n = 1)
12.09.2015	-7.125	-88.451	4151.0	<i>Amperima</i> sp. (n = 1), <i>Benthodytes</i> sp. (n = 1), <i>B.</i> <i>typica</i> (n = 1)

17.09.2015	-7.082	-88.469	4136.3	<i>Benthodytes</i> sp. (n = 2), <i>B. typica</i> (n = 1), <i>Psychronaetes hanseni</i> (n = 1)
18.09.2015	-7.083	-88.470	4429.4	<i>Amperima</i> sp. (n = 1), Elpidiidae gen sp. (n = 1), <i>Psychropotes longicauda</i> (n = 1), <i>Synallactes</i> sp. (morphotype "pink") (n = 1)
22.09.2015	-7.126	-88.451	4150.0	Synallactidae gen sp. (n = 1)
27.09.2015	-7.078	-88.458	4141.9	<i>B. typica</i> (n = 2), <i>Peniagone</i> sp. (n = 1), <i>Psychropotes semperiana</i> (n = 1), <i>Galatheathuria</i> sp. (n = 1), Synallactidae gen sp. (n = 1), <i>Peniagone</i> sp. (n = 1)

144
145 Additionally, the putative holothurians species *Amperima* sp. (n = 3), *Benthodytes* sp. (n = 3), *B.*
146 *typica* (n = 4), *Mesothuria* sp. Ludwig, 1894 (n = 1), *Peniagone* sp. Théel, 1882 (n = 1), and
147 Synallactidae gen sp. Ludwig, 1894 (n = 1) from the study of (Brown et al. 2018) were used.
148 These specimens were collected with the ROV suction sampler in the Peru Basin and transported
149 to respiratory chambers to measure oxygen consumption of individual holothurian specimen over
150 a period of 72 hours. Aboard RV *Sonne*, the holothurians specimens were measured (length,
151 height, width), shock-frozen intact in liquid nitrogen, and stored at. Feces of holothurians that
152 defecated inside the respiratory chambers were sampled and frozen at -21°C.
153 In the shore-based laboratory at NIOZ-EDS (Yerseke, Netherlands), the samples were freeze-
154 dried and finely-ground with mortar and pestle. The organic (org.) C/ $\delta^{13}\text{C}$ and N/ $\delta^{15}\text{N}$ content
155 of the holothurian tissue and of the acidified holothurian gut content were measured with a
156 Thermo Flash EA 1112 elemental analyzer (EA; Thermo Fisher Scientific, USA) which was
157 coupled to a DELTA V Advantage Isotope Ratio Mass Spectrometer (IRMS; Thermo Fisher
158 Scientific, USA). Stable isotope values are presented in δ notation relative to Vienna Pee Dee
159 Belemnite for $\delta^{13}\text{C}$ and relative to air for $\delta^{15}\text{N}$.
160 Sediment grain size of holothurian gut content was determined by laser diffraction on freeze-
161 dried and sieved (<1 mm) sediment samples in a Malvern Mastersizer 2000.

162 163 **Analysis of amino acids**

164 Total hydrolysable amino acids (THAA) from holothurian tissue were extracted following a
165 modified protocol of Veuger et al. (2005): Briefly, THAAs in holothurian tissue were hydrolyzed
166 by adding 0.01 to 0.02 g freeze-dried finely ground tissue to 1.5 ml 6 M HCl in 10 ml screw-cap
167 vials. A N₂-headspace was created in the vials by flushing with N₂-gas for 10 sec before the vials
168 were closed and heated for 20 h at 110°C. After cooling, 10 μL internal L-Norleucine standard
169 per mg dry faunal tissue (stock solution: 2.5 mg mL⁻¹ L-Norleucin acidified with 100 μL 12 M
170 HCl) was added and the solution was evaporated under N₂-flow at 60°C. THAAs from
171 holothurian tissue were derivatized by adding 0.5 ml acidified propan-2-ol to the sample and by
172 heating the closed vials at 110°C for 90 min. Afterwards, the vials were cooled down and the

173 solution was evaporated under N₂-flow at 50°C. After evaporating all solution, 200 µL
174 dichloromethane (DCM) was added and the solution was evaporated again. When the samples
175 were dry, 150 µL DCM and 50 µL pentafluoropropionic anhydride were added, the vials were
176 closed and heated for 10 min at 110°C. The solvent was extracted by adding 0.5 mL
177 chlorophorm and 1 ml phosphorus-buffer to the sample, shaking it until the lower chloroform
178 fraction was clear and centrifuging the vials with 2,000 rpm for 10 min. The chloroform fraction
179 was transferred to GC vials and evaporated again. When the sample was completely dry, it was
180 dissolved in ethyl acetate. Concentrations (µg C g⁻¹ dry mass DM holothurian tissue) and δ¹³C
181 (‰), and δ¹⁵N (‰) of THAAs were measured with a HP 6890 gas chromatograph (Hewlet
182 Packard/ Agilent, USA) coupled with a DELTA-Plus Isotope Ratio Mass Spectrometer (Thermo
183 Fisher Scientific, USA) on a polar analytical column (ZB5-5MS; 60m length, 0.32mm diameter,
184 0.25µm film thickness; Phenomenex, USA).
185 A list with common abbreviation of amino acids and their full name is presented in Table 2.
186

187 **Table 2.** Names and abbreviations of amino acids and fatty acids (PLFAs, NLFAs).

Abbreviation	Full name
Amino acids	
Ala	Alanine
Asp	Aspartic acid
Glu	Glutamic acid
Gly	Glycine
Ile	Isoleucine
Leu	Leucine
Met	Methionine
Phe	Phenylalanine
Pro	Proline
Ser	Serine
Thr	Threonine
Tyr-Lys	Tyrosine and lysine combined
Val	Valine
Fatty acids	
ARA	Arachidonic acid (C20:4ω6)
DHA	Docosahexaenoic acid (C22:6ω3)
EPA	Eicosapentaenoic acid (C20:5ω3)

188
189 **Analysis of fatty acids**
190 Fatty acids (i.e., PLFAs, NLFAs) were extracted from holothurian tissue, feces and gut content
191 following a modified Bligh and Dyer extraction method (Bligh and Dyer 1959; Boschker 2008).
192 Freeze-dried, homogenized powder of holothurian tissue (~50 – 150 mg), feces and gut content
193 (~150 mg – 2.0 g) were mixed with 6 ml MilliQ-water 15 ml methanol (HPLC grade, 99.8%),
194 and 7.5% chloroform (HPLC grade, 99.5%) in pre-cleaned test tubes. The tubes were shaken for
195 2 h, before 7.5 ml chloroform were added and the tubes were shaken again. 7.5 ml MilliQ-water
196 were added and the tubes were stored at -21°C for 12 h for separation of the solvent layers. The
197 lower solvent layer contained the fatty acids extract dissolved in chloroform and was transferred
198 to pre-weighted test tubes. After determining the weight of the chloroform extract, it was
199 fractionated into the different fatty acid classes over an activated silicic acid column (heated at

200 120°C for 2 h; Merck Kieselgel 60) via eluting with 7 ml chloroform, 7 ml acetone, and 15 ml
 201 methanol. The acetone fraction was discarded, whereas the chloroform fraction containing the
 202 NLFAs and the methanol fraction with the PLFAs were collected in separate test tubes and
 203 evaporated to dryness.
 204 PLFAs and NLFAs were derivatized to fatty acid methyl esters (FAMES) by adding 1 ml
 205 methanol-toluene mix (1:1 volume/ volume), 20 µl of an internal standard (1 mg 19:0
 206 FAME mL⁻¹), and 1 ml 0.2 M methanolic NaOH to the test tubes with the PLFAs and NLFAs
 207 extracts. After an incubation at 37°C for 15 min, 2 ml *n*-hexane, 0.3 ml 1 M acetic acid, and 2 ml
 208 MilliQ-water were added. The solution was mixed very well and when the layers had separated,
 209 the (top) *n*-hexane layer was transferred to new test tubes. Additional 2 ml *n*-hexane were added
 210 to the previously used test tubes that contained the acetic acid-MilliQ-water solution, and the
 211 step was repeated. The *n*-hexane layer was transferred again to the new test tubes and 20 µl of a
 212 second internal standard (1 mg 12:0 FAME mL⁻¹) were added. *n*-hexane was evaporated
 213 completely and the FAMES dissolved in 200 µl *n*-hexane were transferred to measuring vials.
 214 The FAMES from holothurian tissues were separated on a BPX70 column (50 m length, 0.32 mm
 215 inner diameter, 0.25 µm film thickness; SGE Analytical Science) with a HP 6890 gas
 216 chromatograph (GC; Hewlett Packard/ Agilent, USA). The FAMES from feces and gut content
 217 were separated on a ZB5-5MS column (60 m length, 0.32 mm diameter, 0.25 µm film thickness;
 218 Phenomenex, USA) on the same GC. Concentrations (µg C g⁻¹ DM holothurian tissue) and δ¹³C
 219 values (‰) of FAMES in holothurian tissue, feces, and gut content were measured on a Finnigan
 220 Delta Plus isotope ratio mass spectrometer (IRMS; Thermo Fisher Scientific, USA) coupled to
 221 the GC via a combustion GC-c-III interface (Thermo Fisher Scientific, USA). Identification of
 222 peaks of the FAME chromatogram were based on equivalent chain length (ECL) and peak areas
 223 were calculated using the two internal standards (12:0 and 19:0) for area correction.
 224 A list with abbreviations and full names of several important fatty acids is presented in Table 2
 225 and Table 3 contains dominant biomarkers.

226
 227 **Table 3.** Fatty acids used as biomarkers of potential food sources of holothurians from the Peru
 228 Basin.

Fatty acid	Main sources of fatty acid	Reference
Organic matter		
C18:1ω9	Highly degraded carrion-derived organic matter	(Graeve et al. 2001)
Bacteria		
<i>i</i> -C14:0, <i>i</i> -C15:0, <i>ai</i> -C15:0, <i>i</i> -C16:0, and C18:1ω7 <i>cis</i> ;	Marine bacteria; gram-positive bacteria; piezotolerant bacteria	(Findlay et al. 1990; Middelburg et al. 2000; Wang et al. 2014)
C18:2ω6		
10-Me-C16:0, <i>ai</i> -C17:0, <i>i</i> -C17:0, and <i>cy</i> -C17:0	Sulfate-reducing and other anaerobic bacteria	(Findlay et al. 1990)
C16:1ω5	Desulfobacteraceae bacteria	(Elvert et al. 2003)
C16:1ω7	Bacteria in fish intestines	(Yano et al. 1997)
C16:1ω9	Deep-water/ benthic bacteria	(Zhao et al. 2015; Choi et al. 2015)
Primary producers		
C16:4ω1, C16:1ω7, and EPA; $\frac{C16:1\omega7}{C16:0} > 1$ or $\frac{DHA}{EPA} < 1$	Diatoms	(Kelly and Scheibling 2012)

C18:4 ω 3 and DHA; $\frac{C16:1\omega7}{C16:0} < 1$ or $\frac{DHA}{EPA} > 1$	Dinoflagellates	(Kelly and Scheibling 2012)
C18:1 ω 9, C18:4 ω 3, and ARA	Kelp	(Kelly and Scheibling 2012)
C16:0, EPA; ARA	Red macroalgae	(Bühning et al. 2002; Kelly and Scheibling 2012)
C18:1 ω 9, C18:4 ω 3, and ARA; C16:1 ω 5	Brown macroalgae	(Khotimchenko 1995; Kelly and Scheibling 2012)
C18:2 ω 6 and C18:3 ω 3	Seagrass	(Kelly and Scheibling 2012)
Consumers		
C20:1 ω 9, C22:1 ω 11	Calanoid copepods	(Falk-Petersen et al. 1987; Dalsgaard et al. 2003)
EPA, DHA ARA; C22:5 ω 5; $\frac{EPA}{ARA}$ -ratio	Hydrothermal vent bivalves Agglutinated foraminifera	(Ben-Mlih et al. 1992) (Larkin et al. 2014; Kharlamenko 2018)

229

230 **Calculations**

231 **Concentration factors**

232 To examine the degree to which PLFAs were concentrated between surface sediment (0 – 2cm
233 layer; 2.32 \pm 0.51 μ g C-PLFA g⁻¹ DM sediment; Stratmann, unpublished) and gut content and
234 feces, a concentration factor CF was calculated:

$$235 \quad CF_{gut\ content} = \frac{[gut\ content_{PLFA}]}{[sediment_{PLFA}]}, \quad (1)$$

$$236 \quad CF_{feces} = \frac{[feces_{PLFA}]}{[sediment_{PLFA}]}, \quad (2)$$

237 where [gut content_{PLFA}] corresponds to the total PLFA concentration in gut content, [feces_{PLFA}]
238 to the total PLFA concentration in feces, and [sediment_{PLFA}] to the average total PLFA
239 concentration in surface sediment.

240

241 **Trophic levels**

242 Trophic levels (TL) of holothurian species were calculated following (Chikaraishi et al. 2009) as

$$243 \quad TL = \frac{(\delta^{15}N_{Glu} - \delta^{15}N_{Phe} - 3.4)}{7.6} + 1. \quad (3)$$

244 $\delta^{15}N_{Glu}$ is the $\delta^{15}N$ of the amino acid glutamic acid (‰) and $\delta^{15}N_{Phe}$ corresponds to the $\delta^{15}N$ of
245 the amino acid phenylalanine (‰).

246 Trophic levels of two different holothurian species are considered robust, when the difference in
247 trophic levels between two species is \geq 0.44. This value corresponds to the average standard
248 deviation of the calculated trophic level (σ_{TL}) across all holothurians that was determined
249 following equation S4 in (Jarman et al. 2017) as:

$$\sigma_{TP} = \sqrt{\left(\frac{1}{\Delta_{Glu-Phe}}\right)^2 \sigma_{\delta^{15}N_{Glu}}^2 + \left(\frac{-1}{\Delta_{Glu-Phe}}\right)^2 \sigma_{\delta^{15}N_{Phe}}^2 + \left(\frac{1}{\Delta_{Glu-Phe}}\right)^2 \sigma_{\beta}^2 + \left(\frac{-1}{\Delta^2_{Glu-Phe}} (\delta^{15}N_{Glu} - \delta^{15}N_{Phe} + \beta)\right)^2 \sigma_{\Delta_{Glu-Phe}}^2}, \quad (4)$$

251 where σ_{β} is 0.9‰ and σ_{Δ} is 1.1‰ (Jarman et al. 2017).

252

253 Heterotrophic re-synthesis of amino acids

254 Total heterotrophic re-synthesis of amino acids ($\sum V$) was approximated as the sum of variance
 255 of individual $\delta^{15}N$ values of the trophic amino acids alanine, aspartic acid, glutamic acid, leucine,
 256 and proline (McCarthy et al. 2007):

$$257 \sum V = \sum_1^n |x_{amino\ acid} - \bar{x}_{amino\ acid}|$$

258 (5)

259 x symbolized each trophic amino acid's $\delta^{15}N$ value, \bar{x} is the average trophic amino acid's $\delta^{15}N$
 260 value and n is the total number of trophic amino acids used in this calculation (McCarthy et al.
 261 2007).

262

263 Statistics

264 The Sørensen–Dice coefficient β_{sor} (Dice 1945; Sørensen 1948; Koleff et al. 2003) was
 265 calculated using the ‘betadiver’ function in the *R* (version 4.1.2; R-Core Team 2017) package
 266 *vegan* (version 2.6-2; (Oksanen et al. 2017)) to compare holothurian species based on their
 267 trophic levels (*TL*), levels of heterotrophic re-synthesis of amino acids ($\sum V$), feeding selectivity
 268 based on concentration factor (*CF*), and food sources/ diet. For this purpose, the quantitative data
 269 *TL* and $\sum V$ were first converted into categories (Table 4) and then converted into binary
 270 (presence/ absence) data; the categorical data ‘feeding selectivity’ (Table 4) and ‘food/sources
 271 diet’ were also converted into binary data. Subsequently, β_{sor} was clustered by average linkage
 272 clustering (unweighted pair-group method using arithmetic averages, UPGMA; Romesburg
 273 1984) using the ‘hclust’ function in *R*. The dendrogram was prepared with *R* package *factoextra*
 274 (version 1.0.7; Kassambara and Mundt 2020).

275

276 **Table 4.** Parameters used to calculate the Sørensen–Dice coefficient β_{sor} presented as
 277 quantitative data (ranges) and categorical data. ‘Food sources/ diet’ includes a list of the main
 278 food sources of the investigated holothurian species which were identified by amino acid and
 279 fatty acid analysis.

Parameter	Quantitative data	Categorical data
Trophic level (<i>TL</i>)	$TL = 2.0-2.5$	$TL_{group\ 1}$
	$TL = >2.5-3.0$	$TL_{group\ 2}$
	$TL = >3.0-3.5$	$TL_{group\ 3}$
Levels of heterotrophic re-synthesis of amino acids ($\sum V$)	$\sum V = 0-1.5$	$\sum V_{group\ 1}$
	$\sum V = >1.5-3.0$	$\sum V_{group\ 2}$
	$\sum V = >3.0-4.5$	$\sum V_{group\ 3}$
	$\sum V = >4.5-6.0$	$\sum V_{group\ 4}$
	$\sum V = >6.0-7.5$	$\sum V_{group\ 5}$
	$\sum V = >7.5-9.0$	$\sum V_{group\ 6}$
Feeding selectivity based on concentration	$CF = 0-10$	no selectivity
	$CF = >10-50$	selective

factor <i>CF</i>	<i>CF</i> = >50–150 <i>CF</i> = >150	very selective extremely selective
Food sources/ diet		<ul style="list-style-type: none">– diatom-derived phytodetritus– dinoflagellate-derived phytodetritus– detritus containing C22:1ω9-<i>cis</i>– secondary consumer of detritus– mix diet (phytodetritus primary consumer, detritus secondary consumer)– mixed diet (phytodetritus, bacteria)– mixed diet (phytodetritus, bacteria, Foraminifera)

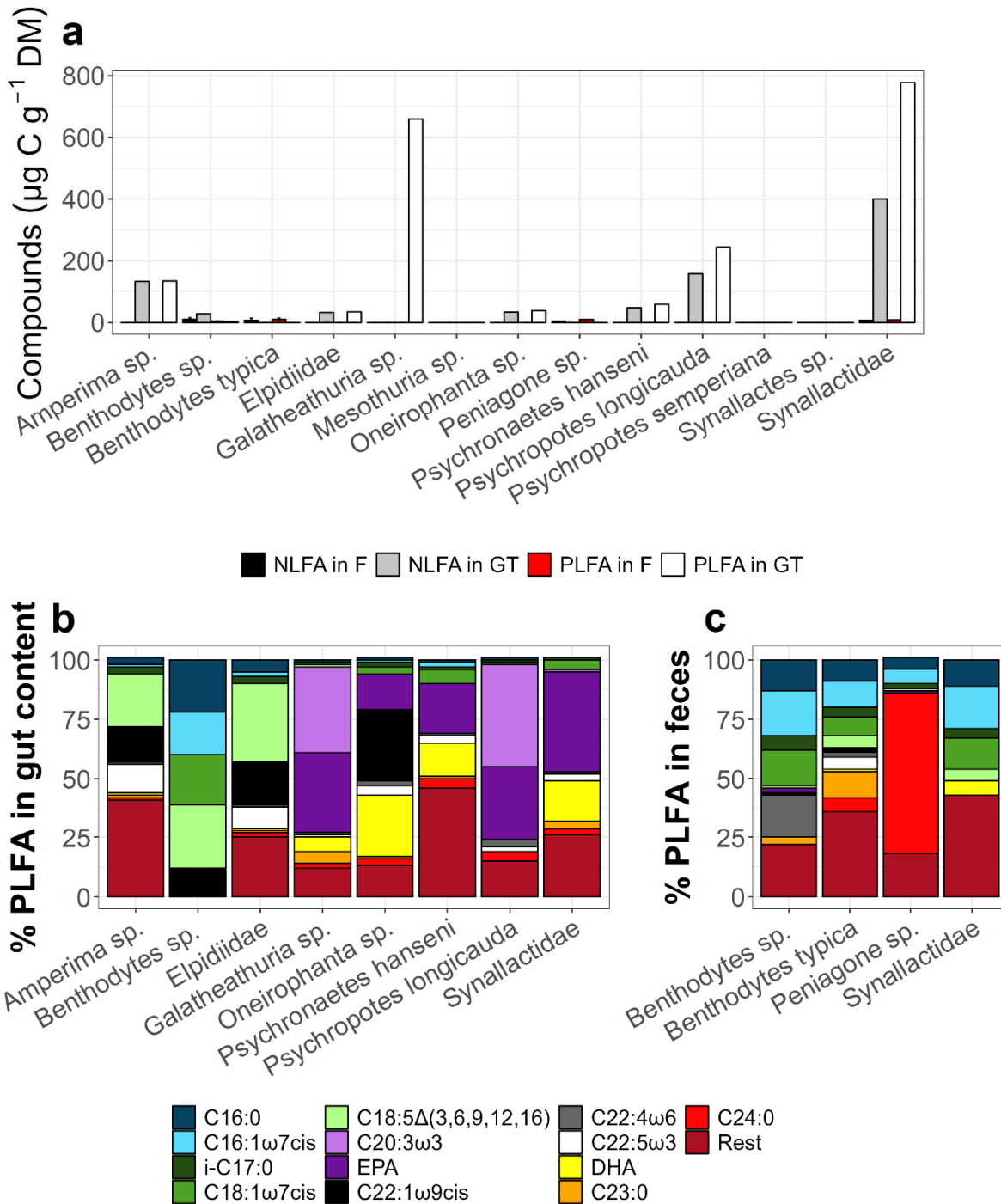
280

281 **Results**

282 **Gut content and feces of holothurians**

283 Gut contents of holothurians in the Peru Basin weighed 1.93 ± 3.56 g dry sediment ($n = 17$) and
284 ranged from 0.11 g dry sediment for *Peniagone* sp. ($n = 1$) to 12.5 g dry sediment for *P. hanseni*
285 ($n = 1$; Table 5). Org. C and TN content of the gut content was $5.34 \pm 4.13\%$ ($n = 17$) and
286 $1.04 \pm 0.87\%$ ($n = 17$), respectively, and it contained 244 ± 304 $\mu\text{g C-PLFA g}^{-1}$ DM gut content
287 ($n = 8$) and 83.3 ± 124 $\mu\text{g C-NLFA g}^{-1}$ DM gut content ($n = 10$) (Fig. 1a). The concentration
288 factor *CF* for PLFAs in holothurian gut content was on average 105 ± 131 ($n = 8$) and ranged
289 from 1.17 to 335 for *Benthodytes* sp. ($n = 1$) and Synallactidae gen sp. ($n = 1$), respectively
290 (Table 5). The average EPA/ARA-ratio for gut content was 3.20 ± 5.58 ($n = 9$), the average
291 DHA/EPA-ratio was 0.62 ± 0.68 ($n = 9$), and the average C16:1 ω 7/C16:0-ratio was 0.81 ± 0.85
292 ($n = 9$) (Fig. 2a).

293



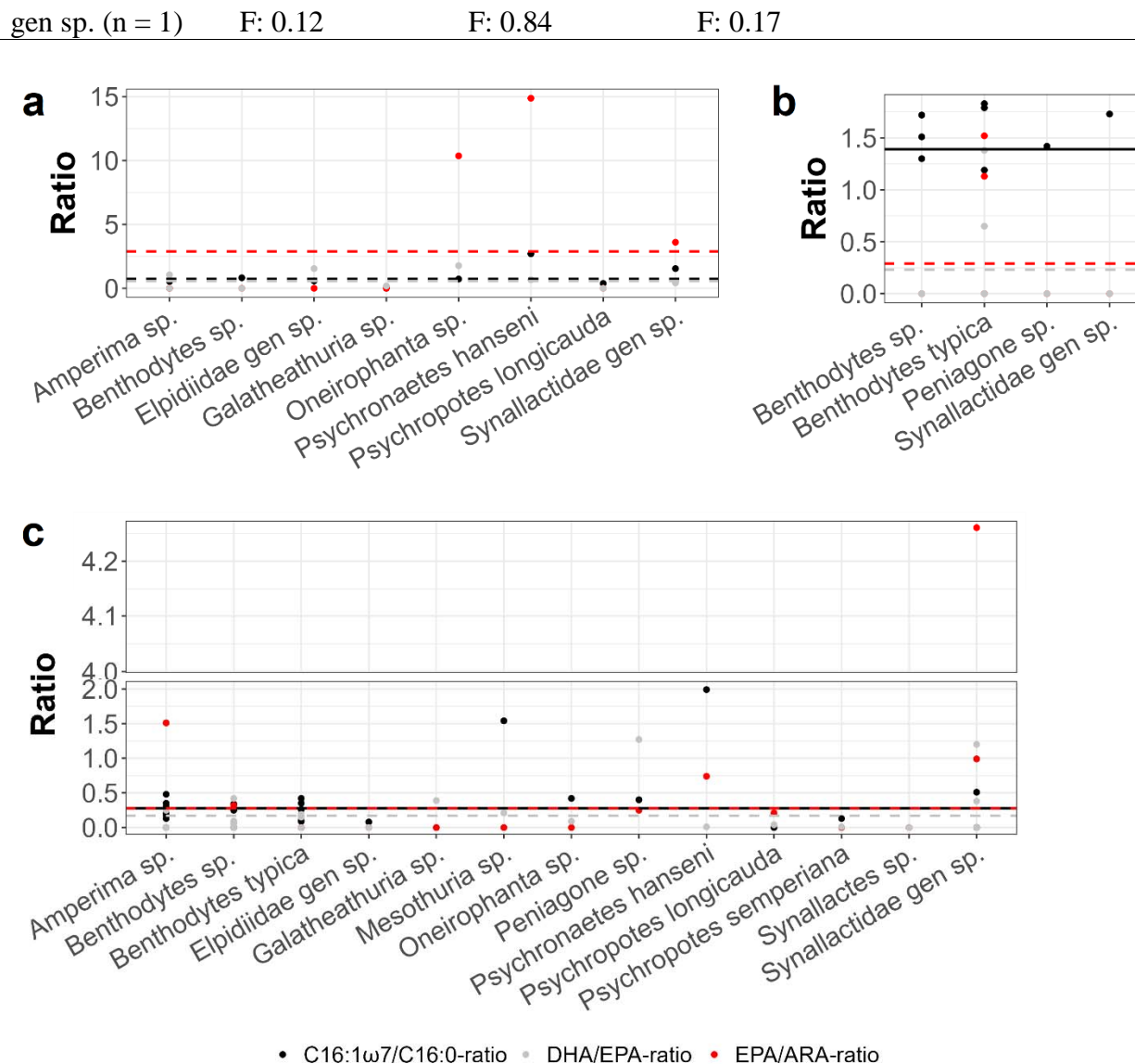
294
295 **Figure 1.** (a) Concentrations ($\mu\text{g C g}^{-1} \text{ DM. C}$) of the compounds PLFA and NLFA in
296 holothurian gut content (GT) and feces (F) and the contribution (%) of individual (b, c) PLFAs
297 and (d, e) NLFAs to the total concentrations. The PLFA and NLFA pools ‘Rest’ include all
298 PLFAs and NLFAs, respectively, that contribute $<2.5\%$ to total % PLFA and NLFA of the
299 average holothurian gut content/ feces.
300 Error bars in (a) indicate SD.
301

302
303
304

Table 5. Sedimentological characteristics of holothurian gut content (GT) and feces (F). Data are presented as mean±SD.

Species	GT/ F (g dry sediment)	GT/ F composition		Concentration factor CF_{gut} content, CF_{feces}
		% org. C;	% TN	
Family: Elpidiidae				
<i>Amperima</i> sp. (GT: n = 5; F: n = 1)	GT: 0.16±0.1; F: 0.54	GT: 7.76±3.85 F: 0.32	GT: 1.60±0.71 F: 0.05	GT: 58.0
Elpidiidae gen sp. (GT: n = 1)	0.88	1.57	0.35	14.9
<i>Peniagone</i> sp. (GT: n = 1; F: n = 1)	GT: 0.11; F: 0.45	GT: 3.27; F: 0.84	GT: 0.66; F: 0.13	F: 4.36
Family: Deimatidae				
<i>Oneirophanta</i> sp. (GT: n = 1)	3.90	2.84	0.51	16.6
Family: Laetmogonidae				
<i>Psychronaetes hanseni</i> (GT: n = 1)	12.5	1.94	0.38	25.6
Family: Psychropotidae				
<i>Benthodytes</i> sp. (GT: n = 2; F: n = 3; sediment grain size: n = 1)	GT: 2.07±2.1 F: 3.01±3.61	GT: 0.95±0.02 F: 0.70±0.16	GT: 0.16±0.01 F: 0.14±0.05	GT: 1.17 F: 1.81±0.53
<i>Benthodytes typica</i> (F: n = 4; sediment grain size: n = 1)	0.85±0.62	1.05±0.47	0.16±0.01	4.44±1.52
<i>Psychropotes longicauda</i> (GT: n = 1)	1.08	1.38	0.29	105
<i>Psychropotes semperiana</i> (GT: n = 1)	1.78	5.25	0.80	
Family: Synallactidae				
<i>Synallactes</i> sp. (morphotype "pink") (GT: n = 1)	6.75	2.22	0.34	
<i>Galatheathuria</i> sp. (GT: n = 1)	0.51	11.4	2.10	285
Synallactidae	GT: 0.14	GT: 11.3	GT: 2.44	GT: 335

305



306

307 **Figure 2.** Ratios of C16:1 ω 7/C16:0, DHA/EPA, and EPA/ARA in (a) holothurian gut content,
 308 (b) holothurian feces, and (c) dried holothurian body walls. Horizontal lines show the average
 309 value of a ratio based on all samples.

310

311 Feces of holothurians weighed 1.36 ± 2.09 g dry sediment (n = 10) and ranged from 0.12 g dry
 312 sediment for *Synallactidae* gen sp. (n = 1) to 3.01 ± 3.61 g dry sediment for *Benthodytes* sp.
 313 (n = 3; Table 5). Org. C and TN content of the feces was $0.83 \pm 0.37\%$ (n = 10) and $0.14 \pm 0.04\%$
 314 (n = 10), respectively, and it contained 7.73 ± 3.60 $\mu\text{g C-PLFA g}^{-1}$ DM sediment (n = 8) and
 315 7.63 ± 5.28 $\mu\text{g C-NLFA g}^{-1}$ DM sediment (n = 15) (Fig. 1a). In holothurian feces (n = 8), PLFAs
 316 were on average still 3.33 ± 1.55 times more concentrated compared to the upper 2 cm of
 317 sediment (CF_{PLFA} range: 1.81 ± 0.53 (n = 3) for *Benthodytes* sp. to 4.44 ± 1.52 for *B. typica* (n = 3))
 318 (Table 5). The average EPA/ARA-ratio for feces was 0.29 ± 0.59 (n = 9), the average DHA/EPA-
 319 ratio was 0.23 ± 0.48 (n = 9), and the average C16:1 ω 7/C16:0-ratio was 1.39 ± 0.57 (n = 9)
 320 (Fig. 2b).

321 Gut content and feces consisted to $81.2 \pm 3.73\%$ of silt (grain size: $<63 \mu\text{m}$; $n = 6$) and to
 322 $10.7 \pm 1.10\%$ of very fine sand (grain size: $62.5 - 125 \mu\text{m}$) (Table 6). The median grain was
 323 $15.5 \pm 2.27 \mu\text{m}$ ($n = 6$).

324

325 **Table 6.** Grain size characteristics of gut content (GT) and feces (FC).

Species	% silt fraction ($<63 \mu\text{m}$)	% very fine sand fraction ($62.5 -$ $125 \mu\text{m}$)	% fine sand fraction ($125 -$ $250 \mu\text{m}$)	% medium sand fraction ($250 -$ $500 \mu\text{m}$)	% coarse sand fraction ($500 -$ $1000 \mu\text{m}$)	Median grain size (μm)
Family: Elpidiidae						
Elpidiidae gen sp. (GT; $n = 1$)	82.3	10.0	4.91	2.05	0.92	14.7
Family: Deimatidae						
<i>Oneirophanta</i> sp. (GT; $n = 1$)	75.5	12.2	7.86	4.01	0.65	19.0
Family: Laetmogonidae						
<i>Psychronaetes</i> <i>hanseni</i> (GT; $n = 1$)	79.0	11.7	6.67	2.48	0.33	17.5
Family: Psychropotidae						
<i>Benthodytes</i> sp. (F; $n = 1$)	86.4	9.20	4.05	0.44	0.00	13.0
<i>Benthodytes</i> <i>typica</i> (F; $n = 1$)	81.0	10.8	6.08	1.77	0.55	14.2
Family: Synallactidae						
Synallactidae gen sp. (F; $n = 1$)	83.0	10.4	4.47	1.58	0.76	14.5

326

327 About $22.3 \pm 15.5\%$ ($n = 8$) of the PLFAs (Fig. 4b) and $23.1 \pm 18.4\%$ ($n = 7$) of the NLFAs
 328 (Fig. 4d) found in holothurian gut content consisted of 'Rest', i.e., the sum of PLFAs and NLFAs
 329 that each contributed $<2.5\%$ to total PLFA and NLFA concentrations. The remaining PLFAs
 330 consisted to $7.00 \pm 6.55\%$ of saturated fatty acids (SFA), $17.0 \pm 17.7\%$ monosaturated fatty acids
 331 (MUFAs, i.e., fatty acids with one double bond), $10.0 \pm 18.4\%$ polyunsaturated fatty acids
 332 (PUFAs, i.e., fatty acids with ≥ 2 double bonds), $41.5 \pm 10.6\%$ highly unsaturated fatty acids
 333 (HUFAs, i.e., fatty acids with ≥ 4 double bonds), and $2.25 \pm 1.36\%$ long-chain fatty acids
 334 (LCFAs, i.e., fatty acids with ≥ 24 C atoms). NLFAs included furthermore $36.9 \pm 11.7\%$ SFAs,
 335 $13.0 \pm 11.0\%$ MUFAs, and $27.0 \pm 17.3\%$ HUFAs. Feces of holothurian consisted to $29.6 \pm 13.3\%$
 336 ($n = 8$) of the PLFAs category 'Rest' (Fig. 4c) and to $31.4 \pm 15.6\%$ ($n = 8$) of the NLFAs category
 337 'Rest' (Fig. 4e). The other PLFAs consisted to $20.1 \pm 9.58\%$ of SFAs, $25.6 \pm 11.7\%$ MUFAs,
 338 $14.1 \pm 12.4\%$ HUFAs, and $10.6 \pm 23.7\%$ LCFAs. The NLFAs included additionally $39.5 \pm 11.1\%$
 339 SFAs, $10.3 \pm 10.5\%$ MUFAs, and $18.8 \pm 24.3\%$ HUFAs.

340

341 **Chemical composition of holothurians**

342 Holothurians in the Peru Basin consisted for $93.0 \pm 10.2\%$ of water ($n = 13$) and their dried body
 343 walls contained $5.87 \pm 3.50\%$ org. C and $1.35 \pm 0.80\%$ total N ($n = 31$), whereas their dried gut
 344 tissues consisted of $16.7 \pm 8.60\%$ org. C and $3.76 \pm 2.18\%$ total N ($n = 15$). The body wall and gut
 345 tissue of the holothurian families Deimatidae and Laetmogonidae had the highest org. C and TN
 346 contents, whereas the families Elpidiidae and Psychropotidae had the lowest org. C and TN
 347 content in body wall tissue (Table 7).

348

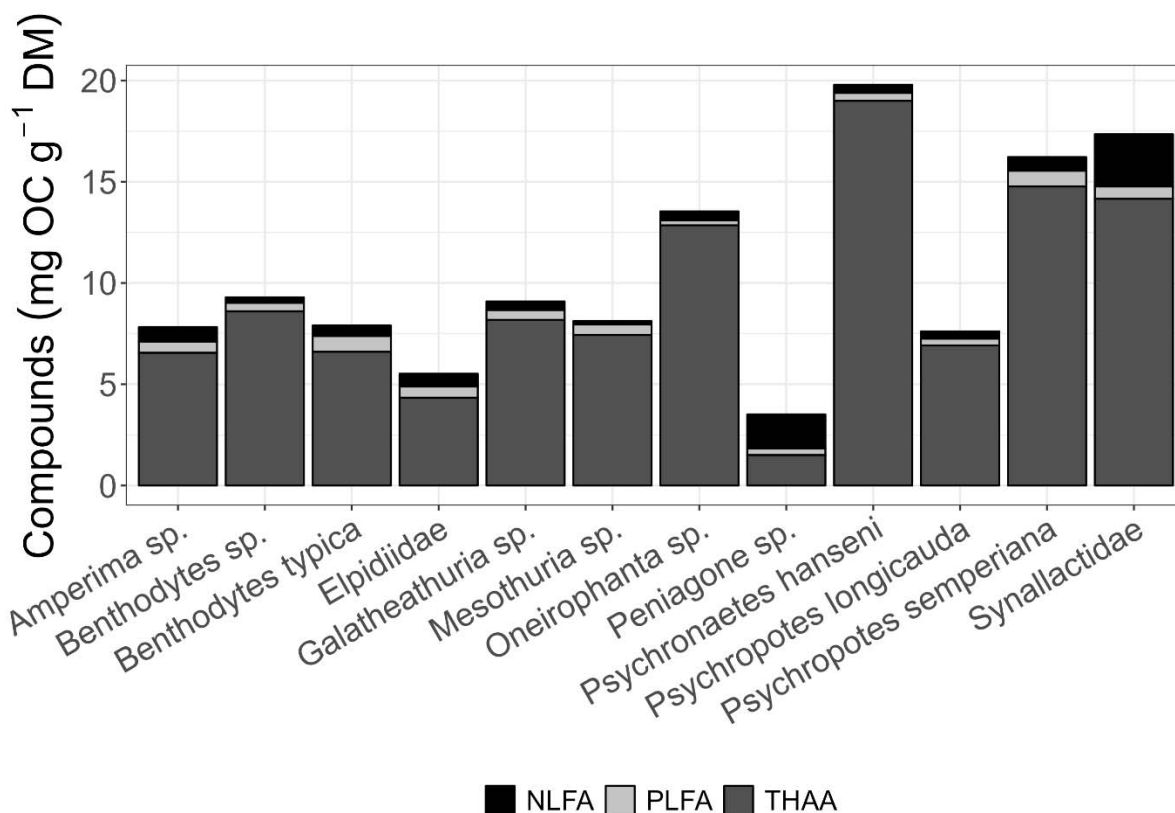
349 **Table 7.** Chemical composition of body wall (BW) and gut tissue (GT) of different holothurian
 350 species collected in the Peru Basin. Data are presented as mean \pm SD.

Species	Body wall (BW)		Gut (G)	
	% org. C	% TN	% org. C	% TN
Family: Elpidiidae (BW: $n = 10$; GT: $n = 4$)	4.48 ± 3.25	1.07 ± 0.86	8.02 ± 2.15	1.63 ± 0.48
<i>Amperima</i> sp. (BW: $n = 9$; GT: $n = 4$)	4.62 ± 3.30	1.10 ± 0.88	8.02 ± 2.15	1.63 ± 0.48
Elpidiidae gen sp. ($n = 1$)	6.94	1.73		
<i>Peniagone</i> sp. ($n = 1$)	0.93	0.17		
Family: Deimatidae ($n = 1$)	16.0	3.25	26.4	6.27
<i>Oneirophanta</i> sp. ($n = 1$)	16.0	3.25	26.4	6.27
Family: Laetmogonidae ($n = 1$)	11.7	2.46	20.6	4.42
<i>Psychronaetes hansenii</i> ($n = 1$)	11.7	2.46	20.6	4.42
Family: Mesothuriidae ($n = 1$)	5.42	1.06		
<i>Mesothuria</i> sp. ($n = 1$)	5.42	1.06		
Family: Psychropotidae (BW: $n = 14$; GT: $n = 4$)	5.19 ± 2.53	1.21 ± 0.60	19.3 ± 7.43	4.15 ± 1.78
<i>Benthodytes</i> sp. ($n = 6$)	4.53 ± 1.54	1.08 ± 0.42		
<i>Benthodytes typica</i> (BW: $n = 5$; GT: $n = 1$)	4.19 ± 1.64	0.95 ± 0.51	28.9 ($n = 1$)	6.55
<i>Psychropotes longicauda</i> ($n = 1$)	4.39	1.15	16.7	3.11
<i>Psychropotes semperiana</i> ($n = 1$)	8.41	2.09	11.2	2.53
Family: Synallactidae (BW: $n = 5$; GT: $n = 4$)	8.60 ± 2.37	1.97 ± 0.53	15.6 ± 8.23	3.61 ± 2.15
<i>Synallactes</i> sp. (morphotype "pink") ($n = 1$)	6.63	1.42	10.5	2.07
<i>Galatheathuria</i> sp. ($n = 1$)	6.98	2.01	8.98	2.23
Synallactidae gen sp. (BW: $n = 3$; GT: $n = 2$)	9.80 ± 2.41	2.14 ± 0.60	21.4 ± 8.15	5.08 ± 2.29

351

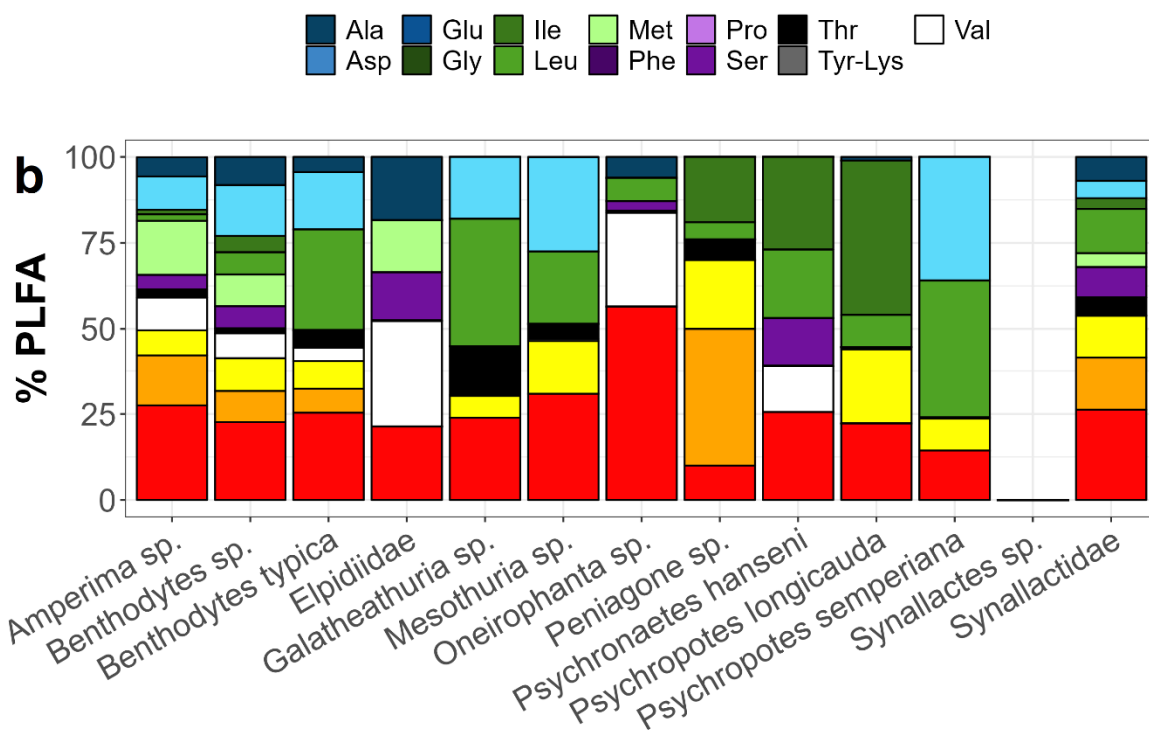
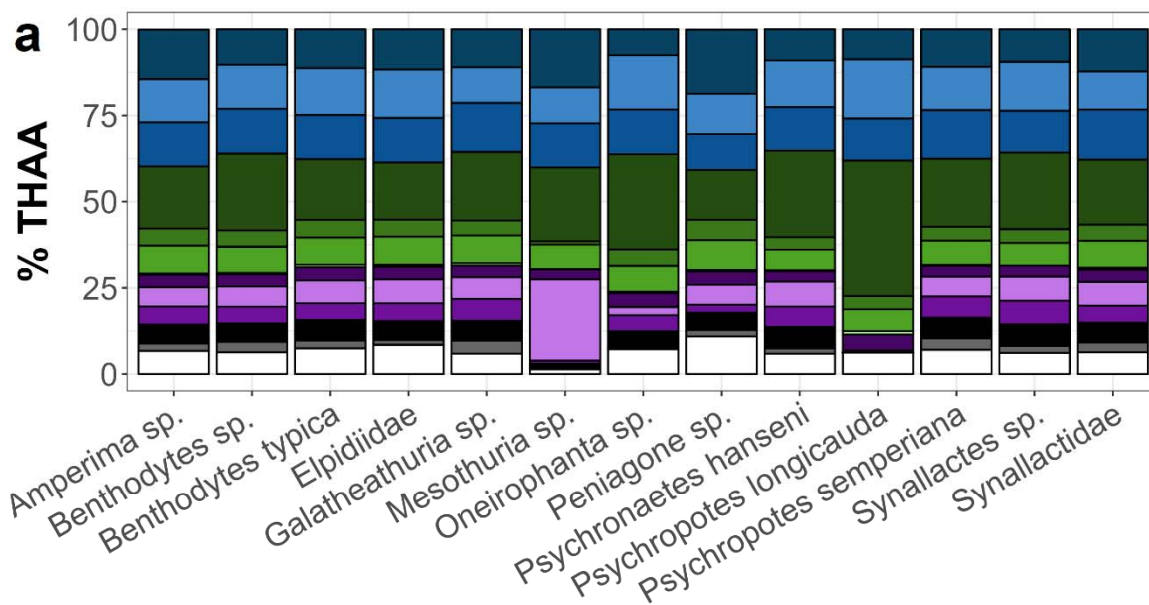
352 THAAs, PLFAs, and NLFAs contributed a total $17.4 \pm 6.11\%$ to the org. C of all specimens
 353 combined ($n = 27$) and ranged from $1.50 \text{ mg C g}^{-1} \text{ DM THAAs}$ (*Peniagone* sp.; $n = 1$) to
 354 $19.0 \text{ mg C g}^{-1} \text{ DM THAA}$ (*P. hansenii*; $n = 1$), $0.24 \text{ mg C g}^{-1} \text{ DM PLFAs}$ (*Oneirophanta* sp.;
 355 $n = 1$) to $0.78 \pm 0.29 \text{ mg C g}^{-1} \text{ DM PLFAs}$ (*B. typica*; $n = 5$), and $0.17 \text{ mg C g}^{-1} \text{ DM NLFAs}$
 356 (*Mesothuria* sp.; $n = 1$) to $2.58 \pm 3.32 \text{ mg C g}^{-1} \text{ DM NLFAs}$ (Synallactidae gen sp.; $n = 3$) (Fig. 3).

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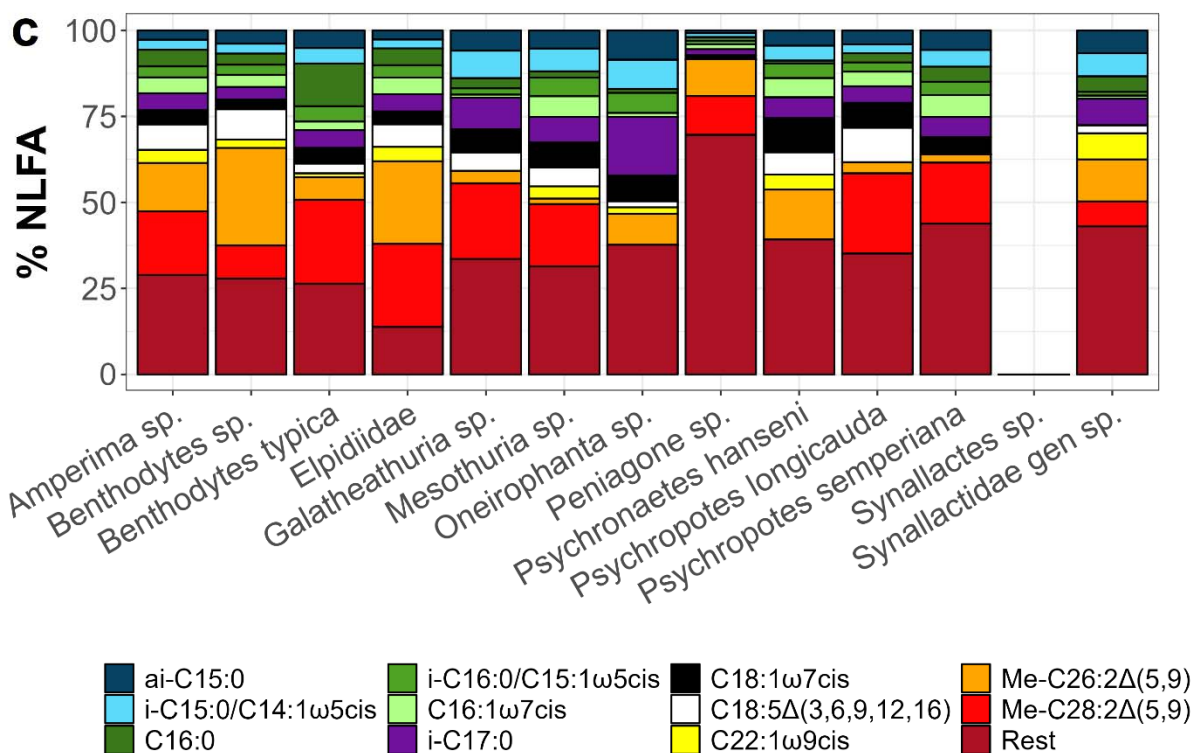


358
 359 **Figure 3.** The compounds neutral lipid-derived fatty acids (NLFA), phospholipid-derived fatty
 360 acids (PLFA), and total hydrolysable amino acids (THAA) in dried holothurian body walls.
 361
 362 The THAA composition did not differ greatly among species, with the main THAAs in
 363 holothurian body walls being alanine, glycine, aspartic acid, and glutamic acid, that contributed
 364 between 55.2% (Elpidiidae; n = 1) and 77.4% (*P. longicauda*; n = 1) to total THAAs of body
 365 walls (Fig. 4a).
 366 In contrast, the PLFA (Fig. 4b) and NLFA (Fig. 4c) composition differed strongly between
 367 species. Between 10.2% (*Peniagone* sp.; n = 1) and 56.5% (*Oneirophanta* sp.; n = 1) of the
 368 PLFAs found in holothurian body walls contributed <2.5% to the total PLFA concentration and
 369 were combined as ‘Rest’ (Fig. 4b). The remaining PLFAs consisted to 4.07±7.57% (n = 32) of
 370 MUFAs, 10.8±12.6% PUFAs, 32.2±14.1% HUFAs, 9.17±14.0% LCFAs, and 15.5±10.1%
 371 methyl-fatty acids. Compared to the average PLFA composition across all holothurian taxa
 372 analyzed, Elpidiidae gen sp. (n = 1) had an above average percentage of MUFAs (12.8% of total
 373 PLFAs) and methyl-fatty acids (48.1% of total PLFAs). *Oneirophanta* sp. (n = 1) had an above
 374 average percentage of SFAs (41.1% of total PLFAs) and *P. longicauda* (n = 1) had an above
 375 average percentage of HUFAs (10.0% of total PLFAs).
 376 The NLFAs consisted for 21.6±10.4% (n = 30) of SFA, 10.1±5.80% MUFAs, 5.36±5.03%
 377 HUFAs, and 28.5±18.4% methyl-fatty acids (Fig. 4c). Between 13.8% Elpidiidae gen sp. (n = 1)
 378 and 69.6% (*Peniagone* sp.; n = 1) of the total NLFAs in holothurian body walls consisted of
 379 NLFAs that individually contributed <2.5% to the total NLFA concentration and were therefore
 380 combined as ‘Rest’. In comparison to the average NLFA composition across all studied

381 holothurian taxa, *Oneirophanta* sp. (n = 1) had an above average percentage of SFA (41.1% of
382 total NLFAs). *P. hanseni* (n = 1) had an above average percentage of MUFAs (19.9% of total
383 NLFAs), *P. longicauda* had an above average percentage of HUFAs (10.0% of total NLFAs),
384 and Elpidiidae gen sp. had an above average percentage of methyl-fatty acids (48.1%).
385



386



387
388 **Figure 4.** Contribution (%) of individual (a) THAAs, (b) PLFAs, and (c) NLFAs to the total
389 concentrations. The PLFA and NLFA pools ‘Rest’ include all PLFAs and NLFAs, respectively,
390 that contribute <2.5% to total % PLFA and NLFA of the average holothurian tissue.

391
392 The ratio of the essential phospholipid-derived PUFAs EPA to ARA, i.e., the EPA/ARA-ratio,
393 ranged from 0.05 ± 0.13 for *Benthodytes* sp. (n = 6) to 1.75 ± 2.23 for Synallactidae gen sp. (n = 3)
394 (Fig. 2c). In comparison, the ratio of DHA to EPA, i.e., the DHA/EPA-ratio, ranged from 0.01
395 for *P. hansenii* (n = 1) to 1.27 for *Peniagone* sp. (n = 1) (Fig. 2c). Due to the absence of the
396 PUFAs ARA and/ or EPA in holothurian body wall tissue, no EPA/ARA-ratios were calculated
397 for *B. typica*, *Elpidiidae* gen sp., *Galatheathuria* sp., *Mesothuria* sp., *Oneirophanta* sp., and *P.*
398 *semperiana*. *Elpidiidae* gen sp. lacked both, DHA and EPA, and therefore no DHA/EPA-ratio
399 could be calculated (Fig. 2c).

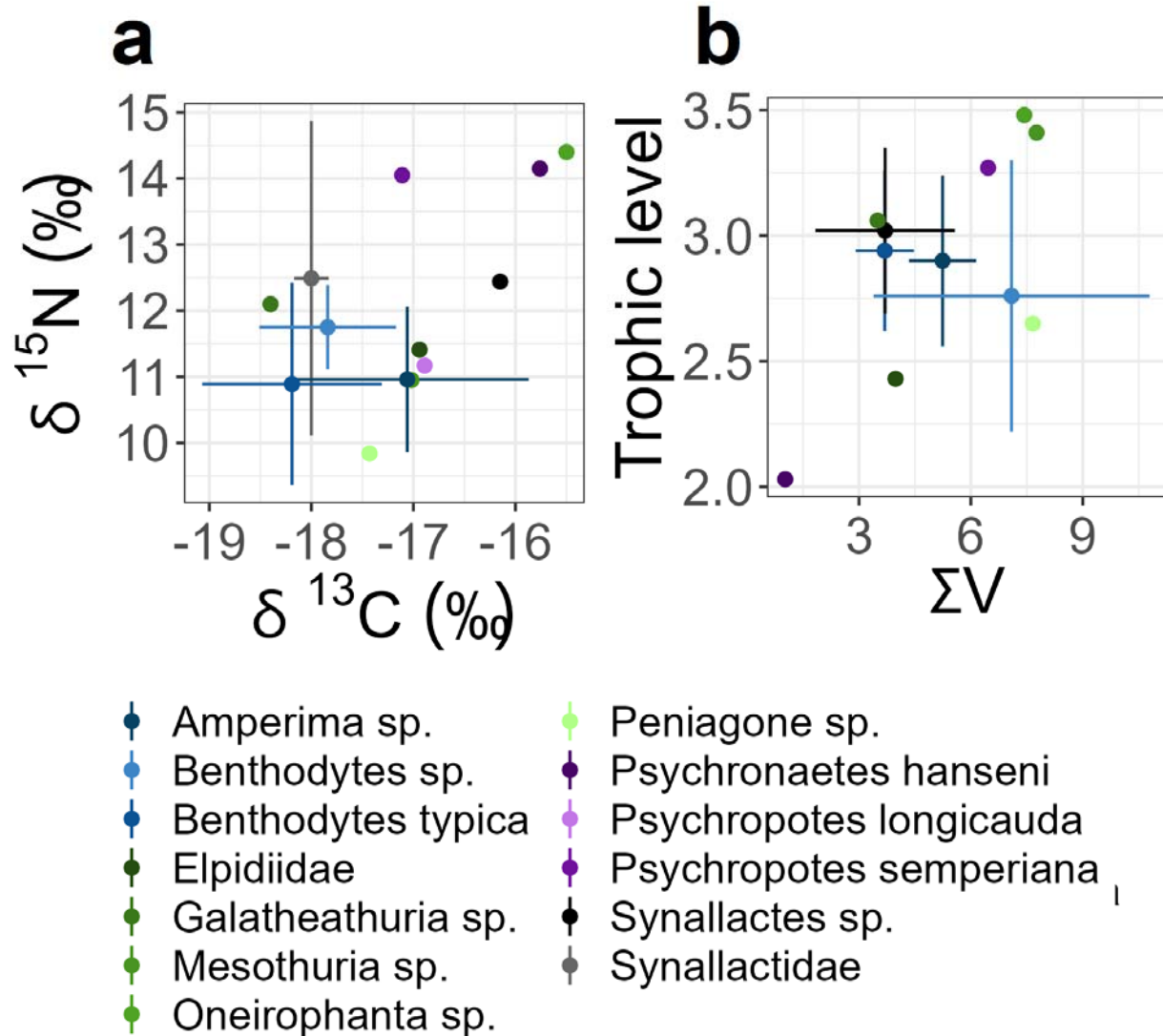
400 401 **Trophic position of holothurians and recycling of amino acids**

402 Holothurians in the Peru Basin had an average $\delta^{13}\text{C}$ -value of $-17.4 \pm 1.02\text{‰}$ (n = 31) with a
403 minimum value of -18.4‰ for *Galatheathuria* sp. (n = 1) and a maximum value of -15.5‰ for
404 *Oneirophanta* sp. (n = 1). The average $\delta^{15}\text{N}$ -value was $11.6 \pm 1.47\text{‰}$ (n = 31) with a minimum
405 value of 9.84‰ for *Peniagone* sp. (n = 1) and a maximum value of 14.4‰ for *Oneirophanta* sp.
406 (n = 1) (Fig. 5a).

407 TL estimates for holothurians in the Peru Basin, based on the $\delta^{15}\text{N}$ values of the THAA glutamic
408 acid and alanine, ranged from 2.0 (*P. hansenii*, n = 1) to 3.5 (*Oneirophanta* sp., n = 1) (Fig. 5b).

409 values ranged from 1.02 to 7.76, with *P. hansenii* having the lowest heterotrophic enrichment
410 and *Mesothuria* sp. having the highest heterotrophic enrichment (Fig. 5b).

411



412
 413 **Figure 5.** (a) Isotopic composition of carbon ($\delta^{13}\text{C}$, ‰) and nitrogen ($\delta^{15}\text{N}$, ‰) of holothurian
 414 body wall tissue from the Peru Basin. (b) Trophic position and heterotrophic enrichment factor
 415 of holothurians. Error bars indicate SD.

416
 417 **Discussion**

418 **Fatty acid composition of holothurians**

419 Deep-sea megabenthic invertebrates consist to 4.5% DM (cnidarians) to 44.9% DM
 420 (crustaceans) of lipids (Drazen et al. 2008a), whereupon holothurians have lipid contents of <1%
 421 DM to 5.8% DM (Drazen et al. 2008b). The largest lipid fraction is phospholipids with 14.5%
 422 total lipids (crustaceans, Drazen et al. 2008a) to 95.2% total lipids (holothurians; Drazen et al.
 423 2008b). The neutral lipids wax esters and triacylglycerol contribute between <1% (polychaetes)
 424 and 83% (crustaceans) to total lipids (Drazen et al. 2008a) and also holothurians consist only of
 425 <1% to 2.6% total lipids wax esters and triacylglycerol (Drazen et al. 2008c). Holothurians from
 426 the Peru Basin contain between 0.16% DM (*Oneirophanta* sp.) and 2.30% DM (*B. typica*)
 427 PLFAs, components of phospholipids, and 0.31% DM (*Oneirophanta* sp.) to 4.71% DM

428 (*Peniagone* sp.) NLFAs, elements of neutral lipids. Hence, they have a relatively high neutral
429 fatty acid content compared to holothurians from Station M (NE Pacific) (Drazen et al. 2008b).
430 This might be related to differences in food availability at the two study sites: The abyssal
431 seafloor at Station M receives on average $22.3 \text{ g C m}^{-2} \text{ yr}^{-1}$ particulate organic carbon (POC)
432 (Baldwin et al. 1998), whereas the POC flux to the Peru Basin is estimated to be $1.49 \text{ g C m}^{-2} \text{ yr}^{-1}$
433 (Haeckel et al. 2001). As a result, holothurians from the Peru Basin might be adapted to a more
434 food-limited environment by building higher concentrations of storage lipids when they
435 encounter fresh phytodetritus than holothurians at Station M.

436

437 **Holothurians trophic level and inferred feeding strategy**

438 Based on the $\delta^{15}\text{N}$ value of body wall tissue, Iken et al., (2001) identified three trophic groups
439 among holothurians from PAP: Group A had $\delta^{15}\text{N}$ values from 10.8 to 12.3‰, group B's $\delta^{15}\text{N}$
440 values ranged from 13.2 to 13.9‰, and group C had $\delta^{15}\text{N}$ values from 15.6 to 16.2‰. The $\delta^{15}\text{N}$
441 values of holothurian tissue from the Peru Basin investigated in this study were lower and ranged
442 from 9.84‰ for *Peniagone* sp. (n = 1) to 14.4‰ for *Oneirophanta* sp. (n = 1). Instead of basing
443 our classification of holothurians from the Peru Basin solely on $\delta^{15}\text{N}$ values, we combined data
444 of trophic level based on compound-specific stable isotope analysis with biomarkers, grain size
445 analysis, and concentration factors for PLFAs.

446

447 **Order Elasipodida**

448 *Psychronaetes hansenii* is a deposit feeder of the **family Laetmogonidae**, which has a trophic
449 level of 2.0, low level of heterotrophic re-synthesis of amino acids ($\sum V = 1.02$) and feeds
450 selectively ($CF_{\text{gut content}} = 25.6$) on sedimentary detritus particles of a medium grain size of
451 $17.5 \mu\text{m}$ which is smaller than the medium grain size of the upper 5 cm of sediment
452 ($20.8 \pm 0.3 \mu\text{m}$; Mevenkamp et al., 2019). Based on the biomarkers present in the body wall tissue
453 of the specimen analysed and in its gut content, parts of the sedimentary detritus likely consists
454 of diatom-derived phytodetritus ($\frac{DHA}{EPA}\text{-ratio}_{\text{body wall}} = 0.01$, $\frac{DHA}{EPA}\text{-ratio}_{\text{gut content}} = 0.67$).

455

456 Elpidiidae gen sp. (**family Elpidiidae**) has a trophic level of 2.4 and medium level of
457 heterotrophic re-synthesis of amino acids ($\sum V = 3.98$). This species is a selective deposit feeder
458 ($CF_{\text{gut content}} = 14.9$) that preferentially feeds upon the PUFA C22:1 ω 9cis which is present in high
459 percentages in its gut content (18%) and in its body tissue (29.0%).

460

461 The benthopelagic *Peniagone* sp. of the **family Elpidiidae** has an estimated trophic level of 2.7
462 and a very high level of heterotrophic re-synthesis of amino acids ($\sum V = 7.66$). This species has
463 a 'sweeping' feeding style (Roberts et al. 2000) and assimilates fresh phytodetritus (Iken et al.
464 2001) with medium efficiency, as the PLFA concentration in its feces (CF_{feces}) is four times
465 higher than in the surface sediment (this study). In the Peru Basin, *Peniagone* sp. seems to feed
466 on diatom-derived phytodetritus ($\frac{C_{16:1\omega7}}{C_{16:0}}\text{-ratio}_{\text{feces}} = 1.42$; this study).

467

468 *Amperima* sp. belongs to the **family Elpidiidae** and its trophic level was estimated to be 2.9 ± 0.3 ,
469 potentially due to a medium level of heterotrophic re-synthesis of amino acids
470 ($\sum V = 5.24 \pm 0.90$). This species is a very selective surface deposit feeder ($CF_{\text{gut content}} = 58.0$)
471 with a 'sweeping' feeding style (Roberts et al. 2000) that grazes on very fresh phytodetritus on
472 the surface sediment (Iken et al. 2001). As a result, the gut content of *A. rosea* at PAP has higher

473 concentrations of chlorophyll-*a* compared to surface sediment or phytodetritus (FitzGeorge-
474 Balfour et al. 2010). A more detailed analysis of the phytopigments in this gut content revealed
475 that *A. rosea* at PAP feeds preferentially on cyanobacteria-derived phytodetritus (Wigham et al.
476 2003). Based on the PLFA composition of its gut content, we found that *Amperima* sp. from the
477 Peru Basin likely feeds on dinoflagellate-derived phytodetritus ($\frac{C_{16:1\omega7}}{C_{16:0}}$ -ratio_{gut content} = 0.53; this
478 study). Also the body wall fatty acid composition in our study differs substantially from
479 specimens from PAP, as the PLFA profile of PAP specimens is dominated by EPA, DHA, ARA,
480 and C18:0 (Hudson et al. 2004), whereas the PLFA profile of Peru Basin specimens is
481 characterized mostly by EPA co-eluted with C22:1 ω 9, Me-C26:2 Δ (5,9), and C28:2 Δ (11,21).
482 Hence, it seems that the feeding niche of the well-studied *Amperima* sp. can differ substantially
483 between ocean basins.

484
485 *Benthodytes* sp. from the **family Psychropotidae** has an estimated trophic level of 2.8 \pm 0.5 and a
486 very high level of heterotrophic re-synthesis of amino acids ($\sum V = 7.09\pm 3.70$). It feeds with a
487 ‘sweeper’ feeding style (Roberts et al. 2000) selectively on smaller sediment particles (medium
488 grain size: 13.0 μ m) from the surface sediment (medium grain size: 20.8 \pm 0.3 μ m; Mevenkamp et
489 al., 2019). However, it likely does not or only moderately selects for specifically detritus-
490 enriched particles ($CF_{gut content} = 1.17$; $CF_{feces} = 1.81\pm 0.53$). In fact, the high percentage of the
491 bacteria-biomarker PLFAs C16:0, C16:1 ω 7*cis*, and C18:1 ω 7*cis* in its gut content and feces, and
492 the very high level of heterotrophic re-synthesis of amino acids indicates *Benthodytes* sp. might
493 host a large biomass of living heterotrophic prokaryotes. Unfortunately, in this study no amino
494 acids from gut content or feces were extracted to assess whether this species concentrates detritus
495 that is highly enriched in amino acids. Such an observation was interpreted by Romero-Romero
496 et al., (2021) as a sign that deep-sea holothurians from Station M are secondary consumers of
497 detritus, whereas the microbial community in their guts are primary consumers of detritus.
498 Therefore, we hypothesize that also *Benthodytes* sp. is a secondary consumer, and its microbial
499 gut community is the primary consumer of detritus.

500
501 *Benthodytes typica* belongs to the **family Psychropotidae** and its trophic level is estimated to be
502 a bit higher (2.9) than the trophic level of *Benthodytes* sp. This species has a medium level of
503 heterotrophic re-synthesis of amino acids ($\sum V = 3.69$) and feeds selectively on smaller particles
504 (medium grain size: 14.2 μ m) from the ambient sediment (20.8 \pm 0.3 μ m; Mevenkamp et al.,
505 2019). These smaller particles contain an at least four times higher concentration of PLFAs than
506 the surrounding sediment ($CF_{feces} = 4.44\pm 1.52$) and consist partially of diatom-derived
507 phytodetritus ($\frac{C_{16:1\omega7}}{C_{16:0}}$ -ratio_{feces} = 1.20 \pm 0.85). Reliance on phytodetritus is confirmed by the
508 PLFA composition of *B. typica* body walls ($\frac{DHA}{EPA}$ -ratio_{body wall} = 0.13 \pm 0.08). In addition, this
509 species either feeds selectively on sediment-bound prokaryotes or hosts prokaryotes as bacteria-
510 specific PLFAs (i.e., C16:0, C16:1 ω 7*cis*, and C18:1 ω 7*cis*) contribute almost 30% to the total
511 PLFA composition in feces, but were not detected in the body wall with >2.5% of total PLFAs.
512 As a medium level of heterotrophic re-synthesis of amino acids was measured, *B. typica* likely
513 has a mixed diet. In this diet, this holothurian species consumes phytodetritus as primary
514 consumer and other types of detritus as secondary consumer following primary processing by a
515 bacterial gut community.

516

517 *Psychropotes longicauda* from the **family Psychropotidae** has a medium level of heterotrophic
518 re-synthesis of amino acids ($\sum V = 5.13$). Feeding selectively was the highest in our data (CF_{gut}
519 $content = 105$), though, surprisingly at PAP this species was found to feed less selectively than
520 *Peniagone diaphana* (FitzGeorge-Balfour et al. 2010). *P. longicauda*'s diet consists likely of
521 diatom-derived phytodetritus ($\frac{DHA}{EPA}$ -ratio_{body wall} = 0.04), but it is also possible that *P. longicauda*
522 consumes filamentous Rhodophyceae. This algae has been found in gelatinous detritus in the
523 deep sea of the NE Atlantic and Bühring et al., (2002) speculated that *P. longicauda* might feed
524 it sporadically, as the body walls of *P. longicauda* from specimens collected at PAP and in the
525 Peru Basin contain EPA, a PLFA typical for Rhodophyceae, at relatively high concentrations
526 (31% of total PLFA, this study; ~24% of total fatty acids at PAP (Ginger et al. 2000)).
527 Additionally, at PAP 70 to 80% of the gut content of this species contained sediment (Iken et al.
528 2001), which might originate from foraminiferans that Roberts and Moore, (1997) found in its
529 guts together with radiolarians, harpacticoids, nematodes, spicules, and diatoms.

530
531 *Psychropotes semperiana* (**family Psychropotidae**) has an estimated trophic level of 3.3, likely
532 related to the high level of heterotrophic re-synthesis of amino acids ($\sum V = 6.46$). This species
533 has been classified as surface deposit feeder (Iken et al. 2001) and based on the biomarkers in the
534 body tissue of a specimen collected in the Peru Basin, it consumes diatom-derived phytodetritus
535 ($\frac{DHA}{EPA}$ -ratio_{body wall} = 0.01, this study).

536 537 **Order Holothuriida**

538 *Mesothuria* sp. belongs to the **family Mesothuriidae** and has an estimated trophic level of 3.4.
539 This species could be a subsurface (Iken et al. 2001) or surface deposit feeder (Miller et al. 2000)
540 with a 'raker' feeding style (Roberts et al. 2000) or feeding with a 'wiping' motion (Hudson et
541 al. 2005). The PLFA composition of its body walls suggests that *Mesothuria* sp. likely consumes
542 diatom-derived phytodetritus ($\frac{DHA}{EPA}$ -ratio_{body wall} = 0.21). Indeed, in a study on the Hawaiian slope,
543 gut contents of *Mesothuria carnosa* had a 2.7 fold enrichment of chlorophyll a pointing towards
544 selective feeding on phytodetritus (Miller et al. 2000). Furthermore, the very high level of
545 heterotrophic re-synthesis of amino acids ($\sum V = 7.76$) from the Peru Basin suggests that
546 *Mesothuria* sp. might also be a secondary consumer of detritus. However, we lack information
547 about its gut content to confirm that it hosts a big(ger) living microbial biomass in its gut that is
548 the primary consumer of detritus.

549 550 **Order Synallactida**

551 *Oneirophanta* sp. as member of the **family Deimatidae** has an estimated trophic level of 3.5 and
552 a very high level of heterotrophic re-synthesis of amino acids ($\sum V = 7.43$). This species feeds
553 selectively ($CF_{gut content} = 16.6$) with a 'raker' feeding style (Roberts et al. 2000) and takes up
554 particles with a median grain size of 19.0 μm , which is slightly smaller than the median grain
555 size of sediment particles in the Peru Basin ($20.8 \pm 0.3 \mu\text{m}$; Mevenkamp et al., 2019). The
556 specimen collected in the Peru Basin likely fed on diatom-derived phytodetritus ($\frac{DHA}{EPA}$ -ratio_{body}
557 $_{wall} = 0.09$) and maybe on bacteria. The very high level of heterotrophic re-synthesis of amino
558 acids and the high trophic level of *Oneirophanta* sp. points to the role of a secondary consumer
559 of detritus, whereas a big biomass of microbial gut community serves as first consumers.
560 However, bacteria-specific PLFAs C16:0, C16:1 ω 7cis, and C18:1 ω 7cis, that were detected in
561 high concentrations in the gut content of *Benthodytes* sp., contribute only 5% to the total PLFA

562 composition in the gut content of *Oneirophanta* sp. Therefore, the diet preferences of this species
563 in the Peru Basin is less clear.

564

565 Synallactidae gen sp. (**family Synallactidae**) has an estimated trophic level of 3.0 ± 1.5 and a
566 medium level of heterotrophic re-synthesis of amino acids ($\sum V = 3.70 \pm 1.87$). It feeds extremely
567 selectively ($CF_{\text{gut content}} = 335$) and consumes particles of a median grain size ($14.5 \mu\text{m}$) that is
568 smaller than the median grain size of the surface sediment in the Peru Basin ($20.8 \pm 0.3 \mu\text{m}$;
569 Mevenkamp et al., 2019). The PLFA composition of the body wall and the gut content of
570 Synallactidae gen sp. indicates that this species predares upon agglutinated foraminiferans ($\frac{EPA}{ARA}$ -
571 $\text{ratio}_{\text{body wall}} = 1.75 \pm 2.23$) and it consumes diatom-derived detritus ($\frac{DHA}{EPA}$ - $\text{ratio}_{\text{body wall}} = 0.13 \pm 0.08$,
572 $\frac{DHA}{EPA}$ - $\text{ratio}_{\text{gut content}} = 0.42$, this study). However, it is not possible to differentiate whether
573 Synallactidae gen sp. is a primary consumer of the phytodetritus or a secondary consumer,
574 whereupon the foraminiferans are the primary consumer. The PLFA composition of the feces
575 shows that this holothurian species is also a bacterivore as bacteria-specific PLFAs (i.e., C16:0,
576 C16:1 ω 7cis, and C18:1 ω 7cis) contribute 42% to the total PLFA composition in the feces. If
577 Synallactes hosts a large community of living bacteria, we would expect to detect a significant
578 amount of bacteria-specific PLFAs in the gut content and a higher level of heterotrophic re-
579 synthesis of amino acids. Therefore we assume that Synallactidae gen sp. has a mixed diet
580 consisting of foraminiferans, bacteria, and phytodetritus.

581

582 *Galatheathuria* sp. from the **family Synallactidae** has an estimated trophic level of 3.1 and a
583 medium level of heterotrophic re-synthesis of amino acids ($\sum V = 3.50$). Similar to Synallactidae
584 gen sp. it feeds extremely selectively ($CF_{\text{gut content}} = 285$) and *Galatheathuria* sp. seems to
585 consume preferably diatom-derived detritus ($\frac{DHA}{EPA}$ - $\text{ratio}_{\text{body wall}} = 0.39$, $\frac{DHA}{EPA}$ - $\text{ratio}_{\text{gut content}} = 0.18$).

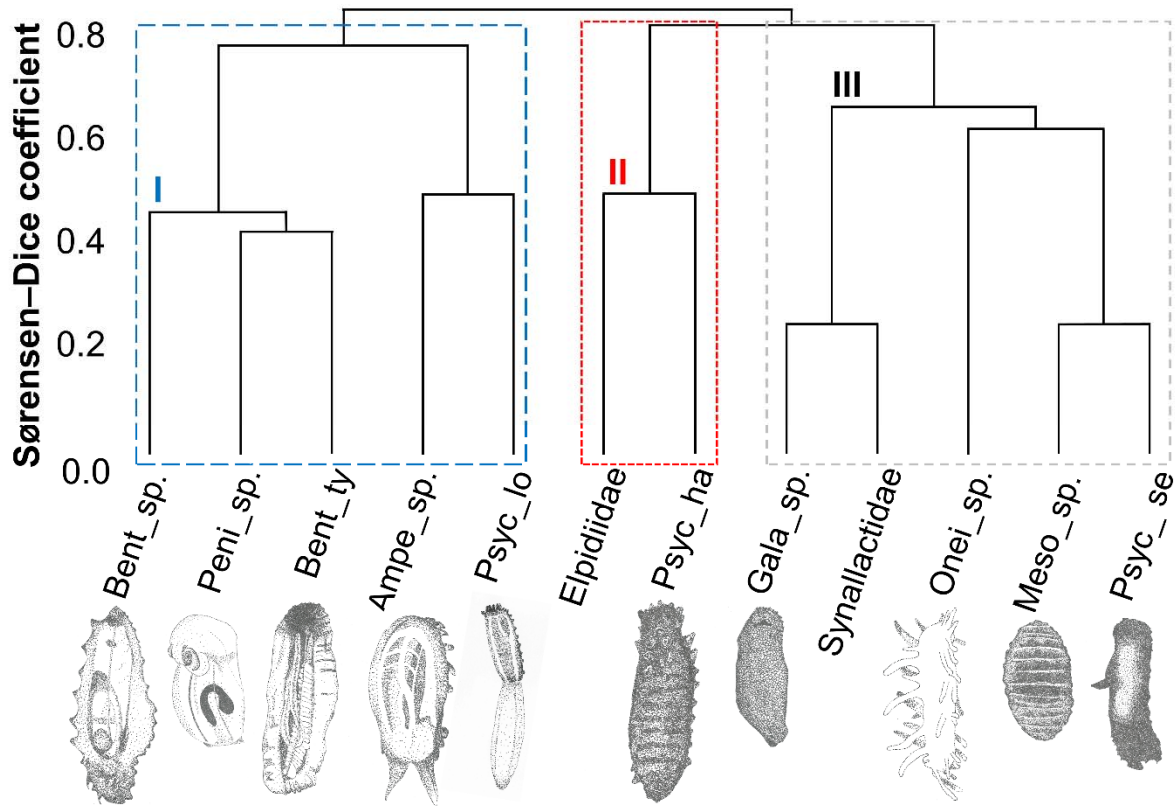
586

587 **Classification of holothurian trophic groups**

588 Here, we propose a classification system of trophic groups for holothurians from the Peru Basin
589 (Fig. 6). It is based on cluster analysis of trophic levels, heterotrophic re-synthesis level of amino
590 acids, feeding selectivity, and diet preferences, instead of on $\delta^{15}\text{N}$ value of body wall tissue.

591 **Trophic group 1** has a trophic level between 2.7 and 3.0, a very diverse diet preference, and
592 includes the species *Amperima* sp., *Benthodytes* sp., *B. typica*, *Peniagone* sp., and *Psychropotes*
593 *longicauda*. **Trophic group 2** has a low trophic level of 2.0 to 2.4 and feeds selectively. It
594 includes Elpidiidae gen sp. and *P. hanseni* and **trophic group 3** has a trophic level between 3.0
595 and 3.5 with a mixed diet and diatom-derived phytodetritus-based diet. It consists of the species
596 *Galatheathuria* sp., *Mesothuria* sp., *Oneirophanta* sp., *P. semperiana*, and Synallactidae gen sp.

597



598
 599 **Figure 6.** Dendrogram of the Sørensen–Dice coefficient calculated for holothurian species from
 600 the Peru Basin based. Trophic group I includes *Amperima* sp. (Ampe_sp.), *Benthodytes* sp.
 601 (*Bent_sp.*), *Benthodytes typica* (*Bent_ty*), *Peniagone* sp. (Peni_sp.), and *Psychropotes*
 602 *longicauda* (Psyc_lo). Trophic group II comprises Elpidiidae gen sp. (Elpidiidae) and
 603 *Psychronaetes hansenii* (Psyc_ha), and trophic group III contains *Galatheathuria* sp. (Gala_sp.),
 604 *Mesothuria* sp. (Meso_sp.), *Oneirophanta* sp. (Onei_sp.), *Psychropotes semperiana* (Psyc_se),
 605 and Synallactidae gen sp. (Synallactidae).
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613
 614 **References**

- 615 Alt CHS, Rogacheva A, Boorman B, et al (2013) Trawled megafaunal invertebrate assemblages from
 616 bathyal depth of the Mid-Atlantic Ridge (48°–54°N). *Deep-Sea Research II* 98:326–340.
 617 <https://doi.org/10.1016/j.dsr2.2013.02.003>
- 618 Baldwin RJ, Glatts RC, Smith KL (1998) Particulate matter fluxes into the benthic boundary layer at a long
 619 time-series station in the abyssal NE Pacific: composition and fluxes. *Deep-Sea Research II* 45:643–
 620 665

- 621 Ben-Mlih F, Marty JC, Fiala-Medioni A (1992) Fatty acid composition in deep hydrothermal vent
622 symbiotic bivalves. *J Lipid Res* 33:1797–1806. [https://doi.org/10.1016/s0022-2275\(20\)41337-9](https://doi.org/10.1016/s0022-2275(20)41337-9)
- 623 Billett DSM (1991) Deep-sea holothurians. *Oceanography and Marine Biology* 29:259–317
- 624 Billett DSM, Bett BJ, Reid WDK, et al (2010) Long-term change in the abyssal NE Atlantic: The “Amperima
625 Event” revisited. *Deep-Sea Research II* 57:1406–1417. <https://doi.org/10.1016/j.dsr2.2009.02.001>
- 626 Billett DSM, Bett BJ, Rice AL, et al (2001) Long-term change in the megabenthos of the Porcupine
627 Abyssal Plain (NE Atlantic). *Prog Oceanogr* 50:325–348. [https://doi.org/10.1016/S0079-
628 6611\(01\)00060-X](https://doi.org/10.1016/S0079-6611(01)00060-X)
- 629 Bligh EG, Dyer WJ (1959) A rapid method of total lipid extraction and purification. *Can J Biochem Physiol*
630 37:911–917. <https://doi.org/10.1139/o59-099>
- 631 Boschker H (2008) Linking microbial community structure and functioning: Stable isotope (¹³C) labeling
632 in combination with PLFA analysis. *Molecular Microbial Ecology Manual II* 1673–1688.
633 https://doi.org/10.1007/978-1-4020-2177-0_807
- 634 Brown A, Hauton C, Stratmann T, et al (2018) Metabolic rates are significantly lower in abyssal
635 Holothuroidea than in shallow-water Holothuroidea. *R Soc Open Sci* 5:172162.
636 <https://doi.org/10.1098/rsos.172162>
- 637 Bühring SI, Koppelman R, Christiansen B, Weikert H (2002) Are Rhodophyceae a dietary component for
638 deep-sea holothurians? *Journal of the Marine Biological Association of the United Kingdom*
639 82:347–348. <https://doi.org/10.1017/S0025315402005556>
- 640 Burdge GC, Calder PC (2014) Introduction to fatty acids and lipids. In: *World Review of Nutrition and*
641 *Dietetics*. pp 1–16
- 642 Chikaraishi Y, Ogawa NO, Kashiyama Y, et al (2009) Determination of aquatic food-web structure based
643 on compound-specific nitrogen isotopic composition of amino acids. *Limnol Oceanogr Methods*
644 7:740–750. <https://doi.org/10.4319/lom.2009.7.740>
- 645 Choi A, Song J, Joung Y, et al (2015) *Lentisphaera profunda* sp. nov., Isolated from deep-sea water. *Int J*
646 *Syst Evol Microbiol* 65:4186–4190. <https://doi.org/10.1099/ijsem.0.000556>
- 647 Dalsgaard J, st. John M, Kattner G, et al (2003) Fatty acid trophic markers in the pelagic marine
648 environment. *Adv Mar Biol* 46:225–340. [https://doi.org/10.1016/S0065-2881\(03\)46005-7](https://doi.org/10.1016/S0065-2881(03)46005-7)
- 649 Dice LR (1945) Measures of the amount of ecological association between species. *Ecology* 26:297–302
- 650 Drazen J, Phleger C, Guest M, Nichols P (2008a) Lipid, sterols and fatty acids of abyssal polychaetes,
651 crustaceans, and a cnidarian from the northeast Pacific Ocean: food web implications. *Mar Ecol*
652 *Prog Ser* 372:157–167. <https://doi.org/10.3354/meps07707>
- 653 Drazen JC, Phleger CF, Guest MA, Nichols PD (2008b) Lipid, sterols and fatty acid composition of abyssal
654 holothurians and ophiuroids from the North-East Pacific Ocean: Food web implications.
655 *Comparative Biochemistry and Physiology - B Biochemistry and Molecular Biology* 151:79–87.
656 <https://doi.org/10.1016/j.cbpb.2008.05.013>

- 657 Drazen JC, Phleger CF, Guest MA, Nichols PD (2008c) Lipid, sterols and fatty acid composition of abyssal
658 holothurians and ophiuroids from the North-East Pacific Ocean: Food web implications. *Comp*
659 *Biochem Physiol B Biochem Mol Biol* 151:79–87. <https://doi.org/10.1016/j.cbpb.2008.05.013>
- 660 Elvert M, Boetius A, Knittel K, Barker Jørgensen BO (2003) Characterization of Specific Membrane Fatty
661 Acids as Chemotaxonomic Markers for Sulfate-Reducing Bacteria Involved in Anaerobic Oxidation
662 of Methane. <https://doi.org/10.1080/01490450390241071>
- 663 Falk-Petersen S, Sargent JR, Tande KS (1987) Lipid composition of zooplankton in relation to the Sub-
664 Arctic food web. *Polar Biol* 8:115–120
- 665 Findlay RH, Trexler MB, Guckerte JB, White DC (1990) Laboratory study of disturbance in marine
666 sediments: response of a microbial community. *Mar Ecol Prog Ser* 62:121–133
- 667 FitzGeorge-Balfour T, Billett DSM, Wolff GA, et al (2010) Phytopigments as biomarkers of selectivity in
668 abyssal holothurians; interspecific differences in response to a changing food supply. *Deep-Sea*
669 *Research II* 57:1418–1428. <https://doi.org/10.1016/j.dsr2.2010.01.013>
- 670 Ginger ML, Santos VLCS, Wolff GA (2000) A preliminary investigation of the lipids of abyssal holothurians
671 from the north-east Atlantic Ocean. *Journal of the Marine Biological Association of the UK* 80:139–
672 146. <https://doi.org/10.1017/s0025315499001654>
- 673 Graeve M, Dauby P, Scailteur Y (2001) Combined lipid, fatty acid and digestive tract content analyses: A
674 penetrating approach to estimate feeding modes of Antarctic amphipods. *Polar Biol* 24:853–862.
675 <https://doi.org/10.1007/s003000100295>
- 676 Haeckel M, König I, Riech V, et al (2001) Pore water profiles and numerical modelling of biogeochemical
677 processes in Peru Basin deep-sea sediments. *Deep-Sea Research I* 48:3713–3736
- 678 Hudson IR, Pond DW, Billett DSM, et al (2004) Temporal variations in fatty acid composition of deep-sea
679 holothurians: evidence of benthic-pelagic coupling. *Mar Ecol Prog Ser* 281:109–120
- 680 Hudson IR, Wigham BD, Solan M, Rosenberg R (2005) Feeding behaviour of deep-sea dwelling
681 holothurians: Inferences from a laboratory investigation of shallow fjordic species. *Journal of*
682 *Marine Systems* 57:201–218. <https://doi.org/10.1016/j.jmarsys.2005.02.004>
- 683 Iken K, Brey T, Wand U, et al (2001) Food web structure of the benthic community at the Porcupine
684 Abyssal Plain (NE Atlantic): a stable isotope analysis. *Prog Oceanogr* 50:383–405
- 685 Jarman CL, Larsen T, Hunt T, et al (2017) Diet of the prehistoric population of Rapa Nui (Easter Island,
686 Chile) shows environmental adaptation and resilience. *Am J Phys Anthropol* 164:343–361.
687 <https://doi.org/10.1002/ajpa.23273>
- 688 Kassambara A, Mundt F (2020) factoextra: Extract and Visualize the Results of Multivariate Data
689 Analyses
- 690 Kelly JR, Scheibling RE (2012) Fatty acids as dietary tracers in benthic food webs. *Mar Ecol Prog Ser*
691 446:1–22. <https://doi.org/10.3354/meps09559>

- 692 Kharlamenko VI (2018) Abyssal foraminifera as the main source of rare and new polyunsaturated fatty
693 acids in deep-sea ecosystems. *Deep-Sea Research II* 154:358–364.
694 <https://doi.org/10.1016/j.dsr2.2017.10.015>
- 695 Khotimchenko S v (1995) Uncommon 16:1(n-5) acid from *Dictyota dichotoma* and fatty acids of some
696 brown algae of Dictyotaceae. *Phytochemistry* 38:1411–1415
- 697 Koleff P, Gaston KJ, Lennon JJ (2003) Measuring beta diversity for presence-absence data. *Journal of*
698 *Animal Ecology* 72:367–382. <https://doi.org/10.1046/j.1365-2656.2003.00710.x>
- 699 Larkin KE, Gooday AJ, Woulds C, et al (2014) Uptake of algal carbon and the likely synthesis of an
700 “essential” fatty acid by *Uvigerina* ex. gr. *semiornata* (Foraminifera) within the Pakistan margin
701 oxygen minimum zone: Evidence from fatty acid biomarker and ¹³C tracer experiments.
702 *Biogeosciences* 11:3729–3738. <https://doi.org/10.5194/bg-11-3729-2014>
- 703 Libes S (2009) *Introduction to marine biogeochemistry*, 2nd edn. Academic Press, Inc., Burlington
- 704 Massin C (1982) Food and feeding mechanisms: Holothuroidea. In: Jangoux M, Lawrence JM (eds)
705 *Echinoderm Nutrition*. Balkema, Rotterdam, pp 43–55
- 706 McCarthy MD, Benner R, Lee C, Fogel ML (2007) Amino acid nitrogen isotopic fractionation patterns as
707 indicators of heterotrophy in plankton, particulate, and dissolved organic matter. *Geochim*
708 *Cosmochim Acta* 71:4727–4744. <https://doi.org/10.1016/j.gca.2007.06.061>
- 709 Mevenkamp L, Guilini K, Boetius A, et al (2019) Responses of an abyssal meiobenthic community to
710 short-term burial with crushed nodule particles in the South-East Pacific. *Biogeosciences* 16:2329–
711 2341. <https://doi.org/10.5194/bg-16-2329-2019>
- 712 Middelburg JJ, Barranguet C, Boschker HTS, et al (2000) The fate of intertidal microphytobenthos
713 carbon: An *in situ* ¹³C-labeling study. *Limnol Oceanogr* 45:1224–1234.
714 <https://doi.org/10.4319/lo.2000.45.6.1224>
- 715 Miller RJ, Smith CR, Demaster DJ, Fornes WL (2000) Feeding selectivity and rapid particle processing by
716 deep-sea megafaunal deposit feeders: A ²³⁴Th tracer approach. *J Mar Res* 58:653–673
- 717 Oksanen J, Blanchet FG, Friendly M, et al (2017) *vegan: Community ecology package*.
- 718 Phillips NW (1984) Role of different microbes and substrates as potential suppliers of specific, essential
719 nutrients to marine detritivores. *Bull Mar Sci* 35:283–298
- 720 Pond D, Dixon D, Bell M, et al (1997) Occurrence of 16:2(n-4) and 18:2(n-4) fatty acids in the lipids of the
721 hydrothermal vent shrimps *Rimicaris exoculata* and *Alvinocaris markensis*: Nutritional and trophic
722 implications. *Mar Ecol Prog Ser* 156:. <https://doi.org/10.3354/meps156167>
- 723 Pond DW, Allen CE, Bell M v., et al (2002) Origins of long-chain polyunsaturated fatty acids in the
724 hydrothermal vent worms *Ridgea piscesae* and *Protis hydrothermica*. *Mar Ecol Prog Ser* 225:219–
725 226. <https://doi.org/10.3354/meps225219>
- 726 R-Core Team (2017) *R: A language and environment for statistical computing*

- 727 Roberts D, Gebruk A v., Levin V, Manship BAD (2000) Feeding and digestive strategies in deposit-feeding
728 holothurians. *Oceanography and Marine Biology: An Annual Review* 38:257–310
- 729 Roberts D, Moore HM (1997) Tentacular diversity in deep-sea deposit-feeding holothurians: implications
730 for biodiversity in the deep sea. *Biodivers Conserv* 6:1487–1505
- 731 Romero-Romero S, Miller EC, Black JA, et al (2021) Abyssal deposit feeders are secondary consumers of
732 detritus and rely on nutrition derived from microbial communities in their guts. *Sci Rep* 11:.
733 <https://doi.org/10.1038/s41598-021-91927-4>
- 734 Romesburg HC (1984) Cluster analysis for researchers. Lifetime Learning Publications
- 735 Ruhl HA (2007) Abundance and size distribution dynamics of abyssal epibenthic megafauna in the
736 northeast Pacific. *Ecology* 88:1250–1262. <https://doi.org/10.1890/06-0890>
- 737 Sørensen T (1948) A method of establishing groups of equal amplitude in plant sociology based on
738 similarity of species content and its application to analysis of the vegetation on Danish commons.
739 Det Kongelige Danske Videnskabernes Selskab, Copenhagen
- 740 Stratmann T, Voorsmit I, Gebruk A v., et al (2018) Recovery of Holothuroidea population density,
741 community composition and respiration activity after a deep-sea disturbance experiment. *Limnol*
742 *Oceanogr* 63:2140–2153. <https://doi.org/10.1002/lno.10929>
- 743 Svensson PA, Wong BBM (2011) Carotenoid-based signals in behavioural ecology: A review. *Behaviour*
744 148:131–189. <https://doi.org/10.1163/000579510X548673>
- 745 Tsushima M (2007) Carotenoids in sea urchins. In: Miller Lawrence J (ed) *Edible Sea Urchins: Biology and*
746 *Ecology*. Elsevier Science B.V., Amsterdam, pp 159–166
- 747 Veuger B, Middelburg JJ, Boschker HTS, Houtekamer M (2005) Analysis of ¹⁵N incorporation into D-
748 alanine: A new method for tracing nitrogen uptake by bacteria. *Limnol Oceanogr Methods* 3:230–
749 240. <https://doi.org/10.4319/lom.2005.3.230>
- 750 Wang J, Li J, Dasgupta S, et al (2014) Alterations in membrane phospholipid fatty acids of gram-positive
751 piezotolerant bacterium *sporosarcina* sp. DSK25 in response to growth pressure. *Lipids* 49:347–
752 356. <https://doi.org/10.1007/s11745-014-3878-7>
- 753 Wigham BD, Hudson IR, Billett DSM, Wolff GA (2003) Is long-term change in the abyssal Northeast
754 Atlantic driven by qualitative changes in export flux? Evidence from selective feeding in deep-sea
755 holothurians. *Prog Oceanogr* 59:409–441. <https://doi.org/10.1016/j.pocean.2003.11.003>
- 756 Yano Y, Nakayama A, Yoshida K (1997) Distribution of polyunsaturated fatty acids in bacteria present in
757 intestines of deep-sea fish and shallow-sea poikilothermic animals. *Appl Environ Microbiol*
758 63:2572–2577
- 759 Zhao JX, Liu QQ, Zhou YX, et al (2015) *Alkalimarinus sediminis* gen. nov., sp. nov., isolated from marine
760 sediment. *Int J Syst Evol Microbiol* 65:3511–3516. <https://doi.org/10.1099/ijsem.0.000446>
- 761