Condition of *Salaria pavo* in the Mar Menor coastal lagoon (SE Iberian Peninsula): potential influence of environmental variables on juveniles

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Summary

The relationship at the local level between fish condition and environmental variables was investigated in *Salaria pavo* (peacock blenny) juveniles. The relationships between fish condition and 18 environmental variables of shallow areas of the lagoon relating to intra- and interspecific fish interactions, water quality and habitat structure, were analysed. The study revealed that water depth and *S. pavo* biomass (both related to intraspecific fish interactions) were the ecological variables accounting for most of the variation in the condition of *S. pavo* juveniles, in this case a negative effect. In this way, the condition of *S. pavo* juvenile populations may be a good indicator of fish density interactions.

Introduction

Distribution of the peacock blenny, *Salaria pavo* (Risso, 1810), runs from the Mediterranean Sea to the Atlantic coasts of Morocco and France. *S. pavo* inhabits rocky and weedy pools and shallow waters and is common in brackish waters. Its prey are benthic invertebrates, especially molluscs. Reproduction occurs in spring and early summer (Zander, 1986). During the breeding season, males build and defend nests and care for the eggs (Gonçalves and Almada, 1997).

Peacock blenny is one of the dominant benthic fish species in the Mar Menor fish assemblage and reproduces within the lagoon (Pérez-Ruzafa et al., 2004). However, no studies exist on its biology and ecology.

In the management of fish populations it is common to analyse fish condition as a measure of both individual and cohort (e.g. age or size group) fitness (Jakob et al., 1996). Such measurements are generally intended to be indicators of tissue energy reserves and may characterize components of the environment in which the fish live (e.g. food and habitat availability, competition, predation, physical factors, parasitic infections and pollution) (Bergeron, 2000; Copp, 2003; Lloret and Planes, 2003; Oliva-Paterna et al., 2003). A low body condition can negatively affect survival, first maturity and the reproductive effort in subsequent phases of fish life history (Rätz and Lloret, 2003; Hoey and McCormick, 2004; Morgan, 2004).

For this reason, the study of fish condition, together with investigations concerning habitat characteristics, will allow a better understanding of the biology and ecology of fish populations. Moreover, fish condition indices are potential indicators for assessing habitat quality in marine ecosystems such as coastal lagoons and estuaries (Lloret et al., 2005).

The purpose of this paper was to determine the relationship between the somatic condition of *S. pavo* juveniles and several environmental variables and indices relating to intra- and interspecific interactions, water quality and habitat structure. These included: fish species richness, size of larger *S. pavo* (total length, TL), *S. pavo* abundance and biomass, biomass/ abundance ratio of *S. pavo*, benthic fish abundance and biomass, biomass/abundance ratio of benthic fish species, total fish abundance and biomass, biomass/abundance ratio of total fish, water temperature (°C), water salinity, depth (cm), submerged vegetation cover (%), submerged vegetation volume, substrate size (SS) and substrate heterogeneity (SH).

Materials and methods

Study area

The Mar Menor is a hypersaline coastal lagoon located in a semiarid region in the south-east of the Iberian Peninsula (Fig. 1). One of the largest coastal lagoons in the Mediterranean region and Europe, with a surface area of 135 km^2 and



Fig. 1. Geographical location of the Mar Menor coastal lagoon and sampling sites

average depth of 3–4 m, it is separated from the Mediterranean Sea by a 22-km-long sandbar, with three narrow channels connecting it with the sea. The lagoon shows a salinity range of 39–45 and a temperature varying from 10°C in winter to 32°C in summer. The bottom is principally covered by dense meadows of the invasive macroalga *Caulerpa prolifera*, although shallow areas are covered by meadows of *Cymodocea nodosa* (Pérez-Ruzafa et al., 2004).

Since the 1970s the Mar Menor has suffered strong environmental changes after the widening of the connecting channels that caused a decrease in salinity from 50–52% to the present levels. Moreover, regular and intermittent watercourses flow into the lagoon, draining a large intensive agricultural area and leading to an important input of agrochemicals (Pérez-Ruzafa et al., 2000, 2002). Finally, the Mar Menor coastal lagoon supports important commercial fisheries and is subject to intensive tourist development (Andreu-Soler et al., 2003).

Sampling sites are located in littoral and shallow areas (maximum ≤ 100 cm) of the lagoon. The bottoms are characterised by soft substrates (principally muddy and sandy bottoms) and isolated patches of submerged vegetation (meadows of *Caulerpa prolifera* and *Cymodocea nodosa*).

Sampling methods

The catches were carried out during the first two weeks of July 2003 as a part of a wider study to examine the effects of human activities on fish communities of the coastal lagoon.

A total of 13 sampling sites was selected in the perimeter coastal shallow areas. Samples were collected using a 10-m-long bag seine net of 0.5 mm mesh size, which allowed the collection of juvenile fish and adults of small size species. Six replicates were collected at each sampling site by adjacent 20 m reaches of shoreline at each site. In each reach, the bag seine was hauled offshore parallel to the shoreline in water <1.0 m for the length of the reach. The area covered by each haul was approximately 160 m² (quantitative sampling).

Additional bag seine hauls and quadrangular $(40 \times 40 \text{ cm})$ hand net sweeps were made along the shoreline in each sampling area (the number of non-quantitative sampling, hauls and sweeps depending on the complexity of the shore). Our goal was to sample all shoreline habitats to detect species richness in a given sampling site.

Thus, seven replicate samples (six quantitative and one nonquantitative) were obtained at each of the 13 sampling sites, enabling us to assess variance within sites and the efficiency of seining. Table 1 Environmental variables and fish community composition at sampling sites with specific emphasis on *Salaria pavo*

Fish from each of the six reaches (quantitative samplings) and non-quantitative samplings were preserved in 7% formaldehyde, before being removed and identified at species level in the laboratory (Whitehead et al., 1986; Arias and Drake, 1990). Relative abundance was expressed as catch per unit effort (CPUEs) and biomass per unit effort (BPUEs):

 $CPUEs = number of specimens/100 m^2$,

BPUEs = fish biomass $(g)/100 \text{ m}^2$.

A total of 584 juvenile *S. pavo* (TL < 4.5 cm) from 13 sampling sites were measured for TL (± 1 mm) and total mass, TM (± 0.1 g). Fish larger than 4.5 cm were excluded from the condition analysis to avoid possible body shape differences between juveniles and adults (Murphy et al., 1990).

	Size of						Benthic	Benthic			Total							
	larger	Fish	S. pavo	S. pavo	S. pavo	Benthic fish	fish	fish	Total fish	Total fish	fish	Water						
	S. pavo	species	abundance	biomass	B/A	abundance	biomass	\mathbf{B}/\mathbf{A}	abundance	biomass	\mathbf{B}/\mathbf{A}	temperature		Vegetation	Vegetation	Depth		
Sampling site	(TL)	richness	(CPUEs)	(BPUEs)	ratio	(CPUEs)	(BPUEs)	ratio	(CPUEs)	(BPUEs)	ratio	(°C)	Salinity	cover (%)	volume	(cm)	SS SF	Н
1 (Tomás Maestre)	6.23	15	4.06	2.78	0.68	20.29	96.77	0.21	128.54	41.19	0.32	27.99	43.36	40.83	2.08	65.00	2.09 0.3	39
2 (Las Brisas)	7.81	15	1.25	1.92	1.54	23.05	87.43	0.26	102.23	30.76	0.30	26.95	43.98	26.67	1.61	67.78	2.39 0.0	69
3 (Los Alíseos)	6.73	12	4.06	2.28	0.56	32.03	178.86	0.18	240.73	54.03	0.22	27.34	44.46	20.00	1.25	53.83	2.36 0.7	71
4 (Isla del Ciervo)	I	10	18.13	11.49	0.63	20.21	54.69	0.37	115.21	57.43	0.50	28.39	44.21	35.83	1.75	42.67	2.22 0.8	81
5 (Playa Paraíso)	6.46	11	1.67	0.99	0.60	9.46	74.58	0.13	90.56	14.73	0.16	30.52	43.89	17.78	1.17	60.11	2.18 0.0	64
6 (Punta del Plome)	8.28	12	1.56	7.19	4.60	10.96	63.13	0.17	91.04	33.91	0.37	29.43	44.28	26.67	1.83	71.67	2.54 0.7	75
7 (Los Nietos)	5.68	10	5.63	3.96	0.70	4.24	19.58	0.22	116.36	35.19	0.30	28.77	44.57	34.17	1.75	61.50	2.11 0.5	58
8 (Los Urrutias)	5.36	13	6.14	4.54	0.74	16.88	53.64	0.31	188.13	68.09	0.36	27.57	44.74	51.67	2.75	41.00	1.83 0.0	63
9 (Los Alcázares)	5.76	12	28.02	14.88	0.53	6.59	50.31	0.13	158.33	58.73	0.37	28.41	42.81	49.17	2.42	43.50	1.65 0.4	46
10 (La Hita)	5.60	6	27.81	9.71	0.35	4.00	25.00	0.16	161.41	45.59	0.28	31.95	42.50	52.50	2.63	43.50	1.44 0.	73
11 (Casablanca)	5.21	7	38.44	28.61	0.74	1.36	8.75	0.16	128.96	63.63	0.49	28.24	44.10	77.50	3.83	52.83	1.70 0.5	57
12 (Villa Nanitos	5.60	11	7.50	3.75	0.50	10.36	84.06	0.12	153.23	36.15	0.24	29.13	43.58	30.83	1.92	62.00	1.95 0.3	38
13 (La Calcetera)	6.26	12	3.23	4.95	1.53	17.48	43.96	0.40	188.64	56.11	0.30	26.86	43.87	62.50	3.17	61.33	2.16 0.7	78
CPUEs, catch per un Submerged vegetatio heterogeneity (stands	nit effort; n volume ard devial	BPUEs, t : 0 (low de tion at eac	iomass per ι snsity of mea	unit effort; dows) to 5 site).	SS, substi (high den	rate size; SH, sity of meado	substrate ws); Subs	heteroge trate size	meity; TL, to (average at e	otal length. each samplii	ng site):	[mud (1), sand	(2), grav	/el (3), pebble	e (4) and bou	lder (5)]	Substra	ate

Each sampling site was characterized by 18 environmental variables and indices (quantified in each reach of every sampling site) relating to intra- and interspecific interactions, water quality (weekly mean values) and habitat structure (local level): size of larger *S. pavo* (TL), fish species richness, *S. pavo* abundance and biomass, biomass/abundance ratio of *S. pavo*, benthic fish abundance and biomass, biomass/abundance ratio of benthic fish species, total fish abundance and biomass, biomass/abundance (°C), water salinity, depth (cm), submerged vegetation cover (%), submerged vegetation volume, SS and SH (Table 1).

Benthic fish or fish species captured at each sampling site that tended to move nearer the bottom and which had carnivorous diets (*sensu* Quignard and Pras, 1986; Dumay et al., 2004) were considered as potential competitors. In this way, total fish, benthic fish and *S. pavo* relative abundance and biomass were assessed as mean CPUEs and BPUEs, respectively, at each sampling site.

Submerged vegetation cover (0-100%) was assessed by average value at each sampling site. Submerged vegetation volume was classified as an ordinate categorical variable from 0 (low density of meadows) to 5 (high density of meadows).We classified substrate *sensu* Bain (1999) [mud (1), sand (2), gravel (3), pebble (4) and boulder (5)] and assessed the SS (average at each sampling site) and SH (standard deviation at each sampling site).

Statistical analyses

The condition of *S. pavo* juveniles was indexed by residuals obtained from the least square regression of TM and TL of all captured individuals (log-transformed data) (Sutton et al., 2000). This residual index (Kr) provides an alternative to the more traditional condition indices, e.g. relative condition factor and Fulton's condition factor, and removes body length effects. The mean condition for *S. pavo* juveniles at each sampling site was determined from the average Kr of individuals at each sampling site. To avoid negative values, the variables were previously multiplied by 10^3 , a procedure considered suitable when variables range from 0 to 1 (García-Berthou and Moreno-Amich, 1993). The existence of significant differences between sampling sites was verified by non-parametric Kruskall–Wallis ANOVA analysis.

A stepwise multiple regression analysis was performed to determine the amount of variation in fish condition (mean Kr value at each sampling site) associated with environmental variables (mean values at each sampling site). This regression procedure firsts selects the most correlated independent variable, and then removes the variance in the dependent variable. It then selects the second independent variable which most correlates with the remaining variance in the dependent variable, and so on until selection of an additional independent does not increase the r^2 by a significant amount (P ≤ 0.05).

The colinearity between environmental variables in the regression model was tested using the Tolerance Index (TI) (Visauta-Vinacua, 1997). Bivariate relationships between environmental variables were also analysed using Pearson's correlation.

Statistical analyses were performed with the SPSS® (SPSS Inc., Chicago, IL, USA) software package and a significance level of $P \le 0.05$ was accepted.

Abundance (number biomass for each spe	of individua cies	ls/100 m ²) ar	nd biomass (g	100 m ⁻²) (in 1	brackets) of	captured spe	cies at each s	ampling site, 1	otal fish abur	idance and bi	omass at eac	h sampling si	te and total a	bundance and
	Sampling site													
Species	1	2	3	4	5	6	7	8	6	10	11	12	13	Total
Aphanias iberus		- 00 00 00	-	0.52 (0.14)	(03 U E) I	0.11 (0.02)	36.46 (7.09)	50.52 (8.22)	9.58 (3.36)	10.16 (2.18)	3.13 (1.04)	2.61 (0.44)	5.00 (0.64)	118.07 (23.14)
Atnerina boyeri Pomatoschistus sp.*	0.07 (10.17) 97.94 (17.04)	0.04 (2.83) 81.46 (19.60)	10.40(11.94) 170.21(23.78)	38.44 (8.99	72.23 (8.02)	4.89 (4.29) 62.61 (10.72)	1.29 (0.08) 18.64 (3.05)	20.51 (17.44) 52.19 (11.04)	49.79 (21.18)	24.53 (3.52)	20.77 (11.48) 8.54 (1.34)	49.17 (18.26) 83.64 (8.46)	108.24 (27.84) 38.02 (7.11	798.23 (128.52)
Liza aurata	0.11 (0.31)	, 	, 	, 	0.069(0.34)	0.21 (1.18)	, , 	0.42 (1.48)	, , ,	0.31 (1.09)	, , 	0.31 (0.73)	, I	1.43 (5.13)
Liza ramado	I	0.07 (0.11)	I	I	I	I	I	I	I	I	I	I	I	0.07 (0.11)
Liza saliens	2.49 (1.64)	0.14 (1.42)	0.42(0.96)	28.86 (19.39)	10.42 (1.86)	19.06 (9.46)	43.96 (12.48)	41.36 (13.54)	30.42 (9.59)	71.56 (21.53)	50.63 (20.72)	8.64 (2.38)	4.23 (0.42)	312.21 (115.39)
Mugil cephalus	- 76 (D 6A)	I	I		I	I	0.31 (0.04)	0.31 (0.63)	I	I	I	I	I	0.63 (0.67)
Linonhrvs dalmatinus*	0.70 (0.04) 0.44 (0.16)	0.63 (0.21)	0.11 (0.04)	0.73(0.39)	0.21 (0.12)	0.31 (0.20)	0.11 (0.08)	(00.0) 20.0	0.11 (0.14)		0.21 (0.02)		0.94 (0.19)	3.77 (1.54)
Salaria pavo	4.23 (2.73)	1.25 (1.92)	4.06 (2.28)	18.13 (11.49)	1.67(0.99)	1.56 (7.19)	5.63(3.96)	6.14 (4.54)	28.02 (14.88)	27.81 (9.71)	38.44 (28.61)	7.50 (3.75)	3.23 (4.95)	147.67 (97.04)
Diplodus sargus sargus*	0.11 (0.05)	0.28 (0.14)	I	1.25 (1.03)	0.90 (0.31	0.21 (0.04)	I	I	I	I	I	I	0.11 (0.03)	2.85 (1.60)
Dicentrarchus punctatus	I	I	I	I	I	I	I	0.11 (0.79	I	I	I	I	0.11(0.63)	0.21 (1.42)
Sygnathus typhle	0.54(0.81)	0.21 (0.07)	Ι	Ι	I	0.11(0.39)	Ι	0.11(0.08)	I	Ι	Ι	Ι	0.11(0.14)	1.07(1.50)
Sygnathus abaster	18.74(4.44)	6.94 (1.23)	40.63 (6.64)	7.39 (1.18)	2.15 (0.57)	1.77 (0.33)	3.13 (0.70)	14.69 (0.36)	10.63 (2.69)	4.69 (1.66)	1.25 (0.42)	0.83 (0.12)	23.44 (4.00)	136.27 (24.34)
Hippocampus guitulatus	0.11 (0.11)	0.07 (0.01)	0.31 (0.17)		- 60 (0 65)	0.11 (0.04)	- 0.72 (1.05)	- 114 (20 00)	0.21 (0.13)	- 47 (0.40)	I	- - 11 0 36)	1 50 (0 06)	0.80 (0.46) 37 46 (37 86)
Gobius voouts Gobius niger*	(11.1) (11.0)	0.07 (0.04)	0.21 (0.25)	-	0.35 (0.27)		(co.i) c/.o	(02.7) ±1.1	0.21 (0.33)	(ct:n) /t:n		(0C'0) 11'0	(07.7) 07. 1	(0.97, (1.99))
Gobius paganellus*	1.52 (1.44)	Í	0.21(0.30)	I	Í	Ι	I	I	Í	I	I	I	I	1.72 (1.74)
Callionymus pusillus*	0.22 (0.22)	0.21 (0.14)	2.29(1.69)	I	0.21 (0.10)	Ι	0.11 (0.05)	Ι	I	Ι	Ι	Ι	I	3.03 (2.19)
Symphodus cinereus*	I	0.07 (0.01)	0.63(1.89)	I	I	I	I	0.31 (2.86)	I	I	I	0.21 (0.19)	0.31(0.18)	1.53 (5.12)
Trachinotus ovatus Total	- 133.69 (41.19)	$^{-}$ 102.14 (30.63)	_ 240.73 (54.01)	- 115.21 (57.41)	- 90.56 (14.73)	0.11 (0.04) 91.04 (33.91)	- 116.36 (35.18)	- 188.13 (64.64)	0.11 (0.08) 158.23 (58.49)	- 161.25 (45.57)	- 128.96 (63.63)	- 153.02 (34.80)	- 188.65 (56.11)	0.21 (0.12) 1867.94 (590.29)
Total	133.69 (41.19)	102.14 (30.63)	240.73 (54.01)	115.21 (57.41)	90.56 (14.73)	91.04 (33.91)	116.36 (35.18)	188.13 (64.64)	158.23 (58.49)	161.25 (45.57)		128.96 (63.63)	128.96 (63.63) 153.02 (34.80)	128.96 (63.63) 153.02 (34.80) 188.65 (56.11)

*Benthic fish species

Results

The fish community structure varied across the 13 sampling sites. Of the 21 taxa captured, *Pomatoschistus* sp. (45.5% of total captures), *Liza saliens* (17.9%) and *Atherina boyeri* (17.4%) were the most abundant species. In turn, the dominant species in biomass terms were *A. boyeri* (24.25% of total biomass), *Pomatoschistus* sp. (21.7%) and *L. saliens* (19.4%). *S. pavo* accounted for 8.5% of the total abundance and 16.3% of the total biomass (Table 2).

Benthic fish species considered as potential competitors due to their benthic and trophic habits (Quignard and Pras, 1986; Dumay et al., 2004) represented 48.0% of the total abundance and 29.6% of the total biomass.

Fish condition data (mean Kr values at each sampling site) are presented in Table 3. Kruskal–Wallis ANOVA analysis showed significant differences in fish condition between sampling sites ($\chi^2 = 166.05$; P < 0.0005). Casablanca (sampling site 11) showed the lowest fish condition value, and Los Alíseos (sampling site 3) showed the highest value.

A stepwise multiple regression model indicated that water depth (P < 0.005) and *S. pavo* biomass (P < 0.05) accounted for most of the variations between sampling sites (59.9%) of Kr (Table 4), showing a negative effect in the condition of *S. pavo* juveniles. The TI value was 0.84, with no colinearity between environmental variables included in the regression model.

Bivariate relationships between Kr and environmental variables, and between the different environmental variables, are presented in Table 5. Of note is the positive and significant relationship between water depth and both the *S. pavo* biomass/abundance ratio and size of larger *S. pavo*.

Table 3

Mean condition (Kr \pm 95% CL) for *Salaria pavo* juveniles at each sampling site obtained from standard residuals of the least square regression of total mass and total length of all captured individuals (log-transformed data) and mean TL (TL \pm 95% CL) for *S. pavo* juveniles at each sampling site

Sampling site	n	$Kr~\pm~95\%~CL$	TL \pm 95% CL
1 (Tomás Maestre)	30	-0.04 ± 0.40	2.7 ± 0.2
2 (Las Brisas)	21	-0.59 ± 0.62	2.9 ± 0.4
3 (Los Alíseos)	24	0.38 ± 1.17	3.1 ± 0.2
4 (Isla del Ciervo)	148	$0.16~\pm~0.32$	3.3 ± 0.1
5 (Playa Paraíso)	23	-0.24 ± 0.35	2.7 ± 0.2
6 (Punta del Plome)	27	-0.51 ± 0.26	3.2 ± 0.2
7 (Los Nietos)	69	$0.02~\pm~0.29$	3.6 ± 0.1
8 (Los Urrutias)	25	0.23 ± 1.19	3.5 ± 0.2
9 (Los Alcázares)	26	$0.37~\pm~0.74$	3.5 ± 0.2
10 (La Hita)	120	-0.02 ± 0.52	2.9 ± 0.1
11 (Casablanca)	31	-0.67 ± 0.24	3.5 ± 0.2
12 (Villa Nanitos	20	-0.13 ± 0.52	3.8 ± 0.4
13 (La Calcetera)	20	-0.33 ± 0.37	$3.4~\pm~0.5$

Discussion

Our results showed that the somatic condition of *S. pavo* juveniles differed among sampling sites. Any differences in Kr were probably caused by differences in habitat conditions. Generally, fish condition varies seasonally due to many factors, including reproductive behaviour and gonadal development (Wootton, 1998). The fact that the sampling period was short (2 weeks) and that the study focused on juvenile specimens (TL < 4.5 cm), ensured that the differences in Kr were not related to seasonal variations on somatic condition.

In our site level analysis of the habitat-fish condition relationship, water depth and *S. pavo* biomass (significant variables in the multiple-regression analyses) were the ecological variables that best correlated with *S. pavo* juvenile condition. Sampling sites with higher water depth and with higher values of *S. pavo* biomass (Las Brisas, Punta del Plome and Casablanca) provided lower juvenile fish condition values.

Competition between fish can be an influential factor in fitness, growth, reproduction and survival. Individuals of the same species within the same population compete for similar resources and are potentially intense competitors (Wootton, 1998).

Intraspecific competition for food resources increases with increasing densities of individuals (Heath, 1992). The juveniles of some fish species suffer a decrease in somatic condition as the density of conspecifics increases (Tupper and Boutilier, 1995). Cowan et al. (2000) showed that, during juvenile stages, a reduction in prey resources due to high population densities is an important factor that reduces cohort growth. In the sampling sites which showed the highest S. pavo biomass values (which are related with high population densities), competition among juvenile fish for empty refuges and/or food resources was particularly strong. Indeed, the sampling sites present a relatively low substrate granulometry (mud-sand), thus agonistic behaviour and competition for refuge was almost certainly higher (Faria et al., 1998). In addition, Faria and Almada (2001) pointed out that intraspecific competition in two blenny species (Lipophrys pholis and Coryphoblennius galerita) was stronger than interspecific competition due to differences in the microhabitat use of each species.

In another context, the competitive superiority of larger individuals may reduce the availability of resources for smaller conspecifics, with the result that the dominant fishes show a higher condition value than subordinate fish (Adams et al., 1998; Sloman et al., 2001). In blennies, the dominant individuals of *Lipophrys pholis* have priority over subordinate fish for access to refuges (Gibson, 1968). In addition, agonistic behaviour is common between larger individuals of blenny (e.g. territorial behaviour, competition for refuge and malemale competition for nests, and parental care) (Faria et al., 1998; Gonçalves and Almada, 1998).

Table 4

Stepwise multiple regression models used to predict condition (Kr) of Salaria pavo juveniles from environmental variables

Environmental variables	Regression equations	Adjusted r^2	F	d.f.	Р
Model 1					
Water depth (cm)	Kr = 1.069 - 0.627 (water depth)	0.338	7.128	1, 11	< 0.05
Model 2	· • • •				
Water depth (cm) S. pavo biomass (BPUEs)	Kr = 1.695–0.856 (water depth) -0.570 (<i>S. pavo</i> biomass)	0.599	9.977	2, 10	< 0.005

 r^2 , proportion of variation explained by the regression models; P, probability levels of the regression models.

Table 5 Pearson's correlation coeff	icients fo	or comp	arisons t	oetween Kr	and envire	onmenta	l variables										
Environmental variables	Kr	Size of larger S. pavo (TL)	Fish species richness	<i>S. pavo</i> abundance (CPUEs)	S. pavo biomass (BPUEs)	<i>S. pavo</i> B/A ratio	Benthic fish abundance (CPUEs)	a Benthic fish biomass (BPUEs)	Benthic fish B/A ratio	Total fish abundance (CPUEs)	Total fish biomass (BPUEs)	Total fish B/A ratio	Water temperature (°C)	Salinity (Vegetation cover %)	Vegetation volume	Depth (cm) SS
Size of larger <i>S. pavo</i> (TL) Fish species richness <i>S. pavo</i> abundance (CPUEs)	0.38 0.11 0.02	0.52 0.57	-0.69*														
S. pavo B/A ratio S. pavo B/A ratio Benthic fish abundance	-0.23 -0.50 0.33	$\begin{array}{c} 0.40 \\ 0.77 \\ 0.43 \end{array}$	-0.68* 0.21 0.55	0.91* -0.35 -0.53	-0.07 -0.55*	-0.02											
(CPUEs) Benthic fish biomass	0.27	0.46	0.65*	-0.57*	-0.53	0.02	0.83										
Benthic fish B/A ratio Total fish abundance	$0.03 \\ 0.57*$	0.08	$0.23 \\ 0.04$	-0.25 0.05	-0.15 -0.08	0.06 -0.36	-0.12 0.41	$0.45 \\ 0.41$	0.15								
(CLUES) Total fish biomass (RPUFs)	0.36	-0.50	-0.21	0.53	0.55	-0.23	-0.16	0.09	0.40	0.61^{*}							
Total fish B/A ratio	-0.12	-0.18	-0.32	0.58*	0.75*	1.05	-0.49	-0.18	0.34	-0.19	0.63*						
water temperature (C) Salinity	-0.08	0.21	0.05	-0.08	-0.17	-0.04 0.26	0.17	0.35	0.39	0.01	0.08	-0.24 0.14	-0.50				
Vegetation cover (%)	-0.19	-0.60*	-0.42	-0.19	0.73^{*}	-0.18	-0.65^{*}	-0.44	0.22	0.22	0.70*	0.57*	-0.12	-0.22	*00 0		
vegetation volume Depth (cm)	-0.63*	+C.U- 0.69*	-0.40 0.38	-0.28 -0.62*	-0.40	-0.07	-0.03* 0.23	-0.44 0.12	-0.19 -0.14	-0.43 -0.43	-0.0/* -0.68*	-0.35 -0.35	-0.12 -0.15	-0.18	-0.40 -0.40	-0.32	
SS	-0.23	0.83*	0.47	-0.77*	-0.52	0.58*	0.53	0.61^{*}	0.27	-0.25	-0.43	-0.17	-0.40	0.60* -	-0.67*	-0.62*	0.68^{*}
SH	-0.14	0.45	-0.16	-0.11	-0.04	0.36	-0.04	0.27	0.55	0.03	0.09	0.15	-0.01	0.32 -	-0.06	0.06	-0.11 0.36
CPUEs, catch per unit eff. *Significance level P < 0.(ort; BPU)5.	JEs, bioı	nass per	unit effort;	SS, substr	ate size:	; SH, substr	ate heterogen	leity; TL,	total length							

In our study, sampling sites with higher water depth presented a higher *S. pavo* biomass/abundance ratio and larger *S. pavo* individuals (TL > 4.5 cm) (Tables 1 and 5). This situation was related to the presence in these deeper sampling sites of adult and dominant fishes, which presumably monopolized refuges and food resources. In addition, because the sampling period coincided with the breeding season, the larger *S. pavo* fish showed territorial behaviour and defended small territories around the nests against intruders (whether of the same or of different species) (Gonçalves and Almada, 1998; Gonçalves et al., 2005). Consequently, adult breeding fish could produce an important level of stress in *S. pavo* juveniles, which would be reflected in lower somatic condition.

We found no relationship between the other variables of habitat structure (except depth) and juvenile fish condition, probably due to the number of sampling sites or perhaps because any relationship was clouded by the very complexity of the ecological interactions (e.g. a non-linear relationship between these variables). To a certain extent, this demonstrates the need for more investigation into the relationships between habitat characteristics, environmental variations and juvenile fish condition in the study area.

In conclusion, the somatic condition of the early phases of the life history of juvenile *S. pavo* in the shallow areas of the Mar Menor was directly affected by their biomass and depth, which are probably related to intraspecific competition, both intracohort and intercohort. Shallow areas with lower *S. pavo* biomass and a lower presence of adult breeding fish produced juvenile fishes with a better somatic condition because they were probably able to avoid competition for refuge and/or food resources with juvenile conspecifics and aggressive interactions with larger males.

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