



**HAL**  
open science

## Experimental shift in diet $\delta^{13}\text{C}$ : a potential tool for ecophysiological studies in marine bivalves

Yves-Marie Paulet, Anne Lorrain, Joëlle Richard, Stéphane Pouvreau

### ► To cite this version:

Yves-Marie Paulet, Anne Lorrain, Joëlle Richard, Stéphane Pouvreau. Experimental shift in diet  $\delta^{13}\text{C}$ : a potential tool for ecophysiological studies in marine bivalves. *Organic Geochemistry*, 2006, 37 (10), pp.1359-1370. hal-00452408

**HAL Id: hal-00452408**

**<https://hal.science/hal-00452408>**

Submitted on 2 Feb 2010

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

2  
3 **Experimental shift in diet  $\delta^{13}\text{C}$ : a potential tool for ecophysiological studies**  
4 **in marine bivalves**

5  
6  
7  
8 Yves-Marie Paulet<sup>1</sup>, Anne Lorrain<sup>2\*</sup>, Joëlle Richard<sup>1</sup> and Stéphane Pouvreau<sup>3</sup>

9  
10  
11  
12 <sup>1</sup>LEMAR, UMR 6539 CNRS, Institut Universitaire Européen de la Mer, Place Nicolas  
13 Copernic, 29 280 Plouzané, France.

14  
15 <sup>2</sup>IRD, UR THETIS, Centre de Recherche Halieutique, 34203 Sète, France.

16  
17 <sup>3</sup>IFREMER, UMR "Physiologie & Ecophysiologie des Mollusques Marins", Station  
18 Expérimentale d'Argenton, Presqu'île du vivier, 29840 Argenton, France.

19  
20  
21 \*Corresponding author: Tel.: +33-4-99 57 32 25; fax: +33-4-99 57 32 95. E-mail address:  
22 [anne.lorrain@ird.fr](mailto:anne.lorrain@ird.fr)

26 **Abstract** - To test the potential of diet switching experiments in ecophysiological studies of  
27 marine invertebrate, stable carbon isotope ratios were measured at different seasons in the  
28 gonad, adductor muscle, digestive gland and gills of scallops (*Pecten maximus*) and oysters  
29 (*Crassostrea gigas*) held for 15 days on a constant diet of phytoplankton depleted in <sup>13</sup>C. The  
30 aim of this study was to determine if differences in carbon incorporation could be detected  
31 among species, seasons and organs, and if so, whether it was consistent with their known  
32 energy-allocation patterns. After offering the new diet, isotope values of the different organs  
33 gradually shifted and significant differences among organs, seasons and species were found.  
34 A carbon incorporation index (CII) was calculated to compare the metabolic activity of each  
35 organ of the two species between day 0 and day 15. For both species, the digestive gland had  
36 the highest CII, the adductor muscle the lowest, while gonad and gills had intermediate  
37 values. The CII was generally much higher in *P. maximus* than in *C. gigas*, suggesting higher  
38 metabolic activity in this species. Seasonal differences in the CII were also observed for the  
39 two species and were interpreted as differences in metabolic activity in accordance with our  
40 energy allocation scenario. Therefore stable isotope diet switching experiments appear to be  
41 of great value for assessing metabolic orientation in bivalves.

42

43

44 Keywords: energy allocation, metabolism, carbon isotopes, *Pecten maximus*, *Crassostrea*  
45 *gigas*

46

## 47 **1. Introduction**

48

49 Knowledge of metabolic activity and energy allocation strategies for bivalves is of  
50 great interest, particularly for aquaculture. Indeed, metabolic pathways are directly or  
51 indirectly linked to processes important for survival or reproduction (Dalhoff 2004).  
52 Therefore, understanding the origin and fate of nutrients provides a mechanistic basis for  
53 successful rearing and reproduction of bivalves in controlled conditions for aquaculture  
54 purposes. For example, the same broodstock conditioning schedule repeated at different  
55 seasons of the year generally gives rise to many different results in terms of fecundity and  
56 hatching success for many mollusc species (e.g. Utting and Millican 1997; Robert and Gérard  
57 1999), thus revealing large seasonal trends in physiology of these organisms and making  
58 knowledge of their metabolic orientation of primary importance.

59 General studies have provided data from single-point measurements of respiration,  
60 assimilation, excretion and organ weights (Vahl 1981a,b; Bayne et al. 1983; MacDonald and  
61 Thompson 1985a,b, 1986, 1987). However, those measures give only instantaneous data, and  
62 the hypothetical energy allocation scenarios produced from these types of measurements offer  
63 only approximations of net production and never depict the real carbon and energy fluxes  
64 from the environment to the animal, nor among organs within the animal. Therefore,  
65 development of new tools in bivalve ecophysiology appears of primary importance.

66 Stable isotope techniques (e.g., C, N, H, S), usually applied in ecological and  
67 population biology studies (reviews by Peterson and Fry 1987; Michener and Schell 1994),  
68 have already been used with success in experimental studies to investigate energy allocation  
69 patterns (O'Brien et al., 2000; Gauthier et al., 2003; Voigt et al., 2003). These techniques are  
70 based on the assumption that the isotopic composition of an organism is linked to that of its  
71 diet. Generally, experimental protocols involve diet switching from one isotopically distinct

72 diet to another. The main principle is that the speed at which the isotopic value of an organ  
73 changes after a diet switch is a function of the metabolic activity of the organ, including both  
74 turnover and growth. For example, such an approach has been successfully applied in  
75 determining energy allocation to reproduction in moths (O'Brien et al., 2000), and tissue  
76 turnover in fishes (Herzka and Holt 2000; Bosley et al. 2002, Suzuki et al., 2005).

77 Previous work on the scallop *Pecten maximus* (Linné 1758) in the field showed that  
78 tissue isotopic composition can be influenced by metabolic activity of the organism (Lorrain  
79 et al. 2002). We therefore expected that carbon incorporation rates would be affected by  
80 changes in bivalve energy demand and allocation. These carbon incorporation rates could then  
81 be followed in several tissues of individuals reared at different periods of the year by  
82 measuring  $\delta^{13}\text{C}$  after a diet switch. To test the potential of stable isotope experiments to  
83 effectively track carbon incorporation in bivalve species, an isotope diet switching experiment  
84 was carried out under controlled conditions. This work was conducted at four different  
85 periods of the year and on two different species, *P. maximus* and the oyster *Crassostrea gigas*  
86 (Thunberg 1793). These two species, intensively studied for aquaculture purposes, are known  
87 to show distinct seasonal behaviour, scallops having a highly regulated annual oscillation of  
88 reserve storage and utilisation (Saout 2000), whereas oysters tend to have a more  
89 opportunistic strategy of energy allocation (Enriquez-Diaz 2004). We therefore expect that the  
90 seasonal patterns of carbon incorporation would be more pronounced in *P. maximus* than in  
91 *C. gigas*. We chose several target tissues because of their different physiological functions  
92 and the likelihood of differences in carbon incorporation: adductor muscle, gonad, digestive  
93 gland and gills for both species; and the remaining tissues (i.e. labial palps, mantle and  
94 perigonadic tissues) for oysters.

95

96

97 **2. Materials and methods**

98

99 *2.1. Collection of bivalves*

100

101 Both scallops and oysters were collected in western Brittany (France). Two-year-old  
102 scallops ( $86 \pm 5$  mm, N = 48) were dredged in the Bay of Brest (Roscanvel,  $48^{\circ}20'N$ ,  
103  $4^{\circ}30'W$ ), whereas oysters of approximately the same age ( $102 \pm 10$  mm, N = 48) were hand  
104 collected at low tide in the Aber Benoît (Landéda,  $48^{\circ}34'N$ ,  $4^{\circ}37'W$ ). Sampling was carried  
105 out in March, May and September 2002, and January 2003 corresponding to the four different  
106 experimental periods (Table 1).

107

108

109 *2.2. Diet switching experiments*

110

111 Experiments were carried out at the IFREMER Shellfish laboratory at Argenton  
112 (Finistère, France) in 2002 and 2003, utilizing cultured unicellular algae with low  $\delta^{13}C$   
113 (caused by bubbling  $CO_2$  from a commercial cylinder into the culture medium) as a food  
114 source.

115 Four experiments were conducted, each one during a different hypothetical temporal  
116 window of energy allocation for scallops and oysters (i.e. in March, June, September and  
117 January; Table 1). Experiments lasted 15 days to minimize the possible effect of laboratory  
118 acclimation and to be sure that they would reflect natural metabolism and windows of energy  
119 allocation. After collection, bivalves were placed in 700 litre tanks with  $1 \mu m$  filtered running  
120 seawater for two days during which time they were not fed to empty the digestive tract.  
121 Afterwards, three individuals were randomly chosen and sacrificed (to represent day 0). The

122 remaining animals were then offered a mixed diet of four unicellular algal species depleted in  
123 <sup>13</sup>C (25% *Chaetoceros calcitrans*, 25% *Skeletonema costatum*, 25% *Isochrysis galbana*  
124 named *T-iso*, 25% *Tetraselmis chui*). During all four experiments, this diet was supplied *ad*  
125 *libitum* using a continuous dripping device with a daily ration equal to 8 % dry weight  
126 algae/dry weight flesh of animal. Algal concentrations were verified each day. A mixed diet  
127 was preferred to a single species diet as tissue production and normal rearing are reduced and  
128 perturbed with single species diets (Utting and Millican 1997; Robert and Gérard 1999). Each  
129 experiment was conducted at the ambient water temperature (Table 1).

130         At each sampling date (in general days 0, 2, 6 and 15), three individuals were taken for  
131 testing. Stomachs of scallops were first rinsed with a few millilitres of 0.2 µm filtered  
132 seawater injected via the mouth, to completely purge digestive tracts (see Lorrain et al. 2002  
133 for more details). Gonad, adductor muscle and digestive gland were then dissected from each  
134 individual. As not considered as organs of predominant role in bivalve energy strategies, gills  
135 of the three individuals were pooled in one sample. For oysters, the remaining tissues (mantle,  
136 labial palps and perigonadic tissues), generally considered as a storage tissue for this species,  
137 were also collected. As these remaining tissues were not sampled in scallop, results from the  
138 remaining oyster tissues will be regarded only as a first attempt to confirm the potential  
139 storage role of these tissues. All samples were frozen at -20°C until analysis.

140         Dietary isotopic composition was monitored by measuring the stable isotope signature  
141 of algae samples (Table 1). These algae were sampled by filtering 15 ml of the mixed algae  
142 through a precombusted Whatman GF/F filter (nominal porosity = 0.7 µm) at different  
143 periods of the experiments. The filters were then stored dried in clean glass vials after 12  
144 hours at 60°C until analysis.

145

146 *2.3. Isotopic analyses*

147 After freeze drying, bivalve tissue samples were ground to a homogeneous powder  
148 and 1 mg samples were folded into 6×4 mm tin cups for continuous flow - isotope ratio mass  
149 spectrometer (IRMS) analysis. Analysis was performed using a Europa Scientific ANCA-NT  
150 20-20 Stable Isotope Analyser with ANCA-NT Solid/Liquid Preparation Module (PDZ  
151 Europa Ltd., Crewz, UK, Scottish Crop Research Institute, Dundee, Scotland). The analytical  
152 precision (SD, N = 5) was 0.2 ‰ for C, estimated from standards analysed along with the  
153 samples. Triplicate analyses performed on some samples confirmed that analytical  
154 reproducibility was very good (0.2 ‰ maximum variation). All isotopic data are given in the  
155 conventional delta notation in units of parts per thousand (‰) relative to the Vienna Pee Dee  
156 Belemnite (VPDB) standard as follows:

$$157 \quad \delta^{13}\text{C}_{\text{sample}} = (R_{\text{sample}} / R_{\text{standard}} - 1) * 1000 \text{ where } R = {}^{13}\text{C} / {}^{12}\text{C}$$

158 Filtered algal samples were exposed to HCl vapour for 4 h at room temperature to  
159 remove carbonates (Lorrain et al. 2003). The filters were then folded, placed into 9×5 tin cups  
160 and kept in closed vials until analysis. The samples were analysed for C content and isotope  
161 ratios by N. Naulet at the University of Nantes (LAIEM, UMR CNRS 6006, France) using a  
162 Carlo Erba NA 2100 elemental analyser coupled to a Finnigan Delta S IRMS. Analytical  
163 reproducibility performed on ten replicate filters was better than 0.2 ‰ (see Lorrain et al.  
164 2003).

165

#### 166 *2.4. Carbon incorporation index*

167

168 To evaluate the differences in carbon incorporation among seasons, organs and  
169 species, we calculated a carbon incorporation index (CII). To express the actual net carbon  
170 change relative to the maximal expected change, such an index should be calculated as:

$$171 \quad \text{CII} = [\delta^{13}\text{C}_{\text{d15}} - \delta^{13}\text{C}_{\text{d0}}] / [\delta^{13}\text{C}_{\text{diet15}} - \delta^{13}\text{C}_{\text{diet0}}] * 100$$



172 where  $\delta^{13}\text{C}_{\text{d}0}$  = the tissue  $\delta^{13}\text{C}$  value at the beginning of the experiment,  $\delta^{13}\text{C}_{\text{d}15}$  = the  
173 tissue  $\delta^{13}\text{C}$  value at day 15;  $\delta^{13}\text{C}_{\text{diet}0}$  = the diet  $\delta^{13}\text{C}$  value before the beginning of the  
174 experiment and  $\delta^{13}\text{C}_{\text{diet}15}$  = the  $\delta^{13}\text{C}$  value of the diet during the experiment.

175 In fact, the food carbon isotopic ratio before the start of the experiment ( $\delta^{13}\text{C}_{\text{diet}0}$ )  
176 remains unknown. We therefore calculated the CII by replacing  $\delta^{13}\text{C}_{\text{diet}0}$  by  $\delta^{13}\text{C}_{\text{d}0}$  *i.e.* the  
177 tissue  $\delta^{13}\text{C}$  value at the beginning of the experiment:

$$178 \quad \text{CII} = [(\delta^{13}\text{C}_{\text{d}15} - \delta^{13}\text{C}_{\text{d}0}) / (\delta^{13}\text{C}_{\text{diet}} - \delta^{13}\text{C}_{\text{d}0})] * 100$$

179 When the tissue  $\delta^{13}\text{C}$  values were not available for day 15, a linear extrapolation based on the  
180 slope obtained from the two preceding values was used. For example, in September, muscle  
181  $\delta^{13}\text{C}_{\text{d}15} = \delta^{13}\text{C}_{\text{d}14} + [(\delta^{13}\text{C}_{\text{d}14} - \delta^{13}\text{C}_{\text{d}6}) / 8]$ .  $\delta^{13}\text{C}_{\text{diet}}$  represents the average  $\delta^{13}\text{C}$  value of the  
182 diet during the 15 days of the experiment (Table 1).

183 This index integrates growth and turn-over processes and therefore gives an idea of the  
184 quantity of metabolites allocated to a specific organ. This CII does not take into account  
185 eventual differential fractionation factors between organs, or some isotopic routing processes,  
186 as is discussed later.

187

## 188 2.5. Data analysis

189

190 Differences in carbon isotope composition between d0 and d15 were tested by  
191 performing a non-parametric Kruskal-Wallis test for each organ (except for the remaining  
192 tissues in March and the gills for which the analyses were conducted on pooled samples) and  
193 each experiment. The same method was applied for CII comparison. When significant  
194 differences were detected, results were classified using the Mann-Whitney non parametric  
195 procedure. Differences were considered significant at  $\alpha = 0.05$ .

196

### 197 **3. Results**

198

199         The decrease of the tissue  $\delta^{13}\text{C}$  over time in both species and for nearly all organs,  
200 indicates successful incorporation of dietary carbon from phytoplankton (Fig. 1 and 2).  
201 Significant carbon isotope change between d0 and d15 was observed for digestive glands,  
202 adductor muscles and gonads of scallops in all seasons (Fig. 1,  $p < 0.05$ ). For oysters,  
203 significant change was also observed in all experiments ( $p < 0.05$ ) except for adductor muscle  
204 during the June experiment and gonad during the September experiment (Fig. 2,  $p > 0.05$ ).  
205 Within this general decreasing trend, some differences between species, seasons, and organs  
206 were apparent. For example, in March, scallop and oyster digestive gland tissues showed the  
207 most rapid decrease of  $\delta^{13}\text{C}$  over time, with an average of 11.3 and 5.8 ‰ change in 12 days,  
208 whereas muscle showed only 1.6 and 0.9 ‰ for scallops and oysters, respectively (Fig. 1 and  
209 2). These differences can be expressed by the carbon incorporation index (CII, see Methods).  
210 The CII clearly reveals a strong difference between scallops and oysters (Fig. 3). Scallops  
211 always had a larger CII than oysters ( $p < 0.01$ ), irrespective of organs and seasons, except for  
212 the digestive gland and muscle in January, when oysters had a higher CII than scallops.

213         Muscle had always the lowest CII, except for oysters in June and September (Figure 3,  
214 Table 3) where muscle did not differ significantly from remaining tissues and gonad,  
215 respectively. Conversely, digestive gland always had the highest CII values, except for  
216 scallops in March and September when gonad and digestive gland did not differ significantly.  
217 Gonads, gills and remaining tissues showed intermediate values between muscle and digestive  
218 gland.

219         Seasonal differences do appear in CII, but differ between species. In scallops, a  
220 general trend toward increasing values from March to January experiments was observed. In

221 oysters, seasonal variations are less pronounced although experiments conducted during  
222 winter months produced significantly higher CII values compared to other months (Table 3).

223

224

225

## 226 **4. Discussion**

227

228 As expected, diet-switching experiments led to changes over time in tissue carbon  
229 isotopic composition, confirming the potential of stable isotope studies to trace carbon within  
230 organisms. Indeed, the carbon incorporation index (CII, see Methods) shows significant  
231 differences among organs, species and seasons. We should stipulate that this CII gives  
232 combined information of tissue growth and turnover (see Gannes et al., 1997), which we  
233 could not separate in this study. Furthermore, as already mentioned in materials and methods,  
234 it would have been better to calculate the CII using the food  $\delta^{13}\text{C}$  value before the diet  
235 switching, instead of using the tissue  $\delta^{13}\text{C}$  value at the beginning of the experiment. Further  
236 experiments should try to evaluate this, even if it is difficult when using bivalves in their  
237 natural environment. However, these global data can still provide valuable information about  
238 metabolite allocation to an organ, which can be very useful for ecophysiological studies.

239 Furthermore, even though the general idea that  $\delta^{13}\text{C}$  variations are directly linked to  
240 organ activity and growth and can help us to study bivalve metabolism, some other processes  
241 may account for the observed differences and reduce the power of this study. According to the  
242 literature, two main processes acting on  $\delta^{13}\text{C}$  variations should be considered: isotopic routing  
243 and differential isotopic fractionation (Gannes et al., 1997).

244 Isotopic routing is a process by which some biochemical components from the diet,  
245 with specific isotopic signatures, are preferentially allocated to certain organs. Most of the  
246 time, lipid accumulation is proposed to explain some decreases in  $\delta^{13}\text{C}$  as lipids are strongly

247 depleted in  $^{13}\text{C}$  (Tieszen et al., 1983). To determine if such a process can fundamentally affect  
248 interpretation of our data, we are able to simulate the effect of a complete isotopic routing,  
249 i.e., only dietary lipids incorporated in one organ. To perform such a simulation, the isotopic  
250 composition of the lipid and non-lipid fraction of the diet has to be known. Data from the  
251 literature allows us to estimate that  $\delta^{13}\text{C}$  of the lipid fraction of a tissue is around 6 to 7 ‰  
252 lower than of its non-lipid fraction (McConnaughey and McRoy, 1979; Kling et al., 1992;  
253 Bearshop et al., 2002). By a mass balance calculation, we estimated the  $\delta^{13}\text{C}$  value of the lipid  
254 fraction of the diet in June (phytoplankton with a maximum 20 % of lipids: Whyte, 1987;  
255 Brown, 1991) to be about -50.8 ‰, whereas the non-lipid fraction is about -42.4 ‰. We  
256 hypothesize that the digestive gland, which is known to be an organ rich in lipids (30 %,  
257 Saout et al., 1999), exclusively incorporates the lipid fraction of the diet (estimated  $\delta^{13}\text{C} =$   
258 -50.8 ‰) and conversely that the muscle (a fat-free organ) incorporates only the non lipid  
259 fraction ( $\delta^{13}\text{C} = -42.4$  ‰). Then using these new values in place of the measured dietary  $\delta^{13}\text{C}$ ,  
260 we can re-estimate the CII. For the muscles of scallops in June, the CII values would shift  
261 from 3.9 to 4.1 %, and for the digestive gland from 45.4 to 34.7 %. In January, recalculated  
262 muscle values would not change significantly whereas digestive gland values would shift  
263 from 17.5 to 14.3 %. Therefore, this estimation of the maximum isotopic routing would  
264 explain only up to 30 % of the observed variations and would not change our conclusions.

265 Differing fractionation factors between organs could also impact the CII results, but  
266 data for differential fractionation between organs of the same organism are scarce (see  
267 Dalerum and Angerbjörn, 2005) and even nonexistent for marine bivalves. However, the  
268 results of Lorrain et al. (2002) suggest that these differing fractionation factors would not  
269 exceed 4 ‰ in scallops (between muscle and digestive gland). Therefore, including differing  
270 fractionation factors (from 1 to 4 ‰) to the  $\delta^{13}\text{C}_{\text{diet}}$  value in the CII calculations would have a  
271 very small impact on CII value differences observed between organs.

272           The CII differed among organs in the following order: digestive gland > gonad >  
273 adductor muscle. Furthermore, seasonal variations are stronger in the digestive gland and in  
274 the gonad than in the muscle, suggesting that the digestive gland and the gonad integrate  
275 shorter time variations as compared to the muscle. On an annual average, the gills presented  
276 intermediate CII values, close to those of the gonad (23 and 9 % for scallops and oysters  
277 respectively) and were constant throughout the year. The “remaining tissues” CII presented an  
278 annual mean value close to 7 % but with large seasonal variations, suggesting that these  
279 tissues might have a storage role, in contrast to the gills.

280           The most striking result of our diet switching experiments is that strong differences in  
281 the CII were observed among seasons. Food, provided ad libitum in all the four experiments,  
282 cannot account for these differences. Temperature differed between experiments, but was not  
283 the cause of these variations as both the maximum and the minimum scallop CII were  
284 observed at the lowest temperature (Fig. 3), in March and January, respectively. To a lesser  
285 extent, the same inference can be drawn for oysters. From this, it can be concluded that  
286 carbon allocation is not driven by immediate thermal conditions, as is suggested for many  
287 bivalves (McDonald and Thomson, 1985b, 1986). Our results constitute the first data set  
288 illustrating this pattern for bivalves obtained using stable isotope diet switching experiments.

289           Variations between the two species CII probably reflect their different energy  
290 allocation strategies. From several works on the same scallop population of the Bay of Brest  
291 (Paulet et al. 1988, 1997; Saout et al. 1999; Saout 2000), a hypothetical schedule of metabolic  
292 activities can be drawn (Fig. 4). Scallops in the Bay of Brest are characterized by a strategy of  
293 storage and postponed use of energy. Basically, energy stored as glycogen in the adductor  
294 muscle and principally as lipids in the digestive gland during spring and summer, is used to  
295 sustain reproductive effort and maintenance during winter. In spring and summer, somatic and  
296 reproductive production is directly fuelled from the available food. Schematically, in terms of

297 energetic allocation priority, the year for an adult scallop can be subdivided in three main  
298 parts (Fig. 4): i) from November to April, metabolic translocation from somatic to  
299 reproductive tissues occurs, ii) from April to May, a transitory period, is characterized by the  
300 simultaneous production of somatic and reproductive tissues mainly from food, and iii) from  
301 June to October, major energetic fluxes originate from the food and are used for reserve  
302 building.

303 In contrast to the scallop, the Japanese oyster, *C. gigas*, from the coast of Brittany,  
304 exhibits an annual cycle by which food can directly sustain growth and reproduction for most  
305 of the year (Chavez-Villalba et al. 2001; Chavez-Villalba et al. 2002a, b; Enriquez-Diaz  
306 2004). Spring and summer are periods of major gonadal production, whereas somatic growth  
307 occurs during this period according to a more opportunistic manner depending on food  
308 availability (Fig. 4). Reproductive activity is sustained either by direct uptake or via  
309 metabolites stored in the reserve tissues ("remaining tissues": mantle, labial palps and  
310 perigonadal tissues). In oysters, seasonally based biological changes are acknowledged to be  
311 less rigid than for scallops, with individuals inhabiting a large range of environmental  
312 conditions (Enriquez-Diaz, 2004).

313 For scallops in the present study, the maximum carbon incorporation in reserve  
314 tissues, i.e. muscle and digestive gland, is observed in March and June. Intense incorporation  
315 into the gonad is limited to March and to a lesser extent to June and September. These results  
316 show that energy allocation to reproduction is observed in March and that reproduction can  
317 still occur in the second temporal window of energy allocation (June to September) in  
318 accordance with the annual schedules shown in figure 4. Therefore, contrary to previous  
319 hypotheses, scallops seem able to assimilate external food as early as March, and not only  
320 after April; this probably reveals the opening of a "receptive window" to food availability  
321 somewhere between the end of January and the beginning of March. The very low CII values

322 observed for somatic tissues in September is contradictory to the prediction of the energy  
323 allocation model (Fig. 4), and probably reveals an overestimation of the storage process in  
324 autumn even though target organs (muscle and digestive gland) are near their maximum  
325 filling. Finally, carbon incorporation is at its minimum in January for all tissues, underscoring  
326 the fact that gonadic activity observed in winter for this species (Paulet et al. 1997) is directly  
327 dependant on the use of storage tissues.

328         In oysters, CII values are generally lower than in scallops and seasonal differences  
329 were less pronounced. In this species, carbon incorporation in the gonad revealed by isotope  
330 results was maximal in March and June, in agreement with the energy allocation model  
331 (Fig.4) and verifying that gonadal tissue develops during this period, at least partially from  
332 food uptake. In September, the gonad CII remains close to zero, corresponding precisely to  
333 the resting stage documented for this species in Brittany (e.g. Lango-Reynoso et al. 1999; Li  
334 et al. 2000; Fig. 4). Seasonal variations of the CII of the digestive gland appear less  
335 pronounced than in scallops, in accordance with its relatively minor role as a storage organ in  
336 oysters compared to scallops. However, the high CII observed for the digestive gland relative  
337 to the other organs in all seasons suggests that this organ would have a more important  
338 storage role in oysters than previously described. The “remaining tissues” exhibited a high CII  
339 value in March, probably due to their predominant role as a reserve compartment. This  
340 heterogeneous tissue composed of the mantle, the labial palps and the perigonadal envelope  
341 will be the object of more extensive studies in the future. For both species, the gills represent  
342 a site of active incorporation during all seasons, which warrants further studies.

343         Our results from carbon isotope tracing agree with previous knowledge on energy  
344 allocation for the studied bivalves. Clearly scallops appear as a species with more rigid and  
345 contrasted temporal allocation windows than oysters, as evidenced by the greater seasonality  
346 in the CII. This apparent highly regulated functioning might be compared with the existence

347 of a putative annual rhythmic physiological oscillation, driven or not by photoperiod, as  
348 proposed by Paulet and Boucher (1991) for this species. One must also consider that scallops  
349 have very distinct periods of energy allocation, using food or tissue reserves, whereas oysters  
350 use both food and tissue reserves simultaneously.

351 Another striking result of these experiments was the generally lower magnitude of the  
352 CII for oysters as compared to scallops. The carbon incorporation, discussed in this study, is a  
353 double source process: i) the renewal of existing tissues (tissue turnover), and ii) the  
354 production of new tissues (growth). This must be considered when discussing differences in  
355 the CII between both species. Although growth was not measured during the course of the  
356 experiment, tissue growth data for oysters in Aber Benoît (Fleury et al., 2001) and for scallops  
357 in the Bay of Brest (Lorrain et al., 2004) reveals that at the same age, a scallop produces  
358 annually at least two times more soft tissues than an oyster. Therefore, a part of the observed  
359 difference in CII values could be due to differences in tissues growth between the two species.  
360 In future studies, the development of methods adapted to precisely assess tissue production at  
361 the individual level would be of primary importance. Secondly, the markedly lower CII  
362 observed for oysters could also reveal a lower metabolism, inducing a lower tissue turnover,  
363 for this species compared to scallops. Such a difference seems consistent with some other life  
364 history traits of these species, such as i) the potential mobility in scallops contrasted with the  
365 sedentary life of the oysters, and ii) the larger pallial cavity, and the greater valve movements,  
366 in scallops compared to oysters of the same size, probably allowing higher pumping rates in  
367 scallops (Møhlenberg and Riisgård, 1979; see also discussion in Bricelj and Shumway, 1991).

368 Finally, for dietary studies, in which stable isotopes are a key tool, this kind of  
369 experiment could be continued over longer time periods to assess turnover and fractionation  
370 factors between food and different tissues. Indeed, the bivalves sampled at the end of our  
371 experiments had not yet reached equilibrium with their new diet. Furthermore, in this study,



372 the elucidation of great differences in carbon incorporation kinetics between organs confirms  
373 the potential of multi-organ analyses to study spatial or temporal variations in diet  $\delta^{13}\text{C}$   
374 (Tieszen et al., 1983; Hobson and Clark, 1992a, b; Hobson et al. 1996). Indeed, to study  
375 trophic dynamics at different time scales, the digestive gland and gonads are more appropriate  
376 than the muscle to detect short-term food source variations, as the muscle only gives an  
377 average value over a long period.

378

## 379 **5. Conclusion**

380

381 Diet switching experiments, conducted under the same diet regime but at different  
382 periods of the year, have revealed differences in carbon incorporation among organs, seasons,  
383 and species. These results are consistent with previous knowledge on energy allocation  
384 strategies for *P. maximus* and *C. gigas*. This study represents an important first step in  
385 establishing the potential of stable isotope diet switching experiments for carbon tracing in  
386 bivalves. In this regard, information from this type of experiment would offer valuable  
387 insights into bivalve ecophysiology and energy allocation patterns. The next stage will be the  
388 coupled study of isotope tracking with a whole carbon budget of the two species, including  
389 consumption, respiration, production (organ by organ) and faecal excretion estimations.

390

391

## 392 **Acknowledgments**

393

394 Financial support of this study was granted by IFREMER and especially by the GDR  
395 “Reproduction of marine bivalves.” We thank all the staff of the IFREMER shellfish  
396 laboratory at Argenton and particularly P. Le Souchu for great help in bivalve conditioning

397 and maintenance during all experiments, and T. Mellado and E. Coquard for their contribution  
398 in sample preparation. Many thanks also to D.P. Gillikin, S. Ford and S. Bouillon for English  
399 corrections and helpful comments; and P. Lopez whose illustrations greatly improved this  
400 manuscript.

401

## 402 **References**

403

404 Bayne, B.L., Salked, P.N., Worrall, C.M., 1983. Reproductive effort and value in different  
405 populations of the marine mussel *Mytilus edulis* L. *Oecologia* (Berl.) 59, 18-26.

406 Bearshop, S., Waldron, S., Votier, S.C., Furness, R.W., 2002. Factors that influence  
407 assimilation rates and fractionation of nitrogen and carbon stable isotopes in avian blood  
408 and feathers. *Physiol Biochem Zool* 75, 451-458.

409 Bosley, K.L., Witting, D.A., Chambers, R.C., Wainright, S.C., 2002. Estimating turnover  
410 rates of carbon and nitrogen in recently metamorphosed winter flounder  
411 *Pseudopleuronectes americanus* with stable isotopes. *Marine Ecology Progress Series*  
412 236, 233-240.

413 Bricelj, V.M., Shumway, S., 1991. Physiology: energy acquisition and utilization. In.  
414 *Scallops: biology, ecology and aquaculture* (ed. S.E. Shumway), pp. 305-346.  
415 Amsterdam: Elsevier (Developments in Aquaculture and Fisheries Science, n°21).

416 Brown, M.R., 1991. The amino-acid and sugar composition of 16 species of microalgae used  
417 in Mariculture. *Journal of Experimental Marine Biology and Ecology* 145, 79-99.

418 Chavez-Villalba, J.E., Mingant, C., Cochard, J.C., Le Pennec, M., 2001. Gamétogenèse chez  
419 l'huître *Crassostrea gigas* de l'Aber Benoît (Bretagne, France), à la limite nord de son aire  
420 de reproduction. *Haliotis* 30, 1-12.

421 Chavez-Villalba, J., Barret, J., Mingant, C., Cochard, J.C., Le Penne, M., 2002a. Autumn  
422 conditioning of the oyster *Crassostrea gigas*: a new approach. *Aquaculture* 210, 171-186.

423 Chavez-Villalba, J., Pommier, J., Andriamiseza, J., Pouvreau, S., Barret, J., Cochard, J.C., Le  
424 Penne, M., 2002b. Broodstock conditioning of the oyster *Crassostrea gigas*: origin and  
425 temperature effect. *Aquaculture* 214, 115-130.

426 Dalhoff, E.P., 2004. Biochemical indicators of stress and metabolism. *Annual Review of*  
427 *Physiology* 66, 183-207.

428 Dalerum, F., Angerbjörn, A., 2005. Resolving temporal variation in vertebrate diets using  
429 naturally occurring stable isotopes. *Oecologia*. DOI: 10.1007/s00442-005-0118-0.

430 Enriquez-Diaz, M.R., 2004. Variabilité et bioénergétique de la reproduction chez l'huître  
431 creuse, *Crassostrea gigas*. Thèse de doctorat, Université de Bretagne Occidentale, Brest  
432 (France), 187 pp.

433 Fleury, P.G., Goyard, E., Mazurié, J., Claude, S., Bouget, J.F., Langlade, A., Le Coguc, Y.,  
434 2001. The assessing of Pacific oyster (*Crassostrea gigas*) rearing performances by the  
435 IFREMER/REMORA network: method and first results (1993-1998) in Brittany (France).  
436 *Hydrobiologia* 465, 195-208.

437 Gannes, L.Z., O'Brien, D.M., Martinez Del Rio, C., 1997. Stable isotopes in animal ecology:  
438 Assumptions, caveats and a call for more laboratory experiments. *Ecology* 78, 1271-1276.

439 Gauthier, G., Bety, J., Hobson, K.A., 2003. Are greater snow geese capital breeders? New  
440 evidence from a stable isotope model. *Ecology* 84, 3250-3264.

441 Herzka, S.Z., Holt, G.J., 2000. Changes in isotopic composition of red drum (*Sciaenops*  
442 *ocellatus*) larvae in response to dietary shifts: potential applications to settlement studies.  
443 *Canadian Journal of Fisheries and Aquatic Sciences* 57, 137-147.

444 Hobson, K.A., Clark, R.G., 1992a. Assessing avian diets using stable isotopes I: turnover of  
445 <sup>13</sup>C in tissues. *The Condor* 94, 181-188.

446 Hobson, K.A., Clark, R.G., 1992b. Assessing avian diets using stable isotopes II: factors  
447 influencing diet-tissue fractionation. *The Condor* 94, 189-197.

448 Hobson, K.A., Schell, D.M., Renouf, D., Noseworthy, E., 1996. Stable carbon and nitrogen  
449 isotopic fractionation between diet and tissues of captive seals: implications for dietary  
450 reconstruction involving marine mammals. *Canadian Journal of Fisheries and Aquatic*  
451 *Sciences* 53, 528-533.

452 Kling, G.W., Fry, B., O'Brien, W.J., 1992. Stable isotopes and planktonic trophic structure in  
453 arctic lakes. *Ecology* 73, 561-566.

454 Lango-Reynoso, F., Devauchelle, N., Le Penne, M., Hatt, P.J., 1999. Elements of  
455 reproductive strategy in oysters, *Crassostrea gigas*, from the "Rade de Brest", France.  
456 *Invertebrate Reproduction and Development* 36, 141-144.

457 Li, K., Osada, M., Mori, K., 2000. Seasonal biochemical variations in Pacific oyster gonadal  
458 tissue during sexual maturation. *Fisheries Science* 66, 502-508.

459 Lorrain, A., Paulet, Y.-M., Chauvaud, L., Savoye, N., Donval, A., Saout, C., 2002.  
460 Differential  $\delta^{13}\text{C}$  and  $\delta^{15}\text{N}$  signatures among scallop tissues: implications for ecology and  
461 physiology. *Journal of Experimental Marine Biology and Ecology* 275, 47-61.

462 Lorrain, A., Savoye, N., Chauvaud, L., Paulet, Y.-M., Naulet, N., 2003. Decarbonation and  
463 preservation method for the analysis of organic C and N contents and stable isotope ratios  
464 of low-carbonated suspended particulate material. *Analytica Chimica Acta* 491, 125-133.

465 Lorrain, A., Paulet, Y.-M., Chauvaud, L., Dunbar, R., Mucciarone, D., Fontugne, M., 2004.  
466  $\delta^{13}\text{C}$  variation in scallop shells: Increasing metabolic carbon contribution with body  
467 size? *Geochimica et Cosmochimica Acta* 68, 3509-3519.

468 MacDonald, B.A., Thompson, R.J., 1985a. Influence of temperature and food availability on  
469 the ecological energetics of the giant scallop *Placopecten magellanicus*. I. Growth rates of  
470 shell and somatic tissue. *Marine Ecology Progress Series* 25, 279-294.

471 MacDonald, B.A., Thompson, R.J., 1985b. Influence of temperature and food availability on  
472 the ecological energetics of the giant scallop *Placopecten magellanicus*. II. Reproductive  
473 output and total production. Marine Ecology Progress Series 25, 295-303.

474 MacDonald, B.A., Thompson, R.J., 1986. Influence of temperature and food availability on  
475 the ecological energetics of the giant scallop *Placopecten magellanicus*. III. Physiological  
476 ecology, the gametogenic cycle and scope for growth. Marine Biology 93, 37-48.

477 MacDonald, B.A., Thompson, R.J., Bayne, B.L., 1987. Influence of temperature and food  
478 availability on the ecological energetics of the giant scallop *Placopecten magellanicus*.  
479 IV. Reproductive effort, value and cost. Oecologia 72, 550-556.

480 Mc Connaughey, T., McRoy, C.P., 1979. Food web structure and the fractionation of carbon  
481 isotopes in the Bering Sea. Marine Biology 53, 257-262.

482 Michener, R.H., Schell, D.M., 1994. Stable isotope ratios as tracers in marine aquatic food  
483 webs. In *Stable isotopes in ecology and environmental science* (ed. K Lajtha and RH  
484 Michener), pp. 138-157. Blackwell.

485 Møhlenberg, F., Riisgård, H.U., 1979. Filtration rate, using a new indirect technique, in  
486 thirteen species of suspension-feeding bivalves. Marine Biology 54, 143-147.

487 O'Brien, D.M., Schrag, D.P., Martinez Del Rio, C., 2000. Allocation to reproduction in a  
488 hawkmoth: a quantitative analysis using stable carbon isotopes. Ecology 81, 2822-2831.

489 Paulet, Y.M., Lucas, A., Gerard, A., 1988. Reproduction and larval development in two  
490 *Pecten maximus* (L.) populations from Brittany. Journal of Experimental Marine  
491 Biology and Ecology 119, 145-156.

492 Paulet, Y.M., Bekhadra, F., Dechauvelle, N., Donval, A., Dorange, G., 1997. Seasonal cycles,  
493 reproduction and oocyte quality in *Pecten maximus* from the Bay of Brest. Annales de  
494 l'Institut océanographique de Paris 73, 101-112.

495 Paulet, Y.M., Boucher, J., 1991. Is reproduction mainly regulated by temperature or  
496 photoperiod in *Pecten maximus*? *Invertebrate Reproduction and Development* 19, 61-70.

497 Peterson, B.J., Fry, B., 1987. Stable isotopes in ecosystem studies. *Annual Review of Ecology*  
498 *and Systematics* 18, 293-320.

499 Robert, R., Gérard, A., 1999. Bivalve hatchery technology: the current situation for the  
500 Pacific oyster *Crassostrea gigas* and the scallop *Pecten maximus* in France. *Aquatic*  
501 *Living Resources* 12, 121-130.

502 Saout, C., Quéré, C., Donval, A., Paulet, Y.M., Samain, J.F., 1999. An experimental study of  
503 the combined effects of temperature and photoperiod on reproductive physiology of  
504 *Pecten maximus* from the bay of Brest (France). *Aquaculture* 172, 301-314.

505 Saout, C., 2000. Contrôle de la reproduction chez *Pecten maximus* (L.): Etudes in situ et  
506 expérimentales. Thèse de doctorat, Université de Bretagne Occidentale, Brest (France),  
507 172 pp.

508 Suzuki, K.W., Kasai, A., Nakayama, K., Tanaka, M., 2005. Differential isotopic enrichment  
509 and half-life among tissues in Japanese temperate bass (*Lateolabrax japonicus*) juveniles:  
510 implications for analyzing migration. *Canadian Journal of Fisheries and Aquatic Sciences*  
511 62, 671-678.

512 Tieszen, L.L., Boutton, T.W., Tesdahl, K.G., Slade, N.A., 1983. Fractionation and turnover of  
513 stable isotopes in animal tissues: Implications for  $\delta^{13}\text{C}$  analyses of diet. *Oecologia* 57, 32-  
514 37.

515 Utting, S.D., Millican, P.F., 1997. Techniques for the hatchery conditioning of bivalve  
516 broodstocks and the subsequent effect on egg quality and larval viability. *Aquaculture*  
517 155, 47-56.

- 518 Vahl, O., 1981a. Energy transformations by the Iceland scallop, *Chlamys islandica* (O.F.  
519 Müller), from 70°N. I. The age-specific energy budget and net growth efficiency. Journal  
520 of Experimental Marine Biology and Ecology 53, 281-296.
- 521 Vahl, O., 1981b. Energy transformations by the Iceland scallop, *Chlamys islandica* (O.F.  
522 Müller), from 70°N. II. The population energy budget. Journal of Experimental Marine  
523 Biology and Ecology 53, 297-303.
- 524 Voigt, C.C., Matt, F., Michener, R., Kunz, T.H., 2003. Low turnover rates of carbon isotopes  
525 in tissues of two nectar-feeding bat species. Journal of Experimental Biology 206, 1419-  
526 1427.
- 527 Whyte, J.N.C., 1987. Biochemical composition and energy content of six species of  
528 phytoplankton used in mariculture of bivalves. Aquaculture 60, 231-241.

529

## Table captions

530

531 Table 1. Details of the experimental protocol of the four experiments (March, June,  
532 September and January): exact dates, water temperature, isotopic composition of the diet  
533 (means  $\pm$  1 S.D, N for number of measurements) and specific composition of the unicellular  
534 algae species that compose the diet given to scallops and oysters.

535

536 Table 2. Results of the multiple range test for differences in CII among organs for the same  
537 experiment (organ effects) and among experiments for the same organ (seasonal effects) for  
538 scallops (A) and oysters (B). ns: non significant, \*\* : significant at the 95% level, nd: not  
539 determined. G: Gonad; DG; Digestive Gland; M: Muscle; R: Remaining tissues.



	March 2002	June 2002	September 2002	January 2003
Dates	8 to 23 March	30 May to 14 June	9 to 24 September	10 to 25 January
Temperature	10°C	14°C	17°C	10°C
Diet $\delta^{13}\text{C}$ (‰)	-42.7 N = 1	-43.9 ± 0.5 N = 2	-50.3 ± 3.1 N = 3	-52.8 ± 2.7 N = 4
Unicellular algal species	<i>Isochrysis galbana</i> , <i>Skeletonema costatum</i> , <i>Tetraselmis chui</i> , <i>Chaetoceros calcitrans</i>			

542 **A. *Pecten maximus***

543  
544 Organ effects

	March		June		September		January	
	G	DG	G	DG	G	DG	G	DG
M	**	**	**	**	**	**	**	**
G		ns		**		ns		**

545  
546  
547 Seasonal effects

	Muscle			Gonad			Digestive Gland				
	March	June	Sept	March	June	Sept	March	June	Sept		
June	**			June	**		June	**			
Sept	**	ns		Sept	**	**	Sept	**	**		
Jan	**	**	**	Jan	**	ns	**	Jan	**	**	ns

548  
549  
550 **B. *Crassostrea gigas***

551  
552 Organ effects

	March			June			September			January		
	G	DG	R	G	DG	R	G	DG	R	G	DG	R
M	**	**	nd	**	**	ns	ns	**	**	**	**	**
G		**	nd		**	**		**	**		**	ns
DG			nd			**			**			**

553  
554  
555 Seasonal effects

	Muscle			Gonad			Digestive Gland			Remaining Tissues		
	March	June	Sept	March	June	Sept	March	June	Sept	March	June	Sept
June	ns			ns			ns			nd		
Sept	ns	ns		**	**		**	ns		nd	**	
Jan	ns	ns	ns	**	**	**	**	**	**	nd	**	ns

556

Figure captions

557

558

559 Figure 1. *Pecten maximus*. Stable carbon isotope values ( $\delta^{13}\text{C}$ , in ‰) of scallops as a function  
560 of time since the diet switch for the four different experiments (March, June, September and  
561 January), each sub-figure corresponding to one of the four organs (adductor muscle, gills,  
562 gonad and digestive gland). Values are means  $\pm$  1 standard deviation (N = 3) except for gills  
563 where the value corresponds to a pool of the three individuals.

564

565 Figure 2. *Crassostrea gigas*. Stable carbon isotope values ( $\delta^{13}\text{C}$ , in ‰) of oysters as a  
566 function of time since diet switch for the four different experiments (March, June, September  
567 and January), each sub-figure corresponding to one of the four organs (adductor muscle, gills,  
568 gonad, digestive gland) and the remaining tissues (mantle, labial palps and perigonadic  
569 tissues). Values are means  $\pm$  1 standard deviation (N = 3) except for gills where the value  
570 corresponds to a pool of the three individuals.

571

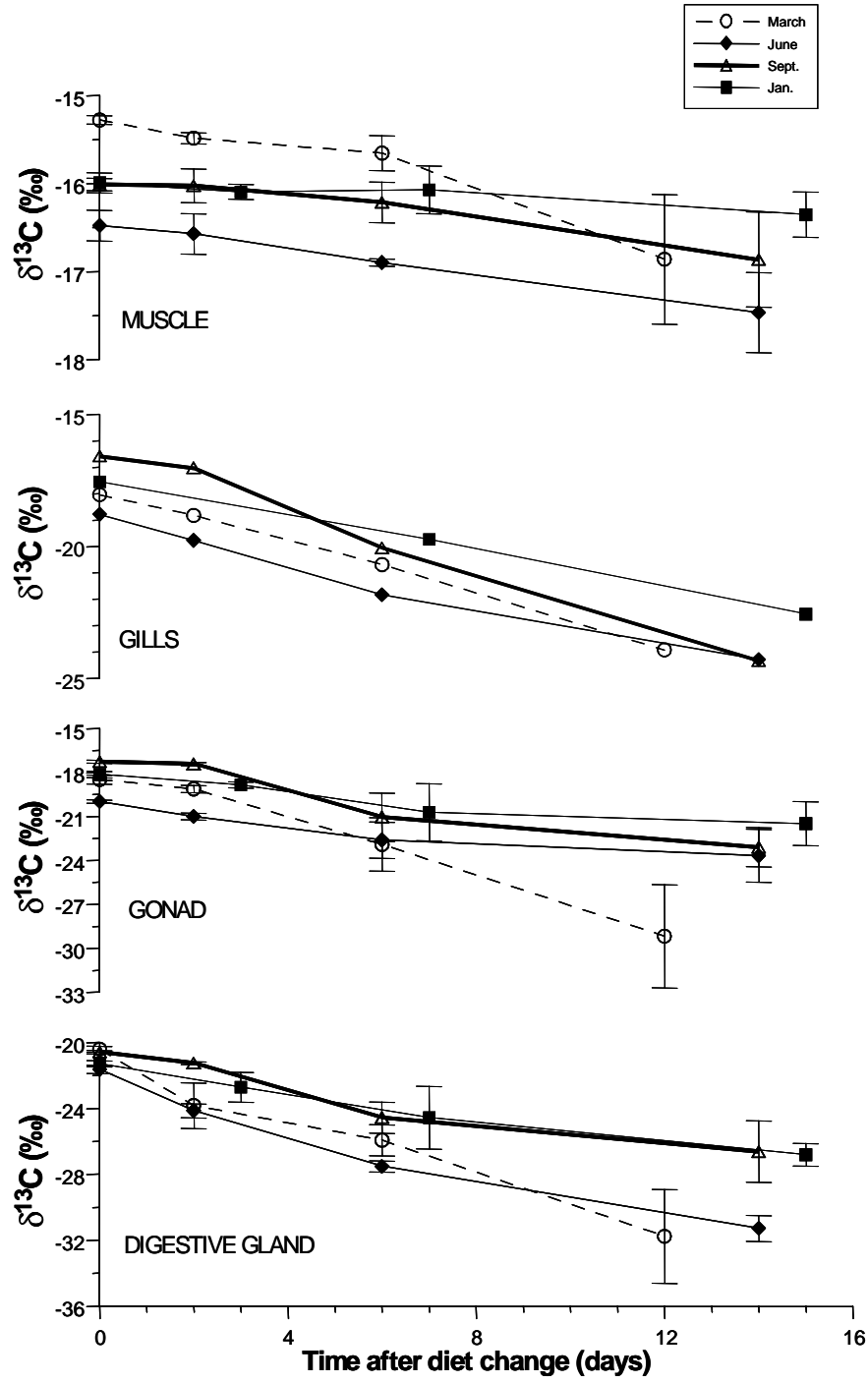
572 Figure 3. Carbon Incorporation Index (CII, in %) in the different organs (adductor muscle,  
573 gonad, digestive gland, gills and remaining tissues) of *P. maximus* (black bars) and *C. gigas*  
574 (grey bars) for the different experiments (March, June, September and January). Temperature  
575 during experiments is also indicated. See materials and methods for calculations of CII.  
576 Standard deviations are indicated when available.

577

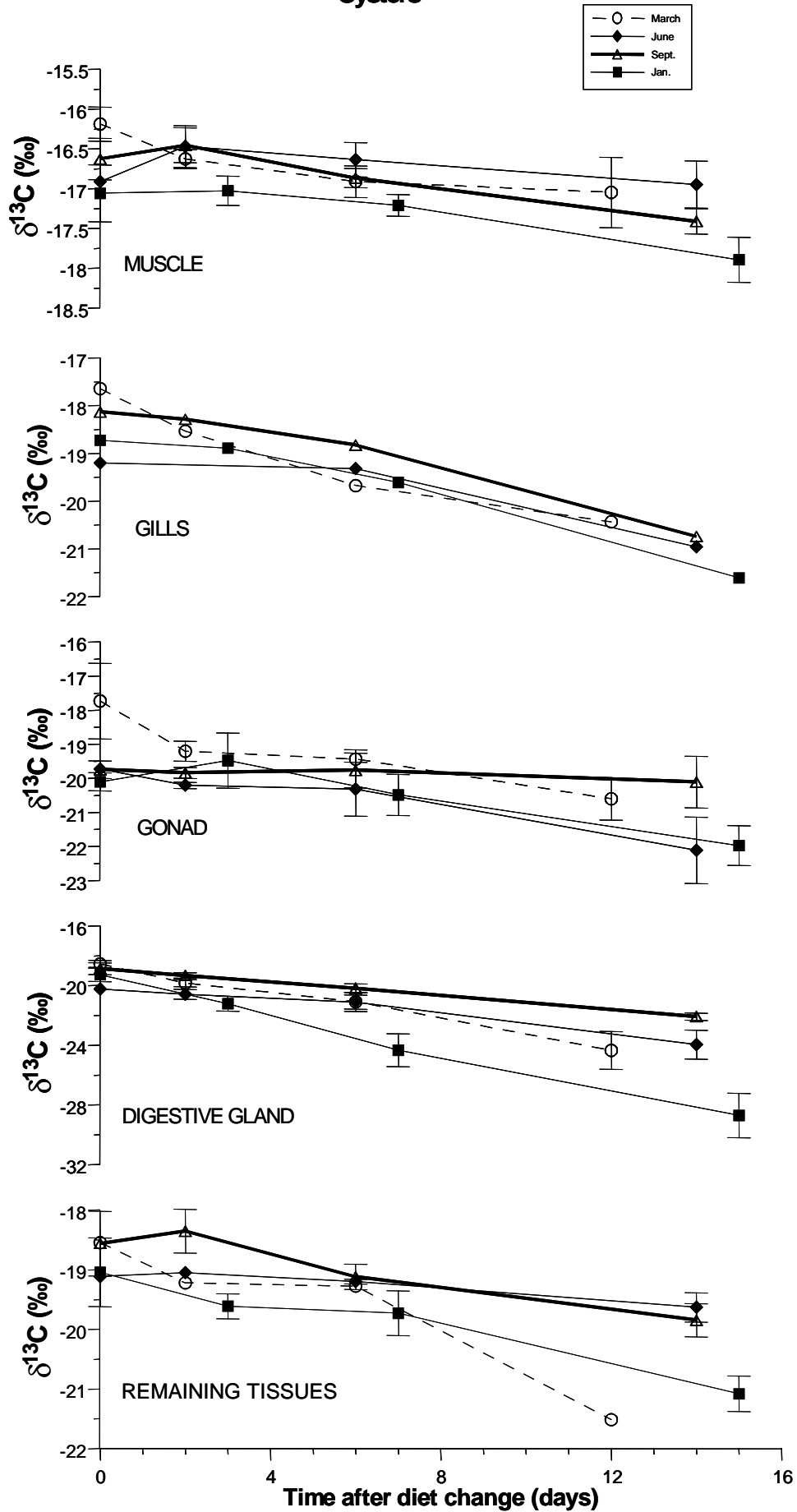
578 Figure 4. Hypothetical annual model of energy allocation for the two bivalves species  
579 developed from previous studies (see discussion) A) *Pecten maximus* and B) *Crassostrea*  
580 *gigas*. Arrows illustrate energy origin (food or reserve tissue) during the three different  
581 periods for each species. R signifies that energy is primarily being allocated to reproduction

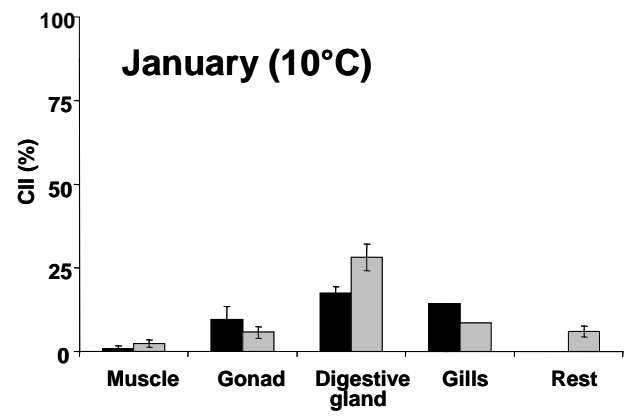
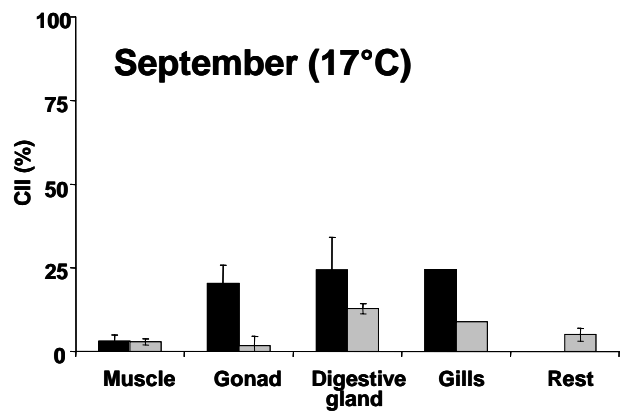
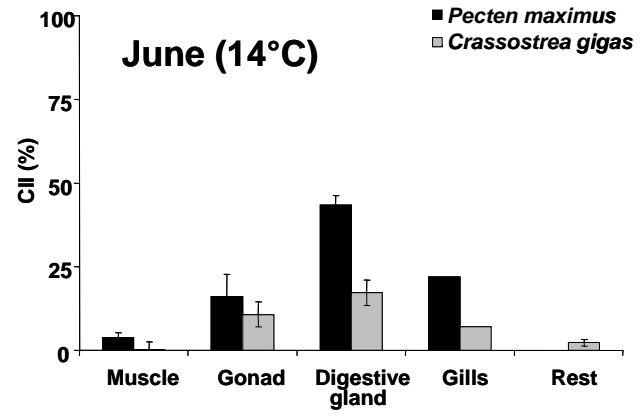
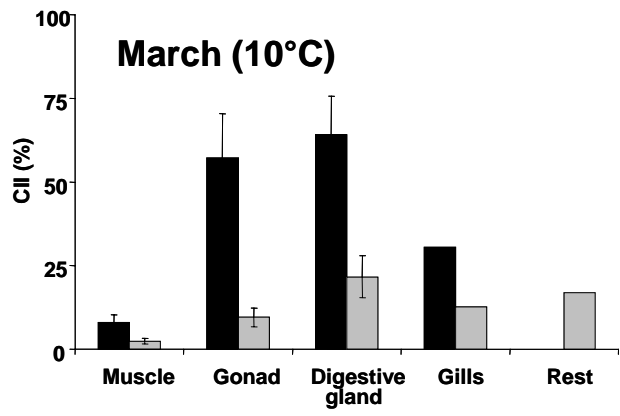
582 and S to somatic growth; small caps indicate secondary processes that can still occur if energy  
583 is in excess. Hachure section represents a resting stage for oysters.

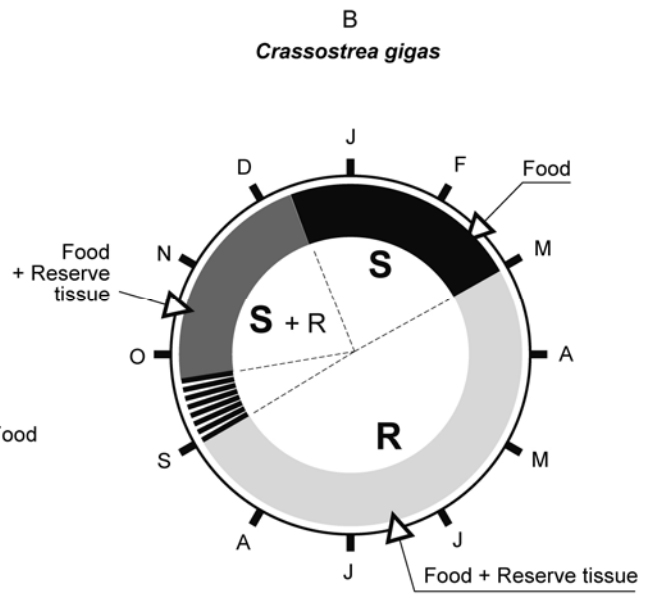
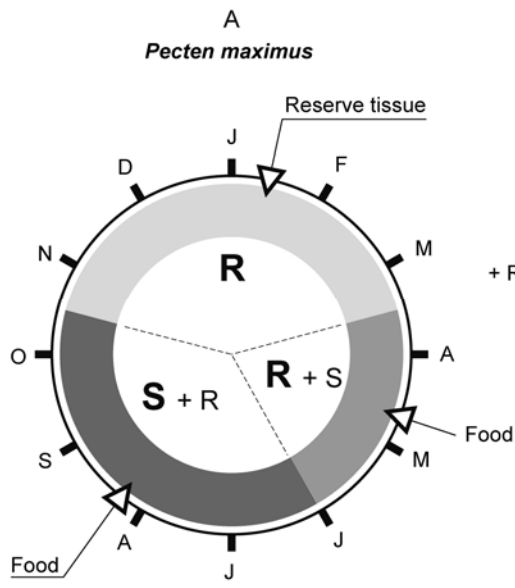
# Scallops



# Oysters







584