Thermal imprinting modifies adult stress and innate immune responsiveness in the teleost sea bream

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Abstract

The impact of thermal imprinting on the plasticity of the hypothalamic–pituitary–interrenal (HPI) axis and stress response in an adult ectotherm, the gilthead sea bream (*Sparus aurata*, L.), during its development was assessed. Fish were reared under 4 thermal regimes, and the resulting adults exposed to acute confinement stress and plasma cortisol levels and genes of the HPI axis were monitored. Changes in immune function, a common result of stress, were also evaluated using histomorphometric measurements of melanomacrophage centers (MMCs) in the head kidney and by monitoring macrophage-related transcripts. Thermal history significantly modified the HPI responsiveness in adult sea bream when eggs and larvae were reared at a higher than optimal temperature (HT, 22°C), and they had a reduced amplitude in their cortisol response and significantly upregulated pituitary *pomc* and head kidney *star* transcripts. Additionally, after an acute stress challenge, immune function was modified and the head kidney of adult fish reared during development at high temperatures (HT and LHT, 18–22°C) had a decreased number of MMCs and a significant downregulation of dopachrome tautomerase. Thermal imprinting during development influenced adult sea bream physiology and increased plasma levels of glucose and sodium even in the absence of an acute stress in fish reared under a high–low thermal regime (HLT, 22–18°C). Overall, the results demonstrate that temperature during early development influences the adult HPI axis and immune function in a teleost fish.

Key Words

- confinement
- HPI axis
- melanomacrophage centers
- plasticity
- thermal imprinting

Journal of Endocrinology
(2017) **233**, 381–394

Introduction

In mammals, the way in which pre-natal or post-natal ‘adverse challenges’ has an impact on the hypothalamic–pituitary–adrenal (HPA) axis is extensively studied and linked with dysfunction during adulthood and associated pathologies (Fahmi *et al.* 2004, Breton 2013, Devlin *et al.* 2013, Johnsen *et al.* 2013, Gonzalez-

Bulnes *et al.* 2014). Early life exposure to stress induces plasticity of the HPA axis and is associated with either hyper-responsiveness to a stress challenge or hypo-responsiveness characterized, respectively, by exaggerated or attenuated levels of glucocorticoids (Elzinga *et al.* 2003, Carpenter *et al.* 2007). Modifications in the HPA axis

during childhood are proposed to provoke dysfunction in adulthood due to epigenetic mechanisms that lead to persistent changes in the expression of genes such as the glucocorticoid receptor (GR), together with lifelong alterations in DNA methylation and histone 3 lysine 9 (H3K9) acetylation (Zhang *et al.* 2010).

Despite differences in stress axis organization between mammals and fish, the overall regulatory loop is conserved and the hypothalamic–pituitary–interrenal (HPI) axis in fish is the homologue of the HPA axis in mammals. Stress in fish triggers the release of hypothalamic, corticotropin-releasing hormone (CRH), which in turn stimulates the synthesis of the prohormone proopiomelanocortin (POMC) and release of adrenocorticotrophic hormone (ACTH) from the pituitary. In the adrenal gland or interrenal tissue of the head kidney in fish, ACTH regulates the production of cortisol by modulating the rate-limiting enzyme, steroidogenic acute regulatory protein (StAR). Cortisol exerts its effect by binding to the GR in target cells and modifies a range of physiological processes such as metabolism and immune function to maintain allostasis (Wendelaar-Bonga 1997).

The ability of juvenile and adult fish to cope with a stressor has been extensively studied, and cortisol levels are generally fairly well correlated with the performance of fish under acute or chronic stress (Wendelaar-Bonga 1997, Mommsen *et al.* 1999, Prunet *et al.* 2008, Aluru & Vijayan 2009, Harper & Wolf 2009, Pankhurst 2011). A notable feature of the stress response in fish is its plasticity, and individuals of the same species can be low- or high-cortisol responders (Barton 2002) and this has led to the suggestion that genetic, developmental and environmental factors determine an individual's stress response (Pottinger & Pickering 1997, Barton 2002, Barton *et al.* 2005, Cnaani 2006). Surprisingly, little attention has been given to understanding the potential lifelong consequences on the stress axis of early gene–environment interactions in teleost fish.

Temperature is a key regulator in ectotherms that strongly influences development and may have a long-term impact on phenotype or genotype and influence adaptive evolution (Somero 2010, Mozes *et al.* 2011). The developmental plasticity of fish means that modifications in the thermal regime during embryonic development can significantly modify their developmental trajectory (Schaefer & Ryan 2006). Skeletal deformities (Boglione & Costa 2011) and modifications in muscle fiber number and growth (Johnston 2006, Garcia de la Serrana *et al.* 2012,

Vieira *et al.* 2012b) are well-established examples of phenotypic plasticity induced by thermal imprinting.

Exposure of early life stages of sea bass (Varsamos *et al.* 2006) and rainbow trout (Auperin & Geslin 2008) to stress results in decreased cortisol production in juveniles exposed to stress. Stress not only acts on the central stress axis but also causes changes in the periphery to re-establish homeostasis, and one well-recognized target is the immune system (Dhabhar 2008). In fish, as in other vertebrates, the majority of stress responses lead to suppression of the immune response (Tort 2011). Stressful conditions including parasitic infections in fish induce modifications in the number, size and pigment content of aggregates of macrophages, the melanomacrophage centers (MMCs) of the piscine innate immune system (Fournie *et al.* 2001, Agius & Roberts 2003, De Vico *et al.* 2008), and this is the predominant immune response during early development (Tort *et al.* 2003).

The present study characterizes the developmental plasticity of the stress axis to thermal imprinting in the gilthead sea bream (*Sparus aurata*), a teleost with a well-studied stress response (Arends *et al.* 1999, Montero *et al.* 1999, Rotllant *et al.* 2001, Barton *et al.* 2005, Caldutch-Giner *et al.* 2010). The hypothesis tested was that thermal imprinting during development would modulate the activity of the HPI axis and lead to a modified stress response in adults. The response to acute confinement stress of young adults that were exposed during development to different thermal regimes was assessed by determining the responsiveness of the HPI axis and elements of the secondary stress response. The influence of thermal history on immune responsiveness to stress at the level of the head kidney, a prominent immune tissue in fish (Whyte 2007) was also evaluated.

Materials and methods

Early life programming

All the procedures for manipulation of embryonic and larval thermal regimes and rearing to juveniles as well as the stress challenge experiments were performed at the Institute for Aquaculture and Food Technology Research (IRTA), St Carles de la Ràpita, Spain, in a temperature-controlled seawater recirculation system (IRTamar). All animal handling procedures were approved by the Ethics and Animal Care Committee (4998-T9900002) and were compliant with the guidelines of the European Union Council (86/609/EU), Spanish and Catalan Governments

legislation. No evidence of infection, modified behavior or mortality was observed during the experiments.

Fertilized eggs of gilthead sea bream (fertilization rate=92%) were maintained during embryogenesis at $18 \pm 0.5^\circ\text{C}$ (low temperature (LT)) or $22 \pm 0.5^\circ\text{C}$ (high temperature (HT)) in duplicate tanks containing 110 mL of fertilized eggs per incubator (30 L). Newly hatched larvae were maintained under four different temperature regimes (2 replicate tanks/group, Fig. 1/Supplementary Fig. 1 and Supplementary Tables 1 and 2, see section on supplementary data given at the end of this article): 18°C from egg incubation through hatching and up until larvae-juvenile transition (LT), 22°C from egg incubation through hatching and up until larvae-juvenile transition (HT), 18°C from egg incubation up until hatching and

then 22°C until larvae-juvenile transition (LHT), and 22°C from egg incubation up until hatching and then 18°C up until larvae-juvenile transition (HLT). Fish from the four temperature regimes were maintained in 2 m^3 tanks connected to a recirculating sea water system (5–10% water renewal per day; IRTAmar) under the same constant temperature regime of $22 \pm 1^\circ\text{C}$ until they were 9 months old. Fish from the four temperature regimes were fed 3% body mass (m/m) with a commercial diet (OptiBream; Skretting, Spain).

Confinement challenge and sampling

Duplicate tanks of fish from each thermal regime were exposed to an acute confinement stress. Sea bream stocked

Figure 1

Schematic representation of the experimental setup. (A) Temperature program imposed to gilthead sea bream from egg fertilization to larvae-juvenile transition. Four temperature treatments (thermal groups) were considered: 2 constant temperatures ($18\text{--}18^\circ\text{C}$ (LT) and $22\text{--}22^\circ\text{C}$ (HT)) and 2 variable temperatures between incubation phase and larval rearing ($22\text{--}18^\circ\text{C}$ (HLT) and $18\text{--}22^\circ\text{C}$ (LHT)). When the fishes' bodies were covered with scales (larvae-juvenile transition), fish from all thermal groups were maintained at 22°C until the onset of the acute stress challenge by confinement ($n=20$ /tank; 2 replicate tanks per thermal history group). Details about differences in the development of fish from different thermal regimes and the time of temperature change/group can be accessed in Supplementary Tables 1 and 2 and also in Supplementary Fig. 1B. Acute stress challenge by confinement and the sampling regime. Adult sea bream ($n=5$ /replicate tank) stocked at 20 kg/m^3 were sampled prior to the application of an acute stress challenge (0h). The remaining fish were exposed to an acute stress by reducing the level of water of each tank, which increased the density to 70 kg/m^3 . Fish were exposed to confinement stress for 30 min before returning them to the initial stocking density (20 kg/m^3) and then sampled ($n=5$ /replicate tank/sampling time) at 1, 4 and 24 h after a stress challenge.

Table 1 General physiological status of the gilthead sea bream with different thermal histories.

Thermal history	Weight (g)	Length (cm)
LT (18–18°C)	155.7 ± 20.2a	15.79 ± 0.86 ^a
HLT (22–18°C)	180.2 ± 28.0b	16.20 ± 0.96 ^a
LHT (18–22°C)	227.1 ± 30.3c	18.03 ± 1.04b
HT (22–22°C)	201.1 ± 20d	16.99 ± 0.73c

Body weight (g) and standard length (cm) of fish from each thermal regime ($n=40$). Data from all sampling times were pooled. Different letters indicate statistical significant difference between thermal groups. Data shown as mean ± s.d.; One-way ANOVA; $P < 0.05$.

at a density of 20 kg/m³ ($n=40$ /group, weight and length are given in Table 1) were acclimated to the experimental circuit for 1 week, which consisted of a recirculating sea water system (IRTamar) with eight 200 L square fiberglass tanks. Acclimating conditions were: 21 ± 1°C, pH 7.5–8, 35–36‰ salinity, >80% oxygen saturation and 12-h light/12-h darkness photoperiod. Fish were fed by hand five times per day using a commercial diet (OptiBream, Skretting) at a ration level of 3% body mass (m/m). Fish ($n=5$ /tank) were sampled before application of the acute stress challenge (time 0). The remaining fish were exposed to an acute stress by increasing their tank density to 70 kg/m³ by lowering the water level for 30 min before increasing the water volume to return them to the initial stocking density. Fish were then sampled ($n=5$ per tank/sampling time) at 1, 4 and 24 h after the stress challenge. For sampling, fish were killed with an overdose (450 ppm) of 2-phenoxyethanol (Sigma-Aldrich) and blotted dry and blood was collected from the caudal vein using a heparinized syringe, centrifuged at 7000g for 4 min at 4°C and the plasma was stored at –80°C. The pituitary was removed and collected into RNAlater (Sigma-Aldrich) and stored at –80°C. The head kidney was dissected out and half of it was placed in RNAlater and the remainder was fixed in Bouin-Hollande for histomorphometric measurements of MMCs in the head kidney.

Biochemical analysis

Plasma cortisol (ng/mL) was measured using a radioimmunoassay as previously described (Rotllant *et al.* 2005). Glucose (mmol/L) and total calcium (mmol/L) in plasma samples were measured with glucose oxidase–peroxidase and o-Cresolphthalein colorimetric commercial kits (Spinreact 1001190 and 1001061, Spain, respectively). Total protein (mg/mL) was measured in plasma using a commercial kit, Protein Assay Dye Reagent Concentrate (#500-0006), a Quick Start BSA Standard

Set (#500-0207, Bio-Rad), and the manufacturer's instructions were followed. Spectrophotometry of colorimetric assays was performed in 96-well plates using a plate reader (Benchmark, Bio-Rad). Osmolality (mmol/kg) was determined using a vapor pressure osmometer (Vapro Wescor 5520, Utah, USA), and sodium (mM) and potassium (mM) concentrations were determined by flame photometry (BWB Technologies, USA).

MMCs morphometric analysis

Head kidney fixed in Bouin-Hollande ($n=3$ /tank/time point) was dehydrated through a graded ethanol series (70–100%) and embedded in low melting point paraffin wax using a tissue processor (Leica TP1020, Leica). Serial longitudinal sections (5 µm) were cut using a rotary microtome (Leica RM 2135) and mounted on glass slides coated with 3-amino-propyltriethoxysilane (APES; Sigma-Aldrich). One section per individual was stained with hematoxylin and eosin to assess general histology, and 3 slides containing 3 serial sections/slide with a gap of 5 sections (25 µm) between consecutive slides were stained with a combination of Alcian blue–Periodic Acid Schiff (PAS)–Orange G. Stained sections were observed using a microscope (Leica DM2000) equipped with a digital camera (Leica DFC480) and 1 section per slide was photographed.

The images obtained were analyzed using Fiji v1.47p software (Schindelin *et al.* 2012). One field of 0.3 mm² per PAS stained section ($n=6$ /group/time point) was randomly selected and semi-automatic analysis of MMCs was carried out by setting a color threshold that highlighted all dark pigment cells as red (Papadopoulos *et al.* 2007). The parameters analyzed were the number of MMCs with an area superior to 0.001 mm² (larger MMCs with a well-defined oval shape); number of MMCs with an area equal or inferior to 0.001 mm² (smaller MMCs, poorly defined); total number of MMCs and total area the MMCs occupied in the defined area.

Analysis of gene expression by quantitative real-time PCR (qPCR)

Total RNA from the pituitary was extracted using PureZOL RNA Isolation Reagent (Bio-Rad), and total RNA from the head kidney was extracted using a Maxwell 16 System (Promega) following the manufacturer's instructions. The concentration and quality of the extracted RNA were determined using NanoDrop 1000 spectrophotometer

(Thermo Fisher Scientific). To eliminate contamination of genomic DNA, total RNA (2–9 µg) was treated with DNase using a DNA-free kit (Ambion). cDNA synthesis was carried out in a 20 µL reaction volume with 500 ng of DNase-treated RNA, 200 ng of random hexamers (Jena Biosciences, Germany), 100 U of RevertAid reverse transcriptase (Fermentas, Thermo Fisher Scientific), 8 U of RiboLock RNase Inhibitor (Fermentas) and 0.5 mM dNTPs. The reaction was incubated for 10 min at 20°C followed by 60 min at 42°C, and the enzyme was inactivated by heating for 5 min at 72°C.

qPCR was used to analyze the mRNA expression of candidate indicator genes of the central stress axis and associated with MMC function before (0) and 4 and 24 h after the acute stress challenge. To monitor how stress modulates the activity of the pituitary gland, the duplicate *pomc* genes ($\alpha 1$ and $\alpha 2$) and *gr* were measured, and in the head kidney, *star* and *gr* transcripts were measured. To assess MMC activity, expression of colony-stimulating factor-1 receptor (*csf1r*) and Tyrosinase (*tyr*) family members *dct* and *tyr* was quantified in the head kidney. Since members of the tyrosinase gene family are not described in sea bream, database mining and phylogenetic analysis were performed to identify them (Supplementary Fig. 2 and Supplementary Table 3). Specific primers for each transcript (Table 2) were designed using Primer Premier 5.0 software (Premier Biosoft Int., CA, USA).

Reactions were performed in duplicate for a 15 µL final volume containing 10 ng of cDNA (25 ng for *dct* and *tyr*

mRNA expression analysis) and 300 nM of specific primers (Table 2) and EvaGreen (SsoFast EvaGreen supermix, Bio-Rad Laboratories). The reaction was carried out in a StepOnePlus qPCR thermocycler, and data were analyzed with StepOne software v2.2 (Applied Biosystems). qPCR cycling conditions were 30 s at 95°C, 40 cycles of 5 s at 95°C and 10 s at 60°C followed by a final melt curve between 60 and 95°C, which gave single product/dissociation curves in all reactions.

Transcripts were quantified using the relative standard curve method (Vieira *et al.* 2012a), generated using serial dilutions of specific qPCR products for each gene (obtained using the same species, tissues and primers for qPCR analysis). All amplicons were sequenced to confirm their identity. Control reactions included a no-template control and a cDNA synthesis control (reverse transcriptase omitted from the reaction). Two candidate reference genes routinely used for sea bream (*rps18* and β -actin) were tested in pituitary and head kidney samples. *Rps18* did not vary significantly in the pituitary samples and was used as the reference gene. β -Actin did not vary significantly in the head kidney and was used as reference gene. Results were calculated by dividing the detected copy number of the target gene by the reference gene in each cDNA sample and expressed as Log2 Fold Change. Results were calculated relative to the LT group at time 0, which was the group reared during the egg and larval stages using the standard temperature regime used for gilthead sea bream (Mozes *et al.* 2011).

Table 2 Primers used for quantitative RT-PCR expression analysis.

Gene	Accession No.	Sequence (5'–3')	Amplicon (bp)	Ta (°C)	Efficiency (%)	R ²
<i>pomc</i> $\alpha 1$	HM584909	F: CTTGAAGAAACCAATGAACATC R: GAAACAGCCAATGAAGACCTAA	162	60	101	0.988
<i>pomc</i> $\alpha 2$	HM584910	F: GCTCGTTAGCAGACCAAT R: CAAAACACTCTCTCTCATCTCT	76	62	91	0.999
<i>gr</i>	DQ486890	F: CCATCACCTCTGCCGATCTG R: CTGGAGGAAGTGTGCTGAACC	195	64	92	0.994
<i>star</i>	EF640987	F: ACATCGGGAAGGTGTTCAAG R: TCTCTGCAGACACCTCATGG	177	62	108	0.999
<i>csf1r</i>	AM050293	F: ACGTCTGGTCTATGGCATC R: AGTCTGGTTGGGACATCTGG	129	62	104	0.997
<i>dct</i>	isotig44494*	F: GCTCTGATCCTGCCAACG R: TCGTCAACATTTCTCAGCCT	112	60	98	0.995
<i>rps18</i>	AM490061	F: AGGGTGTGGCAGACGTTAC R: CTTCTGCCTGTTGAGGAACC	164	60	91	0.997
β -Actin	X89920	F: CCCTGCCACGCCATCC R: TCTCGGCTGTGGTGGTGAAGG	94	60	86	0.996

Gene name, accession number, primer sequence, amplicon length (bp), primer annealing temperature (Ta, °C) and qPCR efficiency (%) and R² are indicated for each primer pair (F = forward and R = reverse).

*Accessed in the database assembly for gilthead sea bream transcripts (Louro *et al.* 2016).

Table 3 Variation of secondary response parameters in gilthead sea bream with different thermal histories.

Thermal history	Glucose (mmol/L)				Potassium (mM)				Sodium (mM)			
	0	1	4	24	0	1	4	24	0	1	4	24
LT	4.3±0.2a	4.2±0.4	4.1±0.6	4.0±0.5	3.2±0.2a	4.2±0.4	4.3±0.3a	4.7±0.4	176.4±2.5a	218.9±15.0	225.8±8.3	233.1±6.7a
LHT	4.2±0.4a	4.3±0.3	3.9±0.3	4.1±0.7	3.4±0.2ab	4.1±0.4	4.2±0.3a	4.7±0.4	173.8±8.7a	223.5±5.2	226.7±9.2	231.3±13.1a
HT	4.3±0.2a	4.3±0.3	3.9±0.6	3.7±0.3	3.4±0.2ab	4.0±0.4	4.5±0.3ab	4.5±0.2	166.2±2.4b	218.7±4.5	219.4±2.9	230.1±7.2a
HLT	4.8±0.5b	4.3±0.4	4.2±0.5	4.1±0.7	3.6±0.3b	4.1±0.6	4.8±0.4b	4.4±0.2	199.6±6.8c	214.4±4.7	228.8±13.7	214.8±1.8b

Statistical analysis

All statistical analyses were performed with SPSS 22.0 software (SPSS, 2013). No significant differences were detected between tank replicates, so all data of fish with a given thermal history at each time point were pooled. Two-way analysis of variance (ANOVA) was used to test the significant differences between fish with different thermal histories at different times after exposure to acute stress. One-way ANOVA was then performed to identify significant differences: (1) in the general physiological status of groups before the application of the acute stress challenge (time 0), (2) between different thermal history groups at the same time point (intergroup comparison) and (3) between the same thermal history group at different time points (intra-group comparison). If significant differences were detected, a Tukey's multiple comparison test was performed. Data were considered significantly different at $P < 0.05$. Log10 transformation of data was used whenever necessary to achieve either normal distribution or equal variance assumptions, and data transformed for analysis are indicated in the respective figure legends. Data are presented as mean ± standard error of the mean (S.E.M.) unless otherwise stated.

Results

Characterization of the general physiological status of experimental fish at time zero

The general physiological status of fish with different thermal histories was assessed before the experimental challenge by measuring body weight and standard length (Table 1). One-way ANOVA revealed that fish from different thermal regimes were significantly ($P < 0.001$) different in most of the biometric parameters assessed. No significant differences in biometric parameters of experimental fish

were detected during the experiments (1, 4 and 24 h). Fish from the LT group were significantly ($P < 0.001$) lighter (155.7 ± 20.2 g) and shorter (15.79 ± 0.86 cm) than LHT and HT fish (Table 1).

Basal plasma glucose concentrations were influenced by thermal imprinting ($P = 0.025$), and before the start of the experimental challenge, the HLT group exhibited significantly ($P < 0.05$) higher glucose concentrations than the other groups. Thermal imprinting also influenced plasma potassium, sodium and total protein concentrations (Table 3). Early thermal regimes did not significantly modify the plasma osmolality or calcium concentration in adult fish (Table 3).

Before the onset of the acute stress challenge, no significant differences were detected in plasma cortisol levels between different groups and cortisol varied from 5.35 ± 1.92 ng/mL (LHT) to 18.40 ± 7.89 ng/mL (HLT, Fig. 2). In the pituitary, expression of *pomca2* was significantly upregulated ($P = 0.04$) in the HLT relative to the LHT group at time 0, but thermal history did not have an impact on *pomca1* transcript abundance. Similarly, *star* transcripts in the head kidney were significantly ($P < 0.05$) upregulated in the HT relative to the LT group. *Gr* transcripts in the head kidney and pituitary of sea bream with different thermal histories were not significantly different at time 0.

Thermal history had no significant impact on the histology (Fig. 3) or transcript abundance of *dct* and *csf1r* in the head kidney (Fig. 4).

Characterization of the fish physiological response to the acute stress challenge

Plasma cortisol Plasma cortisol levels were significantly affected ($P < 0.001$, Two-way ANOVA) after acute stress (Fig. 2). Thermal history did not significantly modify the amplitude of the cortisol response in any of

Osm (mmol/kg)				Protein (mg/mL)				Calcium (mmol/L)			
0	1	4	24	0	1	4	24	0	1	4	24
267.0±7.0	347.6±8.7	347.9±6.5a	350.8±9.0ab	21.8±1.0a	21.9±2.6	22.0±1.4	23.6±2.2ab	3.3±0.1	3.3±0.1a	3.1±0.1a	3.3±0.1
262.6±7.5	344.6±5.2	342.0±5.2ab	358.6±5.8a	21.6±2.1a	21.7±1.6	20.1±2.7	24.9±1.8a	3.4±0.1	3.2±0.1a	3.2±0.1ab	3.3±0.2
270.1±10.6	340.2±5.8	339.0±5.5b	355.6±8.7ab	24.3±2.2b	21.3±2.8	20.5±2.7	23.7±2.1ab	3.5±0.3	2.9±0.1b	3.1±0.1a	3.3±0.1
268.9±6.7	343.6±5.0	347.3±5.8a	346.4±7.5b	21.9±2.4ab	21.3±2.1	21.0±1.8	21.6±1.6b	3.5±0.3	3.1±0.2a	3.3±0.2b	3.4±0.4

Glucose, potassium, sodium, electrolyte-water balance (osmolality), total protein and total calcium of plasma were measured prior (time 0) and after acute stress confinement (1, 4 and 24 h) in the different thermal regimes groups (LT (18–18°C); LHT (18–22°C); HT (22–22°C); HLT (22–18°C)). Statistical differences between the groups are indicated by different letters of the same sampling time. Data is shown as mean±s.d. ($n=10$ individuals per group per sampling time); Two-way ANOVA; $P<0.05$.

HLT, high–low temperature regime; HT, high temperature regime; LHT, low–high temperature regime; LT, low temperature regime.

the experimental groups at any of the time points post stress (intergroup comparison). Intra-group analysis revealed that a short-term hormonal response was observed in all groups irrespective of thermal history, and 1 h after confinement, cortisol levels significantly increased to 113.6 ± 29.26 ng/mL ($P<0.001$) in the LHT group and to 37.2 ± 7.64 ng/mL ($P<0.01$) in the HT group when compared to time 0. Twenty-four hours after application of the stress, only the cortisol levels of the LHT group were significantly ($P=0.01$) increased relative to basal levels.

Pituitary *pomc* transcripts Expression of the duplicate *pomca1* and *pomca2* was modified after acute stress challenge ($P=0.022$ and $P<0.001$, respectively, two-way ANOVA), but only *pomca2* was significantly ($P<0.001$, Two-way ANOVA) modified by thermal history (Fig. 2). Intergroup comparison revealed that at 4 h post stress, significant ($P=0.04$ and $P<0.001$, respectively) upregulation of both *pomca1* and *pomca2* transcripts was observed in the HT fish relative to the LT at time 0. In the same time period, only *pomca2* was significantly ($P=0.01$) upregulated in HLT and LHT relative to LT at time 0. At 24 h post stress, *pomca2* remained significantly ($P<0.05$) upregulated in LHT, HT and HLT and also became significantly $P<0.05$ upregulated in LT fish, all relative to LT at time 0. *Pomca1* was also significantly ($P<0.05$) upregulated in the LHT and HLT fish at 24 h relative to LT at time 0. In contrast, *pomca1* was significantly ($P<0.05$) downregulated in LT relative to HLT 24 h post stress.

Intra-group analysis revealed that *pomca1* was significantly ($P=0.02$) upregulated at 4 h in the LT fish relative to time 0 and then decreased significantly ($P=0.02$) at 24 h (Supplementary Table 4). *Pomca2* was significantly ($P<0.05$) upregulated in LHT and HT at 4 and 24 h, in relation to time 0 (Supplementary Table 4).

Head kidney *star* transcripts Two-way ANOVA revealed that after an acute stress challenge, a significant ($P=0.016$) interaction occurred between thermal history and time in the relative abundance of *star* transcripts (Fig. 2). Intergroup comparisons revealed that the expression of *star* was significantly ($P<0.05$) upregulated in the HLT fish 24 h post stress relative to LT at time 0. Intra-group analysis indicated that in HT fish, the expression of *star* was significantly ($P<0.05$) downregulated at 24 h relative to 0 h (Supplementary Table 4). None of the other groups revealed a significant modification in the expression of *star*.

Pituitary and head kidney *gr* transcripts Abundance of *gr* transcripts in the pituitary was significantly ($P<0.001$, two-way ANOVA) modified by the stress challenge, but no significant effect was observed in the head kidney ($P>0.05$, two-way ANOVA, Fig. 2). Intergroup analysis showed that at 4 h, *gr* expression in the pituitary was significantly ($P<0.05$) downregulated in the LHT fish relative to LT at time 0. At 24 h, *gr* abundance in the head kidney of LHT fish was significantly ($P<0.05$) upregulated relative to LT at time 0 and to HT fish. Intra-group analysis revealed that *gr* expression in the pituitary at 4 h was significantly ($P<0.05$) downregulated in all groups relative to time 0, but by 24 h, returned to basal levels (Fig. 2).

Characterization of the secondary response to acute stress challenge

Plasma glucose concentrations in response to acute stress were significantly influenced by time ($P<0.001$) and thermal history ($P=0.025$, two-way ANOVA, Table 3). Intra-group analysis revealed that plasma glucose significantly ($P<0.05$, one-way ANOVA) decreased

Figure 2

Cortisol levels and expression of molecular indicators (*pomc α1*, *pomc α2*, *gr* and *star*) of the overall stress response of the four experimental thermal groups after acute stress confinement. Cortisol levels were Log10 transformed and measured before (0h) and after acute stress at three time points (1, 4 and 24h). Gene expression was assessed by qPCR at 0, 4 and 24h, and in the pituitary gland, expression of *pomcα1*, *pomcα2* and *gr* was normalized by the mean of *rps18* expression, and in the head kidney, the relative gene expression of *star* and *gr* by the mean of β -actin expression. Normalized data were expressed as Log2 Fold Change relative to LT 18–18°C at 0h. The results are plotted in Tukey box and whiskers graphs: LT (18–18°C); LHT (18–22°C); HT (22–22°C); HLT (22–18°C). '+' represents the mean and dots the outliers ($n=10$ individuals per group per sampling time). Different letters indicate statistical significant differences between the groups of the same sampling time (intergroup analysis). Capital letters under the graph of plasma cortisol indicate if significant differences exist between sampling times for each thermal group (intra-group analysis, ns=not significantly different). Groups with the same letter are not significantly different. Significant upregulation or downregulation relative to LT (18–18°C) at 0h was denoted by: * $P<0.05$, ** $P<0.01$, *** $P<0.001$ using two-way ANOVA.

relative to basal levels in the HLT and HT fish 24h after acute stress (Supplementary Table 5).

Interaction between thermal history and time significantly ($P<0.01$, two-way ANOVA) modified plasma protein together with potassium, sodium, calcium and osmolality that were also significantly ($P<0.001$, two-way ANOVA) changed after confinement stress (Table 3). Intergroup comparisons showed that at 24h post confinement, plasma osmolality and total protein were significantly ($P<0.05$) higher in the LHT fish relative to the HLT fish. The HLT group also had a significantly ($P<0.05$) lower plasma sodium concentration at 24h compared

to all other experimental groups. Calcium levels in the HT group were significantly ($P<0.05$) decreased 1h post confinement in relation to all other experimental groups.

Intra-group analysis revealed that plasma sodium, potassium and osmolality increased significantly ($P<0.05$) in all groups 1h after confinement and remained significantly ($P<0.05$) higher up until 24h, in relation to time 0 (Supplementary Table 5). In all experimental groups, plasma calcium decreased significantly ($P<0.05$) 1h after the onset of acute stress relative to time 0 and subsequently returned to basal concentrations 24h post confinement.

Figure 3

Histological (A) and histomorphometric (B) analysis of gilthead sea bream MMCs in the head kidney before and 1, 4 and 24 h after the acute stress challenge. (A) MMCs in head kidney were stained with Alcian blue–Periodic Acid Schiff (PAS)–Orange G. Different thermal groups (LT (18–18°C); HLT (22–18°C); LHT (18–22°C); HT (22–22°C)) are represented. Erythrocytes enriched area is stained in orange, and lymphoid tissue is stained in purple. MMCs may consist in larger clusters of cells (area >0.001 mm²; arrowhead) or as free cells (area <0.001 mm²; arrow; e). MMCs may present a round (a), oval (c) or irregular shape (g) and is frequently observed near blood vessels (bv, e). Both free and larger MMCs contain dark brown pigmented material. ad, adipocytes. Scale bar = 100 µm. (B) Histomorphometric analysis of MMCs in a randomly selected area of 0.3 mm² in head kidney: number of MMCs with an area equal or inferior to 0.001 mm² (smaller MMCs), total number of MMCs found within the analyzed area and the total area (mm²) occupied by MMCs in the selected area. Analysis of MMCs was performed in unstressed fish (0 h) and during stress response (1, 4 and 24 h). Different thermal regimes (LT (18–18°C); LHT (18–22°C); HT (22–22°C); HLT (22–18°C)) are represented. Different letters indicate statistical difference between thermal groups of the same sampling time. Data shown as mean ± S.E.M.; two-way ANOVA; $P < 0.05$; $n = 10$. A full colour version of this figure is available at <http://dx.doi.org/10.1530/JOE-16-0610>.

Characterization of MMCs following acute stress challenge

Histology of the head kidney MMCs were randomly distributed in the head kidney and were evident as large pigmented masses surrounded by highly cellular tissue stained with a purple or an orange hue due to the accumulation of lymphocytes and erythrocytes, respectively (Fig. 3A). Occasional blood vessels surrounded by cords of cells of interrenal tissue, pale

chromaffin cells and proximal renal tubules were evident in the tissue. The main changes 24 h after acute stress were a decrease in the number of small, highly pigmented masses in the LHT and HT groups and the change from well-delineated spherical MMCs to heterogeneous masses in the LHT group.

Thermal history and time after stress interacted and were associated with a significant ($P < 0.01$, two-way ANOVA) change in the number of small MMCs,

Figure 4

Relative expression of *dct* and *csf1r* in the head kidney before the onset of confinement (0h) and during response (4 and 24h) to stress. Transcript expression levels were obtained by qPCR and were normalized in relation to the mean of β -actin expression and normalized results were expressed as Log2 Fold Change calculated relative to LT 18–18°C at 0h. The results are plotted in Tukey box and whiskers graphs: LT (18–18°C); LHT (18–22°C); HT (22–22°C); HLT (22–18°C) are represented. '+' represents the mean and dots the outliers ($n=10$ individuals per group per sampling time). Different letters indicate statistical significant difference between thermal groups of the same sampling time (intergroup analysis). Data are shown as mean $\pm$ S.E.M. ($n=10$ per group) and significant upregulation or downregulation relative to LT (18–18°C) at 0h was denoted by: * $P<0.05$, ** $P<0.01$, *** $P<0.001$ using two-Way ANOVA.

and consequently, the total number of MMCs, and also in the total area occupied by MMCs in the head kidney at time 0 and during the acute stress response (Fig. 3B). Intergroup comparisons revealed that thermal history had a significant ($P=0.01$, one-way ANOVA) impact on the number of small MMCs and the total number of MMCs and that the HLT fish had significantly ($P<0.05$) fewer MMCs at 1h when compared to the LHT fish. Conversely, 24h after acute stress, the LHT and HT fish exhibited significantly ($P<0.05$) fewer small MMCs relative to the LT group. Although no significant differences were found between groups at 4h post stress in the number of small and total MMCs, the total area occupied by MMCs was significantly ($P=0.01$) smaller in the HLT group compared to LT fish (Fig. 3B). Intra-group comparisons revealed that the number of small MMCs in both LHT and HT groups significantly ($P<0.05$, One-way ANOVA) decreased between time 0 and 24h after acute stress, although the total area of MMCs in the head kidney was not significantly modified in these groups between 0 and 24h post stress (Supplementary Table 6). The number of large MMCs was not significantly modified by confinement stress in any experimental fish at any time point (data not shown).

qPCR of MMC relevant transcripts in the head kidney *Csf1r* expression was highly variable in all experimental groups and was significantly ($P=0.003$, two-way ANOVA) affected by time after acute stress. Intergroup analysis revealed that significant ($P<0.05$) upregulation of *csf1r* transcripts only occurred in the LHT group 4h after acute stress exposure, relative to LT at 0h (Fig. 4). Thermal history and time interacted and had a significant ($P<0.001$, two-way ANOVA) association with

dct downregulation 24h post stress in all experimental groups compared to LT at time 0 (Fig. 4). At the same time, *dct* transcript abundance was significantly ($P<0.05$) downregulated in LHT and HT groups relative to the LT group. No significant changes between groups were identified at any other time point for any of the gene transcripts analyzed. *Tyra* transcripts were below the detection threshold for reliable quantification by qPCR and were excluded from the analysis (data not shown).

Discussion

The results of the present study suggest that, in common with mammals, early life stress modulates the stress response in gilthead sea bream adults (Cottrell & Seckl 2009). Although all the sea bream from each thermal group had a typical response to acute stress with significant differences in plasma cortisol across time (Arends *et al.* 1999, Tort *et al.* 2011), no significant intergroup differences were detected. However, the concentration range of plasma cortisol (37.16 ± 7.64 ng/mL) in fish reared during egg and larval stages at a higher temperature than the standard (HT) was lower than 50–80 ng/mL, typical of stressed Sparidae fish, 1h after an acute stress challenge (Tort *et al.* 2011). These results are reminiscent of the lower cortisol response reported for European sea bass (*Dicentrarchus labrax*) juveniles that as larvae were reared at a higher than optimum water temperature (Varsamos *et al.* 2006). Similarly, when rainbow trout (*Oncorhynchus mykiss*) embryos were stressed, the magnitude of the cortisol response to stress in the subsequent juveniles was lowered

(Auperin & Geslin 2008). Although no significant intergroup differences were observed in plasma cortisol, a temporal shift with delayed resolution in cortisol response was seen in LHT fish at 24 h post stress, which may be explained by a significant upregulation of *gr* in head kidney (Vazzana *et al.* 2010, Teles *et al.* 2013).

Although no significant changes in cortisol response was detected between fish with different thermal histories, a change in the responsiveness of the HPI axis probably resulted from changes at several levels in this axis. Significant differences existed in *pomca1* and *pomca2* pituitary expressions between fish reared at standard temperatures and the fish exposed to acute stress, and this response differed with thermal history. More consistent differences were observed in relation to *pomca2* that is synthesized by corticotrophs and processed and released as ACTH that acts on the interrenal and regulates glucocorticoid production (Cardoso *et al.* 2011). A significantly higher *pomca2* response was observed in the HT and HLT groups, which had a lower median cortisol response, supporting the notion that higher than standard temperatures during embryogenesis (common to both treatment groups) modulate the activity of the HPI axis by increasing the pituitary *pomc* expression (Bizuayehu *et al.* 2015). Increased relative abundance of *Pomc* transcripts have been reported in adult mice exposed to an early life stress (Wu *et al.* 2014).

The responsiveness of the HPI axis was also modified at the level of the head kidney, and basal *star* transcripts that encode the rate-limiting enzyme in the synthesis pathway of cortisol in the head kidney were significantly higher (Stocco 2000) in the HT group, suggesting modified activity of the cortisol producing pathway. An upregulation of *star* expression linked with lower cortisol production was observed in chronically stressed fish and was related to post-translational changes in the pre-existing pool of *star* mRNA, which led to modified biological activity of the enzyme (Geslin & Auperin 2004, Castillo *et al.* 2008). It remains to be established if a similar process occurred in the present study.

All experimental groups of sea bream had a classical stress response with strong downregulation of pituitary *gr* transcripts 4 h after acute stress, which suggests that the central activation of negative feedback on the stress axis was maintained irrespective of thermal history (Alderman *et al.* 2012). These results suggest that thermal imprinting during development in sea bream increased the responsiveness of the pituitary gland to stress but in a *gr*-independent manner. This contrasts with the

situation in mammals, in which exposure to stress during early development leads to reduced expression of *GR* in the hippocampus and reduced negative feedback on the HPA axis relative to unstressed animals (Cottrell & Seckl 2009). Future studies will be required to establish if increased *pomc* expression in sea bream is a consequence of reduced expression of hypothalamic *gr* or a result of epigenetic mechanisms at the level of the pituitary gland (Wu *et al.* 2014).

The mechanism by which early thermal regimes modified stress was not established, but it may be most likely linked to epigenetic modulation of the genome by temperature (Bizuayehu *et al.* 2015). It has been shown that epigenetic mechanisms cause a reduction in hypothalamic *GR* expression (Cottrell & Seckl 2009) and the increase in the expression of *POMC* in the pituitary gland of amniotes (Wu *et al.* 2014) and that these changes in gene expression are associated with hypermethylation or hypomethylation of DNA, respectively (Jaenisch & Bird 2003).

Thermal imprinting during early ontogeny modified the subsequent growth trajectory of the sea bream and changed the whole animal physiology even in the absence of a stress challenge, as plasma parameters related to hydromineral balance and osmoregulation were modified. It was notable that a drop in temperature at hatch (HLT group) had the most profound impact on the adult physiology, and HLT fish displayed significantly higher levels of glucose, sodium and potassium in relation to LT fish (the reference). Although in the present study the mechanism by which temperature after hatching modified the plasma glucose and ions in adult fish was not established, it may be linked to a time-specific window for temperature-induced epigenetic modulation (Bizuayehu *et al.* 2015).

The overall organization of the gilthead sea bream interrenal tissue was not modified by thermal history, and the MMCs had a typical morphology (large encapsulated, small clusters or free pigmented cells; Fig. 3; Meseguer *et al.* 1994). Different thermal histories were associated with a significantly modified immune response to stress in adults, and the number of small MMCs was reduced 24 h post stress in the LHT and HT groups relative to LT group. The reduction in the number of MMCs in the teleost fish head kidney has been proposed as a sign of immunosuppression and decreased phagocytic activity (Gregori *et al.* 2014). However, the expression of the well-established macrophage marker in gilthead sea bream *csf1r* (Roca *et al.* 2006) in the present study was highly variable

and did not corroborate the histology that indicated less MMCs, and presumably less macrophages, were present 24 h post stress in the LHT and HT fish.

The melanosynthetic activity of the macrophages in MMCs has been demonstrated in Atlantic salmon (*Salmo salar*, Fagerland *et al.* 2013) and DCT, which is involved in conversion of L-tyrosine into melanin, is expressed exclusively in cells capable of pigment production (Steel *et al.* 1992). In addition, DCT has antioxidant and phagocytic functions in the innate immunity of ectothermic vertebrates (Kovacs *et al.* 2012). In the present study, a highly significant downregulation in *dct* expression occurred 24 h after the acute stress challenge in all fish. However, LHT and HT had a significantly greater downregulation compared to LT suggesting stronger suppression of innate immunity after a stress challenge in these fish (Kovacs *et al.* 2012). In a short-term study in sea bass exposed to thermal imprinting, juveniles had depressed serum IgM and higher susceptibility to a viral challenge (Varsamos *et al.* 2006). Our results demonstrate the effect of thermal imprinting not only on the HPA axis but also on innate immune elements of adults exposed to stress. It will be important to establish if these modifications are a consequence of the bioenergetics/metabolic cost of a modified stress response or the result of temperature-induced epigenetic modifications of immune genes (Bizuayehu *et al.* 2015).

The present study revealed that the temperature regime during early development of sea bream caused thermal imprinting with long-term consequences for physiological and homeostatic responses to a stressor in adults. Clearly, a change in temperature between egg and larval stages (HLT and LHT) and an increase in temperature (HT) above the optimum (16–19°C) modified the whole adult physiology (HLT) and attenuated the plasma cortisol response to an acute stress in an over-stimulated HPI axis (HT). Additionally, rearing larvae at higher temperatures (HT and LHT) resulted in a suppression of elements of the innate immunity in adults, but the reason for this change remains to be established; it may arise from changed bioenergetics/metabolic costs of a modified stress response or from epigenetic modifications of immune genes. The present results provide valuable information for future research about the epigenetic mechanisms underlying the plasticity of the stress axis and immune function.

Supplementary data

This is linked to the online version of the paper at <http://dx.doi.org/10.1530/JOE-16-0610>.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

Funding

This study was funded by the project Lifecycle EU-FP7 222719 and national funds through FCT—Foundation for Science and Technology through project CCMAR/Multi/04326/2013. R A C is funded by FCT SFRH/BD/81625/2011 grant and J C R C is supported by an auxiliary research contract under the project UID/Multi/04326/2013.

Acknowledgments

Authors are extremely grateful to G Macià, M Matas, S Molas and M Monllaó (IRTA) for their technical assistance in live prey production and gilthead sea bream larval and juvenile rearing.

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Received in final form 22 March 2017

Accepted 18 April 2017

Accepted Preprint published online 18 April 2017