Metabolism of the mussel *Mytilus galloprovincialis* from two origins in the Ría de Arousa (north-west Spain)

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Mussel seed *Mytilus galloprovincialis* (Bivalvia: Mytilidae) from two origins (rocky shore and collector ropes) was cultivated on a raft in the Ría de Arousa (north-west Spain), from seeding to thinning out, for 226 d (November 1995–July 1996) and two aspects of metabolism, oxygen consumption rate (VO_2) and ammonia excretion rate ($\text{VNH}_4\text{-N}$) were studied *in situ*.

The model derived from multiple analysis of oxygen consumption accounted for 91.9% of the variance, based on dry weight of the mussels and the environmental factors quality of food (organic content) and mainly chlorophyll-*a*. Seed origin also showed significant influence. The seasonal pattern of the oxygen consumption can be attributed mainly to the variation of chlorophyll-*a*, which showed a higher range of values in the spring months.

Origin of seed did not show a homogeneous effect on oxygen consumption throughout the cultivation period. Collector rope mussels showed higher oxygen consumption values at the beginning of the cultivation period and after the first 15 d, but the rocky shore mussels showed a higher oxygen consumption between days 22 and 110. Energy-conserving patterns and lower condition index at the onset of the experiment for rocky shore mussels could explain these initial differences.

Multiple analysis on the variation of ammonia excretion rate provided a model that accounted for 72.6% of the variance based on dry weight of mussels, seed origin and the environmental parameters chlorophyll-*a* and total particulate matter. The rocky shore mussels showed a significantly higher excretion values for most of the cultivation period, although there was no constant tendency throughout. High excretion values were recorded between January and March, whilst for the rest of the cultivation period values were low.

The O:N index was higher in collector rope mussels for most of the cultivation period, which may suggest a more favourable energy metabolism and/or a more appropriate nutritional state for these specimens.

INTRODUCTION

In studies of physiological energetic, oxygen consumption and nitrogen excretion rates represent the costs and losses of energy associated with metabolism. The physiological responses of bivalve molluscs to changes in the environment are extremely variable (Bayne & Widdows, 1978; Widdows et al., 1984; Tedengren et al., 1990; Navarro et al., 1991), and the adaptation of each population to the specific characteristics of their habitats have on occasion made it possible to establish effects linked to origin when specimens are subjected or transplanted to new environmental and/or nutritional conditions. These responses have been collectively defined by Mallet et al. (1987) as the 'ecological memory' of the specimens with regard to their original habitat when they are placed in a different environment. To this effect, differences in growth linked to origin (collector rope and rocky shore mussels) have been observed for mussels cultivated on a raft (Pérez Camacho et al., 1995; Babarro et al., 2000a). In the case of physiological parameters for *Mytilus edulis* and *Mytilus galloprovincialis*, Okumus & Stirling (1994), in a natural environment, and Labarta et al. (1997) under laboratory conditions respectively, observed a significant effect of origin on ammonia excretion, but not on oxygen

consumption. Widdows et al. (1984) did however find differences in both physiological rates, but at specific times during the annual cycle.

In this study we analyse the physiological rates of oxygen consumption and ammonia excretion in mussels from collector rope and rocky shore origins when both are cultivated on a raft from a length of 20 to 60 mm, and evaluate their behaviour in terms of endogenous and environmental factors.

MATERIALS AND METHODS

Harvesting and maintenance of mussels

In November 1995 seed of *Mytilus galloprovincialis* Lmk of approximately 20 mm in length was gathered from the rocky shore and collector ropes on a raft, both in the mid-outer area of the Ría de Arousa (Galicia, north-west Spain). Both groups of mussels were detached from rocks and ropes respectively and subsequently allowed re-attaching to cultivation ropes on the raft by using the standard procedure of the mussel culture in Spain (linen net which degraded itself after some days in seawater). Both types of seed, from the same year class, came from spawning period in the previous spring–summer and the

Table 1. Characteristics of the natural seston (means $\pm$ SD).

Date	TPM (mg/l)	POM (mg/l)	PIM (mg/l)	Vol (mm ³ /l)	T (°C)	S (‰)	Chl- <i>a</i> (µg/l)	Q ₁ (POM/TPM)	Q ₂ (POM/Vol)	Chl- <i>a</i> /POM
27/11/95	0.68 $\pm$ 0.18	0.37 $\pm$ 0.11	0.30	0.57 $\pm$ 0.29	15.7 $\pm$ 0.7	32.8 $\pm$ 1.9	0.75 $\pm$ 0.18	0.55	0.65	2.02
05/12/95	0.49 $\pm$ 0.15	0.29 $\pm$ 0.04	0.20	0.39 $\pm$ 0.21	15.4 $\pm$ 1.1	31.3 $\pm$ 4.7	0.36 $\pm$ 0.27	0.59	0.75	1.25
13/12/95	0.83 $\pm$ 0.11	0.31 $\pm$ 0.03	0.52	0.52 $\pm$ 0.09	15.6 $\pm$ 0.4	35.2 $\pm$ 0.5	0.48 $\pm$ 0.15	0.38	0.60	1.52
20/12/95	0.74 $\pm$ 0.21	0.37 $\pm$ 0.08	0.36	0.45 $\pm$ 0.08	13.7 $\pm$ 0.4	34.8 $\pm$ 0.4	0.61 $\pm$ 0.15	0.51	0.82	1.63
03/01/96	2.56 $\pm$ 0.68	1.00 $\pm$ 0.27	1.56	1.29 $\pm$ 0.09	14.3 $\pm$ 0.3	30.1 $\pm$ 2.9	0.36 $\pm$ 0.36	0.39	0.78	0.36
17/01/96	0.79 $\pm$ 0.06	0.46 $\pm$ 0.02	0.33	0.51 $\pm$ 0.08	13.9 $\pm$ 0.6	28.0 $\pm$ 4.8	0.29 $\pm$ 0.28	0.59	0.90	0.62
31/01/96	0.55 $\pm$ 0.07	0.28 $\pm$ 0.03	0.27	0.29 $\pm$ 0.05	13.2 $\pm$ 0.3	30.1 $\pm$ 1.4	0.64 $\pm$ 0.27	0.51	0.95	2.32
15/02/96	1.34 $\pm$ 0.20	0.39 $\pm$ 0.06	0.95	0.71 $\pm$ 0.06	13.9 $\pm$ 0.6	31.1 $\pm$ 2.2	1.43 $\pm$ 0.35	0.29	0.54	3.70
28/02/96	1.31 $\pm$ 0.11	0.57 $\pm$ 0.08	0.74	1.07 $\pm$ 0.14	12.9 $\pm$ 0.4	34.1 $\pm$ 1.4	1.16 $\pm$ 0.18	0.43	0.53	2.05
13/03/96	0.88 $\pm$ 0.22	0.46 $\pm$ 0.10	0.42	0.90 $\pm$ 0.29	13.1 $\pm$ 0.1	34.9 $\pm$ 0.4	1.19 $\pm$ 0.37	0.52	0.51	2.59
27/03/96	0.96 $\pm$ 0.17	0.52 $\pm$ 0.08	0.44	1.26 $\pm$ 0.26	13.8 $\pm$ 0.2	33.1 $\pm$ 0.4	1.71 $\pm$ 0.28	0.54	0.41	3.27
10/04/96	1.38 $\pm$ 0.12	0.64 $\pm$ 0.04	0.74	1.66 $\pm$ 0.15	14.4 $\pm$ 0.5	32.5 $\pm$ 2.7	1.96 $\pm$ 1.60	0.47	0.39	3.04
24/04/96	0.74 $\pm$ 0.09	0.39 $\pm$ 0.02	0.35	0.65 $\pm$ 0.15	15.0 $\pm$ 0.6	33.7 $\pm$ 1.7	1.59 $\pm$ 0.35	0.53	0.60	4.05
05/06/96	0.62 $\pm$ 0.09	0.29 $\pm$ 0.04	0.33	0.76 $\pm$ 0.10	15.0 $\pm$ 1.2	35.0 $\pm$ 0.9	1.11 $\pm$ 0.59	0.46	0.38	3.86
03/07/96	0.72 $\pm$ 0.13	0.43 $\pm$ 0.08	0.29	0.64 $\pm$ 0.16	14.3 $\pm$ 0.7	35.6 $\pm$ 0.2	1.03 $\pm$ 0.30	0.60	0.67	2.38

TPM, total particulate matter; POM, particulate organic matter; PIM, particulate inorganic matter; Vol, total particulate volume; T, temperature; S, salinity; Chl-*a*, chlorophyll-*a*; Q₁, organic content (by weight). Q₂, organic content (by volume); Chl-*a*/POM, chlorophyll-*a* content (by total organic weight).

sampling locations are 2 km away from each other. Experimental cultivation, which was carried out in a raft usually employed for the culture in Ría de Arousa (500 m²), began in winter with the aim of minimizing any possible advantages accruing to the collector ropes seed as a result of its better adaptation to raft cultivation. The experiment continued until July 1996 (226 d) and covered the first stage of mussel cultivation from seeding to thinning out (50–60 mm). Sixteen cultivation ropes (12 m) were used, eight for each type of seed, hung alternately and with a density of 19 kg of seed per rope (1.6 kg/m of rope or 2600 ind/m of rope). Sampling was performed by removing mussels from adjacent ropes at an average depth of 2–4 m for both groups of seed.

The initial length (mean $\pm$ SD) of the seed was 22.55 $\pm$ 1.55 mm for collector ropes and 19.02 $\pm$ 1.93 mm for the rocky shore mussels. Mean total dry weight (shell plus soft tissues) was 0.36 $\pm$ 0.06 and 0.27 $\pm$ 0.06 g/ind for each type of seed, respectively. No significant differences in length and total dry weight were observed between both types of seed at the beginning of the experiment (ANOVA; $P > 0.05$; $N = 300$). The final length (mean $\pm$ SD) achieved for both groups of mussels in July 1996 was 55.61 $\pm$ 1.60 mm for collector ropes and 50.11 $\pm$ 1.96 mm for the rocky shore mussels. Mean total dry weight (shell plus soft tissues) at the end of the experimental period was 4.74 $\pm$ 0.23 and 3.08 $\pm$ 0.30 g/ind for each type of seed, respectively (ANOVA; $P < 0.001$; $N = 300$).

Measurements

Natural seston was characterized as total particulate matter (TPM mg/l), particulate organic matter (POM mg/l), particulate inorganic matter (PIM mg/l), particulate volume (Vol mm³/l) and chlorophyll-*a* (chl-*a* µg/l). Chlorophyll-*a* values as well as temperature (T°C) and salinity (S psu) values of the water column were provided by the Centro de Control da Qualidade do Medio Mariño da Consellería de Pesca, Marisqueo e Acuicultura

Figure 1. Fluctuation of condition index values (mg cm⁻³) for rocky shore and collector ropes mussels and chlorophyll-*a* concentration throughout the cultivation time.

(Xunta de Galicia). Seston quality was expressed as Q₁ (POM/TPM), Q₂ (POM/Vol) and the chl-*a*/POM index. Seawater samples were filtered onto pre-ashed (450°C for 4 h) and weighed GFC filters and rinsed with isotonic ammonium formate (0.5M). Total dry matter was established as the weight increment determined after drying the filters to constant weight at 110°C for 12 h with an accuracy of 0.001 mg. Organic matter corresponded to the weight loss after ignition at 450°C for 4 h in a muffle furnace. Particulate volume per litre of seawater was determined by counting in the range of 2–56 µm using a Coulter Counter Multisizer II fitted with a 100-µm aperture tube. The variation in these environmental and/or nutritional parameters over the cultivation period is shown in Table 1 and was described in a previous paper (Babarro et al., 2000a).

Metabolic rate (VO₂)

The metabolic rate was obtained by measuring the oxygen consumption (VO₂). Mussels were placed in

Table 2. Values of physiological parameters (mean $\pm$ SD; $N=30$) of two sources of seed mussel standardized to tissue dry weight (DW mg) according to growth of *Mytilus galloprovincialis* Lmk during the experiment.

Date	Days of cultivation	Source of seed mussel	DW mg	VO ₂ (ml O ₂ /h)	VNH ₄ -N (μ g NH ₄ -N/h)	O:N Index
27/11/95	0	Collector ropes	50	0.050 $\pm$ 0.007***	1.57 $\pm$ 0.42**	39.84
		Rocky shore		0.040 $\pm$ 0.011	2.33 $\pm$ 0.48	21.48
05/12/95	8	Collector ropes	50	0.038 $\pm$ 0.006	0.50 $\pm$ 0.19	95.54
		Rocky shore		0.038 $\pm$ 0.005	0.71 $\pm$ 0.27	67.35
13/12/95	15	Collector ropes	50	0.036 $\pm$ 0.007*	0.84 $\pm$ 0.23*	53.74
		Rocky shore		0.032 $\pm$ 0.006	1.11 $\pm$ 0.23	35.96
20/12/95	22	Collector ropes	50	0.039 $\pm$ 0.003***	0.68 $\pm$ 0.19	71.98
		Rocky shore		0.053 $\pm$ 0.004	0.86 $\pm$ 0.27	76.83
03/01/96	36	Collector ropes	50	0.021 $\pm$ 0.009	0.37 $\pm$ 0.17**	70.35
		Rocky shore		0.015 $\pm$ 0.003	0.81 $\pm$ 0.42	23.28
17/01/96	50	Collector ropes	50	0.027 $\pm$ 0.004***	0.71 $\pm$ 0.25**	47.52
		Rocky shore		0.042 $\pm$ 0.003	1.17 $\pm$ 0.20	44.85
31/01/96	64	Collector ropes	70	0.061 $\pm$ 0.004***	2.02 $\pm$ 0.79	37.92
		Rocky shore		0.066 $\pm$ 0.005	1.64 $\pm$ 0.76	50.41
15/02/96	80	Collector ropes	100	0.113 $\pm$ 0.006***	2.19	64.47
		Rocky shore		0.137 $\pm$ 0.008	2.99	57.25
28/02/96	95	Collector ropes	100	0.122 $\pm$ 0.013**	2.05 $\pm$ 1.35	74.51
		Rocky shore		0.133 $\pm$ 0.009	2.87 $\pm$ 1.73	57.86
13/03/96	110	Collector ropes	200	0.184 $\pm$ 0.019**	2.21 $\pm$ 0.55	104.03
		Rocky shore		0.200 $\pm$ 0.018	2.56 $\pm$ 0.91	97.66
27/03/96	125	Collector ropes	300	0.240 $\pm$ 0.026	2.05 $\pm$ 0.46**	146.28
		Rocky shore		0.244 $\pm$ 0.028	2.92 $\pm$ 0.81	104.59
10/04/96	140	Collector ropes	400	0.361 $\pm$ 0.086	3.13	144.11
		Rocky shore		0.335 $\pm$ 0.071	4.10	102.09
24/04/96	155	Collector ropes	600	0.488 $\pm$ 0.104	7.00	87.11
		Rocky shore		0.409 $\pm$ 0.102	8.30	61.57
05/06/96	197	Collector ropes	1000	0.701 $\pm$ 0.139	8.10	108.13
		Rocky shore		0.658 $\pm$ 0.176	9.24	88.97
03/07/96	226	Collector ropes	1000	0.585 $\pm$ 0.088	6.25 $\pm$ 1.88	116.90
		Rocky shore		0.538 $\pm$ 0.075	7.32 $\pm$ 1.94	91.81

VO₂, oxygen consumption rate; VNH₄-N, ammonia excretion rate. (*, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$).

closed 780 ml cylindrical chambers (height 85 mm, diameter 115 mm) sealed with oxygen probes (YSI® 5730). The oxygen concentration was recorded at regular intervals with oxygen meters (YSI® 58) connected to the probes for approximately 60 min, until the oxygen concentration fell below 30% of the initial value. One parallel control chamber without animals was always used to correct for bacterial respiration, electrode drift, etc. Metabolic rate was then calculated:

$$\text{VO}_2 = 60[\text{Ct}_0 - \text{Ct}_1][V/(\text{t}_1 - \text{t}_0)] \quad (1)$$

where Ct₀–Ct₁ are the concentration of oxygen in the water (mg O₂/l) at start and finish times (min) respectively; V is the volume of the chamber; t₁–t₀ represent start and finish times (min) of the measured period, respectively.

Ammonia excretion rate (VNH₄-N)

Rate of ammonia excretion was determined by placing mussels in closed chambers filled with 250 ml of air-saturated water previously filtered through 0.2 μ m Millipore membranes. After 150 min, 10 ml samples of the water were extracted from each experimental chamber to be analysed following Solorzano's method (1969):

$$\mu\text{gNH}_4 - \text{N/h} = [(\text{test} - \mu\text{M control}) / (14/(1000/V))(1/t)] \quad (2)$$

where V is the volume of the experimental container (250 ml) and t is the time of incubation (2.5 h).

The number of specimens taken into consideration for the physiological measurements varied according to their size (i.e. with the length of cultivation period). At the beginning, eight specimens of 50 mg of tissue dry weight class were placed in each chamber in a set of ten, and this number was also employed for 70 and 100 mg of tissue dry weight classes. From 200 mg of tissue dry weight class onwards the number of animals used progressively decreased and at the end of the experiment there was only one specimen of 1000 mg of tissue dry weight class in each chamber. Physiological measurements were taken weekly from November to December, fortnightly from January to May, and monthly in June and July.

The ratio of oxygen consumed to nitrogen excreted (O:N index) was computed by atomic equivalents according to Widdows (1985).

Condition index of the animals was calculated as the individual dry weight per volume. As an index of (shell) volume the length to the third power was used. Variation of this condition index throughout the experimental time

in relation to chlorophyll-*a* concentration is shown in the Figure 1.

Size standardization

To preclude variability in physiological rates caused by size differences, these rates were corrected to a standardized individual according to growth of the animals. Once physiological measurements were completed, shell length of each individual was recorded to the nearest 0.1 mm with vernier callipers and the soft tissues excised from the shell, dried at 110°C for 12 h and weighed with an accuracy of 0.01 g. Oxygen consumption and ammonia excretion rates were obtained according to the formula:

$$Y_s = Y_e (DW_{st}/DW_e)^b \quad (3)$$

where Y_s and Y_e are the corrected and non-corrected physiological rates respectively, DW_{st} is the range 50–1000 mg of tissue dry weight, DW_e is the experimental weight of the mussels (DW in mg) and b is the power value that scales physiological rate to body weight: $b=0.75$ (Bayne & Newell, 1983) and $b=0.72$ (Hawkins et al., 1985) for the oxygen consumption and ammonia excretion rates respectively. In order to show fluctuation of the physiological rates vs time of culture for a standard size class, 1 g of tissue dry weight was chosen for both physiological rates.

Data analysis

Standard analysis of variance (ANOVA) was used to compare means for both physiological rates (VO_2 and VNH_4-N). Kruskal–Wallis and Friedman non-parametric tests were employed when homogeneity was not obtained (Bartlett's test) after data transformation.

Multiple stepwise regression analysis was used to determine the effect of a variety of factors, both endogenous and environmental, on the variation of oxygen consumption and ammonia excretion rates. The seed origin factor was added to this analysis with values of 0 and 1 for collector rope and rocky shore mussels respectively. Both physiological rate values were transformed by $\log_{10}(x+1)$ to stabilize variances.

All the analyses were carried out in accordance with the statistical methods described by Zar (1984).

RESULTS

Oxygen consumption (VO_2)

Oxygen consumption values for both groups of mussels (collector rope and rocky shore) throughout the experimental cultivation period are shown in Table 2.

Oxygen consumption rate was 25% higher in collector rope mussels than those of rocky shore origin at the beginning of the culture period ($P<0.001$; Table 2). After 8 d, VO_2 was lower for both types of seed, although this decrease was greater for collector rope mussels, which meant no difference between them ($P>0.05$). The initial differences observed between both types of seed reappeared after 15 d of cultivation on the raft ($P<0.05$; Table 2). However, between days 22 and 110 of the culture period the differences in VO_2 were inverted, and this rate was

Figure 2. Variation of oxygen consumption (VO_2) and ammonia excretion (VNH_4-N) rates throughout the cultivation time. Both physiological rates have been standardized to 1 g of tissue dry weight.

significantly higher in the rocky shore mussels when compared to the collector rope specimens (Table 2). After day 110 of cultivation on the raft, VO_2 values were similar for both groups of mussels ($P>0.05$).

Oxygen consumption vs cultivation time

Variation of oxygen consumption for 1 g of tissue dry weight class over the experimental period is shown in Figure 2A. During the first 15 d, VO_2 showed an acclimation response to the experimental cultivation conditions, as has been described above. After this initial period, oxygen consumption fluctuation clearly corresponded to a seasonal model: low values during the winter months and a rise of this physiological rate in the spring and summer months (Figure 2A). There was a strong variation in VO_2 during wintertime (a range of 0.123–0.497 ml O_2 /h for both types of mussels). On the other hand, there was greater uniformity in the spring–summer period with values ranging from 0.538–0.772 ml O_2 /h for both groups of mussels.

Oxygen consumption vs dry weight and environmental parameters

The effect of mussel growth (weight) and the variation of environmental factors on oxygen consumption, as well as the seed origin factor, was studied by means of a multiple regression analysis (Table 3). The model thus

Table 3. Stepwise multiple regression of log physiological rates of *Mytilus galloprovincialis* (VO_2 , oxygen consumption rate and VNH_4-N , excretion rate) with logarithm of tissue dry weight (log DW), quality of seston ($Q_s=POM/TPM$), temperature ($T^{\circ}C$), chlorophyll-*a* ($\mu g/l$), total particulate matter ($TPM\ mg/l$) and origin of seed.

Parameter	VO_2			
	Coefficient	SE	F-ratio	P
Constant	-2.505			
Log DW	0.647	0.014	2081.016	<0.001
Q_s	0.191	0.065	8.706	<0.01
T	-0.013	0.006	5.123	≈ 0.05
Chl- <i>a</i>	0.255	0.015	278.525	<0.001
Origin	0.069	0.010	46.018	<0.001

$r^2=0.919$; $N=720$; $F_{5,174}=1620.962$; $P<0.001$.

Parameter	VNH_4-N			
	Coefficient	SE	F-ratio	P
Constant	-1.098			
Log DW	0.567	0.047	147.610	<0.001
Chl- <i>a</i>	0.191	0.049	15.233	<0.001
Origin	0.108	0.031	11.903	<0.01
TPM	-0.110	0.029	13.951	<0.001

$r^2=0.726$; $N=180$; $F_{4,175}=139.610$; $P<0.001$.

derived accounted for 91.9% of the variance, which in turn was mostly explained by the logarithm of the weight ($P<0.001$). The origin factor also showed a highly significant effect whereas temperature, quality of seston ($Q_s=POM/TPM$) and in particular chlorophyll-*a* were the environmental parameters that presented significant influence on the oxygen consumption rate as well (see Table 3).

Oxygen consumption vs chlorophyll-*a* concentration

A significant positive linear relationship was established between oxygen consumption rate and chlorophyll-*a* concentration for the whole experimental time (Figure 3). There was no differences between equations of both groups of mussels (collector ropes and rocky shore), therefore it can be shown as a common linear equation (Figure 3):

$$VO_2 = 0.279 \pm 0.035 \text{ Chl-}a + 0.243; \quad N=30; \quad r^2=0.694; \quad P<0.001 \quad (4)$$

Ammonia excretion (VNH_4-N)

Ammonia excretion values of mussels from both origins during the experimental culture period are shown in Table 2.

At the beginning of the experiment, VNH_4-N was 48% higher for rocky shore mussels than those from collector ropes ($P<0.01$; Table 2), and no significant differences were observed after 8 d of cultivation

Figure 3. Relationship between oxygen consumption rate and chlorophyll-*a* concentration throughout the cultivation time for both groups of mussels. The common linear equation corresponding to both groups of mussels together is shown in the text.

Figure 4. Relationship between O:N index and chlorophyll-*a* concentration during the experimental time for rocky shore and collector ropes mussels. The equations corresponding to the figure are shown in the text.

($P>0.05$). After 15 d, VNH_4-N continued to be higher in the rocky shore specimens ($P<0.05$), and this difference persisted over most of the cultivation period, although significant differences could only be observed at certain times (Table 2).

Ammonia excretion vs cultivation time

Variation of ammonia excretion for 1 g of tissue dry weight class over the experimental period is shown in Figure 2B. As was the case for oxygen consumption, ammonia excretion decreased between the beginning of the experimental period and day 36. Excretion values increased during January and February, but unlike VO_2 , dropped again during the spring and summer months, returning to values close to those observed during winter.

Ammonia excretion vs dry weight and environmental parameters

The effect of mussel growth (weight), their origin, and environmental factors on the ammonia excretion variation was studied by means of a multiple regression analysis (Table 3). The model accounted for 72.6% of the variation in VNH_4-N , which in turn mostly corresponded to the logarithm of the weight ($P<0.001$). As was the case for VO_2 , origin factor showed significant influence whereas environmental parameter importance was

represented by the effect of chlorophyll-*a* concentration and total particulate matter.

O:N index

The ratio between the oxygen consumption and the ammonia excretion rates, O:N index calculated according to their atomic equivalents, indicates the proportion of protein in relation to lipids or carbohydrates that is catabolised for the energy requirements of the metabolism. Thus, a high rate of catabolism of proteins as compared to lipids or carbohydrates is expressed by a low O:N ratio.

The highest O:N index values corresponded to the collector rope mussels for most of the experimental period (Table 2), which may indicate that mussels with this origin had a more favourable energy metabolism.

The O:N index values increased in both types of mussel throughout the experimental period, coinciding with a greater availability of food in the spring and summer months, and one of the effects due to this factor is a higher accumulation of reserve tissues in the mussels. When the O:N index is plotted against a factor of food quality like chlorophyll-*a* concentration, good concordance was achieved throughout the experiment (Figure 4). The corresponding functions for these relationships are:

Collector ropes: $O:N = 40.71 + 44.42 \pm 13.23 \text{ Chl-}a$; $N = 15$
 $r^2 = 0.464$ $P < 0.01$

Rocky shore: $O:N = 31.70 + 34.14 \pm 11.04 \text{ Chl-}a$; $N = 15$
 $r^2 = 0.424$ $P < 0.01$

The covariance analysis performed for the linear equations showed no significant differences between the slopes of both groups ($t = 0.213$; $df = 26$; $P > 0.05$; common slope: 39.281). However, the intercept obtained for collector rope mussels was significantly higher than that for the rocky shore specimens ($t = 2.181$; $df = 27$; $P < 0.05$), which means higher O:N values for the former group when chlorophyll-*a* is low, as was the case in wintertime at the onset of the experiment.

DISCUSSION

Oxygen consumption

The oxygen consumption rate showed a clearly seasonal pattern, with low values being recorded in winter and maximums during spring and early summer. A similar seasonal pattern to that of the present study has been observed on many previous occasions (Bayne & Widdows, 1978; Widdows, 1978; Navarro & Thompson, 1996; Smaal et al., 1997). These fluctuations have been related to such factors as temperature, the availability of food, and above all, reproductive condition.

Multiple regression analysis provided a model that accounted for 91.9% of the variation in oxygen consumption based on weight, origin of seed and environmental factors quality of food, temperature and chlorophyll-*a*. The important role of multifactorial analyses in order to explained physiological variations should be noted; such changes are difficult to explain unless they are associated

with specific major factors such as temperature and gametogenic activity (Iglesias & Navarro, 1991).

In this study, temperature fluctuated in a very narrow range (2.7°C between maximum and minimum values), though oxygen consumption rate showed a clear seasonal variation. The significant effect of temperature in the multiple model can be considered residual (Table 3). However, the importance of quality of food ($Q_1 = \text{POM/TPM}$) and greatly chlorophyll-*a* (Figure 3) on VO_2 variation was quite clear. High respiration values throughout spring and early summer can be related to an increase of food quality, which in turn should be interpreted according to the findings of Navarro & Thompson (1996). These authors reported that low values of food quality (corresponding to a higher proportion of inorganic matter in the total particulate matter) lead to a lessening of metabolic costs, either as a result of a drop in digestive activity and growth when food quality is low (Bayne & Widdows, 1978; Bayne et al., 1989), or due to the mechanical response of valve closure when there is a high concentration of particles in the water column.

The influence of gametogenesis processes on the oxygen consumption has been widely reported (see Bayne & Widdows, 1978; Widdows, 1978; Smaal et al., 1997 for some examples). In this survey we observed that mussels from both origins became reproductively active in February–March 1996 when they reached 30–40 mm size class (100–200 mg of dry weight of tissue). In terms of chlorophyll-*a*, the first peak in February (Figure 1) coincided with a high increase of oxygen consumption rate after wintertime. With our experimental design, it is difficult to distinguish between both factors (reproductive stage and chlorophyll-*a*) on their importance because reproductive condition was roughly assessed by sight observations.

The inversion of oxygen consumption differences between both groups of seed from day 22 up to day 110 of culture with higher values for rocky shore mussels, relative to first 15 days, may be interpreted according to the increase of clearance and ingestion rates for this type of seed (Babarro et al., in press, b), which would lead to a higher metabolic expenditure. Shick et al. (1988) observed that animals that inhabit rocky shore environments exhibit a characteristic compensatory trait related to energy conservation, given that the environment of these specimens is characterized by desiccation, osmotic stress and temperature changes that would all affect the oxygen consumption rate. Additionally condition index values for the rocky shore mussels were clearly lower for the first two weeks on the raft (Figure 1) which could partly account for these initial differences in VO_2 between both groups of mussels.

Other factors that might have showed a significant influence on initial differences like detachment–reattachment of seed mussel from the experimental cultivation ropes, can be considered negligible with our experimental design. Although we did not quantify this energetic demand to re-establish byssal attachment, it is possible to consider that this factor affected both groups of animals in a similar magnitude by removing both seed from their original habitats (collector ropes and rocky shore) and allowing them to re-attach again by standard technique used by the mussel farmers (see Material and Methods).

Ammonia excretion

The high initial values of ammonia excretion for both types of seed should be attributed to a state of unsettlement that might cause an initial increase in protein catabolism that quickly fell again after 8 d. A more sustained increase in $\text{VNH}_4\text{-N}$ was observed between January and March, which indicates a similar trend to that established by Bayne & Widdows (1978) for *Mytilus edulis* between the winter and spring periods.

Importance has traditionally been attached to the effect that factors such as temperature and energy reserves cycle have on the variation of $\text{VNH}_4\text{-N}$, since they are closely linked to and have a direct bearing on the reproductive cycle (Bayne & Scullard, 1977; Bayne & Widdows, 1978; Bayne & Newell, 1983; Smaal et al., 1997). The relationship reproductive cycle vs ammonia excretion involves an increase in the protein metabolism, and therefore, in excretion when the gonads reach maturity. Additionally, when glucose reserves are scarce, which can occur in wintertime, an additional energy contribution from protein catabolism may well be needed in the January–March period when the energy produced by the metabolism of glucose could prove to be insufficient, since it could be mainly absorbed by gametogenesis (see above, oxygen consumption). This proposed explanation for the increase in the excretion rate (shortage of glucose reserves and the use of protein metabolism) is in agreement with the findings of other studies under conditions of food scarcity or nutritional stress (Gabbott & Bayne, 1973; Bayne & Widdows, 1978).

The significant effect of the seed origin in the present study is in agreement with results obtained under laboratory conditions for collector rope and rocky shore mussels (Labarta et al., 1997). In the present study the rocky shore mussels also showed higher ammonia excretion values than those for collector rope specimens, and the extension of these differences over the most part of the experimental time may indicate, as has been suggested by Okumus (1993), a slow process of adaptation for this particular physiological rate, which would include osmotic adjustments in which the pool of free amino acids plays an important role. This may be especially important in littoral habitats, with periods of exposure to the air, and for this reason the disturbances caused by transplanting these specimens to the cultivation conditions that exist when suspended from a raft may well last for some considerable time.

O:N index

As an integration of oxygen consumption and ammonia excretion rate fluctuations, the O:N index showed a seasonal variation, with low values in winter and a gradual increase in spring. Low O:N index values have been associated with situations of limited availability of food (Kreeger, 1993) and this could be associated in our study with wintertime. In spring and early summer, a quite considerable increase in chlorophyll-*a* content implied higher O:N index values (Figure 4).

Widdows (1985) pointed out that O:N values below 30, which correspond to those of rocky shore origin at the beginning of the experimental period, lie on the threshold of stress. The reason for this is that whilst

the oxygen consumption rate has been described as diet-dependent and drops dramatically within inanition situations (Bayne et al., 1989), the excretion rate increases when food is scarce, as a result of the increase in protein catabolism (Gabbott & Bayne, 1973; Bayne & Scullard, 1977). Bayne & Scullard (1977) have suggested that the use of the O:N index as a stress indicator is only appropriate if the seasonal cycles of gametogenesis/energy reserves are taken into account, since although feeding conditions may be excellent, the O:N index can decrease during the spawning period (Widdows, 1985).

In this study, collector rope mussels showed higher values than the rocky shore specimens for this index, which can be interpreted as an indicator of the better nutritional condition of the former specimens. Kreeger (1993) described low O:N index values in states of nutritional deprivation, which are characteristic of areas where food is scarce, as is the case of the original habitat for rocky shore mussels. The availability of food is the major differential factor with regard to the characteristics of the original habitats in the present study (collector rope and rocky shore), and the differences observed in the O:N index at the beginning of the experimental period (42–85% higher for the collector rope mussels for the first two weeks), together with the wide variation in these differences throughout the experiment, which are nevertheless almost always in favour of the collector rope mussels, may be accounted for by the persistence of ammonia excretion differences in spite of a long period of acclimatization. The present experiment began in winter, a period in which the possible differences in protein metabolism between the two groups under study are thought to be at their lowest. However, with the arrival of a period with greater availability of food, the differences in this metabolism may be enhanced, especially once the carbohydrate reserves have been deviated to reproductive processes.

The differences in the O:N index throughout a large part of the experimental period related to ammonia excretion behaviour are indicative of slow process of adaptation for this physiological rate when specimens are subjected to new environmental and/or nutritional conditions, as occurs when rocky shore mussels are transferred to a raft for cultivation.

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