



Different Fish Meal and Fish Oil Dietary Levels in European Sea Bass: Welfare Implications After Acute Confinement Stress

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To provide practical feeding management guidelines preceding a stressful episode during farming practices, European sea bass juveniles (initial weight: 72.3 g) were fed for 60-days different fish meal (FM) and fish oil (FO) dietary levels [high (30% FM, 15% FO, FM30/FO15), intermediate (20% FM, 7% FO, FM20/FO7), and low (10% FM, 3% FO, FM10/FO3)] in triplicate conditions. Fish were then fasted for 36 h and exposed to a 2-h acute crowding (80 kg m⁻³ biomass). Plasma biochemistry, skin mucus parameters and gene expression of stress and immune-related genes were performed before, at 2 and 24 h after crowding. At the end of the trial, the FM10/FO3 group showed lower final body weight, weight gain, and specific growth rate compared to the other treatments. Most of the plasma parameters were mainly affected by crowding condition rather than diet; however, after stress, lactate was higher in the FM30/FO15 group compared to the other treatments. Similarly, protease, antiprotease, peroxidase and lysozyme in skin mucus were mostly affected by crowding conditions, while fish fed FM10/FO3 displayed higher skin mucosal IgM and bactericidal activity against *Vibrio anguillarum* and *V. harveyi*. Most of the stress-related genes considered (*hsp70* and *gr-1* in the brain; *hsp70*, *gr-1* and *gr-2* in the head kidney), showed an overall expression pattern that increased over time after stress, in addition, *hsp70* in the head kidney was also up-regulated in fish fed FM30/FO15 after stress. Higher plasmatic lactate together with the up-regulation of some stress-related transcripts suggest a higher reactivity to acute crowding of the stress-response mechanism in fish fed high FM and FO dietary levels. Otherwise, the higher skin mucosal IgM and bactericidal activity observed in fish fed FM10/FO3 dietary levels seems to indicate that acute crowding was able to activate a higher pro-inflammatory response in this treatment. Overall, the results of the present study seem to indicate that 10% FM and 3% FO dietary levels might affect stress and immune responses.

Keywords: aquaculture, feeding strategy, dietary fishmeal and fishoil, crowding stress, growth, skin mucus, plasma biochemistry, gene expression

INTRODUCTION

Nowadays, fish farmers are urged to apply feeding management practices that implement feed formulations which address both optimal growth and fish welfare issues, especially for coping with exposures to various types of stress (Boyd et al., 2020; Reverter et al., 2021). Certainly, under current aquaculture intensive rearing conditions, fish are continuously exposed to many stressors both biotic and abiotic; among them, husbandry-related stressors such as handling, crowding, transportation, or confinement are the most important (Tort and Teles, 2012). When experienced, these stressors are well-known to cause a wide range of physiological alterations, thus, worsening overall fish performance (Barton et al., 2005; Bagni et al., 2007; Castro et al., 2018; Balasch and Tort, 2019). In recent years, for several fish species including Mediterranean carnivorous ones, significant amounts of dietary fishmeal (FM) and fish oil (FO) have been successfully replaced with alternative ingredients mainly derived from commodity agricultural crops (Kaushik et al., 2004; Sitjà-Bobadilla et al., 2005; Médale and Kaushik, 2009; Hardy, 2010; Sales and Glencross, 2011; Nasopoulou and Zabetakis, 2012; Bonvini et al., 2018c; Sutuli et al., 2018; Parma et al., 2019; Kwasek et al., 2020). Among Mediterranean fish species, European sea bass (*Dicentrarchus labrax*) feed formulation has been evolving constantly, successfully reaching very low contents of FM and FO while providing for optimal rearing conditions; thus, demonstrating their potential for commercial use (Torrecillas et al., 2017a; Bonvini et al., 2018a; Parma et al., 2019). However, at the commercial farm scale this species is highly susceptible to stressful rearing-related problems such as netting, handling, sorting, transportation, and crowding giving rise to a wide range of negative physiological responses affecting their rearing performance and welfare (Gornati et al., 2004; Poltronieri et al., 2007; Fanouraki et al., 2011; Maricchiolo et al., 2011; Ellis et al., 2012; Tsalafouta et al., 2015; Samaras et al., 2017; Serradell et al., 2020). To counteract the negative impact of these stressors in fish, several feeding strategies have been developed to boost animal welfare, most of them including the utilization of functional feed additives (Volpatti et al., 2013; Gisbert et al., 2018; Serradell et al., 2020). However, only limited studies have investigated the interaction between very low dietary levels of FM and FO and animal welfare under suboptimal rearing conditions (Gisbert et al., 2018; Machado et al., 2019; Serradell et al., 2020). The effects of chronic stress on the immune system of fish are known to be clearly immunosuppressive; however, acute stress-induced immunomodulation is still poorly understood (Guo and Dixon, 2021). For instance, a recent study in European sea bass juveniles exposed to acute stress (netting and 1 min air exposure) and fed high levels of vegetable oils (VO) and carbohydrates resulted in depression of the immune response in both undisturbed and stressed fish (Machado et al., 2019). In order to find practical feeding management guidelines for fish farmers, the aim of this study was to evaluate the effect of different FM and FO dietary levels on growth performance, feed efficiency and, stress and immune-related biomarkers in European sea bass exposed to acute crowding. Fish responses to acute confinement were evaluated by analysis of different humoral immune biomarkers

from skin mucus, plasma biochemistry, and gene expression of stress and immune-related genes in order to provide insight into the overall effects of experimental diets on fish welfare.

MATERIALS AND METHODS

Experimental Diets

Three isonitrogenous (46% protein) and isolipidic (18% lipid) extruded diets (pellet size = 4.0 mm) were formulated to contain high, medium and low dietary levels of FM and FO (30% FM, 15% FO, FM30/FO15; 20% FM and 7% FO, FM20/FO7 and 10% FM and 3% FO, FM10/FO3, respectively). Diets were formulated with both, FM and a mixture of vegetable ingredients currently used for European sea bass aquafeeds (Parma et al., 2019), and were produced by SPAROS Lda (Portugal). Ingredients, proximate composition and fatty acid profile of the experimental diets are presented in **Tables 1, 2**.

TABLE 1 | Ingredients and proximate composition of the experimental diets.

Ingredients, % of the diet	FM30/FO15	FM20/FO7	FM10/FO3
Fishmeal LT70	30.0	20.0	10.0
Soy protein concentrate	10.0	12.0	20.0
Wheat gluten	5.0	10.0	11.0
Corn gluten	10.0	11.7	13.5
Soybean meal 48	9.0	9.0	8.1
Rapeseed meal	5.0	5.0	4.5
Sunflower meal	5.0	5.0	4.5
Wheat meal	9.5	9.0	7.9
Fish oil	15.0	7.0	3.0
Rapeseed oil		8.5	13.0
*Vitamin and mineral premix	1.0	1.0	1.0
Lutavit C35	0.1	0.1	0.1
Betaine HCl	0.1	0.1	0.1
Antioxidant	0.2	0.2	0.2
Sodium propionate	0.1	0.1	0.1
MCP	–	0.8	1.8
L-Lysine	–	0.4	0.6
L-Threonine	–	–	0.2
L-Tryptophan	–	–	0.1
D,L-Methionine	–	0.10	0.3
Proximate composition, % on a wet basis			
Moisture	7.92	6.85	6.26
Protein	44.84	45.75	45.25
Lipid	17.90	18.48	18.10
Ash	8.05	7.12	6.26
Gross energy (Cal g ⁻¹)	21.2	21.2	21.1

*Vitamins and mineral premix (IU or mg kg⁻¹ diet; in vivo NSA: Portugal); DL alpha tocopherol acetate, 200 mg; sodium menadione bisulfate, 10 mg; retinyl acetate, 16,650 IU; D,L-cholecalciferol, 2,000 IU; thiamine, 25 mg; riboflavin, 25 mg; pyridoxine, 25 mg; cyanocobalamin, 0.1 mg; niacin, 150 mg; folic acid, 15 mg; L-ascorbic acid monophosphate, 750 mg; inositol, 500 mg; biotin, 0.75 mg; calcium pantothenate, 100 mg; choline chloride, 1,000 mg; betaine, 500 mg; copper sulfate heptahydrate, 25 mg; ferric sulfate monohydrate, 100 mg; potassium iodide, 2 mg; manganese sulfate monohydrate, 100 mg; sodium selenite, 0.05 mg; zinc sulfate monohydrate, 200 mg; yttrium oxide, 100 mg.

TABLE 2 | Fatty acids composition of the experimental diets.

Fatty acids composition (g/100 g)	FM30/FO15	FM20/FO7	FM10/FO3
Lauric acid (12:0)		0.01 ± 0.00	0.02 ± 0.00
Myristic acid (14:0)	1.23 ± 0.15	0.40 ± 0.06	0.18 ± 0.04
Pentadecanoic acid (15:0)	0.08 ± 0.02	0.03 ± 0.01	0.02 ± 0.00
Palmitic acid (16:0)	3.15 ± 3.32	1.54 ± 0.18	1.10 ± 0.14
Isoheptadecanoic acid (17:0 iso)	0.07 ± 0.01	0.02 ± 0.00	0.01 ± 0.00
Hexadecenoic acid (16:1)	1.49 ± 0.45	0.50 ± 0.42	0.24 ± 0.07
14-methylhexadecanoic acid (17:0 anteiso)	0.05 ± 0.01	0.02 ± 0.00	0.01 ± 0.00
Margaric acid (17:0)	0.07 ± 0.01	0.03 ± 0.01	0.02 ± 0.00
Heptadecenoic acid (17:1)	0.05 ± 0.01	0.03 ± 0.00	0.02 ± 0.00
Stearic acid (18:0)	0.58 ± 0.08	0.35 ± 0.05	0.30 ± 0.05
Octadecenoic acid (18:1)	3.05 ± 0.27	4.80 ± 0.41	5.72 ± 0.49
Octadecadienoic acid (18:2)	1.34 ± 0.15	2.56 ± 0.27	3.12 ± 0.31
Arachidic acid (20:0)	0.03 ± 0.01	0.05 ± 0.01	0.05 ± 0.01
Octadecatrienoic acid (18:3)	0.43 ± 0.06	0.56 ± 0.07	0.67 ± 0.08
Eicosenoic acid (21:0)	0.35 ± 0.05	0.24 ± 0.04	0.18 ± 0.04
Stearidonic acid (18:4 n-3)	0.39 ± 0.06	0.13 ± 0.03	0.05 ± 0.00
Behenic acid (22:0)	0.01 ± 0.00	0.03 ± 0.00	0.04 ± 0.01
Docosanoic acid (22:1)	0.45 ± 0.06	0.24 ± 0.04	0.12 ± 0.02
Lignoceric acid (24:0)	0.01 ± 0.00	0.02 ± 0.00	0.01 ± 0.00
Polyunsaturated fatty acids (> C20)	2.19 ± 0.21	0.76 ± 0.08	0.30 ± 0.04
Polyunsaturated fatty acids	5.38 ± 0.37	2.53 ± 0.20	1.76 ± 0.16
Monounsaturated fatty acids	5.50 ± 0.53	5.87 ± 0.59	6.30 ± 0.65
Saturated fatty acids	7.84 ± 0.42	5.12 ± 0.32	4.59 ± 0.33
Fatty acids' ratios			
Polyunsaturated fatty acids/monounsaturated fatty acids	1.43 ± 0.16	0.87 ± 0.11	0.73 ± 0.09
Polyunsaturated fats/saturated fatty acids	1.46 ± 0.13	2.02 ± 0.21	2.61 ± 0.30

FM30/FO15 = 300 g kg⁻¹ fishmeal (FM), 150 g kg⁻¹ fish oil (FO); FM20/FO7 = 200 g kg⁻¹ fishmeal (FM), 70 g kg⁻¹ fish oil (FO); FM10/FO3 = 100 g kg⁻¹ fishmeal (FM), 30 g kg⁻¹ fish oil (FO).
SD, standard deviation.

Fish and Rearing Conditions

The experiment was carried out at the Laboratory of Aquaculture, Department of Veterinary Medical Sciences of the University of Bologna (Cesenatico, Italy). European sea bass (*Dicentrarchus labrax*) specimens were obtained from an Italian fish farm (Orbetello, GR) and adapted to the laboratory facilities for 1 week before the beginning of the trial. Afterward, 50 fish tank⁻¹ (initial weight: 72.2 ± 1.3 g, mean ± standard deviation, SD) were randomly distributed in nine 800 L volume tanks. Each diet was randomly assigned and administered to triplicate tanks over 60 days. Tanks were provided with natural seawater and connected to a closed recirculation system (overall water volume: 13 m³). The rearing system consisted of a mechanical sand filter (PTK 1200, Astralpool, Barcelona, Spain), ultraviolet lights (SH-88, BLUGEO S.r.l., Parma, Italy) and a biofilter (PTK 1200, Astralpool, Barcelona, Spain). The water exchange rate within each tank was 100% every hour, while the overall

water renewal amount in the system was 5% daily. During the trial, the temperature was kept at 23.0 ± 0.5°C and an artificial photoperiod of 12 h light and 12 h dark. The oxygen level was kept constant (8.0 ± 1.0 mg L⁻¹) through a liquid oxygen system regulated by a software program (B&G Sinergia snc, Chioggia, Italy). Ammonia (total ammonia nitrogen ≤ 0.1 mg L⁻¹) and nitrite (≤ 0.2 mg L⁻¹) were daily monitored spectrophotometrically (Spectroquant Nova 60, Merck, Lab business, Darmstadt, Germany), while salinity (30 g L⁻¹) was measured by a salt refractometer (106 ATC, Giorgio Bormac S.r.l., Carpi, Italy). Sodium bicarbonate was added on a daily basis to keep pH at 7.8–8.2. Fish were fed twice a day (8:30 and 16:30 h) for 6 days a week by automatic feeders to provide 10% overfeeding as reported by Bonvini et al. (2018a). Each meal lasted 1 h, after which the uneaten pellets of each tank were collected, dried overnight at 105°C, and weighed for further calculations of feed intake.

Sampling

At the beginning and at the end of the experiment, all the fish in each tank were anesthetized by MS222 at 100 mg L⁻¹ and individually weighed (BW, g). Proximate composition of carcasses was determined at the beginning of the trial on a pooled sample of 10 fish and on a pooled sample of five fish per tank at the end of the trial. Furthermore, wet weight of body, viscera, liver, perivisceral fat and total length were individually recorded for five fish per tank to determine viscerosomatic (VSI), hepatosomatic (HSI) and mesenteric fat (MFI) indices, and condition factor (CF).

At the end of the 60-day period of feeding, animals were fasted for 36 h. Subsequently, a crowding stress challenge (80 kg m⁻³ biomass) was conducted by lowering the water level in tanks for 2 h. At the end of this time frame, the water level was brought back to the initial level, allowing the animals to recover from the stressful event for 24 h. Before (time 0, T0), at the end of the 2-h exposure (time 1, T1) and 24 h after confinement at high density (time 2, T2), eight fish per tank were euthanized with an overdose of MS222 (300 mg L⁻¹) and sampled at each considered time point (T0, T1, and T2) in order to explore the capacity of the different diets to provide any advantage or disadvantage in front of the challenge test.

Skin mucus samples were collected from eight fish per tank according to Parma et al. (2020). Briefly, skin mucus was collected by gently scraping the dorsolateral surface of specimens using a cell scraper, taking care to avoid contamination with urinogenital and intestinal excretions. Collected mucus samples (3 mL each) were then stored at -80°C until analysis.

For the assessment of plasma biochemistry, blood from five fish per tank ($n = 15$ fish per diet treatment, same fish used for skin mucus samples) was collected from the caudal vein and placed in 4 mL heparin tubes (Vacutest Kima, Arzergrande, Italy). Samples were then centrifuged (3,000 × g, 10 min, 4°C) and plasma aliquots were stored at -80°C until analysis (Bonvini et al., 2018b).

Similarly, two fish per tank ($n = 6$ fish per diet treatment at T0, T1, and T2) were randomly sampled, euthanized with an overdose anesthetic as previously described, then head kidney

(both sides) and brain (whole) were dissected to gain 100 mg of tissue in order to evaluate the impact of dietary treatments on genetic markers of stress-response in fish that had been submitted to crowding stress. The brain and head kidney samples were kept at 4°C overnight into RNA Later (Sigma-Aldrich), and then stored at -80°C until further analysis.

All experimental procedures were evaluated and approved by the Ethical-Scientific Committee for Animal Experimentation of the University of Bologna (protocol ID 942/2019) in accordance with European directive 2010/63/UE on the protection of animals used for scientific purposes.

Growth Performance, Feed Efficiency, and Body Condition Indices

In order to evaluate the effect of experimental diets, the following formulae were employed to calculate growth performance, somatic and nutritional index values:

Weight gain (WG, g) = BWf - BWi (where BWf and BWi represent the final and the initial body weight).

Specific growth rate (SGR, % day⁻¹) = 100 * (ln BWf - ln BWi)/days

Feed intake (FI, g kg ΔBW⁻¹ day⁻¹) = ((1000*total feed ingestion)/(ΔBW))/days.

Feed conversion ratio (FCR) = FI/WG.

Condition factor (CF) = 100 * (BWf/total body length³).

Viscerosomatic index (VSI, %) = 100 * (viscera weight/BWf).

Hepatosomatic index (HSI, %) = 100 * (liver weight/BWf).

Mesenteric fat index (MFI, %) = 100 * (mesenteric fat weight/BWf).

Protein efficiency rate (PER) = (BWf - BWi)/total protein intake.

Gross protein efficiency (GPE, %) = 100 * [(% final body protein content * BWf) - (% initial body protein * BWi)]/total protein intake fish.

Lipid efficiency rate (LER) = (BWf - BWi)/total lipid intake.

Gross lipid efficiency (GLE, %) = 100 * [(% final body lipid content * BWi) - (% initial body lipid * BWi)]/total lipid intake fish.

Proximate Composition Analyses

Diets and whole bodies of fish were analyzed for proximate composition. Moisture content was obtained by weight loss after drying samples in a stove at 105°C overnight. Crude protein was determined as total nitrogen (N*6.25) using the Kjeldahl's method according to the AOAC International (AOAC, 2010). Total lipids were determined according to Bligh and Dyer's (1959) extraction method. Ash content was estimated by incineration in a muffle oven at 450°C overnight. Gross energy was determined by a calorimetric bomb (Adiabatic Calorimetric Bomb Parr 1261; PARR Instrument, IL, United States). Fatty acid analyses composition of diets was performed according to ISO16958:2015.

Stress and Metabolic Biomarkers in Plasma

Plasma samples (500 μL) were analyzed by automated analyzer (AU 480; 220 Olympus/Beckman Coulter, Brea,

CA, United States) using specific methods (Olympus system 221 reagent, OSR) detecting the levels of glucose (GLU), triglycerides (TRIG), lactate, total protein, aspartate aminotransferase (AST), alanine transaminase (ALT), alkaline phosphatase (ALP), creatine kinase (CK), lactate dehydrogenase (LDH), calcium (Ca²⁺), inorganic phosphorus (P), potassium (K⁺) sodium (Na⁺), magnesium (Mg) according to Pelusio et al. (2021). In addition, plasma cortisol (CORT) concentration was determined using a chemiluminescence immunoassay (Immulite cortisol, Diagnostic Product Corporation, Los Angeles, CA, United States) using an automated analyzer (Immulite XP2000, Siemens) (Pelusio et al., 2021).

Humoral Immunity Markers in Skin Mucus

The peroxidase activity was measured by oxidation of 3,3',5,5'-Tetramethylbenzidine (TMB) according to Quade and Roth (1997). Aliquots of 10 μL of mucus solution were diluted with Hanks' buffer (HBSS) to a final volume of 50 μL in a flat-bottomed 96-well plate. As substrate, H₂O₂ was added, and the color change reaction was halted by adding 50 μL 2 M H₂SO₄. The optical density (OD) was read at λ = 450 nm in a plate reader (FLUOstar Omega, BMG Labtech, Germany). Samples without mucus solution were used as blanks. One unit was defined as the amount producing an absorbance change of 1, and the activity was expressed as U mg protein⁻¹.

Protease activity was quantified using the azocasein hydrolysis assay according to the method described by Ross et al. (2000). Briefly, 100 μL of skin mucus were incubated with 100 μL of ammonium bicarbonate buffer and 125 μL of 0.7% azocasein (Sigma, United States) in sterile Eppendorf vials. The reaction was then halted by adding 250 μL of 4.6% trichloroacetic acid (TCA, Sigma, United States) and the OD was read at λ = 450 nm using a plate reader. Skin mucus was replaced by trypsin (5 mg mL⁻¹, Sigma) for the positive controls or by ammonium bicarbonate buffer for the negative one. The activity for each sample was expressed as % protease activity in relation to the controls.

Antiprotease activity was determined by the capacity of skin mucus to inhibit trypsin activity (Hanif et al., 2004). Briefly, 100 μL of skin mucus were incubated (10 min, RT) with 10 μL of trypsin solution (5 mg mL⁻¹, Sigma, United States). Subsequently, 100 μL of ammonium bicarbonate buffer and 125 μL of 0.7% azocasein (Sigma, United States) were added and mixtures incubated for 2 h at RT. Afterward, 250 μL of 4.6% trichloroacetic acid (TCA; Sigma, United States) were added and samples incubated for an additional 30 min. Later 100 μL of 0.5 N NaOH were added and the OD was read at λ = 450 nm using a plate reader. Buffer without sample or trypsin was used as positive controls and a combination of buffer and trypsin solution was used for the negative controls. The activity for each sample was expressed as % trypsin inhibition in relation to the controls.

Lysozyme activity was measured according to the turbidimetric method described by Parry et al. (1965). Volumes of 10 μL of skin mucus diluted 1:2 with 10 mM PBS, pH 6.2, were placed in a flat-bottomed 96-well plate. To each well, 200 μL

of freeze-dried *Micrococcus lysodeikticus* in the above buffer (0.3 mg mL^{-1} , Sigma) were added as lysozyme substrate. The reduction in absorbance at $\lambda = 450 \text{ nm}$ was measured. One unit of lysozyme activity was defined as a reduction in absorbance of $0.001 \text{ units min}^{-1}$. The units of lysozyme present in skin mucus were obtained from a standard curve made with hen's egg-white lysozyme (HEWL, Sigma), and the results were expressed as U mg protein^{-1} .

Alkaline phosphatase activity was measured by incubating $100 \mu\text{L}$ of skin mucus with an equal volume of 4 mM *p*-nitrophenyl liquid phosphate (Sigma, United Kingdom) in 100 mM ammonium bicarbonate buffer containing 1 mM MgCl_2 ($\text{pH } 7.8$, 30°C) (Ross et al., 2000). The OD was continuously measured at 1 min intervals over 3 h at 405 nm in a plate reader and one unit of activity was defined as the amount of enzyme required to release 1 mmol of *p*-nitrophenol product in 1 min . The activity was expressed as U mg protein^{-1} .

Esterase activity was measured by incubating aliquots of $100 \mu\text{L}$ of skin mucus with an equal volume of 0.4 mM *p*-nitrophenylmyristate substrate (Sigma, United Kingdom) in 100 mM ammonium bicarbonate buffer containing 0.5% Triton X-100 (Sigma, United Kingdom) ($\text{pH } 7.8$, 30°C) in a flat-bottomed 96-well plate, as described by Ross et al. (2000). The OD and activity were determined and expressed as previously described for the alkaline phosphatase.

Total skin mucus immunoglobulin M (IgM) levels were analyzed by ELISA method as described by Hanif et al. (2004). Samples of $100 \mu\text{L}$ per well of skin mucus diluted 1:5 with 50 mM carbonate-bicarbonate buffer (35 mM NaHCO_3 and 15 mM Na_2CO_3 , $\text{pH } 9.6$), the proteins were adsorbed during an overnight incubation at 4°C and blocked for 2 h at room temperature with $200 \mu\text{L}$ blocking buffer containing 3% BSA in PBS-T, followed by three new rinses with PBS-T. Plates were then incubated (1 h with $100 \mu\text{L}$ per well) with mouse anti-European sea bass IgM monoclonal antibody (Aquatic Diagnostics Ltd., 1/100 in blocking buffer) and with $100 \mu\text{L}$ per well of secondary antibody anti-mouse IgG-HRP (1/1000 in blocking buffer, Sigma-Aldrich). After exhaustive rinsing with PBS-T the plates were developed using $100 \mu\text{L}$ of a 0.42 mM solution of 3,3',5,5'-tetramethyl benzidine hydrochloride (TMB, Sigma-Aldrich). The reaction was allowed to proceed for 10 min and halted by the addition of $50 \mu\text{L}$ 2 M H_2SO_4 and the plates were then read at $\lambda = 450 \text{ nm}$. Negative controls consisted of samples without skin mucus or without primary antibody, whose OD values were subtracted from each sample value.

Two bacteria, pathogenic for marine fish species (*Vibrio harveyi* and *Vibrio anguillarum*), were used in the bactericidal assays. Bacteria were cultured for 48 h at 25°C in Tryptic Soy Agar (TSA, Difco Laboratories), and then single colonies were inoculated in Tryptic Soy Broth (TSB, Difco Laboratories), both supplemented with NaCl to a final concentration of 1% (w/v), and grown with continuous shaking (100 rpm) for 24 h . Bactericidal activity was determined according to the method of Stevens et al. (1991). For this, aliquots of $100 \mu\text{L}$ of bacteria were added to wells of a flat-bottomed 96-well plate and incubated with the same volume of skin mucus. The absorbance of the dissolved formazan was measured at $\lambda = 570 \text{ nm}$. Bactericidal activity was expressed

as a percentage of non-viable bacteria (the difference between the absorbance of surviving bacteria in test samples compared with the absorbance of bacteria from positive controls).

Soluble total protein concentration in mucus samples was determined by the Bradford dye binding method, using bovine serum albumin (BSA, Sigma-Aldrich) as a standard (Bradford, 1976).

Gene Expression in Brain and Head-Kidney

RNA from tissues (50 mg per sample) was extracted using the RNeasy Mini Kit (Qiagen, Spain) following the instructions of the manufacturer. RNA concentration and quality were determined by spectrophotometry ($\lambda = 260/280$) (Nanodrop-2000® spectrophotometer; Thermo Scientific, United States) and denaturing electrophoresis in TAE agarose gel (1.5%) (Masek et al., 2005). For each tissue sample, ca. $1 \mu\text{g}$ of RNA was reverse transcribed into cDNA using Super-Script™ First-Strand Synthesis System for RT-PCR (Invitrogen™, Life Technologies, Warrington, United Kingdom) primed with oligo (dT)_{12–18} and random hexamers. Primers for quantitative PCR analyses were obtained for most genes by searching the literature for previously reported gene expression studies in this species (Table 3). When genes of interest were not previously published, an alignment of marine teleost homologs for the gene of interest obtained from GenBank, was made using BioEdit Sequence Alignment Editor ver 7.0.5.2 (Hall, 1999). Consensus primers designed from conserved regions identified in these alignments were used for *D. labrax*-specific qPCR assays (Table 2). Quantitative real-time polymerase chain reaction (qPCR) was performed using a C1000 Touch Thermal Cycler (BioRad, Spain). Beta actin (*β-actin*) was used as the reference gene. All cDNA samples were diluted 1:20 in molecular biology grade water. An aliquot of $5 \mu\text{L}$ from each diluted sample was pooled to provide a “calibrator sample” that was included as a calibrator reaction (*β-actin*) sample on each plate to correct for variation among experimental plates. A standard curve was prepared by amplification of a dilution series of this pooled cDNA for calculation of the efficiency (%E) for each assay to ensure that efficiency was close to 100%.

Quantitative PCR analyses for each gene were performed in triplicate in a total volume of $20 \mu\text{L}$ containing $2 \mu\text{L}$ cDNA, $20 \mu\text{mol}$ of each primer, $10 \mu\text{L}$ SYBR™ Green (Life Technologies, Warrington, United Kingdom) and $7 \mu\text{L}$ of sterile water. A negative control was included (NTC = no template control) for each set of reactions on each 96-well plate. All reactions were performed using the following conditions: 1 cycle at 95°C for 10 min ; 40 cycles at 95°C for 15 s , $T_a^\circ\text{C}$ (primer-specific annealing temperature) for 30 s . A final melt curve stage completed the program, enabling confirmation of the amplification of a single product in each reaction. The final relative mRNA levels were then calculated following Pfaffl's method (Pfaffl, 2004).

Statistical Analysis

All data are presented as mean \pm SD. A tank was used as the experimental unit for analyzing growth performance, whereas

TABLE 3 | Primer sequences used for gene expression analyses.

Gene	Abbreviation	GenBankID	Primer sequence (5'-3')	Amplicon (bp)	References	Annealing T°
β -actin	β -act	AJ537421.1	Fw: 5'-ATGTGGATCAGCAAGCAGG-3' Rv: 5'-AGAAATGTGTGGTGTGGTCG-3'	144	El Aamri et al., 2015	5860
Glucocorticoid receptor-1	<i>gr-1</i>	AY549305	Fw: 5'-GAGATTTGGCAAGACCTTGACC-3' Rv: 5'-ACCACACCAGGCGTACTGA-3'	401	Reyes-López et al., 2018	6660
Glucocorticoid receptor-2	<i>gr-2</i>	AY619996	Fw: 5'-GACGCAGACCTCCACTACATTC-3' Rv: 5'-GCCGTTCACTCTCAACCAC-3'	403	Reyes-López et al., 2018	6864
Melanocortin receptor	<i>mr</i>	JF824641.1	Fw: 5'-CCTGTCTCCTCTATGAATGG-3' Rv: 5'-AATCTGGTAATGGAATGAATGTC-3'	173	Samaras et al., 2018	6062
Caspase 3	<i>csp3</i>	DQ345774.1	Fw: 5'-ACGAAGCAGGTCAATCATCC-3' Rv: 5'-GCAGTTAAGGGTATCCAGAGC-3'	98	El Aamri et al., 2015	6066
Heat shock protein 70	<i>hsp70</i>	AY423555	Fw: 5'-CAACCTCCTGGGAAGTTTG-3' Rv: 5'-AGAATCTTCTGCTTGTCGTCG-3'	353	Serradell et al., 2020	56.356.3
Transforming growth factor	<i>tgf</i>	AM421619.1	Fw: 5'-ACGGGCTACCATGCCAATC-3' Rv: 5'-AGAGGCTCCTGGGTTGTGATG-3'	110	Designed	6670
Tumor necrosis factor α	<i>tnf-α</i>	DQ200910	Fw: 5'-ACAGCGGATATGGACGGTG-3' Rv: 5'-GCCAAGCAAACAGCAGGAC-3'	76	El Aamri et al., 2015	6060
Interleukin 10	<i>il10</i>	AM268529	Fw: 5'-ACCCCGTTTCGCTTGCCA-3' Rv: 5'-CATCTGGTGACATCACTC-3'	163	El Aamri et al., 2015	5654
Dicentrarin	<i>dcn</i>	AY303949.1	Fw: 5'-GTTTACGTCGGCAAGTCCATCC-3' Rv: 5'-GATGAACCGCCATAGATCACTC-3'	175	Designed	7068
Cyclooxygenase 2	<i>cox2</i>	AJ630649	Fw: 5'-AGCACTTACCCACCAGTTC-3' Rv: 5'-AAGCTTGCCATCCTTGAAGA-3'	148	Cordero et al., 2016	6258
Lysozyme	<i>lys</i>	KJ433681.1	Fw: 5'-TGATGCAGGTTGTTGATGTTAATC-3' Rv: 5'-TCCATCCCCATATTGTAGGC-3'	194	Reyes-López et al., 2018	6664

a pool of five sampled fish was considered the experimental unit for analyzing carcass composition and somatic indices. Individual fish were used for analyzing plasma biochemistry, skin mucus parameters and relative expression of stress and immune-related transcripts. At the end of the feeding trial with diets differing in the FM and FO levels (60 days), data of growth performance, nutritional indices and somatic indices were analyzed by a one-way analysis of variance (ANOVA) and in case of significance ($P \leq 0.05$) Tukey's *post hoc* test was performed. In order to provide the effect of diets on the basal status of skin mucus parameters, plasma and gene expression before the stress episode; results were analyzed by a two-tier statistical approach analyzing first the data at T0 by one way ANOVA and followed by a two-way ANOVA considering sampling points and diets as factors. When statistically significant differences were found among groups ($P \leq 0.05$), Tukey's *post hoc* test was performed to determine differences among experimental groups. Prior to ANOVA analyses, all data were checked for normality and homogeneity of variance by means of the Shapiro Wilk and Levene tests, respectively. All the statistical analyses were conducted using GraphPad Prism and Statistical Package for Social Science (SPSS) for Windows version 20.0.

RESULTS

Key Performance Indicators and Body Proximate Composition

Results on growth performance and feeding efficiency parameters are summarized in **Table 4**. Significant differences were found in terms of BWf, WG, and SGR, while the lowest values were found in animals fed the FM10/FO7 diet. In particular,

European sea bass fed FM30/FO15 and FM20/FO7 diets were 11.2 and 9.2% heavier than those from the FM10/FO3 dietary treatment. However, FCR, FI, and survival values displayed no significant differences among experimental groups.

Results on proximate body composition and, nutritional and somatic indices are reported in **Table 5**. No significant differences were found on any of the analyzed parameters.

Plasma Biochemistry

Plasma biochemistry results are summarized in **Supplementary Table 1** and **Table 6**. At the end of the growth trial (T0) dietary treatments did not affect all the parameters analyzed except cortisol which was higher in FM30/FO15 than the other treatments (One-way ANOVA – **Supplementary Table 1**).

TABLE 4 | Growth performance of European seabass fed experimental diets over 60 days before crowding stress exposure.

	FM30/FO15	FM20/FO7	FM10/FO3	P-value
BWi	72.40 \pm 1.66	71.75 \pm 1.02	72.42 \pm 1.73	0.832
BWf	210.72 \pm 3.00 ^b	203.65 \pm 6.33 ^b	187.00 \pm 5.00 ^a	0.003
WG	138.32 \pm 4.20 ^b	131.90 \pm 5.33 ^b	114.58 \pm 4.32 ^a	0.002
SGR	1.78 \pm 0.07 ^b	1.74 \pm 0.04 ^b	1.58 \pm 0.05 ^a	0.003
FCR	1.25 \pm 0.01	1.29 \pm 0.06	1.37 \pm 0.08	0.098
FI	20.41 \pm 0.53	20.56 \pm 0.67	20.16 \pm 0.84	0.782
Survival	100.00 \pm 0.00	99.30 \pm 1.20	99.30 \pm 1.20	0.630

Data are given as the tanks mean ($n = 3$) \pm SD. In each line, different superscript letters indicate significant differences among treatments ($P \leq 0.05$). FM30/FO15 = 300 g kg⁻¹ fishmeal (FM), 150 g kg⁻¹ fish oil (FO); FM20/FO7 = 200 g kg⁻¹ fishmeal (FM), 70 g kg⁻¹ fish oil (FO); FM10/FO3 = 100 g kg⁻¹ fishmeal (FM), 30 g kg⁻¹ fish oil (FO).

TABLE 5 | Body composition and, nutritional and somatic indices of European seabass fed experimental diets over 60 days before crowding stress exposure.

	FM30/FO15	FM20/FO7	FM10/FO3	P-value
Whole body composition, %				
Protein	16.71 ± 0.68	17.02 ± 0.11	17.02 ± 0.61	0.722
Lipid	17.50 ± 0.33	17.31 ± 0.75	17.26 ± 1.42	0.949
Ash	5.85 ± 0.08	3.76 ± 0.15	3.75 ± 0.08	0.994
Moisture	61.66 ± 0.12	61.57 ± 0.47	61.60 ± 1.00	0.984
Nutritional indices				
PER	1.78 ± 0.02	1.70 ± 0.08	1.62 ± 0.09	0.080
GPE	29.96 ± 1.93	29.45 ± 1.64	28.11 ± 2.61	0.573
LER	4.46 ± 0.04	4.20 ± 0.20	4.04 ± 0.24	0.079
GLE	104.31 ± 3.75	98.13 ± 6.36	97.80 ± 9.44	0.475
Somatic indices				
HSI	1.38 ± 0.06	1.35 ± 0.05	1.40 ± 0.10	0.363
MFI	7.52 ± 0.54	7.78 ± 0.44	7.29 ± 0.72	0.942
VSI	12.00 ± 0.84	12.24 ± 0.30	11.63 ± 0.50	0.980
CF	1.28 ± 0.06	1.29 ± 0.06	1.24 ± 0.04	0.447

Data are given as the mean ($n = 15$ per diet for proximate composition and nutritional indices; $n = 12$ per diet for somatic indices) ± SD. FM30/FO15 = 300 g kg⁻¹ fishmeal (FM), 150 g kg⁻¹ fish oil (FO); FM20/FO7 = 200 g kg⁻¹ fishmeal (FM), 70 g kg⁻¹ fish oil (FO); FM10/FO3 = 100 g kg⁻¹ fishmeal (FM), 30 g kg⁻¹ fish oil (FO).

PER = Protein efficiency ratio = ((FBW-IBW)/protein intake); Gross protein efficiency (GPE, %) = 100*[(% final body protein*FBW) - (%initial body protein*IBW)]/total protein intake fish; Gross lipid efficiency (GLE, %) = 100*[(% final body lipid*FBW) - (%initial body lipid*IBW)]/total lipid intake fish; Lipid efficiency ratio (LER) = ((FBW-IBW)/lipid intake); Hepatosomatic index (HSI, %) = 100*(liver weight/FBW); Mesenteric Fat Index (MFI, %) = 100*(mesenteric fat weight/FBW); Viscerosomatic index (VSI, %) = 100*(viscera weight/FBW); Condition factor (CF) = 100*(FBW/length³).

SD, standard deviation.

Table 6 reports plasma results in relation to dietary treatments and time after stress (Two-way ANOVA). Glucose was affected by diet and time after stress where FM30/FO15 showed higher values than FM20/FO7 and values were higher at T1 compared to T0 and T2. Triglyceride were affected by time where T0 groups displayed higher values than those of T1 and T2. Similarly, plasma protein was affected by time with higher values found at T1 and T2 compared to T0. Concerning AST and ALP, no significant diet- nor time-dependant effects were found among experimental groups while a significant time effect was found in ALT, with higher values found in T0 than T2 groups. CK and LDH were affected by time with lower values found in T2 compared to T0. Ca²⁺ was affected by time where T1 and T2 showed higher values than T0. Both significant diet and time effects were found in P where FM30/FO15 and T0 displayed higher values than the other diets and time groups, respectively. K⁺ was affected by time with higher values at T0 compared to the other sampling times. Concerning Na⁺, significant diet, time and interaction effects were found and the combined effect of diet and time resulted in an increasing pattern from T0 to T1 only in FM10/FO3 diets. In addition, at T1, FM10/FO3 displayed higher values than the other dietary treatments. Plasma Mg²⁺ levels were influenced by time; specifically, values were lower at T2 in comparison to T0 and T1. Cortisol level was affected by time; specifically, values were higher at T1 in comparison to T0

and T2. A significant diet and interaction effect were observed in lactate, in particular the lactate levels recorded at T1 were higher in the FM30/FO15 group in comparison to those observed in FM20/FO7 and FM10/FO3 at the same time point. In addition, at T0 FM30/FO15 displayed higher lactate values at T1 than the other time points, while in FM20/FO7 values were higher at T0 than T1. In FM10/FO3 no significant difference within time points were observed.

Humoral Immune Biomarkers in Skin Mucus

Results concerning humoral immune biomarkers in skin mucus are presented in **Supplementary Figures 1A–I** and in **Figures 1A–I**. At the end of the growth trial (T0) dietary treatments did not affect all the parameters analyzed except esterase which was lower in FM20/FO7 compared to the other treatments (One-way ANOVA – **Supplementary Figure 1**). In addition, bactericidal activities against *V. anguillarum* and *V. harveyi* were higher in FM10/FO3. **Figure 1** reports humoral immune biomarkers results in relation to dietary treatments and time after stress (Two-way ANOVA). Protease activity was affected by time after stress displaying higher values in T0 than T1 and T2 (**Figure 1A**). Antiprotease activity was affected by time with significant diet × time effect which resulted to increasing values at T1 and T2 in FM10/FO3. In addition, at T0, FM20/FO7 was higher than FM10/FO3 (**Figure 1B**). Peroxidase activity was significantly affected by time and diet, showing higher values at T2 compared to T0 and T1 (**Figure 1C**). In addition, peroxidase activity was also lower in fish fed FM10/FO3 than FM30/FO15. Esterase activity was affected by time with higher values at T2 than T1 and T0 (**Figure 1D**). Alkaline phosphatase activity showed a diet, time and interaction effect with higher values at T2 in the FM30/FO15 and FM10/FO3 compared to the same dietary groups recorded at T0 and T1. In addition at T2, FM20/FO7 was lower than the other diets while FM10/FO3 was lower than FM30/FO15 (**Figure 1E**). Lysozyme was affected by time with general increasing values at T1 even if not specific differences were detected in the *post hoc* test (**Figure 1F**). IgM levels showed a diet, time and interaction effect; specifically, values increased in animals fed the FM10/FO3 diet and sampled at T1 and T2 compared to those sampled at T0 within the same treatment. In contrast, fish from the FM20/FO7 group were characterized by lower IgM values at T2 compared to T0, while those from the FM30/FO15 group showed lower values at T1 compared to T0 and T2 sampling points (**Figure 1G**). In addition, at T1 FM30/FO15 displayed lower values than FM20/FO7 and FM10/FO3, while at T2, FM10/FO3 showed higher values than the other diets and FM30/FO15 was higher than FM20/FO7.

Bactericidal activity against *V. anguillarum* was affected by time, diet and interaction; specifically, in fish fed the FM10/FO3 values were higher at T2 than T0 within the same dietary group, while in fish fed FM20/FO7 values were higher at T1 compared to T0. At each considered time point, FM10/FO3 showed higher values than the other dietary treatments.

Similarly, bactericidal activity against *V. harveyi* was also affected by time and diet with higher values at T1 and T2 than

TABLE 6 | Plasma biochemistry values for European seabass fed the experimental diets and exposed to crowding stress and recovery after small fasting period.

Analyte	Time	FM30/FO15	FM20/FO7	FM10/FO3	P-value		
					Diet	Time	Inter
Glucose (mg dL ⁻¹)	T0	188.25 ± 66.73	166.07 ± 40.07	170.14 ± 43.81	<0.001(B/A/AB)	<0.001(A/B/A)	0.382
	T1	293.30 ± 110.62	213.12 ± 48.48	252.24 ± 65.27			
	T2	223.15 ± 69.39	181.04 ± 44.81	188.14 ± 38.49			
TRIG (mg dL ⁻¹)	T0	730.39 ± 284.73	851.80 ± 347.61	773.87 ± 240.28	0.170	<0.001(B/A/A)	0.625
	T1	436.07 ± 92.14	471.81 ± 115.19	389.09 ± 43.56			
	T2	451.37 ± 123.85	442.58 ± 84.36	387.00 ± 69.65			
TP (mg dL ⁻¹)	T0	4.38 ± 0.45	4.31 ± 0.42	4.12 ± 0.38	0.051	<0.001(A/B/B)	0.910
	T1	4.97 ± 0.54	4.90 ± 0.44	4.85 ± 0.34			
	T2	4.78 ± 0.28	4.79 ± 0.55	4.52 ± 0.37			
AST (U L ⁻¹)	T0	273.61 ± 275.40	196.80 ± 157.33	153.07 ± 104.89	0.065	0.293	0.545
	T1	137.00 ± 79.34	132.80 ± 105.08	139.64 ± 73.54			
	T2	283.26 ± 443.76	142.58 ± 94.39	105.50 ± 82.14			
ALT (U L ⁻¹)	T0	8.61 ± 15.94	6.40 ± 7.84	4.07 ± 3.86	0.323	0.008(B/AB/A)	0.730
	T1	3.00 ± 2.25	4.25 ± 5.00	2.36 ± 1.29			
	T2	2.47 ± 2.25	2.70 ± 2.66	1.69 ± 1.01			
ALP (U L ⁻¹)	T0	79.00 ± 82.14	51.87 ± 17.24	52.20 ± 15.32	0.173	0.146	0.323
	T1	53.93 ± 10.65	52.88 ± 10.05	53.00 ± 5.04			
	T2	50.95 ± 13.20	50.32 ± 13.46	44.88 ± 7.76			
CK (U L ⁻¹)	T0	3530.93 ± 2783.44	4206.50 ± 4362.01	3190.43 ± 3469.04	0.274	0.003(B/AB/A)	0.460
	T1	3940.69 ± 3519.65	2364.88 ± 2969.64	2049.73 ± 830.50			
	T2	1544.67 ± 1106.93	2282.00 ± 1608.62	1169.62 ± 748.82			
LDH (U L ⁻¹)	T0	232.78 ± 252.62	167.80 ± 113.83	174.33 ± 169.45	0.481	0.024(B/AB/A)	0.770
	T1	154.93 ± 107.26	126.27 ± 108.60	143.45 ± 62.59			
	T2	120.95 ± 84.84	134.74 ± 109.52	95.75 ± 88.39			
Ca ²⁺ (mg dL ⁻¹)	T0	12.96 ± 3.17	13.69 ± 0.89	13.35 ± 0.73	0.912	<0.001(A/B/B)	0.217
	T1	15.18 ± 1.65	14.47 ± 0.94	15.09 ± 1.30			
	T2	14.62 ± 1.04	14.28 ± 0.89	13.99 ± 0.71			
P (mg dL ⁻¹)	T0	12.47 ± 4.44	11.75 ± 1.25	10.86 ± 1.29	0.004(B/A/A)	<0.001(B/A/A)	0.317
	T1	11.54 ± 2.03	9.53 ± 0.91	10.28 ± 1.05			
	T2	9.90 ± 1.14	9.39 ± 1.12	9.26 ± 0.97			
K ⁺ (mEq L ⁻¹)	T0	6.17 ± 2.67	6.01 ± 2.48	6.29 ± 1.83	0.512	<0.001(B/A/A)	0.869
	T1	1.73 ± 0.64	2.32 ± 1.33	2.20 ± 1.16			
	T2	2.59 ± 1.35	2.66 ± 1.46	3.18 ± 1.56			
Na ⁺ (mEq L ⁻¹)	T0	188.40 ± 4.03 ^a	188.13 ± 4.00 ^a	187.71 ± 3.60 ^a	0.041	<0.001(A/A/B)	0.009
	T1	187.56 ± 2.34 ^{a#}	185.16 ± 12.5 ^{a#}	194.56 ± 6.63 ^{b†}			
	T2	193.84 ± 3.73 ^b	195.65 ± 3.27 ^b	195.63 ± 9.93 ^b			
Mg (mg dL ⁻¹)	T0	4.50 ± 0.82	4.23 ± 0.73	4.48 ± 1.17	0.103	<0.001(B/B/A)	0.171
	T1	4.63 ± 0.94	4.44 ± 0.70	5.10 ± 1.00			
	T2	3.94 ± 0.61	3.52 ± 0.36	3.46 ± 0.55			
Cortisol (μg dL ⁻¹)	T0	35.61 ± 12.33	26.16 ± 5.86	25.56 ± 12.42	0.055	<0.001 (A/B/A)	0.123
	T1	41.34 ± 12.30	43.19 ± 5.96	40.86 ± 8.43			
	T2	34.98 ± 9.19	34.26 ± 10.26	31.59 ± 8.42			
Lactate (mmol L ⁻¹)	T0	53.37 ± 18.29 ^a	49.13 ± 17.46 ^b	48.98 ± 11.56	<0.001(B/A/A)	0.420	0.001
	T1	68.90 ± 39.29 ^{b†}	33.60 ± 7.07 ^{a#}	45.87 ± 13.55 [#]			
	T2	48.79 ± 8.28 ^a	43.19 ± 7.21 ^{ab}	46.77 ± 9.84			

Data are given as the mean ($n = 15$ diet⁻¹) each time (T) sampling ± SD. The uppercase letters in the parenthesis after P-values of two-way ANOVA represent significant differences in two-way ANOVA. The three sets of letters in a parenthesis correspond to the three time (T0/T1/T2) or diet (FM30/FO15/FM20/FO7/FM10/FO3) groups, respectively, and data value corresponding to "A" are lower than those corresponding to "B." Different symbols stand for significant differences among dietary treatments for the same time while lowercase letters stand for significant differences between times for the same diet. FM30/FO15 = 300 g kg⁻¹ fishmeal (FM), 150 g kg⁻¹ fish oil (FO); FM20/FO7 = 200 g kg⁻¹ fishmeal (FM), 70 g kg⁻¹ fish oil (FO); FM10/FO3 (100 g kg⁻¹ fishmeal (FM), 30 g kg⁻¹ fish oil (FO). Pre-stress time point after 36 h fasting (T0); post-stress time point after 2 h confinement stress exposure (T1); recovery time point 24 h after the end of confinement stress exposure (T2).

TP, total protein; TRIG, triglycerides; AST, aspartate aminotransferase; ALT, alanine transaminase; ALP, alkaline phosphatase; CK creatine kinase (U L⁻¹); LDH, lactate dehydrogenase; Ca²⁺ calcium; P, inorganic phosphorus; K⁺ potassium; Na⁺, sodium; Mg, magnesium; SD, standard deviation.

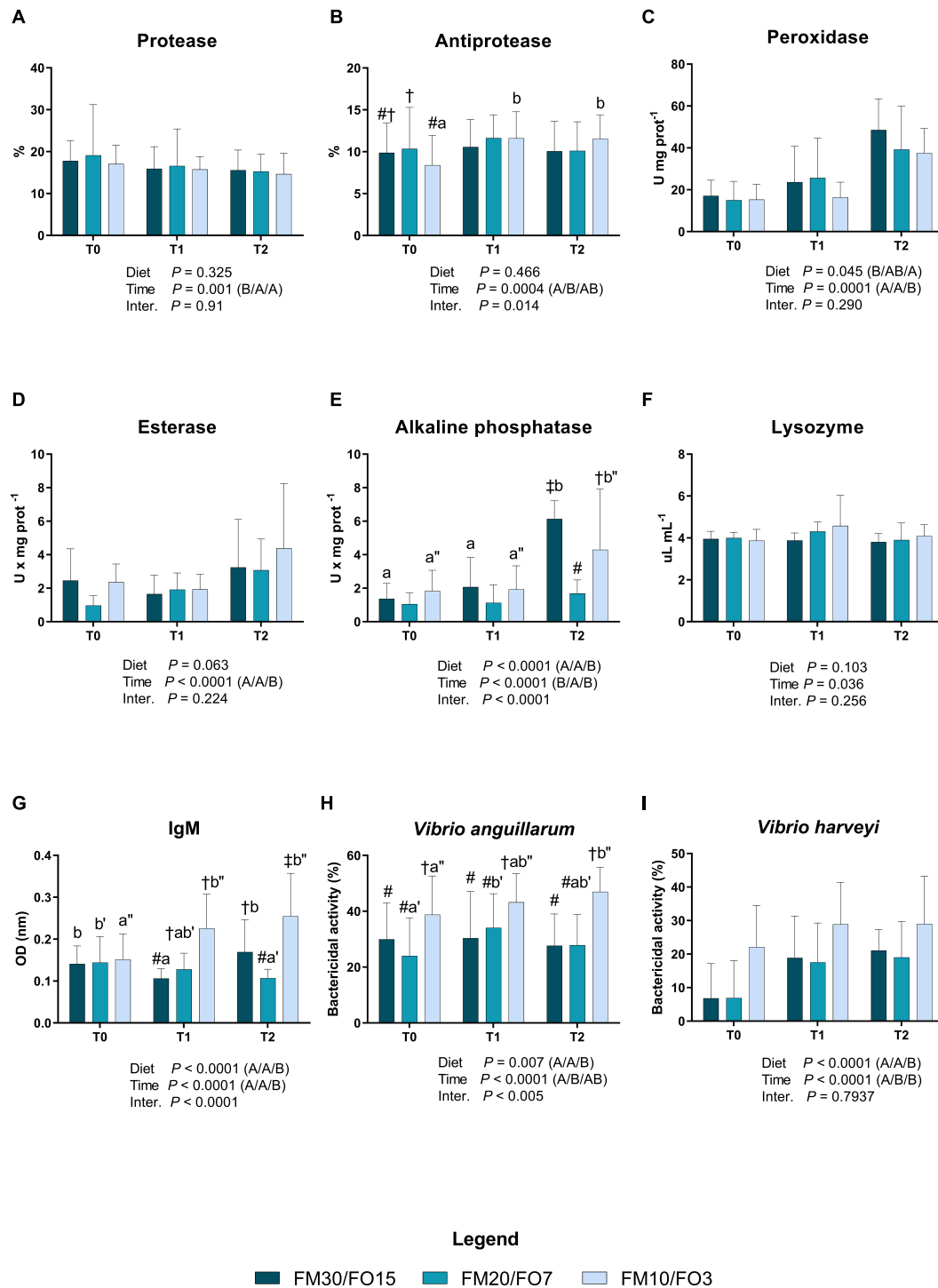


FIGURE 1 | Humoral immune biomarkers in skin mucus of European sea bass fed the experimental diets and exposed to crowding stress (2 h T1) and recovery (24 h T2) after fasting period (36 h, T0). FM30/FO15 = 300 g kg⁻¹ fishmeal (FM), 150 g kg⁻¹ fish oil; (FO); FM20/FO7 = 200 g kg⁻¹ FM; 70 g kg⁻¹ FO; FM10/FO3 = 100 g kg⁻¹ FM; 30 g kg⁻¹ FO. Data represent the mean \pm S.D. ($N = 24$). The uppercase letters in the parenthesis after P -values of two-way ANOVA represent significant differences in two-way ANOVA ($P < 0.05$). The three sets of letters in a parenthesis correspond to the three times (T0/T1/T2) or diets (FM30/FO15/FM20/FO7/FM10/FO3) groups, respectively, and data value corresponding to "A" are lower than those corresponding to "B." Different symbols stand for significant differences among dietary treatments for the same time while lowercase letters (a b c, FM30/FO15; a'b'c' FM20/FO7; a''b''c'' FM10/FO3) stand for significant differences between times for the same diet. **(A)** Protease activity (%); **(B)** antiprotease activity (%); **(C)** peroxidase activity ($U^* \text{ mg prot}^{-1}$); **(D)** esterase activity ($U^* \text{ mg prot}^{-1}$); **(E)** alkaline phosphatase activity ($U^* \text{ mg prot}^{-1}$); **(F)** lysozyme activity ($\mu\text{L}^* \text{ mg prot}^{-1}$); **(G)** IgM (OD nm); **(H)** bactericidal activity (%) against *Vibrio anguillarum*; **(I)** bactericidal activity (%) against *Vibrio harveyi*.

T0 and higher values reported in FM10/FO3 compared to the other treatments.

Gene Expression of Stress and Immune-Related Genes in Brain and Head-Kidney

Supplementary Figure 2 and Figure 2 show the relative expression values of the four target genes involved in the stress response measured in the brain and the head kidney.

At the end of the growth trial (T0) brain *gr-1* and *gr-2* relative expression decreased at decreasing FMFO level, while in the head kidney *hsp70* was lower in FM20/FO7 than FM30/FO15 and *mr* was lower in FM20/FO7 and FM10/FO3 compared to FM30/FO15 (One-way ANOVA – Supplementary Figure 2).

Figure 2 reports the same stress target genes in relation to dietary treatments and time after stress (Two-way ANOVA).

In the brain, *hsp70* were affected by time displaying lower values at T0 than T1 and T2 while brain *gr-1* was lower in T0 than T2. Brain *gr-2* was affected by diet supported mainly by the higher values observed in FM30/FO15 than FM10/FO3. No significant time nor diet effects were reported for the relative expression of *mr* in the brain.

In the head kidney, *hsp70* was up-regulated at T2 in fish fed the FM30/FO15 diet with higher values compared to the other time points within dietary treatment and in comparison to the other diets at T2. In addition, head kidney *gr-1* and *gr-2* were affected by time; specifically, *gr-1* showed higher values at T2 than T1 and T0 while *gr-2* showed higher values at T0 and T2 compared to T1. In the same tissue, *mr* was affected by time, diet and interaction; specifically, in FM30/FO15 transcripts decreased from T0 to T1, increasing again at T2 while in FM20/FO7 and FM10/FO3 values were higher at T0 than T1. At T0, FM30/FO15 displayed higher values than the other diets.

Concerning *gr-1/mr* a significant interaction effect was observed for brain where FM20/FO7 displayed higher values at T1 than T0. The *gr-1/mr* in the head kidney showed a time effect with values higher at T2 compared to the other time points.

The expression of the selected genes involved in the immune response for the brain and head kidney are depicted in Supplementary Figure 3 and Figure 3. At the end of the growth trial (T0) brain *tgf* was lower in FM20/FO7 compared to FM10/FO3 while brain *tnf- α* was lower in FM10/FO3 than FM30/FO15 and FM20/FO7. At the same time brain *cox2* was lower in FM10/FO3 compared to FM30/FO15, while brain *csp3* was lower in FM10/FO3 than FM20/FO7 (One-way ANOVA – Supplementary Figure 3). Concerning head kidney, at the end of the growth trial (T0) FM20/FO7 and FM10/FO3 showed *tnf- α* , *cox2* and *csp3* values lower than FM30/FO15, while *lys* and *il10* values of FM20/FO7 were lower than FM30FO15 (One-way ANOVA – Supplementary Figure 3).

Figure 3 reports the same immune response genes in relation to dietary treatments and time after stress (Two-way ANOVA).

In the brain, *tgf* was affected by time and interaction; specifically, values were up-regulated at T2 only in fish fed FM10/FO3. A similar trend occurred for *dcn*, where the relative expression in fish fed FM10/FO3 was higher than FM20/FO7

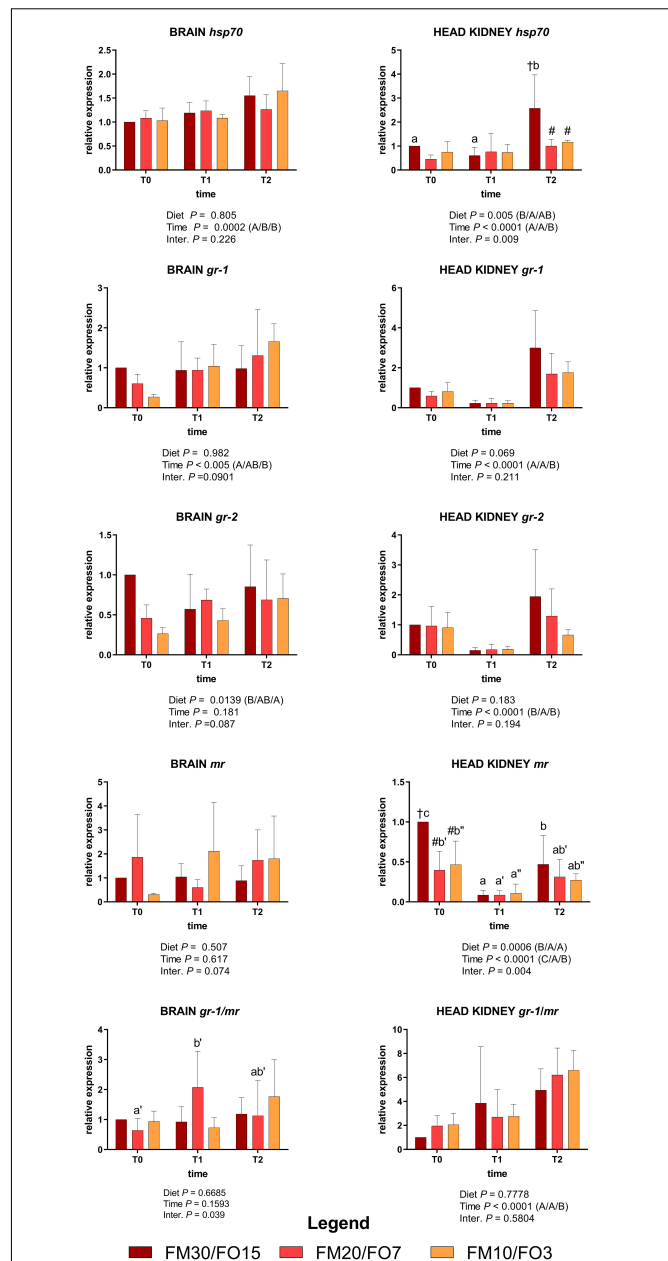


FIGURE 2 | Relative expression of stress-related genes in brain and head kidney of European sea bass fed the experimental diets and exposed to crowding stress (2 h T1) and recovery (24 h T2) after fasting period (36 h, T0). FM30/FO15 = 300 g kg⁻¹ fishmeal (FM), 150 g kg⁻¹ fish oil; (FO); FM20/FO7 = 200 g kg⁻¹ FM; 70 g kg⁻¹ FO; FM10/FO3 = 100 g kg⁻¹ FM; 30 g kg⁻¹ FO. Data represent the mean \pm S.D. (N = 6). The uppercase letters in the parenthesis after P-values of two-way ANOVA represent significant differences in two-way ANOVA (P < 0.05). The three sets of letters in a parenthesis correspond to the three times (T0/T1/T2) or diets (FM30/FO15/FM20/FO7/FM10/FO3) groups, respectively, and data value corresponding to "A" are lower than those corresponding to "B." Different symbols stand for significant differences among dietary treatments for the same time while lowercase letters (a b c, FM30/FO15; a'b'c' FM20/FO7; a"b" c" FM10/FO3) stand for significant differences between times for the same diet. Genes are reported as follows: *gr-1*, glucocorticoid receptor-1; *gr-2* glucocorticoid receptor-2; *mr*, melanocortin receptor; *hsp70*, heat shock protein 70.

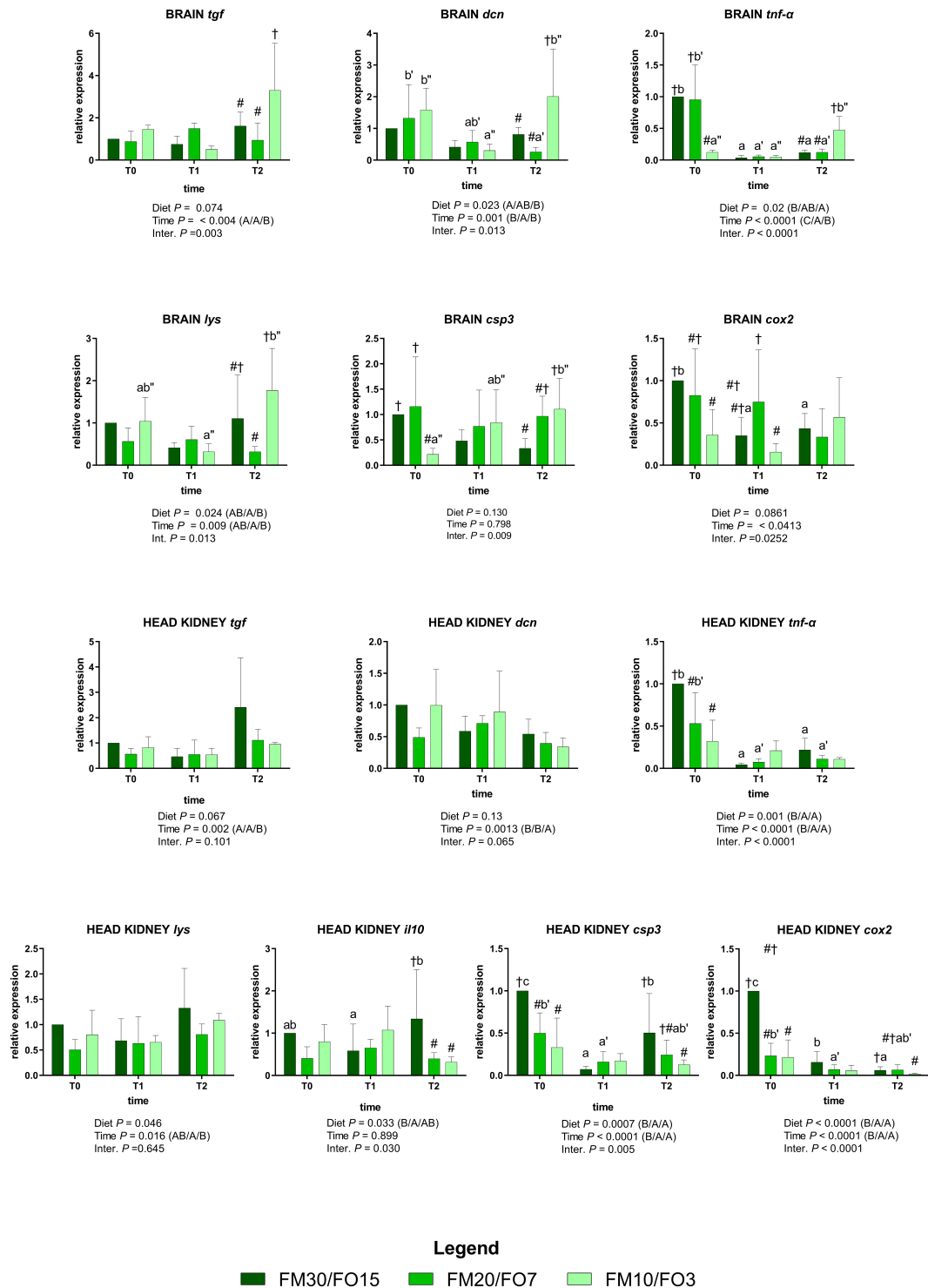


FIGURE 3 | Relative expression of immune-related genes in brain and head kidney of European sea bass fed the experimental diets and exposed to crowding stress (2 h T1) and recovery (24 h T2) after fasting period (36 h, T0). FM30/FO15 = 300 g kg⁻¹ fishmeal (FM), 150 g kg⁻¹ fish oil; (FO); FM20/FO7 = 200 g kg⁻¹ FM; 70 g kg⁻¹ FO; FM10/FO3 = 100 g kg⁻¹ FM; 30 g kg⁻¹ FO. Data represent the mean ± S.D. (N = 6). The uppercase letters in the parenthesis after P-values of two-way ANOVA represent significant differences in two-way ANOVA (P < 0.05). The three sets of letters in a parenthesis correspond to the three times (T0/T1/T2) or diets (FM30/FO15/FM20/FO7/FM10/FO3) groups, respectively, and data value corresponding to “A” are lower than those corresponding to “B.” Different symbols stand for significant differences among dietary treatments for the same time while lowercase letters (a b c, FM30/FO15; a’b’c’ FM20/FO7; a’’b’’c’’ FM10/FO3) stand for significant differences between times for the same diet. Genes are reported as follows: *csp3*, caspase 3; *hsp70*, heat shock protein 70; *tgf*, transforming growth factor; *tnf-α*, tumor necrosis factor α; *il10*, interleukin 10; *dcn*, dicentrarin, *cox2*, cyclooxygenase 2, *lys*, lysozyme.

and FM30/FO15 at T2 and higher in comparison to the same dietary treatment at T1. In addition, values in FM20/FO7 were lower at T2 compared to the values from the same treatment recorded in T1. The relative expression of *tnf- α* was affected by time, diet and interaction; specifically, values decreased at T1 and T2 only in FM30/FO15 and FM20/FO7, while values increased at T2 compared to T1 in FM10/FO3. In addition, at T0 values were lower in FM10/FO3 than the other dietary treatments while the same diet displayed higher values at T2 than the other dietary treatments.

The expression of *lys* was affected by diet, time and interaction; specifically, values were higher in FM10/FO3 at T2 compared to fish fed the FM20/FO7 diet and compared to the same dietary treatment at T1.

The relative expression of *csp3* displayed a significant diet \times time interaction where fish from the FM10/FO3 group displayed an increasing trend from T0 to T2 sampling points. In addition, at T0 values were lower in FM10/FO3 than the other diets while at T2 the same diet displayed higher values than FM30/FO15. Brain *cox2* reported significant time and interaction effects; specifically, values within FM30/FO15 were higher at T0 compared to T1 and T2. Data at T0 reported values in FM30/FO15 higher than FM10/FO3 while at T1 FM20/FO7 values were higher than FM10/FO3.

Regarding the assessment of gene expression patterns in the head kidney, *tgf* was affected by time with higher values at T2, than T1 and T0 while *dcn* displayed lower values in T2 than T0 and T1. The relative expression of *tnf- α* showed significant diet, time and interaction effect; specifically, values in FM30/FO15 and FM20/FO7 were higher at T0 than T1 and T2, while at T0 FM30/FO15 values were higher the other diets.

Head kidney *lys* gene expression showed higher values at T2 than T1. The pro-inflammatory cytokine *il10* displayed significant diet \times time interaction effect with higher values in fish fed FM30/FO15 diet at T2 compared to the same diet at T1, while at T2 FM30/FO15 values were higher compared to the other dietary treatments. Caspase 3 displayed a significant diet, time and interaction effects. In particular FM30/FO15 displayed higher values at T0 compared to the same diet at T1 and T2, while the same diet was higher at T2 than T1. In addition, *csp3* values at T0 were higher in FM30/FO15 than the other diets, while at T2 FM30/FO15 values were higher than FM10/FO3. The relative expression of *cox2* was higher in FM30/FO15 group at T0 compared to the same treatment at T1 and T2, while FM20/FO7 at T0 was higher than T1 and T2. At T0 *cox2* values were higher in FM30/FO15 than the other treatments, while at T2 FM30/FO15 values were higher than FM10/FO3.

DISCUSSION

Key Performance Indicators and Body Proximate Composition

Recently, several studies have explored the feasibility of using very low FM and FO diets in European sea bass, as their effects on fish performance, gut morphology and microbiome, plasma biochemistry and filet quality have been described

(Torrecillas et al., 2017b; Bonvini et al., 2018c; Parma et al., 2019; Henry et al., 2020). Although it has been shown that the FM and FO reduction may affect stress resistance, cellular immunity and oxidative status in different marine fish species (Sitjà-Bobadilla et al., 2005; Montero and Izquierdo, 2010; Geay et al., 2011; Gisbert et al., 2018), this topic still deserves further attention especially regarding this species, as it is highly susceptible to common stressors under intensive farming conditions.

In the present study, despite the absence of significant differences on feed intake and FCR values among diets, fish from the FM10/FO3 group showed a tendency for higher FCR; thus, leading to a negative outcome on BWf and SGR, values that were 11.3% lower than the FM30/FO15 group individuals. These findings are in contrast with those previously reported by Parma et al. (2019), where juveniles (22.8–68.2 g in BW) fed 10% FM as their sole animal protein source showed equal growth, feed and protein utilization in comparison to a control containing 20% FM. These differences could be related to the different feed formulation (i.e., lower FO inclusion) in the present trial. Similar results were reported also by Coutinho et al. (2017) where juveniles (7–28 g in BW) fed diets containing 15% FO and 8 and 2% (on dry weight basis) of FM and soluble fish protein concentrate, respectively. On the other hand, Bonvini et al. (2018a) reported a trend toward lower growth performance and protein efficiency at higher FM replacement levels in 70–200 g European sea bass when fed a diet containing 10% FM compared to their congeners fed diets containing 20 and 30% FM.

In FM10/FO3 the reduced level of marine ingredients with a consequence reduction of LC-PUFA could have affected growth performance as previously described in this species where the supplementation of n-3 and n-6 LC-PUFA in fish fed free FM and FO diet led to increased growth (Torrecillas et al., 2017a, 2018).

Concerning body composition and nutritional uptake, all treatments were found to be comparable under the present experimental conditions, results which agreed with those previously reported by Bonvini et al. (2018a) and Parma et al. (2019).

Plasma Biochemistry

Studying a farmed fish species blood biochemistry profile can be a valuable tool to assess nutritional and health status (Peres et al., 2014; Bonvini et al., 2015). Overall, the present results at T0 were within the range of normal values reported for this species under proper nutritional and non-stressing conditions (Bonvini et al., 2018b). Furthermore, different FM/FO dietary levels did not affect the basal status before stress even if cortisol level in FM30/FO15 was higher than the other treatments.

Data on plasma cortisol coupled with glucose levels is one of the first and main metabolites being released into the blood stream as a marker to acute stress response (Barton, 2002). European sea bass has been well characterized to have high basal and post-stress plasma cortisol concentrations (Fanouraki et al., 2011; Maricchiolo et al., 2011; Ellis et al., 2012) after even the minimum ordinary handling that occurs in fish farms (Tsalafouta et al., 2015; Samaras et al., 2017). In accordance with Di Marco et al. (2008), in our study the cortisol level peaked after stress

exposure with a tendency to return to values similar to those reported in the pre-stress condition.

Plasma lactate values reflect metabolic and hormonal responses and can be considered an indicator of secondary stress responses (Saccol et al., 2018). The observed higher plasma lactate after stress exposure only in fish fed the FM30/FO15 diet may indicate a stronger response to crowding by increasing anaerobic muscle activity, as also reported in other fish species exposed to stressors such as chasing, netting, and air exposure (Ruane et al., 2002; Fanouraki et al., 2007).

Under current experimental conditions, confinement stress induced a significant increase of plasma glucose, which tended to return to initial basal values after a recovery time of 24 h. Notably, glucose was found to be slightly higher in groups fed FM30/FO15 at 2 h (T1) after confinement stress. Similar trends in plasmatic glucose levels and recovery time have also been reported in European sea bass exposed to 100 kg/m³ for 15 min (Di Marco et al., 2008). Basal TRIG levels before stress are within the range of values reported by Peres et al. (2014) on this species; however, they were lower than those reported by Bonvini et al. (2018a,b). The different basal levels among these studies might be attributed to different dietary formulations regarding lipid sources as well as to different post-prandial time at which analyses were conducted. In this work, TRIG levels decreased after 2 h post-stress exposure in all the treatments. A similar decrease was reported in European sea bass after 48–72 h post-stress challenge and in animals exposed to starvation (Pérez-Jiménez et al., 2007; Di Marco et al., 2008; Peres et al., 2014). The authors suggested that this decrease is probably due to its use as an energetic substrate and to the reduction in lipogenesis, which may have contributed to the maintenance of plasma glucose levels after an acute stress event (Pérez-Jiménez et al., 2007). Total protein level is a good potential indicator to predict the nutritional status of fish, since it is usually very stable in well-nourished animals. Under present experimental conditions, total protein increased after stress in all treatments. In contrast to our results, previous studies evidenced a plasma protein decline under fasting or stress conditions as a consequence of amino acid oxidation or peripheral proteolysis (Mommensen et al., 1999; Di Marco et al., 2008; Coeurdacier et al., 2011; Peres et al., 2014). However, Montero et al. (1999), described an increase of cortisol, glucose, and plasma protein in gilthead sea bream held at high stocking density as a strategy for increasing oxygen-carrying capacity of blood during periods of high energy demand. Such differences between studies may be related to the duration and intensity of the stressor, which would affect host metabolism and plasmatic protein levels.

Non-specific enzymes ALT, ALP, LDH, and CK are plasma indicators that may provide important information on fish health status, whose elevated levels in blood may indicate tissue damage in many organs including liver, muscle, spleen and kidney (Peres et al., 2013; Guardiola et al., 2018; Parma et al., 2020). Under current experimental conditions, ALT levels dropped after stress exposure and also continued to decrease after recovery time in all the treatments. An increasing level of plasma ALT may imply liver damage and less efficient absorption and transportation of lipids and carbohydrates from the intestine as described

by other authors (Peres et al., 2014; Bonvini et al., 2018b), while ALT reduction may also be associated with changes in amino acid metabolism, since ALT degrades alanine for cellular energetic purposes.

The enzyme CK is particularly active in heart and skeletal muscle, while LDH converts lactate to pyruvate in cells; and their activity was described as being lowered by 30 and 50% respectively after 1 week fasting in European sea bass (Peres et al., 2014). Accordingly, CK and LDH concentration were found to be reduced at 24 h post-stress in a similar manner for all the treatments under current experimental conditions. Furthermore, plasma electrolytes are indicators of the secondary phase of stress response in fish, providing an indirect indication of altered plasma cortisol levels (Peres et al., 2014). Although fish may absorb some inorganic nutrients from the aquatic environment, when starvation or elevated stocking density occurs, a significant reduction in plasma phosphorus and calcium levels take place, confirming their importance as a general indicator of stress in fish (Lall, 2002; Hrubec and Smith, 2010; Guardiola et al., 2018). In this study, plasmatic calcium levels tended to increase after confinement stress and return to basal values at 24 h after stress in all dietary groups. Contrariwise, phosphorus concentration continued to decrease at T2, with generally lower values in animals fed FM20/FO7 and FM10/FO3 diets. This is consistent with a calcium and phosphorus drop and eventual resilience after starvation or stress described for this species (d'Orbcastel et al., 2010; Peres et al., 2014). Other plasma electrolytes such as potassium, sodium and magnesium alterations may be linked to acid-base balance and ions-transport disturbance (Roque et al., 2010; Bonvini et al., 2018b), which in the present study were found to tend to return to the initial values after the recovery period. However, at 2 h after stress only fish fed FM10/FO3 were characterized by an increase in the level of plasma sodium, finding which may be related to the reduced dietary LC-PUFA such as arachidonic acid which was able to attenuate the stressor-induced increase in plasma osmolality induced by acute crowding in gilthead sea bream (Van Anholt et al., 2004).

Humoral Immune Biomarkers in Skin Mucus

The study of the innate immune parameters of skin mucus (lysozyme, proteases, anti-proteases, peroxidase, esterase, alkaline phosphatase, and bactericidal activities) is of great importance in the evaluation of mucosal immunity of fish as this is the first line of the host's defense. Their evaluation has been studied in relation to pathogens, environmental stress, feeding conditions, contaminants, crowding or handling in several fish species. However, data concerning the specific response of mucosal components to different dietary formulations and stressors such as crowding or handling in European sea bass are still scarce (Guardiola et al., 2014b, 2016; Sanahuja et al., 2019).

In the present study, acute crowding slightly reduced protease activity in the skin mucus with a similar response for all the dietary treatments. In contrast, a different response of skin mucus protease activity to acute crowding was observed in gilthead sea bream, which was characterized by a decrease and increase after

2 and 24 h stress exposure, respectively. In addition, protease activity in gilthead sea bream was reduced when fish were fed low fish meal content (10% FM and 3% FO vs. 30 FM%, 15% FO), while chronic high density rearing conditions did not affect its activity (Parma et al., 2020). Antiprotease activity was significantly affected by crowding condition in fish fed the FM10/FO3 diet, which resulted in increasing values at 2 and 24 h after stress. In contrast, antiprotease activity was not affected by acute crowding in gilthead sea bream and, in agreement with our findings, antiprotease and protease were negatively correlated as a physiological response to compensate for the variations in protease activity (Guardiola et al., 2016). Peroxidase activity peaked at 24 h after stress for all the treatments. No data on skin mucus peroxidase are available for European sea bass exposed to acute crowding; however, in gilthead sea bream a similar response to crowding was observed with increasing values obtained at 48 h after stress. These differences between species could be attributed to species-specific factors, fish health status or exposure time to stress (Guardiola et al., 2014a; Sanahuja et al., 2019). In addition, in the present study a lower response of peroxidase activity was observed in fish fed the FM10/FO3 diet in comparison to those animals from the FM30/FO15 group, data that may indicate that these low FM/FO levels may reduce the specific action of this skin mucus component. Esterase activity in fish skin mucus has been previously studied mostly in terms of defensive activity against pathogens or in response to environmental chemical compounds, although their role is still not fully understood (Guardiola et al., 2014a; Sanahuja et al., 2019). No data are reported on esterase activity in fish skin mucus in response to acute crowding; however, our results indicated a positive response to stress of this mucus component in all dietary groups. Interestingly, in olive flounder (*Paralichthys olivaceus*) esterase activity of skin mucus was negatively correlated with protease activity. A similar trend occurred in the present trial, supporting the hypothesis that these enzymes may have a direct or indirect role in immune function either by acting alone or in association with other biologically active substances present in the skin mucus (Jung et al., 2012). Alkaline phosphatase is an antibacterial agent in mucus due to its hydrolytic activity, and increases in fish after physical or chemical stress, including skin regeneration (Sheikhzadeh et al., 2012a,b). In our study, ALP peaked at 24 h post-stress with a similar time-related effect as that observed for peroxidase and esterase, indicating possible synergistic effects of these three parameters in response to stress (Jung et al., 2012). Increased activity of ALP was also demonstrated in tiger barb (*Puntius tetrazona*) mucus during crowding stress as response to the increase in mucus secretion or improved mucosal immune responses (Hoseinifar et al., 2014; Roosta and Hoseinifar, 2016). Lysozyme activity was marginally affected by crowding conditions with an increasing level at 2 h post stress. Recent findings showed that acute crowding conditions in Atlantic salmon (*Salmo salar*) induced an increase in lysozyme activity of skin mucus only at 1 h post-stress, while it returned to basal values at 3 h after stress (Djordjevic et al., 2021). Conversely, high rearing density during 98 and 120 days in gilthead sea bream and turbot (*Scophthalmus maximus*), respectively, reduced skin mucus lysozyme activity as a possible

immunodepressive effect due to chronic stress caused by high stocking density (Jia et al., 2016).

In addition, the bactericidal activity of IgM production in skin mucus reached a relevant post-stress peak mainly in fish fed the FM10/FO3 diet, which displayed higher values in comparison to other treatments. Our values were higher than those reported for skin mucus IgM in gilthead sea bream after crowding stress (transfer from 10 to 50 kg m⁻³ over 48 h) (Guardiola et al., 2016). The crowding stress employed in the present study seemed sufficient to activate a further excretion of IgM in fish fed the FM10/FO3 diet. Such a response may be interpreted as a mechanism to avoid possible invasion by pathogens when animals are highly challenged, subject to frequent surface abrasion, or vulnerable to infections. A further reinforcement for the idea of there being an enhanced humoral response in low FM/FO diet was the more elevated *in vitro* bactericidal activity against *V. anguillarum* and *V. harveyi* observed in fish fed the FM10/FO3 diet. These results may be the consequence of the sum of many intrinsic antimicrobial factors such as skin antimicrobial peptides and other innate immune components to fight pathogens. In particular, recent findings have shown the importance of LC-PUFA fractions in the composition of the skin mucus of gilthead seabream in relation to different FMFO dietary level, highlighting the needs for further studies to better understand the role of feed composition on skin mucus protective functions (Torrecillas et al., 2019).

Gene Expression of Stress and Immune-Related Genes in Brain and Head-Kidney

Glucocorticoid (*gr-1*, *gr-2*) and mineral corticoid receptors (*mr*) are receptors for cortisol found in many tissues including brain and immune cells, which are related to the regulatory role of cortisol in immune defense such as the secretion of pro- and anti-inflammatory cytokines (Reyes-López et al., 2018). In addition, *hsp70* is responsible for repairing and preventing cellular damage from stress-induced protein denaturation and is considered an important indicator for stress in many species including European sea bass (Islam et al., 2021). In the present study, most of the stress-related genes considered, specifically *hsp70* and *gr-1*, in the brain, and *hsp70*, *gr-1*, and *gr-2* in head kidney, showed an overall time-dependent increasing pattern in expression after stress, findings which are in line with a positive response of the marker genes under consideration to crowding and handling stress in European sea bass and other fish species (Eissa and Wang, 2016).

Additionally, concerning *gr-1* and *gr-2*, it was previously explained that different expression patterns among *gr-1* and *mr* function as differential feed-forward and feed-back mechanisms, respectively, of cortisol on these receptor targets (Samaras et al., 2018). The *gr1/mr* ratio has been considered an appropriate indicator of stress load in fish species including European sea bass, as also reported in mammalian vertebrates (De Kloet et al., 2005, 2008; Samaras et al., 2018). In the present study, the *gr1/mr* ratio observed in the kidney increased after stress as result of the increasing, decreasing patterns of expression for *gr-1* and *mr*,

respectively. Interestingly, these genes were similarly expressed in the brain of European sea bass exposed to a combination of stresses including confinement, chasing and air exposure (Samaras et al., 2018).

Although not all the transcripts analyzed in the brain and head kidney showed a clear dietary effect, data reported for *hsp70*, *gr1*, *gr2* in the head kidney seemed to indicate a more acute response to crowding stress for fish fed the FM30/FO15 diet compared to lower FM/FO levels. In general, these results are in agreement with previous findings in European sea bass fed high levels of VOs, since dietary lipids can alter the ability of fish to cope with different types of stressors (Montero et al., 2003; Machado et al., 2019). At the biochemical level, these data are consistent with the higher glucose and lactate plasma level found in FM30/FO15, while no evident relation with plasma cortisol level was found. The absence of a direct relation between cortisol plasma level and the gene expression of glucocorticoid receptors was also previously observed in this species (Pavlidis et al., 2011; Reyes-López et al., 2018). In addition, the effect of the diet on the expression of stress-related genes was less evident in the brain than in the head kidney, possibly indicating a difference in the degree to which there is involvement of the diet in different target tissues.

While fish fed the high FM/FO (FM30/FO15) diet showed a higher response to stress, a low FM diet (FM10/FO3) resulted in a higher up-regulation of several immune-related genes at 24 h post stress, including *dcn*, and *lys* in the brain. These data were in agreement with the higher level of IgM and bactericidal activity against *V. anguillarum* and *V. harveyi* observed in skin mucus from animals fed the FM10/FO3 diet at 24-h post-stress; collectively these data all indicate a stronger activation of the immune response in fish fed this diet. Several antimicrobial molecules including lysozyme and dicentracin are known to play a key role in the innate immune response as they possess powerful microbiocidal activity against a wide range of pathogens. In European sea bass particularly, it has been suggested that dicentracin is involved in the antimicrobial function against bacteria such as *V. anguillarum* and *Photobacterium damsela* and its expression was up-regulated in head kidney at 5h post-infection with nodavirus (Mosca et al., 2014; Meloni et al., 2015; Valero et al., 2015, 2020). The higher pro-inflammatory activity observed at mucosal immune effectors (IgM and bactericidal activity) and gene expression levels (lysozyme and dicentracin) in fish from the FM10/FO3 group could also be related to the lower stress response observed after crowding, as evidenced by the lower lactate, glucose and gene expression of stress-related genes (*hsp70*, *gr-1*, *gr-2*) obtained in fish from this treatment. It is well documented that stress and immune response are closely connected due to the existing crosstalk between these two systems, since a stressor induces alterations in immune response. For instance, in fish species the inhibition of the pro-inflammatory response related to high cortisol production is well known (Castillo et al., 2009; Reyes-López et al., 2018). However, in the head kidney a different pattern of expression of immune-related genes compared to the brain was observed: in particular *tgf* and *il10* were up-regulated in fish fed the FM30/FO15 diet compared to the other treatments. Both of these

cytokines have been shown to have significant autocrine activities for normal kidney function in mammals (Sinuani et al., 2013). Some types of nephropathy in mammals have been shown to benefit from increased expression of *il10*. Therefore, the up-regulation of anti-inflammatory cytokines such as *il10* and *tgf* may support the beneficial dietary effect of high FM/FO levels enhancing anti-inflammatory response after exposure to stress (Busti et al., 2020).

CONCLUSION

In conclusion, 10% FM, 3% FO dietary inclusion levels showed a moderate reduction in growth performance, an effect that seemed to be related to the higher FCR reported in fish fed this diet. Although most of the plasma and skin mucus parameters analyzed were not affected by dietary treatments before stress, acute crowding induced different dietary-mediated responses. Specifically, a higher level of plasma lactate together with the up-regulation of some stress-related transcripts (*hsp70*, *gr-2*) suggests a higher reactivity to acute crowding of the stress-response mechanism, but a lower inflammatory response, as evidenced by higher expression of the anti-inflammatory *tnf-α* and *il10* genes in fish fed high FM and FO dietary levels (FM30/FO15). Otherwise, the higher IgM levels and bactericidal activity observed in skin mucus of fish fed the FM10/FO3 diet seems to indicate that acute crowding was able to activate a higher pro-inflammatory response in this treatment. The results of the present study seem to indicate that the low FM/FO dietary level employed could affect stress and immune response. Repeated or chronic exposure to stressors would be needed in order to further evaluate the dietary effects on fish welfare.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/**Supplementary Material**, further inquiries can be directed to the corresponding author/s.

ETHICS STATEMENT

The animal study was reviewed and approved by Ethical-Scientific Committee for Animal Experimentation of the University of Bologna (protocol ID 942/2019) in accordance with European directive 2010/63/UE on the protection of animals used for scientific purposes.

AUTHOR CONTRIBUTIONS

AB, NFP, LP, and PPG conceived and designed the experiment. NFP, AB, and LP wrote the first draft of the manuscript. NFP and LP carried out fish maintenance and sample collection. EG and KBA carried out the gene expression analysis and data processing. MAE carried out the skin mucus analysis and data

processing. FD and MCS carried out plasma biochemistry analysis. All authors reviewed, improved the writing and approved the final manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fmars.2021.779053/full#supplementary-material>

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