ORIGINAL RESEARCH



A sensory and nutritional validation of open ocean mussels (*Mytilus galloprovincialis* Lmk.) cultured in SE Bay of Biscay (Basque Country) compared to their commercial counterparts from Galician Rías (Spain)

K. Azpeitia · Y. Ríos · I. Garcia · J. Pagaldai · D. Mendiola

Received: 4 January 2017/Accepted: 17 March 2017/Published online: 27 April 2017 © The Author(s) 2017. This article is an open access publication

Abstract Mussel shell biometry, nutritional quality as well as consumer sensory evaluation of experimental open ocean cultured mussel *Mytilus galloprovincialis* (Lamarck 1819) were analyzed and compared to that of commercial mussels from Galician Rías available in the local market. Both mussel products were of the same commercial size. In this study, open ocean mussels were significantly higher and wider than those of Galician Rías. In addition, with the exception of ash content, both mussel products showed similar biochemical composition. Regarding fatty acid profiles, however, statistical differences were detected. These differences were not fully reflected in the sensory assessment. In terms of consumer acceptability, both mussel products were considered equally satisfactory.

Keywords Fatty acids · Mussel · Mytilus galloprovincialis · Open ocean aquaculture · Proximate composition · Sensory analysis

Introduction

Aquaculture is a seafood supply that has increased gradually over the last few decades (Langan and Horton 2003). This growth is due to progress in technology, changes in policy, and an increased societal awareness of sustainability (Bushek et al. 2004; Coen et al. 2007; Costa-Pierce 2008; Forrest et al. 2009; Bostock et al. 2010). In particular, mollusk aquaculture represents more than 75% (13.9 million tons) of the world's aquaculture, with mussel production representing around 13% (1.8 million tons) of its annual production (FAO 2014). Spain, France, and the Netherlands produce one-third of the global mussel aquaculture production (Buck et al. 2010). Specifically, Spain, with an average production near 200,000 tons per year, is the second largest producer worldwide and the first in Europe (FAO 2014). A large part (95%) of its annual production is produced in the Galician Rías from the SW Bay of Biscay (Galician region). Since 2008, mussel products from that area have had a Protected Designation of Origin certifying quality and traceability within the standards of the EU seafood policy.

Shellfish species are a valuable food source in the socioeconomic context of the Basque Country (SE Bay of Biscay) (Gracia 1996), yet mussel aquaculture is limited. Indeed, over the last few decades, every

K. Azpeitia · D. Mendiola (⋈)

Marine Research Division, AZTI-Tecnalia, Herrera Kaia; Portualdea, s/n, 20110 Pasaia, Spain e-mail: dmendiola@azti.es

Y. Ríos · I. Garcia · J. Pagaldai AZTI-Tecnalia, Astondo Bidea, Edificio 609, Parque Tecnológico de Bizkaia, 48160 Derio, Bizkaia, Spain





mollusk (especially bivalves) consumed in this area has been imported. The lack of development of an aquaculture or shellfish sector in the region of the Bay of Biscay is mainly linked to high wave-energetic scenarios (Galparsoro et al. 2012) and previous issues related to estuarine water quality (Chust et al. 2009). A further restriction is current legislation, which aims to prevent conflict between users of marine habitats. These conflict-prevention methods hinder the development of near-shore aquaculture as the limited inshore spaces are protected marine habitats (Galparsoro et al. 2010; Borja et al. 2011; Pascual et al. 2011).

In addition to these local limitations, the main obstacles in mussel aquaculture in general are those regarding infectious diseases and parasites (Bower et al. 1994), marine phycotoxins (Svensson 2003; Moroño et al. 2003) as well as carrying capacity and competition for the use of marine space (Martínez et al. 2007; Costa-Pierce 2008; Byron et al. 2011). Limited marine space coupled with commercial objectives often results in non-sustainable practices for commercial mussel cultivation (Smaal 2002; Waite et al. 2005; Duarte et al. 2008). It is encouraging, however, that ecosystem-based approachmanagement strategies have been recommended to promote the sustainable development of mussel cultivation (Soto et al. 2008).

One such strategy is open ocean aquaculture which represents a promising alternative for the sustainable cultivation of bivalve species (Polk 1996; Hesley 1997; Stickney 1998, Bridger and Costa-Pierce 2003). However, this activity is still in development stages (Hickman 1992), and main mussel culture techniques (i.e., rafts, pole racks and/or longline systems) have been established almost exclusively in protected near-shore waters (Pérez-Camacho et al. 1991; Myrand et al. 2009) or in estuarine habitats (Penney et al. 2002; LeBlanc et al. 2005; Drapeau et al. 2006). Therefore, few studies have reported successful biological and/or engineering results on open ocean mussel aquaculture to date (Langan 2001; Langan and Horton 2003; Buck 2007).

Open ocean aquaculture is not only a financially attractive but also an environmentally sound one with a highly nutritious end product for the consumer. Indeed, there is an increasing demand for products sourced from open ocean aquaculture methods, resulting in increased revenue in international markets (Orban et al. 2002; Fuentes et al. 2009; Pogoda et al. 2013). Further, developing open ocean aquaculture methods will be environmentally beneficial (Shumway et al. 2003), as mussels and other bivalves are filter feeders, capturing and eating microscopic particles, plants, and nutrients by filtering, and thus cleaning ocean water (Shpigel et al. 1993). For the consumer, as most farmed mussels have fast growth, they provide year-round nutritious food, much in demand (Seed and Suchanek 1992; Gosling 2003). Mussels are appreciated by humans for their organoleptic properties (Grienke et al. 2014) as well as for their health benefits. They provide high-quality protein, minerals, essential trace elements, fat-soluble vitamins (i.e., vitamin D), and essential fatty acids EFA, especially long-chained polyunsaturated fatty acids: omega-3 (ω3 LC-PUFAs) (Stankovic et al. 2011; James 2013).

Regarding consumption, mussel shape and appearance are decisive factors for consumers' purchase (Fuentes et al. 2009), whereas meat quality (and hence, apparent health benefits) is regulated by biochemical composition (Orban et al. 2002; Filgueira et al. 2006; Fuentes et al. 2009). Mussel meat quality has been described by consumers as the result of not only chemical and biological characteristics, but also organoleptic properties, such as the appearance of the muscle, the intrinsic flavour, and the absence of undesirable components (Vernocchi et al. 2007). As reported by Fuentes et al. (2009) mussel organoleptic characteristics can be origin dependent. This means that different production sites, with different conditions and culture technologies, are expected to promote changes in growth performance and biochemical composition of mussel individuals, which could in turn reflect different product qualities at consumer level (Oliveira et al. 2015).

To date, no studies have investigated organoleptic properties on open ocean mussel products from the SE Bay of Biscay. Both biochemical composition and organoleptic properties are of special importance to anticipate consumer acceptance and market feasibility (Gökoglu 2002; Oliveira et al. 2015) before considering any implementation of mussel activity in the region.

The specific goals of this study were to characterize and compare the biometric parameters, nutritional content, and sensory aspects between experimental mussel product from the SE Bay of Biscay (open ocean) and commercial mussel product from the SW Bay of Biscay (Galician Rías).



Materials and methods

Location and sampling

An experimental suspension longline culture system (Langan 2000) was developed to cultivate shellfish in the open waters of the south eastern corner of the Bay of Biscay (Basque region; Spain). This region extends over 150 km and it is oriented E–W. The climate is temperate, oceanic, with moderate winters and warm summers. According to Köppen's classification, the area is associated with a marine temperate climate (Cfb) (Collins and Borja 2004). Seeds $(17.40 \pm 5.07 \text{ mm})$ were naturally collected during June 2013 and cultivated at the experimental site $(43^{\circ}21.39'\text{N}; 02^{\circ}26.90'\text{W})$ over a 1-year period. The open ocean site is already classified as a Class A production area (according to EU regulation (EC) 854/2004). Therefore, those mussels are not subjected to depuration requirements. The site is located at 2 nautical miles off the coast without any protection from land masses (Ryan 2004). Galparsoro et al. (2012) described strong hydrodynamic conditions on Basque Country's coast. A detailed description of current and waves in the experimental site can be found in Azpeitia et al. (2016).

Mussel individuals were cultured within 17 m water column depth, on mussel-growing ropes of 12 m in length. Mussel ropes hung from 5 m headline depth, with a distance between ropes of 0.5 m and an initial stocking density of 681 ± 37 individuals per lineal meter. After 13 months of cultivation, in July 2014, 200 mussel specimens of "commercial size" (Peteiro et al. 2006, i.e., 70 mm) were carefully detached from the ropes by cutting the byssal threads and cleaned of encrusting organisms. Selected mussel individuals were transported in plastic self-draining boxes covered by flaked ice. Later in the laboratory, mussels were conserved in plastic boxes, stored at 4 °C and covered with flaked ice for 24 h previous to analysis to preserve them. Similarly, 200 mussel individuals from Ría de Arosa (Galician Region; Spain) with a comparative commercial size (69.65 ± 6.11 mm) were obtained at early hours on the morning of the day of the sensory analysis. Commercial mussels were bought from a local seafood retailer to ensure a comparable size product with the same conservation procedure (plastic box and stored at 4 °C). These mussels were also stored at 4 °C and covered with flaked ice for some hours previous to analysis. Subsequently, two stocks of 50 mussel individuals from Galicia and from the experimental open ocean site were frozen at -80 °C to await morphometric and biochemical analysis.

Morphometry

Twenty mussel individuals from each origin (experimental open ocean and commercial) were carefully detached by cutting the byssal threads, cleaned of encrusting organisms and measured for morphometric analysis. The individuals were measured for their shell length (L; maximum anterior—posterior axis), shell height (H; maximum dorso-ventral axis), and shell width (W; maximum lateral axis) using a digital caliper to the nearest 0.1 mm. The biometric terminology used in this study is that described by Seed (1968). Mussel shape was analyzed by linear allometric relation between log-transformed mussel shell dimensions [length (L), height (H) and width (W)].

Biochemical composition and fatty acids

Three replicate samples of 10 pooled mussel individuals from each origin were considered for biochemical characterization (including proximate and glycogen analysis) and lipids determination (including total lipids and fatty acids), respectively.

The fresh individuals were first minced in a food processor (IKA® M 20 universal mill, IKA 1603601, Germany). Moisture and ash were determined according to the gravimetric determination. Briefly, minced tissue pools were placed in pre-weighed porcelain trays for drying at 80 °C for 24 h and then weighed to the nearest 0.001 g. Subsequently, dry mussels' tissue pools were ashed at 450 °C for 4 h in a muffle furnace, cooled in a desiccator and weighed again to determine the ash-free dry weight content.

Total protein was determined according to the Kjeldahl's method (AOAC 1975). Samples were digested in a digestor BUCHI B-435, distilled using a Rotavapor[®] R-210 (BÜCHI Labortechnik AG, Switzerland) and automatic volumetric titration developed through a Mettler Toledo T- 50 (Mettler Toledo, United States).



Glycogen concentration (mg·g⁻¹ mussels' tissue) was calculated based on a mussel glycogen standard (Sigma, Saint Louis, MO) following the methodology reported by Gallardi et al. (2014). Briefly, 30% KOH was added to 0.5 g of homogenized mussels' tissue (ratio 2:1). The samples were then heated in a shaking-water bath at 100 °C for 20 min, vortexed for 30 s and subsequently chilled on ice for 5 min. After cooling, 200 μL of each sample was transferred to a 4 mL glass vial followed by 200 μL of 95% ethanol. The solution was vortexed again briefly and then placed in a boiling-water bath for 15 min followed by the addition of 1.2 mL of lukewarm water. The samples were again briefly vortexed and then allowed to stand at room temperature for 5 min before measuring the glycogen content. Glycogen content was measured by colorimetric reaction using 25 μL of prepared sample. 10 μL of 80% aqueous phenol and 200 μL of sulphuric acid were added to the samples on 96-well plates. Absorbance was measured using a multi- detection microplate reader (Synergy HT, BIO-TEK) at 490 nm. The concentration of glycogen in the samples was calculated based on a mussel glycogen standard (Sigma, Saint Louis, MO).

Total lipids were determined based on Blight and Dyer method (Hanson and Olley 1963). In this way, total lipids from a known weight of tissue was extracted and purified by homogenization in a suitable excess of chloroform/methanol/water (2:2:1.8 v/v/v). Following this, total lipids in chloroform solution were determined by weighting the eluate after removal of the solvent by rotary evaporation. All lipid extracts were then stored in chloroform at -30 °C to await analysis.

Fatty acid methyl esters (FAME) were separated and quantified using an Agilent 7890 Gas Chromatographer equipped with auto-injector (Agilent Technologies, United States). The gas chromatograph is equipped with a flame-ionization detector (290 °C) and a DB-WAS 122-7032 Agilent Technologies capillary column (30 m _ 0.25 mm I.D., film 0.25 μm). Helium was used as a carrier gas and initial oven temperature was 50 °C, followed by an increase at a rate of 10 °C·min⁻¹ to a final temperature of 240 °C for 9 min. Individual FAME were identified by comparison of retention times with four commercial fatty acid standards (PUFA 1, PUFA 3, BAME and 37-component) supplied by Supelco (Bellefonte, PA) for area percent normalization. At each sample relative quantities were expressed as weight percent of total fatty acids.

Sensory evaluation/consumer testing

Thirty regular domestic consumers of mussel were randomly recruited by email from AZTI's database. They were segmented by gender (50% men) and age range (18-70 years). To remove debris and any remaining byssal thread, 150 mussels were washed and scrubbed for the sensory analysis. Samples were steamed at 100 ± 1 °C for 1 min and then served at 65 ± 1 °C. Mussels were presented to consumers on white plastic plates labeled with 3-digit random numbers, containing two mussels from each origin (experimental from open ocean and commercial from Galician Rías). Consumers tasted one mussel from each origin. The consumers received oral and written instructions and were provided with water and unsaltedcrisp bread for palate cleansing after each tasting. Each consumer tasted the samples in randomized order and evaluated five attributes (appearance, odor, taste, texture, and global acceptance) with a structured 9-point scale ranging from 1 (equivalent to I dislike it very much) to 9 (equivalent to I like it very much). The 9-point hedonic scale (Peryam and Girardot 1952; Peryam and Pilgrim 1957) is the most commonly used scale for testing consumer preference and acceptability of foods (Schutz and Cardelo 2001). This kind of scales produce central tendency effects on panelist, which reduces the options to a 7-point scale (Villanueva et al. 2005). Despite this possible reduction, a 9-point scale with an arithmetic adjustment can always be straightforward re-scaled to a 7-point scale (Epler et al. 1998). This enables the comparisons with different scale formats with less response options (Dawes 2012), which are common in current scientific literature. Consumers' purchase intention was also assessed with a structured 5-point scale ranging from 1 (equivalent to I would not buy it) to 5 (equivalent to I will buy it). The same test was repeated three times with each panelist. Consumers tasting was performed in the sensory laboratory of AZTI- Tecnalia specifically designed according to ISO 8587:2006.

Statistical analysis

Relative shell morphology was investigated through the general linear function, where y and x are size-related measures and a and b are constants (i.e., intercept and slope; which represents the coefficient of relative shape)



(Pauly 1983). Similarly, allometric equations were generated separately for mussels from each origin, using linear regression and logarithmic transformation. The differences between allometric coefficients (b) from each origin were evaluated using ANCOVA (Rohlf and Sokal 1981).

Means and standard deviations (mean \pm SD) of data on morphometrics, proximate composition (humidity, ash, protein, glycogen, and lipid content), as well as fatty acid profiles, of mussels were plotted using Sigmaplot (11.0) statistical and graphical software (Systat software). Statistical analyses were carried out using a statistical package (SPSS 20.0 for Windows®). Sensory attributes and evaluation data were analyzed using the R-project software (R Development Core Team 2008). A significance level of 0.05 was used. Normality was tested by Shapiro–Wilk test and the homoscedasticity was checked using Leven's test. The differences between mussel products type (experimental and commercial products) were compared using independent sample t test. Arcsine transformation (Zar 1984) was applied to the data expressed as percentage.

Results

Morphometry

The biometric results of the two mussel products (experimental open ocean and commercial from Galician Rías) tested within the present study are shown in Table 1. No statistical differences between the shell lengths (L) of both mussel products were detected. Conversely, significant differences were detected for the measurements of shell height (H) and shell width (W) between both mussel products. As observed, the open ocean mussel individuals from the present study displayed significantly higher morphometric values than those from the commercial samples.

The allometric relationships between shell length (mm) and shell width (mm), shell length and shell height (mm), or between shell width (mm) and shell height (mm) of both mussel products are shown in Table 2. Based on ANCOVA analysis (Table 3), the factor of product type significantly affected both the linear relations between log-transformed shell length (mm) and log-transformed shell height (mm), as well as the linear relation between log-transformed shell length (mm) and log-transformed shell width (mm). Higher values were found in the experimental mussels. Finally, no ANCOVA analysis was performed to compare the linear relation between log-transformed shell height (mm) and log-transformed shell width (mm) as significant interaction between the factor (product type) and the covariable [log-transformed shell width (mm)] was observed. Hence, no direct relation between the change in shell width (mm) and the change in shell height (mm) among mussel products could be assessed.

Table 1 *M. galloprovincialis.* Biometric characteristic of the mussel shell and proximate composition of mussel tissue pools from experimental open ocean (SE the Bay of Biscay; Basque

region) and sheltered Rías (SW Bay of Biscay; Galician region) conditions

	Experimental	Commercial	One-way AN	OVA
	Open ocean	Galician Rías	\overline{F}	P value
L (mm)	70.29 ± 7.96	69.65 ± 6.11	0.037	0.849
H (mm)	$37.56 \pm 4.59*$	33.58 ± 2.48	10.471	0.003
W (mm)	$24.58 \pm 4.04*$	21.18 ± 4.59	5.538	0.024
Moisture (%)	78.77 ± 1.54	76.57 ± 1.77	2.661	0.178
Protein (dry weight %)	49.52 ± 4.29	44.79 ± 4.02	1.962	0.234
Lipid (dry weight %)	6.27 ± 1.47	6.30 ± 1.34	0.001	0.976
Ash (dry weight %)	13.00 ± 1.16 *	9.40 ± 0.82	19.253	0.012
Carbohydrates (dry weight %)	31.21 ± 4.87	39.51 ± 6.07	3.356	0.141

Mean \pm standard deviation, n = 20



^{*} Statistical significance was set up at (P < 0.05) (one-way ANOVA)

Table 2 Mytilus galloprovincialis. Parameter of shell shape (length (L), width (W) and height (H)) regression equations used to relate shell length (mm) to shell height (mm) or shell width (mm), and shell width (mm) to shell height (mm) of different mussel products (experimental vs. commercial);

equation fitted to $\log_{10} X = \log_{10} a + b \log_{10} Y$, where a intercept and b slope, and X and Y shell measurements. Error of correlation coefficient (r), number of specimens (n) used to derive equation are also given

	Mussel product		SE	t	Sig	95% CI		R^2	N
						Upper	Lower		
Experimental	Intercept	-0.185	0.224	-0.823	0.421	_	0.287	0.773	20
	Slope	0.952	0.122	7.834	0.000	0.697	1.208		
Commercial	Intercept	0.390	0.241	1.620	0.123	-0.116	0.897	0.552	20
	Slope	0.616	0.131	4.709	0.000	0.341	0.891		
Experimental	Intercept	-0.874	0.284	-3.075	0.007	_	_	0.882	20
	Slope	1.225	0.154	7.951	0.000	0.901	1.549		
Commercial	Intercept	-0.501	0.243	-2.066	0.053	-1.011	0.008	0.873	20
	Slope	1.000	0.132	7.595	0.000	0.724	1.277		
Experimental	Intercept	0.735	0.161	4.557	0.000	0.396	1.073	0.775	20
	Slope	0.604	0.116	5.197	0.000	0.360	0.848		
Commercial	Intercept	0.790	0.149	5.287	0.000	0.476	1.104	0.757	20
	Slope	0.548	0.111	4.917	0.000	0.314	0.782		
	Commercial Experimental Commercial Experimental	Commercial Slope Experimental Intercept Slope Commercial Intercept Slope Commercial Intercept Slope Experimental Intercept Slope Commercial Intercept Slope Commercial Intercept	Slope 0.952	Slope 0.952 0.122	Slope 0.952 0.122 7.834 Commercial Intercept 0.390 0.241 1.620 Slope 0.616 0.131 4.709 Experimental Intercept -0.874 0.284 -3.075 Slope 1.225 0.154 7.951 Commercial Intercept -0.501 0.243 -2.066 Slope 1.000 0.132 7.595 Experimental Intercept 0.735 0.161 4.557 Slope 0.604 0.116 5.197 Commercial Intercept 0.790 0.149 5.287	Slope 0.952 0.122 7.834 0.000	Experimental Intercept Slope -0.185 0.224 -0.823 0.421 - Commercial Intercept Slope 0.952 0.122 7.834 0.000 0.697 Commercial Intercept Slope 0.390 0.241 1.620 0.123 -0.116 Experimental Intercept -0.616 0.131 4.709 0.000 0.341 Experimental Intercept -0.874 0.284 -3.075 0.007 - Slope 1.225 0.154 7.951 0.000 0.901 Commercial Intercept -0.501 0.243 -2.066 0.053 -1.011 Slope 1.000 0.132 7.595 0.000 0.724 Experimental Intercept -0.735 0.161 4.557 0.000 0.396 Slope 0.604 0.116 5.197 0.000 0.360 Commercial Intercept -0.790 0.149 5.287 0.000 0.476	Experimental Intercept -0.185 0.224 -0.823 0.421 - 0.287 Slope 0.952 0.122 7.834 0.000 0.697 1.208 Commercial Intercept 0.390 0.241 1.620 0.123 -0.116 0.897 Slope 0.616 0.131 4.709 0.000 0.341 0.891 Experimental Intercept -0.874 0.284 -3.075 0.007 - - Slope 1.225 0.154 7.951 0.000 0.901 1.549 Commercial Intercept -0.501 0.243 -2.066 0.053 -1.011 0.008 Slope 1.000 0.132 7.595 0.000 0.724 1.277 Experimental Intercept 0.735 0.161 4.557 0.000 0.396 1.073 Slope 0.604 0.116 5.197 0.000 0.360 0.848 Commercial Intercept 0.790<	Experimental Intercept -0.185 0.224 -0.823 0.421 - 0.287 0.773 Commercial Intercept 0.952 0.122 7.834 0.000 0.697 1.208 Commercial Intercept 0.390 0.241 1.620 0.123 -0.116 0.897 0.552 Slope 0.616 0.131 4.709 0.000 0.341 0.891 Experimental Intercept -0.874 0.284 -3.075 0.007 - - 0.882 Slope 1.225 0.154 7.951 0.000 0.901 1.549 Commercial Intercept -0.501 0.243 -2.066 0.053 -1.011 0.008 0.873 Experimental Intercept 0.735 0.161 4.557 0.000 0.396 1.073 0.775 Experimental Intercept 0.604 0.116 5.197 0.000 0.360 0.848 Commercial Intercept 0.790

Table 3 *Mytilus galloprovincialis.* Mussel product type effects on shell length (mm) to shell width (mm) and on shell length (mm) to shell height (mm), allometric relations

analyses. Regression equation exponent b (slope) were compared using ANCOVA (Rohlf and Sokal 1981), with α set at 0.05

Allometric relation	Source of variation	ANCOVA			R^2
		df	F	Sig	
Log H/Log L	Log length (covariable)	1	83.561	0.000	0.759
	Product type (factor)	1	29.992	0.000	
	Error	37			
Log W/Log L	Log length (covariable)	1	121.922	0.000	0.803
	Product type (factor)	1	19.953	0.000	
	Error	37			

Biochemical composition and fatty acids

The proximate composition of both mussel products is shown in Table 1. Moisture levels were relatively high within both samples. Mussels from commercial origin (Galician Rías) showed slightly higher moisture values (78.77% vs. 76.57%) than mussels from the open ocean conditions of the present study, yet significant differences were not detected.

Significantly lower ash levels were detected within the commercial individuals (13.00% vs. 9.40%) compared to the study ones.

Lipid contents were slightly higher in the commercial samples (6.27% vs. 6.30%), although no significant differences were found. Protein contents did not show any significant differences between mussel products. Finally, glycogen levels (71.06 \pm 16.71 mg·g⁻¹ and 100.59 \pm 25.05 mg·g⁻¹ at experimental and commercial mussels, respectively) were slightly higher in commercial samples, although differences were not statistically significant.

The FA composition of both mussel products are shown in Table 4. The saturated fatty acids (SAFA) accounted for 29% in experimental mussels and significantly lower content (25%) for commercial mussels. Saturated fatty acid predominated over the monounsaturated fatty acid (MUFA) with 12% for the experimental and significantly higher level (22%) for the commercial samples. Both mussel products showed the



Table 4 *M. galloprovincialis.* Fatty acid compositions (percentage of total fatty acids) of mussel tissue pools analyzed during the comparative study between the experimental open

ocean mussels from the SE Bay of Biscay and commercial mussels obtained from the sheltered Rías from Galicia

Fatty acids	Experimental open ocean	Commercial Galician Rías	
14:0	2.52 ± 0.80*	4.14 ± 0.60	
15:0	$0.57 \pm 0.07*$	0.36 ± 0.02	
16:0	$18.91 \pm 1.29*$	15.50 ± 0.43	
17:0	$1.90 \pm 0.32*$	1.12 ± 0.20	
18:0	4.70 ± 0.93	3.74 ± 0.52	
20:0	0.06 ± 0.05	0.02 ± 0.04	
22:0	0.03 ± 0.03	$0.18 \pm 0.03*$	
∑SAFA	28.69*	25.05	
14:1	0.42 ± 0.07	$0.99 \pm 0.17*$	
16:1ω7	2.40 ± 0.87	$11.14 \pm 1.42*$	
16:1ω5	0.30 ± 0.05	0.25 ± 0.02	
17:1	0.19 ± 0.16	0.40 ± 0.10	
18:1ω9	2.22 ± 0.68	1.14 ± 0.05	
18:1ω7	1.22 ± 0.15	$3.19 \pm 0.07*$	
20:1 ω11	1.21 ± 0.23	1.32 ± 0.17	
20:1ω9	$3.48 \pm 0.28*$	1.59 ± 0.19	
20:1ω7	0.47 ± 0.05	$2.03 \pm 0.16*$	
22:1	0.06 ± 0.06	0.24 ± 0.14	
∑MUFA	11.98	22.28*	
16:4ω3	8.32 ± 2.25	3.04 ± 4.70	
18:2ω6	$2.03 \pm 0.32*$	0.87 ± 0.02	
18:3ω3	$1.82 \pm 0.44*$	0.77 ± 0.09	
18:4ω3	$3.74 \pm 1.23*$	1.35 ± 0.24	
20:2 α	$2.76 \pm 0.36*$	1.57 ± 0.31	
20:2 β	0.25 ± 0.04	$1.14 \pm 0.35*$	
20:2ω6	$0.71 \pm 0.12*$	0.38 ± 0.03	
20:4ω6	1.30 ± 0.21	$2.61 \pm 0.12*$	
$20:4\omega 3$	0.34 ± 0.13	0.64 ± 0.23	
$20.5\omega 3$	9.85 ± 1.06	$24.71 \pm 2.21*$	
22:2 NMIDa	$0.73 \pm 0.10*$	0.12 ± 0.10	
22:2 NMIDb	1.22 ± 0.13	$4.54 \pm 0.43*$	
21:5ω3	1.20 ± 0.16	1.09 ± 0.06	
22:4ω6	0.20 ± 0.04	1.90 ± 1.54	
22:5ω6	$0.61 \pm 0.03*$	0.15 ± 0.15	
22:5ω3	1.12 ± 0.11	$2.13 \pm 0.21*$	
22:6ω3	$23.13 \pm 2.20*$	5.66 ± 0.47	
∑PUFA	59.33*	52.67	
NMID	1.95	4.66*	
Terrestrial $(18:2\omega6 + 18:3 \omega3)$	3.85*	1.64	
$\Omega 3/\Omega 6$	10.21	6.67	
DHA/EPA	2.35*	0.23	

Values (mean \pm SD, n = 10)

NMID Non-methylene-interrupted dienoic fatty acid



^{*} Statistical significance was set up at (P < 0.05) (one-way ANOVA)

majority of fatty acids consisted of polyunsaturated fatty acids (PUFA), with significantly higher level (59%) for experimental mussels than for commercial mussels (52%).

In the case of SAFA, palmitic acid (16:0) was predominant (18.91 and 15.50%) in both products, followed by stearic acid 18:0 in the open ocean mussels (4.70%), and by myristic acid 14:0 in the commercial mussels (4.14%). Significant differences (one-way ANOVA; P < 0.05) in the SAFA profile (i.e., the sum of pentadecanoic 15:0, palmitic 16:0 and behenic 22:0 acids) were found between both mussel products.

Regarding MUFA, the commercial mussel from Rías displayed abundant related palmitoleic acid $16:1\omega7$ (11.4%). This was followed by vaccenic acid $18:1\omega7$ (3.19%) and eicosenoic acid $20:1\omega7$ (2.03%). Conversely, the observed trend of MUFAs in the open ocean experimental individuals was not equivalent; the major acid was gondoic acid $20:1\omega9$ with 3.48%, followed by palmitoleic acid $16:1\omega7$ with 2.40% and oleic acid $18:1\omega9$ with 2.20%. Thus, significant differences (one-way ANOVA; P < 0.05) were found between MUFA of both mussel products.

Regarding PUFAs, eicosapentaenoic (EPA) ($20:5\omega3$) and docosahexaenoic (DHA) ($22:6\omega3$) acids were the most important. Commercial mussels showed higher EPA ($20:5\omega3$) with 24.71% and less DHA ($22:6\omega3$) with 5.66%, whereas the experimental open ocean mussels showed less EPA ($20:5\omega3$) content with 9.85% and higher (DHA ($22:6\omega3$) with 23.13%. Thus, the DHA/EPA ratios were significantly different (one-way ANOVA; P < 0.05) between mussel products.

Non-methylene-interrupted dienoic fatty acid (i.e., 22:2 NMIDa and 22:2 NMIDb) were higher in the commercial samples (4.66%) than in the experimental ones (1.95%). Differences at NMID level were statistically significant (one-way ANOVA; P < 0.05) between mussel products.

Finally, terrestrial fatty acids (i.e., $18:2\omega6 + 18:3\omega3$) accounted for the 3.85% in experimental mussels and 1.64% in the commercial ones; differences between mussel products were statistically significant (one-way ANOVA; P < 0.05).

Sensory evaluation/consumer testing

The panelist male:female ratio (%) participating in the present study was 37.7:63.3. The range of ages between participants was distributed as follows: 18–30 years (13.3%); 31–40 years (40%); 41–50 years (33%); 51–60 years (13.3%). The 77% of participants were regular mussel consumers (3.3% more than once a week; 26.7% once a week; 46.7% once a month), whereas the 23.3% declared to be occasional mussel consumers.

The resulted acceptance for both mussel products was around 7 points out of 9 (Fig. 1). No significant differences (t test; P > 0.05) were detected for any of the sensory attributes analyzed between both products (Table 5). The results of the qualitative assessment showed that both mussel products showed good appearance and odor. The results of the qualitative assessment showed that both mussel products showed good appearance and odor. Particularly, the experimental ones showed to provide more intense sea-flavour with a slightly softer and more juicy texture. Some other aspects such as the whitish color on the muscle tissue and/or the salty taste were criticized when observed. A large proportion of the panelists (70%) expressed potential interest in purchasing the experimental product, while a small minority (17%) did not. The remaining 13% of panelists did not express a purchase preference. For the commercial mussels, 83% of the panelists showed purchase intention, while 10% showed uncertainty (or doubt) and 7% would not buy it (Fig. 2).

Discussion

All mussels consumed in the Basque region, during recent decades, are imported (Gracia 1996). Currently, the only way to cultivate mussels in the Basque Country is offshore and with submerged longline method (Azpeitia et al. 2016). The aim of the study was to compare if experimental open ocean cultured mussels of commercial size were as competitive as local market leader of the same commercial size. No significant differences were observed in shell length between both products, which validated the analysis of quality differences among mussel products as well as the analysis of consumer perception among products. Although panelists were not able to inspect mussel shells, only mussels without visible damage (open valves or broken shell) were used in the study. Direct influences on the appearance of the product should be derived from the differences found at biometric levels between both mussel (*M. galloprovincialis* Lmk.) origins. Different shell



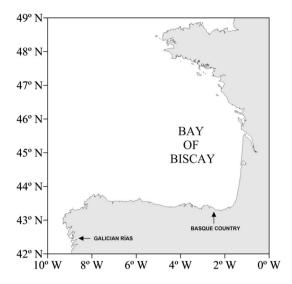


Fig. 1 Map of the Bay of Biscay showing the specific location of the open ocean experimental site at the SE Bay of Biscay (Basque Country) and the commercial mussel product from the SW Bay of Biscay (Galician Region)

Table 5 *M. galloprovincialis.* Student's *t* analysis carried out for all sensorial attributes analyzed in experimental mussel from the open ocean conditions of the SE Bay of Biscay and mussels cultured in at sheltered Rías from Galicia

	Experimental open ocean	Commercial Galician Rías	Student's t	
			t	P value
Appearance	6.467 ± 2.097	7.033 ± 1.066	1.32	0.194
Odor	6.967 ± 1.402	7.3 ± 1.088	1.029	0.308
Flavor	7.533 ± 1.106	7.4 ± 1.303	-0.427	0.671
Texture	7.033 ± 1.586	7.533 ± 0.973	1.472	0.148
Global impression	6.9 ± 1.517	7.333 ± 1.269	1.2	0.235

Values (mean \pm SD, n = 30)

^{*} Statistical significance was set up at (P < 0.05) (pairwise comparison)

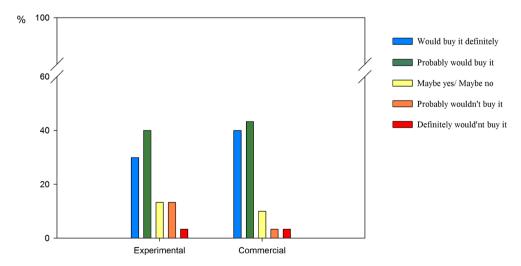


Fig. 2 M. galloprovincialis. Buying-intention results carried out on the comparison between mussels from the open ocean in the SE Bay of Biscay ("experimental") and from sheltered estuaries in the SW Bay of Biscay ("commercial")



shapes among mussel product were observed. Mussels cultured in Galician Rías were found to be thinner (reduced W. L^{-1} ratio) and lower (reduced H. L^{-1} ratio) than experimental open ocean mussels. Commercial mussels may have been cultured at higher densities than experimental mussels. Stocking density on experimental mussels was 681 individuals/m, whereas commercial label does not provide the consumer with those culture characteristics. Nevertheless, literature value for standard mussel density on ropes cultured in sheltered Galician Rías is between 700 and 1000 mussels·m⁻¹ (Irisarri et al. 2014a). Cubillo et al. (2012) observed that mussels cultured at higher densities displayed a lower shell height to shell length ratio compared to mussels cultured at higher density displayed lower shell width to shell length ratio compared to those cultured at low densities.

Regarding differences in oceanographic conditions, mean current speed in the experimental site was significantly higher than values reported from previous studies carried out in Galician sheltered Rías (Pérez-Camacho et al. 2014a, b). Similarly, highly significant wave height scenarios were also found (with peaks of 796 m) at the experimental site. Indeed, only open ocean mussel growth studies of Buck and Buchholz (2004) and Langan (2000), from the German Bight and the Gulf of Maine, respectively, showed similar wave height scenarios (with peaks of 7.96 m). The mussel shells of the experimental site were characteristically round, wide and great in thickness. Although these observations were not validated through specific comparisons, they are consistent with the findings from previous studies developed with mussels from wave beaten shores (Akester and Martel, 2000, Lewis and Powel 1961, Seed 1968).

Despite the limited information available on the culture method and culture characteristic of commercial mussel product, the new product cultured at the experimental site would compete with current commercial products. No differences regarding mussel appearance of cooked samples were pointed out by the group of local panelists during the sensory test.

In this study, no relevant differences at biochemical composition level other than ash content were observed between the open ocean experimentally cultured mussels and the commercially purchased mussels from Galician Rías. Biochemical composition of mussels is affected by water temperature, nutrient availability, and the reproductive cycle of individuals (Fernández-Reiriz et al. 2007). These multiple factors change seasonally, which, in turn, cause seasonal changes in biochemical composition of mussels (De Zwaan and Zandee 1972; Dare and Edwards 1975; Ruano et al. 2012). As reported by previous authors, major differences in biochemical composition relied on: (i) feeding behavior, in terms of filtration capacity and clearance rate (Vilela 1950; Heral et al. 1980); (ii) quality and quantity of seston available in the environment; (iii) state of sexual maturity of individuals (Fernández–Reiriz et al. 2007).

This study showed similarities on proximate composition between both mussel products, consistent with previous findings from mussels from Galician waters as reported by Fuentes et al. (2009). This may be explained by similarities occurring within the main factors regulating growth and biochemical performance in both culture conditions. Both culture conditions have a close annual average in environmental conditions, the same seasonality in reproductive cycles, and/or the same genetic heritage due to geographic proximity. However, open ocean mussels exhibited a higher amount of ash when compared to commercial ones from Galician Rías. Fuentes et al. (2009) observed similar values of ash content in commercial product from Galician Rías when comparing mussel product from different Spanish regions. Higher ash content than open ocean mussels of this study were also observed in the same study in mussels from Ebro Delta and from Valencia (NE and SE Spain, respectively). Mussels cultured in the exposed environmental conditions (e.g., phytoplankton variability, strong hydrodynamics, and very marked seasonal changes in seston composition) may experience situations of stress or may have greater energetic expenditure (Azpeitia et al. 2016) than mussels cultured in sheltered conditions. Oliveira et al. (2015) found similar results when comparing open ocean mussels to sheltered cultured mussels. The authors related this difference to the hydrodynamic conditions found by mussels in the offshore culture areas, which would interfere with mussel metabolism in a set of complex interactions between temperature, food availability, growth, and reproductive cycle (Gabbot 1976). Thus, ash content is influenced by many abiotic factors such as salinity and presence of metals on the water column (Okumus and Stirling 1998), as well as by the nutritional state and the reproductive cycle of the animal (Beninger and Lucas 1984). Regarding nutritional state, Mayzaud (1976) associated an increase in ash content of zooplankton with a state of starvation, and Wilkins (1967) observed the same in herrings. Nevertheless, this higher ash percentage may account for the decrease in other constituents (e.g., lipids, proteins) rather than an increase in the amount of ash, as data analysis was performed with relative percentages and not



in absolute terms. Despite these differences in ash content, similar general properties at biochemical level should be expected when comparing the open ocean mussels from the SE Bay of Biscay to their commercial counterparts from another northern Spanish region.

Depending on the time of the year, levels of nutrients, both in Galician Rías and in the open ocean of the SE Bay of Biscay, differ. In Galician Rías, upwelling events typically take place during spring–summer. Highest chlorophyll levels have been recorded during spring (<5 μg·l⁻¹)(Irisarri et al. 2014a). However, upwelling events are usually followed by thermal stratification periods during the summer which might result in a significant reduction of chlorophyll levels up to 0.4 μg·l⁻¹ (Irisarri et al. 2014a). The thermocline layer prevents nutrients from fertilizing the surface and chl-a eventually becomes depleted. The author corroborated this with a negative scope for growth (SFG) observed in mussel, when abnormally high temperatures are coupled with aforementioned low energy intake in Galician Rías in summer. In the SE Bay of Biscay, there is also a thermal stratification during summer, which decreases phytoplankton biomass to lower values than those described in Galician Rías (0.2 μg·l⁻¹ in surface waters) (Morán et al. 2012). However, during summer, short periods of atmospheric instability (storms) may act as modulation factors of this bloom decay in the SE Bay of Biscay. Although inputs of nutrient-rich continental waters can be observed, the intensified aforementioned current velocities may also resuspend inorganic sediments. Thus, in storm conditions, a dilution of available particulate organic matter (POM) for bivalve filter-feeders can be expected (Cranford et al. 2011; Irisarri et al. 2013; Zuñiga et al. 2014) in the experimental site.

The biochemical composition of the present contribution are consistent with previous studies reporting that mussels are high in proteins and low in calories and fat (Shahidi 2004; Fuentes et al. 2009; Gallardi et al. 2014). Indeed, some authors have reported that a daily protein intake of 0.75 g·kg body weight (which means that with 100 g of mussels for a person of 70 kg) is recommended to satisfy approximately 23% of the daily protein requirements (Garcia-Gabarra 2006; Stankovic et al. 2011; James 2013), thus demonstrating the rich nutritional value of mussels for human consumption.

In addition, it is also generally accepted that mussels represent an important source of essential fatty acids and particularly for long-chain polyunsaturated variety (PUFAs) (James 2013). In this study, not only concentration levels but also their representativeness on the profile of the major fatty acid groups (PUFA, MUFA and SAFA) is consistent with previous mussel studies (King et al. 1990; Freites et al. 2002b; Orban et al. 2002). Irisarri et al. (2014b) observed a selective accumulation of PUFA in different tissues of mussels. Mussels have a limited capability for de novo synthesis of PUFAs (Alkalani et al. 2007) and acquire them from diet. The unsaturation degree of the FA increases during the colder seasons, as mussels require these FA to maintain membrane fluidity (Hall et al. 2002). The contrary is observed in warmer seasons and thus, the PUFA level varies in opposite fashion to SAFA level. Mean seawater temperature in open ocean experimental site in summer was slightly higher (Azpeitia et al. 2016) than the typical summer temperature in Galician Rías (Pazos et al. 1997), which could have promoted a faster decrease in the unsaturation degree. This may have caused SAFA level to be higher in experimental mussel than commercial ones. Several authors have reported that higher proportions of SAFA may represent bacterial loads or organic-detritus-rich environments (Galap et al. 1999; Freites et al. 2002b). However, open ocean conditions are related to high hydrodynamic scenarios which favor marine water recycling and consequently less detritus accumulation compared to sheltered sites. Despite this higher water recycling, resuspension of material may have occurred at the experimental site. Moreover, regarding bacterial biomarker FA (15:0, 17:0, $18:1\omega7$ and $18:1\omega7/18:1\omega9 > 1$; Budge et al. 2001), higher values were observed in mussels from Galician Rías compared to experimental mussels. This may indeed support the aforementioned notion that temperature differences may well lead to higher SAFA levels in experimental mussels.

Major fatty acid levels in marine bivalves are generally known for being highly dependent on biochemical and environmental conditions (Lucas and Beninger 1985; De Moreno et al. 1980; Fernández-Reiriz et al. 1989; Fuentes et al. 2009). Nevertheless, the aforementioned results are consistent with current existing knowledge on mussel biochemistry and could be of use in further research in the same field at a local level.

Palmitic acid (16:0) has been reported in numerous studies (King et al. 1990; Freites et al. 2002a; Orban et al. 2002; Alkalani et al. 2007; Fuentes et al. 2009) as the major saturated fatty acid (SAFA) present in mussels. The experimental open ocean mussels herein displayed not only the same trend, but also, consumers reported a more intense sea-flavor after tasting the product. This would be consistent with Oliveira et al. (2015), who found that palmitic acid along with DHA are mainly responsible for mussel flavour.



PUFA levels of both mussel products herein were abundant and consistent with results from previous mussel studies from other localities (King et al. 1990; Orban et al. 2002; Freites et al. 2002b; Dridi et al. 2007; Alkalani et al. 2007; Fuentes et al. 2009), where healthy mussels always contain PUFA levels near to 50% of the total fatty acids proportion. As reported by Musa-Veloso et al. (2011) a dose of 250 mg·day⁻¹ of the long-chain omega-3 fatty acids EPA and DHA (PUFA) is strongly recommended for human health. In this way, although mussels are low in lipids, these lipids are of high quality and provide a healthy source of omega-3 fatty acids with numerous nutritional benefits (Shahidi 2004; Byrd-Bredbenner et al. 2009; James 2013). A high ω 6/ ω 3 ratio has also been reported as harmful to human health (i.e., over values of 12/50; Ackman 1990; Shahidi and Miraliakbari 2004; Fuentes et al. 2009). The results on the EPA and DHA of the present study were suitable within all ranges for both types of mussels examined.

Suspension feeding bivalves feed on living and non-living available material in their habitat (i.e., generalist filter feeders) (Ward and Shumway 2004). They select particles from seston, separating the grain from the chaff, and acquire the optimum energetic budget (Ward and Shumway, 2004). Seston consists of plankton of a wide range of sizes and palatability, material resuspended from the benthos, aggregates, detritus, fecal pellets as well as microorganisms (Ward and Shumway 2004, Karayücel et al. 2010). Fatty acids (FA) have been successfully used as trophic markers of bivalves to identify food sources, habitat preferences, feeding strategies and trophic links (Kharlamenko et al. 1995, 2001; Prato et al. 2010). Several studies on mussel Mytilus galloprovincilais Lmk. (Freites et al. 2002a, b; Ezgeta-Balić et al. 2012; Irisarri et al. 2014a) found that variability in nutritional quality of seston was explained by the variance of FAs found in mussels. Bivalves have some capability to synthesize FAs independently of diet (Ventrella et al. 2013), but they are not capable of biosynthesizing de novo polyunsaturated FAs (PUFAs) (Fernández-Reiriz et al. 2011). In addition, fatty acids transferred from primary producers, when incorporated by bivalves, do not undergo major changes (Graeve et al. 1994; Xu and Yang 2007), and thus, fatty acid measurements are a qualitative measurement of both energy transfer and assimilation of nutrients in mussel tissue. FA signature in mussel tissues can reveal the food source of the bivalve (i.e., several weeks prior) (Ezgeta-Balić et al. 2012) and thus, certain FA and their ratios can be used as biomarkers of the diet of the mussel. With these FA ratios, the seasonal contribution of bacteria, phytoplankton classes, as well as microzooplankton to the diet of the mussel can be assessed (Budge et al. 2001; Handå et al. 2012). Such analysis is an important factor in understanding the feeding physiology of mussels (Xu and Yang 2007).

In spite of presenting healthy EPA and DHA proportions, the experimental open ocean mussels of the present study showed less EPA and higher DHA contents than the commercial ones from Galician Rías. EPA level could be related to the ingestion of dinoflagellates (Freites et al. 2002a), whereas DHA could be a consequence of consumption and accumulation of diatoms (Handå et al. 2012). Sidari et al. (1998) observed that Mytilus galloprovincialis Lmk. as a preference selected dinoflagellates rather than diatoms. DHA/EPA ratio was >1 in experimental mussels, whereas, it was <1 in commercial mussels. These FA trophic markers (Budge et al. 2001; Dalsgaard et al. 2003) suggest mussels from the experimental site feed on dinoflagellates. Additionally, the ratio 16:1ω7/16:0 was <1 in experimental mussels and >1 in commercial mussels. Thus, all these FA trophic markers suggest commercial mussels feed on diatoms rather than dinoflagellates (Budge et al. 2001; Dalsgaard et al. 2003). No water samples were collected for phytoplankton analysis. However, as described by Alvarez et al. (2008) in the western Iberian Peninsula (i.e., Portugal and Galicia) during spring summer the continental shelves are strongly affected by upwelling pulses. These natural events are caused by the prevailing winds and the coastline orientation, causing an inflow of deep nutrient-rich waters which boosts phytoplankton productivity (Varela et al. 2005). Moreover, as described by Figueiras et al. (2002) and Álvarez-Salgado et al. (2008), these upwelling pulses result in the transportation of dinoflagellates to the continental shelf and leave the embayment dominated by diatoms. Similarly, Irisarri et al. (2014b) found that diatom FA biomarkers were significantly higher in mussel tissues from Galician Rías during spring-summer although those FA trophic markers were >1 throughout the entire year. Oliveira et al. (2015) also reported higher levels of EPA in mussels (M. galloprovincialis Lmk.) from the Galician Rías compared to individuals of the same species but different culture origin [S Atlantic Sea (Portugal)]. The authors suggested that the resultant DHA/EPA ratios would reflect the higher relative ingestion of diatoms. The contribution, however, of phytoplankton (as diatoms and dinoflagellates) to the diet (derived by 16:1ω7/16:0 and 20:5ω3/22:6ω3 ratios) was proportionally low and moderate in open ocean and commercial (M. galloprovincialis Lmk.) mussels, respectively. In the present study mussels were harvested in July along the north Spanish coast. There is a high



likelihood that previous poor environmental conditions (Prato et al. 2010) or post-spawning stages (Garmendia et al. 2010; Ortiz-Zarragoitia et al. 2011; Múgica et al. 2015; Cuevas et al. 2015) may have affected mussels' fatty acids composition in summer.

Fatty acid trophic markers of terrestrial organic sources [i.e., $\Sigma(18:2\omega6 + 18:3\omega3)$] higher than 2.5 were described by Budge and Parrish (1998) as indicative of a terrestrial organic diet source (i.e., terrestrial plants). A significant input of terrestrial markers was observed in the present study, where the experimental open ocean mussels showed to exceed the threshold. This may be related to the aforementioned differences in seston dynamics in Galician Rías and in the SE Bay of Biscay during summer season.

Finally, the presence of 22:2NMID acid was lower in experimental mussels compared to commercial ones. Sánchez-Lazo and Martínez-Pita (2012) also reported the presence of this fatty acid, although accounting for very little proportion.

The sensory analysis herein was consistent with previous mussel studies (Kryzynowek and Wiggin 1979; Ablett et al. 1986; Gökoglu 2002). The panelists were able to determine the mussel product qualitative characteristics on appearance, odor, flavor, texture, and global perception. Both types of mussel (open ocean and commercial) reached scores of 7 over 9 points at hedonic scale for all sensory attributes. In general terms, even though no triangle test was done, the sensory analysis session was performed stably and consistently among panelists.

Regarding appearance and odor, both characteristics were scored slightly lower for the open ocean samples when compared to the commercial ones from Galician Rías. Observed discrepancies on mussel appearance could be related to inherent biological and/or operational factors (e.g., higher variability in the coloration of mussel meat from open ocean conditions; previous industrial procedures applied on the commercial samples, etc.). The results of the present study showed no significant differences in tasting levels between open ocean and commercial products. However, most panelists also identified some distinctive attributes such as intense sea-flavour, softer and more juicy textures, and salty tastes in the open ocean samples.

These results may specifically be related to the absence of previous depuration requirement on the open ocean samples as a result of having an origin with a class A production area according to EU and local public legislation. During depuration processes, mussels are fasted and animals excrete their metabolic waste products, which include salts (Lee et al. 2008). Despite depuration of commercial mussels, the organoleptic characteristics did not differ greatly between both mussel products. The results on texture, at the time of the sensory analysis, also revealed softer meat in experimental mussels. Nevertheless, differences were not statistically significant and thus it can be concluded that softer texture does not imply mussels are less desirable. It is noteworthy that desired quality among fishery products relies on regional preferences, consumers' attitudes and methods of preservation and/or consumption (Haard 1992). Both mussel products reached 7 points out of 9 in qualitative category; a positive result considering the potential of open ocean mussels of the present study to be marketable in the near future. The tested consumers also expressed an intention to purchase both products (>70%). In general terms, this prescreening hedonic analysis showed that experimental open ocean mussels were similar in attributes to those routinely imported from the Galician Rías to the Basque Country. From a marketing perspective, the mussels cultured in the open ocean of the SE Bay of Biscay may be well accepted by consumers and fulfill the minimum requirements to fit and compete in the existing local market. The results of the present study provide baseline information to validate nutritional value characteristics and consumers' acceptance of mussels (M. galloprovincialis Lmk.) cultured in the open ocean of the Bay of Biscay. In addition, as a commercial validation was positive, it is now worth investigating how changes in the culture method can optimize production and commercial activity of future aquaculture in the Basque Country.

Conclusion

Space available in coastal areas for aquaculture facilities is growing more and more limited and is subjected to strict jurisdictions and limited leases. In addition, the geography of the coast in many regions might not be sheltered enough for mooring rafts or longlines, while limited seston supply and/or polluted waters can also represent limiting factors for mussel coastal aquaculture. Hence, this study investigates an interesting alternative to traditional suspended mussel culture in coastal inlets.



This study is the first sensory assessment of mussels produced in open ocean from the SE Bay of Biscay. First, this new information will be valuable for the development of strategies that favor origin specifications linked to health benefits which, in turn, would contribute to consumers' decision-making processes (i.e., purchase intention and preferences). Secondly, it will help to predict cost-effectiveness during mussel production and commercialization.

The approach of this study aimed to validate consumer acceptance as a meaningful factor in market feasibility studies. In addition, it also aimed to investigate if mussels cultured in the open ocean could compete with other mussels currently available on the market. However, future market needs will ultimately decide the final viability of mussel open ocean aquaculture.

Acknowledgements We wish to thank Gaizka Bidegain, Pilar Martín, Ainhoa Bikandi, and AZTI staff for their capable assistance during the laboratory and field works. Aileen Logan and Emma Ruizdelgado critically reviewed the manuscript. This work was partially supported by the Agriculture, Fisheries and Foods Department, and also through a PhD grant by the Department of Education, Linguistic Policy and Culture of the Basque Government.

Open Access This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (http://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made.

References

Ablett RF, Gould SP, Sherwood DA (1986) Frozen storage performance of cooked cultivated mussels (*Mytilus edulis* L.)—influence of ascorbic acid and chelating agents. J Food Sci 51(5):1118–1121

Ackman RG (1990) Seafood lipids and fatty acids. Food Rev Int 6(4):617-646

Akester RJ, Martel AL (2000) Shell shape, dysodont tooth morphology, and hinge-ligament thickness in the bay mussel *Mytilus trossulus* correlate with wave exposure. Can J Zool 78(2):240–253

Alkanani T, Parrish CC, Thompson RJ, McKenzie CH (2007) Role of fatty acids in cultured mussels, *Mytilus edulis*, grown in Notre Dame Bay, Newfoundland. J Exp Mar Biol Ecol 348(1):33–45

Álvarez I, Gómez-Gesteira M, DeCastro M, Dias JM (2008) Spatio temporal evolution of upwelling regime along the western coast of the Iberian Peninsula. J Geophys Res Oceans 113:14

Álvarez-Salgado XA, Labarta U, Fernández-Reiriz MJ, Figueiras FG, Rosón G, Piedracoba S, Filgueira R, Cabanas JM (2008) Renewal time and the impact of harmful algal blooms on the extensive mussel raft culture of the Iberian coastal upwelling system (SW Europe). Harmful Algae 7(6):849–855

AOAC (Association of Official Analytical Chemists) (1975) Official Methods of Analysis, 12th edn. Association of Official Analytical Chemists, Whashington, D.C

Azpeitia K, Ferrer L, Revilla M, Pagaldai J, Mendiola D (2016) Growth, biochemical profile, and fatty acid composition of mussel (*Mytilus galloprovincialis* Lmk.) cultured in the open ocean of the Bay of Biscay (northern Spain). Aquaculture 454:95–108

Beninger PG, Lucas A (1984) Seasonal variations in condition, reproductive activity, and gross biochemical composition of two species of adult clam reared in a common habitat: *tapes decussatus* L. (Jeffreys) and *Tapes philippinarum* (Adams & Reeve). J Exp Mar Biol Ecol 79(1):19–37

Borja Á, Galparsoro I, Irigoien X, Iriondo A, Menchaca I, Muxika I, Pascual M, Quincoces I, Revilla M, Germán Rodríguez J, Santurtún M, Solaun O, Uriarte A, Valencia V, Zorita I (2011) Implementation of the European Marine Strategy Framework Directive: a methodological approach for the assessment of environmental status, from the Basque Country (Bay of Biscay). Mar Pollut Bull 62:889–904

Bostock J, McAndrew B, Richards R, Jauncey K, Telfer T, Lorenzen K, Corner R (2010) Aquaculture: global status and trends. Philos Trans R Soc B Biol Sci 365(1554):2897–2912

Bower SM, McGladdery SE, Price IM (1994) Synopsis of infectious diseases and parasites of commercially exploited shellfish. Annu Rev Fish Dis 4:1–199

Bridger CJ, Costa-Pierce BA (2003) Open ocean aquaculture: from research to Commercial reality. The World Aquaculture Society, Baton Rouge, Louisiana

Buck BH (2007) Experimental trials on the feasibility of offshore seed production of the mussel *Mytilus edulis* in the German Bight: installation, technical requirements and environmental conditions. Helgol Mar Res 61(2):87–101

Buck BH, Buchholz CM (2004) The offshore-ring: a new system design for the open ocean aquaculture of macroalgae. J Appl Phycol 16(5):355–368

Buck BH, Ebeling MW, Michler-Cieluch T (2010) Mussel cultivation as a co-use in offshore wind farms: potential and economic feasibility. Aquac Econ Manag 14(4):255–281

Budge SM, Parrish CC (1998) Lipid biogeochemistry of plankton, settling matter and sediments in Trinity Bay, Newfoundland: II. Fatty acids. Organ Geochem 29:1547–1559

Budge SM, Parrish CC, Mckenzie CH (2001) Fatty acid composition of phytoplankton, settling particulate matter and sediments at a sheltered bivalve aquaculture site. Mar Chem 76(4):285–303



- Bushek D, Richardson D, Bobo MY, Coen LD (2004) Quarantine of oyster shell cultch reduces the abundance of *Perkinsus marinus*. J Shellfish Res 23(2):369–374
- Byrd-Bredbenner C, Moe G, Beshgetoor D, Berning J (2009) Wardlaw's perspectives in nutrition. In: Weatley CH (ed) McGraw-Hill, New York
- Byron C, Link J, Costa-Pierce B, Bengtson D (2011) Calculating ecological carrying capacity of shellfish aquaculture using mass-balance modeling: Narragansett Bay, Rhode Island. Ecol Model 222(10):1743–1755
- Chust G, Borja Á, Liria P, Galparsoro I, Marcos M, Caballero A, Castro R (2009) Human impacts overwhelm the effects of sealevel rise on Basque coastal habitats (N Spain) between 1954 and 2004. Estuar Coast Shelf Sci 84:453–462
- Coen LD, Brumbaugh RD, Bushek D, Grizzle R, Luckenbach MW, Posey MH, Tolley S (2007) Ecosystem services related to oyster restoration. Mar Ecol Prog Ser 341:303–307
- Collins M, Borja Á (2004) Preface (to: Oceanography and Marine Environment of the Basque Country). Oceanogr Mar Environ Basqu Ctry. http://eprints.soton.ac.uk/11051/
- Costa-Pierce BA (2008) Epilogue: aquaculture, innovation and social transformation. Aquac Innov Soc Transf 17:315-326
- Cranford PJ, Ward JE, Shumway SE (2011) Bivalve filter feeding: variability and limits of the aquaculture biofilter. Shellfish Aquac Environ 81:124
- Cubillo AM, Peteiro LG, Fernández-Reiriz MJ, Labarta U (2012) Density-dependent effects on morphological plasticity of *Mytilus galloprovincialis* in suspended culture. Aquaculture 338:246–252
- Cuevas N, Zorita I, Costa PM, Franco J, Larreta J (2015) Development of histopathological indices in the digestive gland and gonad of mussels: integration with contamination levels and effects of confounding factors. Aquat Toxicol 162:152–164
- Dalsgaard J, John MS, Kattner G, Müller-Navarra D, Hagen W (2003) Fatty acid trophic markers in the pelagic marine environment. Adv Mar Biol 46:225-340
- Dare PJ, Edwards DB (1975) Seasonal changes in flesh weight and biochemical composition of mussels (*Mytilus edulis* L.) in the Conwy Estuary, North Wales. J Exp Mar Biol Ecol 18(2):89–97
- Dawes JG (2012) Do data characteristics change according to the number of scale points used? An experiment using 5 point, 7 point and 10 point scales. Int J Mark Res 51(1):2008
- De Moreno JEA, Pollero RJ, Moreno VJ, Brenner RR (1980) Lipids and fatty acids of the mussel (*Mytilus platensis* d'Orbigny) from South Atlantic waters. J Exp Mar Biol Ecol 48:263–276
- De Zwaan A, Zandee DI (1972) Body distribution and seasonal changes in the glycogen content of the common sea mussel *Mytilus edulis*. Comp Biochem Physiol A Physiol 43(1):53–58
- Drapeau A, Comeau LA, Landry T, Stryhn H, Davidson J (2006) Association between longline design and mussel productivity in Prince Edward Island, Canada. Aquaculture 261(3):879–889
- Dridi S, Romdhane MS, Elcafsi MH (2007) Seasonal variation in weight and biochemical composition of the Pacific oyster, Crassostrea gigas, in relation to the gametogenic cycle and environmental conditions of the Bizert lagoon, Tunisia. Aquaculture 263(1):238–248
- Duarte P, Labarta U, Fernández-Reiriz MJ (2008) Modelling local food depletion effects in mussel rafts of Galician Rías. Aquaculture 274(2):300–312
- Epler S, Edgar Chambres IV, Kemp KE (1998) Hedonic scales are better predictor than jus about right scales of optimal sweetness in lemonade 1. J Sens Stud 13(2):191–197
- Ezgeta-Balić D, Najdek M, Peharda M, Blažina M (2012) Seasonal fatty acid profile analysis to trace origin of food sources of four commercially important bivalves. Aquaculture 334:89–100
- FAO (2014) The state of world Fisheries and Aquaculture_2014 (Rome). FAO Fisheries and Aquaculture Department 176
- Fernández-Reiriz MJ, Pérez-Camacho A, Ferreiro MJ, Blanco J, Planas M, Campos MJ, Labarta U (1989) Biomass production and variation in the biochemical profile (total protein, carbohydrates, RNA, lipids and fatty acids) of seven species of marine microalgae. Aquaculture 83:17–37
- Fernández-Reiriz MJ, Duarte P, Labarta U (2007) Modelos de comportamiento alimentario en el mejillón de las rías de Galicia. In: Figueras AJ (ed) Biología y cultivo del mejillón, vol 26. Editorial CSIC-CSIC Press, pp 197–223
- Fernández-Reiriz MJ, Pérez-Camacho A, Peteiro LG, Labarta U (2011) Growth and kinetics of lipids and fatty acids of the clam Venerupis pullastra during larval development and postlarvae. Aquac Nutr 17(1):13–23
- Figueiras FG, Labarta U, Reiriz MF (2002) Coastal upwelling, primary production and mussel growth in the Rías Baixas of Galicia. In: Sustainable increase of marine harvesting: fundamental mechanisms and new concepts. Springer, Netherlands, pp 121–131
- Filgueira R, Labarta U, Fernández-Reiriz MJ (2006) Flow through chamber method for clearance rate measurements in bivalves. Design and validation of individual chambers and mesocosm. Limnol Oceanogr Methods 4:284–292
- Forrest BM, Keeley NB, Hopkins GA, Webb SC, Clement DM (2009) Bivalve aquaculture in estuaries: review and synthesis of oyster cultivation effects. Aquaculture 298(1):1–15
- Freites L, Labarta U, Fernández-Reiriz MJ (2002a) a Evolution of fatty acid profiles of subtidal and rocky shore mussel seed (Mytilus galloprovincialis, Lmk.). Influence of environmental parameters. J Exp Mar Biol Ecol 268(2):185–204
- Freites L, Fernández-Reiriz MJ, Labarta U (2002b) b Lipid classes of mussel seeds *Mytilus galloprovincialis* of subtidal and rocky shore origin. Aquaculture 207(1–2):97–111
- Fuentes A, Fernandez-Segovia I, Escriche I, Serra JA (2009) Comparison of physico-chemical parameters and composition of mussels (*Mytilus Galloprovincialis* Lmk.) from Different Spanish Origins. Food Chem 112:295–302
- Gabbot PA (1976) Chapter 8: energy metabolism. In Bayne BL (ed) Marine mussels: their ecology and physiology. International biological programme. Cambridge University Press, Cambridge, vol 10, pp 293–355
- Galap C, Netchitaïlo P, Leboulenger F, Grillot JP (1999) Variations of fatty acid contents in selected tissues of the female dog cockle (*Glycymeris glycymeris* L, Mollusca, Bivalvia) during the annual cycle. Comp Biochem Physiol A122:241–254



- Gallardi D, Hobbs K, Mills T, Couturier C, Parrish CC, Murray HM (2014) Effects of extended ambient live holding on cultured blue mussels (*Mytilus edulis* L.) with reference to condition index, lipid profile, glycogen content and organoleptic testing. Aquaculture 430:149–158
- Galparsoro I, Borja Á, Legorburu I, Hernández C, Chust G, Liria P (2010) Morphological characteristics of the Basque continental shelf (Bay of Biscay, northern Spain); their implications for Integrated Coastal Zone Management. Geomorphology 118(3):314–329
- Galparsoro I, Liria P, Legorburu I, Bald J, Chust G, Ruiz-Minguela P, Pérez G, Marqués J, Torre-Enciso Y, González M, Borja A (2012) A Marine Spatial Planning approach to select suitable areas for installing wave energy converters on the Basque continental shelf (Bay of Biscay). Coast Manag 40:1–9
- García-Gabarra A (2006) Nutrient intakes: concepts and international recommendations (part two) (Review). Nutrición Hospitalaria 21(4):437–447
- Garmendia L, Soto M, Cajaraville MP, Marigómez I (2010) Seasonality in cell and tissue-level biomarkers in *Mytilus galloprovincialis*: relevance for long-term pollution monitoring. Aquat Biol 9(3):203–219
- Gökoglu N (2002) A descriptive method for sensory evaluation of mussels. LWT Food Sci Technol 35(7):563-567
- Gosling E (2003) Bivalve Molluscs: biology, ecology and culture. Fishing News Books-Blackwell Publishing, MPG Books, Bodmin
- Gracia J (1996) El sector pesquero en la historia del País Vasco: esbozo de los conocimientos actuales y problemas abiertos a la futura investigación historiográfica. Itsas Memoria Revista de Estudios Marítimos del País Vasco 1:209–214
- Graeve M, Kattner G, Hagen W (1994) Diet-induced changes in the fatty acid composition of Arctic herbivorous copepods: experimental evidence of trophic markers. J Exp Mar Biol Ecol 182(1):97–110
- Grienke U, Silke J, Tasdemir D (2014) Bioactive compounds from marine mussels and their effects on human health. Food Chem 142:48–60
- Haard NF (1992) Control of chemical composition and food quality attributes of cultured fish. Food Res Int 25(4):289-307
- Hall JM, Parrish CC, Thompson RJ (2002) Eicosapentaenoic acid regulates scallop (*Placopecten magellanicus*) membrane fluidity in response to cold. Biol Bull 202(3):201–203
- Handå A, Min H, Wang X, Broch OJ, Reitan KI, Reinertsen H, Olsen Y (2012) Incorporation of fish feed and growth of blue mussels (*Mytilus edulis*) in close proximity to salmon (*Salmo salar*) aquaculture: implications for integrated multi-trophic aquaculture in Norwegian coastal waters. Aquaculture 356:328–341
- Hanson SWF, Olley J (1963) Application of the Bligh and Dyer method of lipid extraction to tissue homogenates. Biochem J 89(3):101–102
- Heral M, Razet D, Maestrini S, Garnier J (1980) Composition de la matière organique particulaire dans les eaux du bassin de Marennes-Oléron: apport énergétique pour la nutrition de l'huître. In: CIEM Conseil International pour l'Exploration de la mer
- Hesley C (1997) Open ocean aquaculture: chartering the future of ocean farming. In: Proceedings of an international conference. UNIHI-Seagrant-CP-98–08, University of Hawaii Sea Grant College Program, Maui, 23–25 April 1997
- Hickman RW (1992) Mussel cultivation. The mussel Mytilus: ecology, physiology, genetics and culture. Elsevier, New York, pp 465–510
- Irisarri J, Fernández-Reiriz MJ, Robinson SM, Cranford PJ, Labarta U (2013) Absorption efficiency of mussels *Mytilus edulis* and *Mytilus galloprovincialis* cultured under integrated multi-trophic aquaculture conditions in the Bay of Fundy (Canada) and Ría Ares-Betanzos (Spain). Aquaculture 388:182–192
- Irisarri J, Fernández-Reiriz MJ, De Troch M, Labarta U (2014a) Fatty acids as tracers of trophic interactions between seston, mussels and biodeposits in a coastal embayment of mussel rafts in the proximity of fish cages. Comp Biochem Physiol B Biochem Mol Biol 172:105–115
- Irisarri J, Fernández-Reiriz MJ, Cranford PJ, Labarta U (2014b) b Effects of seasonal variations in phytoplankton on the bioenergetic responses of mussels (*Mytilus galloprovincialis*) held on a raft in the proximity of red sea bream (*Pagellus bogaraveo*) net-pens. Aquaculture 428:41–53
- James D (2013) Risks and benefits of seafood consumption. GLOBEFISH Research Programme, FAO, Rome, p 28
- Karayücel S, Çelik MY, Karayücel İ, Erik G (2010) Growth and production of raft cultivated Mediterranean mussel (Mytilus galloprovincialis Lamarck, 1819) in Sinop, Black sea. Turk J Fish Aquat Sci 10(1). doi:10.4194/trjfas.2010.0102
- Kharlamenko VI, Zhukova NV, Khotimchenko SV, Svetashev VI, Kamenev GM (1995) Fatty acids as markers of food sources in a shallow-water hydrothermal ecosystem (Kraternaya Bight, Yankich Island, Kurile Islands). Mar Ecol Prog Ser 120:231–241
- Kharlamenko VI, Kiyashko SI, Imbs AB, Vyshkvartzev DI (2001) Identification of food sources of invertebrates from the seagrass Zostera marina community using carbon and sulfur stable isotope ratio and fatty acid analyses. Mar Ecol Prog Ser 220:103–117
- King PA, McGrath D, Britton W (1990) The use of artificial substrates in monitoring mussel (*Mytilus edulis* L.) settlement on an exposed rocky shore in the west of Ireland. J Mar Biol Assoc U K 70(02):371–380
- Kryzynowek J, Wiggin K (1979) Seasonal variation and frozen storage stability of blue mussel (*Mytilus edulis*). J Food Sci 44(6):1644–1645
- Langan R (2000) Submerged longline culture of blue mussels (*Mytilus edulis*) at an open ocean site in the Gulf of Maine. J Shellfish Res 19:575
- Langan R (2001) Advances in submerged longline culture of blue mussels *Mytilis edulis* in the open ocean. In Bridger CJ, Reid TH (eds.) Open ocean aquaculture IV, symposium program and book of abstracts. Mississippi-Alabama Sea Grant Consortium, Ocean Springs, MS MASGP-01–006, St Andrews, pp 73–74
- Langan R, Horton F (2003) Design, operation and economics of submerged longline mussel culture in the open ocean. Bull Aquac Assoc Can 103:11–20



LeBlanc N, Landry T, Stryhn H, Tremblay R, McNiven M, Davidson J (2005) The effect of high air and water temperature on juvenile *Mytilus edulis* in Prince Edward Island, Canada. Aquaculture 243(1):185–194

Lee R, Lovatelli A, Ababouch L (2008) Bivalve depuration: fundamental and practical aspects. Food and Agriculture Organization of the United Nations

Lewis JR, Powell HT (1961) The occurrence of curved and ungulate forms of the mussel *Mytilus edulis* L. in the British Isles and their relationship to *M. galloprovincialis* Lamarck. Proc Zool Soc Lond 137(4):583–598

Lucas A, Beninger PG (1985) The use of physiological condition indices in marine bivalve aquaculture. Aquaculture 44(3):187-200

Martínez ML, Intralawan A, Vázquez G, Pérez-Maqueo O, Sutton P, Landgrave R (2007) The coasts of our world: ecological, economic and social importance. Ecol Econ 63(2):254–272

Mayzaud P (1976) Respiration and nitrogen excretion of zooplankton. IV. The influence of starvation on the metabolism and the biochemical composition of some species. Mar Biol 37(1):47–58

Morán XAG, Barbosa AB, Revilla M, Bode A, Orive E, Soudant D (2012) Phytoplankton and microbial plankton of the Bay of Biscay and western Iberian shelf. In: O'Brien TD, Li WKW, Morán XAG (eds.) ICES Phytoplankton and Microbial Plankton Status Report 2009/2010. ICES Cooperative Research Report 313, pp 114–135

Moroño A, Arévalo F, Fernández ML, Maneiro J, Pazos Y, Salgado C, Blanco J (2003) Accumulation and transformation of DSP toxins in mussels *Mytilus galloprovincialis* during a toxic episode caused by *Dinophysis acuminata*. Aquat Toxicol 62(4):269–280

Múgica M, Sokolova IM, Izagirre U, Marigómez I (2015) Season-dependent effects of elevated temperature on stress biomarkers, energy metabolism and gamete development in mussels. Mar Environ Res 103:1–10

Musa-Veloso K, Binns MA, Kocenas A, Chung C, Rice H, Oppedal-Olsen H, Lemke S (2011) Impact of low v. moderate intakes of long-chain n-3 fatty acids on risk of coronary heart disease. Br J Nutr 106(08):1129–1141

Myrand B, Tremblay R, Sévigny JM (2009) Decreases in multi-locus heterozygosity in suspension-cultured mussels (*Mytilus edulis*) through loss of the more heterozygous individuals. Aquaculture 295(3):188–194

Okumuş I, Stirling HP (1998) Seasonal variations in the meat weight, condition index and biochemical composition of mussels (*Mytilus edulis L.*) in suspended culture in two Scottish sea lochs. Aquaculture 159(3–4):249–261

Oliveira RA, Sykes AV, Hachero-Cruzado I, Azeiteiro UM, Esteves E (2015) A sensory and nutritional comparison of mussels (*Mytilus* sp.) produced in NW Iberia and in the Armona offshore production area (Algarve, Portugal). Food Chem 168:520–528

Orban E, Di Lena G, Nevigato T, Casini I, Marzetti A, Caproni R (2002) Seasonal changes in meat content, condition index and chemical composition of Mussels (*Mytilus galloprovincialis*) cultured in two different Italian Sites. Food Chem 77(1):57–65

Ortiz-Zarragoitia M, Garmendia L, Barbero MC, Serrano T, Marigómez I, Cajaraville MP (2011) Effects of the fuel oil spilled by the Prestige tanker on reproduction parameters of wild mussel populations. J Environ Monit 13(1):84–94

Pascual M, Borja A, Eede SV, Deneudt K, Vincx M, Galparsoro I (2011) Marine biological valuation mapping of the Basque continental shelf (Bay of Biscay), within the context of marine spatial planning. Estuar Coast Shelf Sci 95(1):186–198

Pauly D (1983) Some simple methods for the assessment of tropical fish stocks. FAO Fish. Tech. Pap. 234

Pazos AJ, Román G, Acosta CP, Abad M, Sánchez JL (1997) Seasonal changes in condition and biochemical composition of the scallop *Pecten maximus* L. from suspended culture in the Ria de Arousa (Galicia, NW Spain) in relation to environmental conditions. J Exp Mar Biol Ecol 211(2):169–193

Penney RW, Hart MJ, Templeman N (2002) Comparative growth of cultured blue mussels, *Mytilus edulis*, *M. trossulus* and their hybrids, in naturally occurring mixed-species stocks. Aquac Res 33(9):693–702

Pérez-Camacho A, Gonzalez R, Fuentes J (1991) Mussel culture in Galicia (NW Spain). Aquaculture 94(2):263-278

Pérez-Camacho A, Aguiar E, Labarta U, Vinseiro V, Fernández-Reiriz MJ, Álvarez-Salgado XA (2014a) Ecosystem-based indicators as a tool for mussel culture management strategies. Ecol Ind 45:538–548

Pérez-Camacho A, Aguiar E, Labarta U, Vinseiro V, Fernández-Reiriz MJ, Álvarez-Salgado XA (2014b) Ecosystem-based indicators as a tool for mussel culture management strategies. Ecol Ind 45:538–548

Peryam DR, Girardot NF (1952) Advanced taste-test method. Food Eng 24:58-61

Peryam DR, Pilgrim FJ (1957) The hedonic scale method of measuring food preference. Food tech 11(4):9-14

Peteiro LG, Babarro JM, Labarta U, Fernández-Reiriz MJ (2006) Growth of *Mytilus galloprovincialis* after the Prestige oil spill. ICES J Mar Sci J du Cons 63(6):1005–1013

Pogoda B, Buck BH, Saborowski R, Hagen W (2013) Biochemical and elemental composition of the offshore-cultivated oysters Ostrea edulis and Crassostrea gigas. Aquaculture 400–401:53–60

Polk M (1996) Open ocean aquaculture. Proceedings of an international conference, May 8–10, 1996, Portland, Maine. In: Polk M (ed) New Hampshire-Maine Sea Grant College Program No. UNHMP-CP-SG-96-9, pp 640

Prato E, Danieli A, Maffia M, Biandolino F (2010) Lipid and fatty acid compositions of *Mytilus galloprovincialis* cultured in the Mar Grande of Taranto (Southern Italy): feeding strategies and trophic relationships. Zool Stud 49(2):211–219

R Development Core Team (2008) R: A language and environmental for statistical computing. R Foundation fro Statistical Computing, Viena, Austria. http://www.R-project.org

Rohlf FJ, Sokal RR (1981) Comparing numerical taxonomic studies. Syst Zool 30:459-490

Ruano F, Ramos P, Quaresma M, Bandarra NM, Da Fonseca IP (2012) Evolution of fatty acid profile and condition index in mollusc bivalves submitted to different depuration periods. Revista Portuguesa da Ciências Veterinárias 111(581–582):75–84

Ryan J (2004) Farming the Deep Blue. Board Iascaigh Mhara. Technical Report 82

Sánchez-Lazo C, Martínez-Pita I (2012) Biochemical and energy dynamics during larval development of the mussel *Mytilus galloprovincialis* (Lamarck, 1819). Aquaculture 358–359:71–78



Schutz HG, Cardello AV (2001) A labeled affective magnitude (LAM) scale for assessing food liking/disliking. J Sens Stud 16(2):117–159

Seed R (1968) Factors influencing shell shape in Mytilus Edulis. J Mar Biol Assoc U K 48(03):561-584

Seed R, Suchanek TH (1992) Population and community ecology of *Mytilus*. Mussel Mytilus Ecol Physiol Genet Cult 25:87–170 Shahidi F (2004) Functional seafood products. ACS Symp Ser 702:29–49

Shahidi F, Miraliakbari H (2004) Omega-3 (n-3) fatty acids in health and disease: part 1—cardiovascular disease and cancer. J Med Food 7(4):387–401

Shpigel M, Neori A, Popper DM, Gordin H (1993) A proposed model for "environmentally clean" land-based culture of fish, bivalves and seaweeds. Aquaculture 117(1–2):115–128

Shumway SE, Davis C, Downey R, Karney R, Kraeuter J, Parsons J, Wikfors G (2003) Shellfish aquaculture—in praise of sustainable economies and environments. World Aquac 34(4):8–10

Sidari L, Nichetto P, Cok S, Sosa S, Tubaro A, Honsell G, Della Loggia R (1998) Phytoplankton selection by mussels, and diarrhetic shellfish poisoning. Mar Biol 131(1):103–111

Smaal AC (2002) European mussel cultivation along the Atlantic coast: production status, problems and perspectives. Dev Hydrobiol 167:89–98

Soto D, Aguilar-Manjarrez J, Brugère C, Angel D, Bailey C, Black K, Wainberg A (2008) Applying an ecosystem-based approach to aquaculture: principles, scales and some management measures. Build Ecosyst Approach Aquac 14

Stanković S, Jović M, Milanov R, Joksimović D (2011) Trace elements concentrations (Zn, Cu, Pb, Cd, As and Hg) in the Mediterranean mussel (*Mytilus galloprovincialis*) and evaluation of mussel quality and possible human health risk from cultivated and wild sites of the southeastern Adriatic Sea, Montenegro. J Serb Chem Soc 76(12):1725–1737

Stickney RR (1998) Joining forces with industry—open ocean aquaculture. In: Proceedings of the third annual international conference. TAMU-SG-99–103, Corpus Christi, Texas Sea Grant College Program

Svensson S (2003) Depuration of Okadaic acid (Diarrhetic Shellfish Toxin) in mussels, *Mytilus edulis* (Linnaeus), feeding on different quantities of nontoxic algae. Aquaculture 218(1):277–291

Varela M, Prego R, Pazos Y, Moroño Á (2005) Influence of upwelling and river runoff interaction on phytoplankton assemblages in a Middle Galician Ria and Comparison with northern and southern rias (NW Iberian Peninsula). Estuar Coast Shelf Sci 64(4):721–737

Ventrella V, Pagliarani A, Nesci S, Trombetti F, Pirini M (2013) Dietary enhancement of selected fatty acid biosynthesis in the digestive gland of *Mytilus galloprovincialis* Lmk. J Agric Food Chem 61(4):973–981

Vernocchi P, Maffei M, Lanciotti R, Suzzi G, Gardini F (2007) Characterization of Mediterranean mussels (*Mytilus galloprovincialis*) harvested in Adriatic Sea (Italy). Food Control 18:1575–1583

Vilela H (1950) Vida bentónica de Tapes decussatus (L.). Arquivos do Museu Bocage 21:1-120

Villanueva ND, Petenate AJ, Da Silva MA (2005) Performance of the hybrid hedonic scale as compared to the traditional hedonic, self-adjusting and ranking scales. Food Qual Prefer 16(8):691–703

Waite L, Grant J, Davidson J (2005) Bay-scale spatial growth variation of mussels *Mytilus edulis* in suspended culture, Prince Edward Island, Canada. Mar Ecol Progress Ser 297:157

Ward JE, Shumway SE (2004) Separating the grain from the chaff: particle selection in suspension-and deposit-feeding bivalves. J Exp Mar Biol Ecol 300(1):83–130

Wilkins NP (1967) Starvation of the herring, Clupea harengus L.: survival and some gross biochemical changes. Comp Biochem Physiol 23(2):503–518

Xu Q, Yang H (2007) Food sources of three bivalves living in two habitats of Jiaozhou Bay (Qingdao, China): indicated by lipid biomarkers and stable isotope analysis. J of Shellfish Res 26(2):561–567

Zar JH (1984) Biostatistical analysis. Prentice-Hall, Englewood Cliffs, p 718

Zúñiga D, Castro CG, Aguiar E, Labarta U, Figueiras FG, Fernández-Reiriz MJ (2014) Biodeposit contribution to natural sedimentation in a suspended *Mytilus galloprovincialis* Lmk mussel farm in a Galician Ría (NW Iberian Peninsula). Aquaculture 432:311–320

