A novel hepcidin-like in turbot (*Scophthalmus maximus* L.) highly expressed after pathogen challenge but not after iron overload

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Abstract

Hepcidins are antimicrobial peptides with an important role in the host innate immunity. Moreover, it has been reported that mammalian hepcidins present a dual-function being a key regulator in the iron homeostasis. Here, we describe the coding sequence of a novel hepcidin-like peptide in turbot, *Scophthalmus maximus*. This molecule presents several differences with regard to the previously characterized hepcidin in this flatfish species and it has not the hypothetical iron regulatory sequence Q-S/I-H-L/I-S/A-L in the N-terminal region. Therefore we propose the existence of at least two types of hepcidin in turbot. Moreover, results revealed a higher variability in the mRNA sequences of the novel hepcidin compared with the other form. Constitutive expression of turbot hepcidins (Hepcidin-1 and Hepcidin-2) was analyzed in several tissues and as expected, both molecules were highly represented in liver. On the other hand, the effect of three different stimuli (bacterial or viral infection and iron overloading) in the level of hepcidin mRNA was also examined and a differential response to pathogens and iron was observed. Whereas both hepcidins were affected by pathogen challenge, only Hepcidin-1 was up-regulated after iron overloading. Therefore, this and other evidences suggest that these peptides could be involved in different functions covering the dual role of mammalian hepcidins.

Keywords: Turbot (*Scophthalmus maximus*), Fish, Innate immunity, Hepcidin, LEAP (liver-expressed antimicrobial peptide), Antimicrobial peptide, Bactericidal activity, *Aeromonas salmonicida*, VHSV, Iron metabolism

1. Introduction

Antimicrobial peptides (AMPs) are cationic and amphipathic host defense peptides generally composed of 12–50 amino acids in length and produced by many organisms, including plants, invertebrates, and vertebrates [1]. A wide variety of AMPs have been described to date, presenting diversity in their sequence and structure [2], but all AMPs have in common their capacity to disrupt lipidic membranes in order to kill or inhibit the proliferation of microbes [3] and in some cases, inhibit DNA or protein synthesis [4]. Moreover, several AMPs possess immunomodulatory properties. They can act as chemokines or regulate the chemokine production, participate in the recruitment, maturation and activation of immune cells, as well as act as powerful anti-inflammatory or pro-inflammatory molecules [5-8].

Hepcidin is one of these multifunctional peptides. It was discovered for the first time as a cysteine-rich peptide composed by 25 amino acids in human blood and named LEAP-1 (liver-expressed antimicrobial peptide) because its expression was predominantly detected in liver [9]. In addition to its bactericidal activity, hepcidin is also implicated in immunomodulatory functions and iron metabolism [10-18]. To our knowledge, the only mammal presenting two hepcidin genes in the genome is the mouse [10]. However, two or more hepcidin genes have been identified in several bony fish [19-26], with different degrees of sequence divergence. There are evidences suggesting that murine hepcidins (hepc1 and hepc2) perform different functions, given that only hepc1 seems to be involved in iron metabolism [27]. In fish, the different roles of hepcidin peptides still remain less clear.

Previously to our study, one hepcidin gene from turbot (*Scophthalmus maximus*) had been characterized [28]. In the present work, several Expressed Sequence Tags (ESTs) presenting homology to an “Antimicrobial peptide precursor” were analyzed and a novel hepcidin-like antimicrobial peptide was characterized in turbot. In addition, the constitutive expression of both hepcidins was analyzed in several tissues and the response to bacterial or viral infections and iron overload was also examined. On the basis of the obtained results, a potential role of turbot hepcidins is proposed.

2. Materials and Methods

2.1. Animals

Juvenile turbot (average weight 2.5 g) were obtained from a commercial fish farm (Insuiña S.L., Galicia, Spain). Animals were maintained in 500 L fiberglass tanks with a re-circulating saline-water system with a light–dark cycle of 12:12 h at 18 °C and fed daily with a commercial diet (LARVIVA-BioMar, France). Prior to experiments, fish were acclimatized to laboratory conditions for 2 weeks. Fish care and challenge experiments were reviewed and approved by the CSIC National Committee on Bioethics.

2.2. ESTs analysis

ESTs presenting similarity to an “Antimicrobial Peptide precursor” were obtained from a turbot sequencing assays performed by Pardo *et al.* [29] and from a high-throughput sequence analysis of turbot transcriptome using 454-pyrosequencing performed in our laboratory. The ESTs alignment was performed using ClustalW [30] and a contig was constructed using CAP3 sequence assembly program [31]. The sequence was translated from nucleotide to amino acid sequence using the translation tool from ExPasy [32] and was submitted to *Blastp* (<http://blast.ncbi.nlm.nih.gov/Blast.cgi>) for detecting homologies with other proteins. A comparison with other hepcidin proteins was performed for determining if this sequence represents the full-length coding sequence. According to the classification of hepcidin peptides performed by Hilton & Lambert [33] we propose the nomenclature of Hepcidin-1 (Hep-1) for the molecule described by Chen *et al.* [28] and Hepcidin-2 (Hep-2) for the novel AMP identified in the present work, and they will be named accordingly in this paper.

2.3. Analysis of sequence variability of both turbot hepcidins

Total RNA was extracted from a pool composed by head kidney from 4 turbot intraperitoneally stimulated with the pathogen bacteria *Aeromonas salmonicida* subsp. *salmonicida* (5.5×10^5 CFU/fish) using *TRIzol*® (Invitrogen) in accordance with instructions provided by the manufacturer in combination with the RNeasy mini kit (Qiagen) for RNA purification after DNase treatment (RNase-free DNase set, Qiagen). Quantity of the total purified RNA was determined using the spectrophotometer Nanodrop ND-1000. The reverse transcription was performed with the SuperScript II Reverse Transcriptase (Invitrogen) using 0.5 µg of RNA and following the manufacturer indications.

In order to determine if Hepcidin-1 and Hepcidin-2 presented variations in the sequence, specific primers were designed to amplify the open reading frame (ORF) (Table 1). The amplifications were performed in a 25 µl total volume containing 10µl of ultrapure water (*Sigma-Aldrich*), 12 µl of 2x PCR Master Mix (*Fermentas*), 1 µl of each specific primer (10 µM) and 1 µl of cDNA. PCR conditions consisted on an initial denaturation for 5 min at 94°C, 30 cycles of denaturation at 94°C for 30 s, annealing at 54°C for 30 s, elongation at 72°C for 1 min and a final extension for 7 min at 72°C in a GeneAmp® PCR System 2700 thermocycler (Applied Biosystems). The PCR products were analyzed on a 1.5% agarose gel stained with ethidium bromide under UV light using the Gel Doc XR system and the Quantity one program (Bio-Rad, Hercules, Ca). 3 µl of PCR product were linked into pCR® 2.1 plasmid vector (Invitrogen) for its cloning following the protocol instructions and One Shot® TOP10F' competent cells (Invitrogen) were transformed using the plasmid. Bacteria were cultured on LB/Ampicillin/IPTG/X-Gal plates during 24 hours at 37°C and the bacterial colonies were selected by the blue-white screening method. 25 positive clones for each hepcidin were selected and the insert was amplified using the M13 vector specific primers (M13F: 5'-GTAAAACGACGGCCAG-3', M13R: 5'-CAGGAAACAGCTATGAC-3). PCR was performed as mentioned above. cDNA sequencing was conducted using an automated ABI 3730 DNA Analyzer (Applied Biosystems, Inc. Foster City, CA, USA). For each hepcidin, the 25 sequences were compared using ClustalW [30] in order to detect sequence variations.

The consensus sequences for both hepcidin were obtained using CAP3 software [31] that corresponded to the most prevalent clone. In each case, sequences were aligned for identifying the conserved regions between both turbot hepcidins. Sequence similarity and identity scores were calculated with the software BioEdit [34] using the BLOSUM62 matrix. Prediction of cleavage sites for the signal peptide and the prodomain was performed using ProP 1.0 server [35]. The molecular weight and isoelectric point of the mature peptide were determined using the *Compute pI/Mw tool* from *ExPASy* [32] and the estimated half-life of both peptides was predicted using ProtParam tool too from *ExPASy* [32].

2.4. Genomic organization

Genomic DNA was extracted from healthy turbot muscle using the Phenol-Chloroform method [36]. In order to determine the presence of introns in the genomic sequence of

the novel Hepcidin-2 the amplification was performed with the same primers used for the confirmation of the full-length coding sequence (Table 1). One µg of DNA was used as template for PCR amplification. The elongation step on each cycle was 90 seconds for the genomic sequence. Four positive clones were sequenced. Exon-intron junctions were deduced using Wise2 software (<http://www.ebi.ac.uk/Wise2/programming>).

2.5. Phylogenetic analysis

Amino acid sequences of turbot hepcidins were compared with those from other fish and vertebrates (whole prepropeptides). Sequences employed for the alignment and their GenBank accession numbers were: *Ictalurus punctatus* (NP_001187130), *Siniperca chuatsi* (ACO88905), *Salmo salar* (AAO85553), *Oncorhynchus mykiss* (ADU85830), *Sparus aurata* (CAO78619), *Morone chrysops* (AAM28440), *Danio rerio* (AAN10302), *Dicentrarchus labrax* (AAZ85124), *Paralichthys olivaceus* (BAE06234) and (BAE06235), *Puntius sarana* (CAZ68137), *Crocodylus siamensis* (ADA68357), *Xenopus tropicalis* (NP_001090729) and (ABL75284), *Homo sapiens* (NP_066998), *Pan troglodytes* (ABU75211), *Equus caballus* (NP_001161799), *Mus musculus* (NP_115930) and *Rattus norvegicus* (NP_445921). Multiple sequence alignment was performed using the *T-Coffee* server [37] in regular computation mode and using the *t_coffee_msa* multiple alignment method and phylogenetic tree was drawn using Mega 4.0 software [38] Neighbor-Joining algorithm [39] was used as clustering method, the distances matrix was computed using Poisson correction method and all positions containing alignment gaps and missing data were eliminated. Statistical confidence of the phylogenetic analysis was assessed by performing 1,000 bootstrap replicates.

2.6. Constitutive expression of hepcidins

Eight different tissues (kidney, spleen, gill, liver, intestine, heart, brain and muscle plus skin) were removed from 20 healthy fish in order to examine the constitutive expression of both hepcidins. Equal amounts of the same tissue from four fish were pooled, obtaining 5 biological replicates for each tissue (4 turbot/replicate). Total RNA was extracted and cDNA was synthesized as mentioned above. Hepcidin expression profiles were determined using Real-time quantitative PCR. Specific PCR primers (Table 1) were designed using the *Primer3* program [40] and their amplification efficiency was calculated using seven serial five-fold dilutions of head kidney cDNA from

unstimulated turbot with the Threshold Cycle (C_T) slope method [41]. Individual real-time PCR reactions were carried out in 25 μ l reaction volume using 12.5 μ l of SYBR® GREEN PCR Master Mix (Applied Biosystems), 10.5 μ l of ultrapure water (Sigma-Aldrich), 0.5 μ l of each specific primer (10 μ M) and 1 μ l of five-fold diluted cDNA template in MicroAmp® optical 96-well reaction plates (Applied Biosystems). All reactions were performed using technical triplicates in a 7300 Real-Time PCR System thermocycler (Applied Biosystems) with an initial denaturation (95°C, 10 min) followed by 40 cycles of a denaturation step (95°C, 15 s) and one hybridization-elongation step (60°C, 1 min). An analysis of melting curves was performed for each reaction. Relative expression of Hepcidin-1 and Hepcidin-2 was normalized using the Elongation Factor-1 alpha as reference gene, which was constitutively expressed and not affected by the experimental treatments, and calculated using the Pfaffl method [41]. Fold-change units were calculated by dividing the normalized expression values for each tissue by the normalized expression values obtained in the organ presenting the lower expression level for each hepcidin. The relative rates of turbot hepcidin types expressed in the different tissues were calculated taking into account the normalized values of each hepcidin in the same tissue.

2.7. Expression of hepcidins under different stimuli

In order to study the transcriptional induction of turbot hepcidins several experimental stimulations were performed. Three kinds of stimuli were used: bacterial or viral infections and iron overloading. Fish were divided into five groups, composed of 80 fish each. Turbot belonging to one group were injected intraperitoneally with 50 μ l of an *Aeromonas salmonicida* subsp. *salmonicida* (strain VT 45.1 WT) suspension (5.5×10^5 CFU/fish), other group was inoculated with a Viral Haemorrhagic Septicaemia Virus (VHSV- strain UK-860/94) suspension (1.2×10^5 TCID₅₀/fish) and eighty turbot using Iron-dextran (Sigma Chemical Co., St. Louis, MO, USA) (0.1 mg/fish). The Gram-negative bacteria and Iron-dextran were resuspended in 1x Phosphate Buffered Saline (PBS 1x) and the viral suspension using Eagle's minimum essential medium (MEM, Gibco) supplemented with 2% fetal bovine serum (FBS), penicillin (100 IU/ml) (Invitrogen) and streptomycin (100 μ g/ml) (Invitrogen). Therefore, two control groups were necessary, one injected intraperitoneally with 50 μ l of PBS 1x and another one using MEM + 2% FBS + P/S. Head kidney and liver from twenty fish belonging to each

experimental group were removed at different sampling points (3, 8, 24 and 72 hours). For each sampling point and treatment, equal amounts of each tissue from four turbot were pooled, constituting five biological replicates for head kidney and other five for liver (4 turbot/replicate). Moreover, in order to determine if hepcidins expression was iron dose-dependent, thirty-six fish were divided into 3 batches of 12 individuals each and every group was injected with a different Iron-dextran dose (0.1 µg/fish, 0.1 mg/fish and 0.5 mg/fish). Head kidney and liver were removed at 24 hour after stimulation and the tissues from four turbot were pooled, obtaining 3 biological replicates. Total RNA was extracted and cDNA was synthesized as mentioned above and amplification was carried out using the same protocol previously described.

2.8. Statistical analysis

Expression results were represented graphically as the mean $\pm$ the standard deviation of the biological replicates. In order to determine statistical differences, data were analyzed using the computer software package SPSS v.19.0. One-way ANOVA followed by Tukey's multiple comparison test of the means among the different time groups and between each time group and its corresponding control was performed. Differences were considered significant when $p < 0.05$.

3. Results

3.1. Molecular characterization of Hepcidin-2

Hepcidin-1, previously identified in turbot [28], was composed of 90 amino acids, however, the deduced amino acid sequence of the new hepcidin form characterized in the present work consisted of 84 amino acids with a full-length coding sequence of 255 bp in length (Figure 1). Turbot Hep-2 exhibited cleavage sites for processing the immature peptide. A potential cleavage site for the signal peptide was predicted to be located between positions 24 and 25 (SAA-TF) and a motif for the propeptide cleavage was predicted between the residues Arg⁶² and Gly⁶³. Consequently, the predicted prepropeptide consisted of a 24 amino acids signal peptide, a prodomain of 38 residues and a mature processed peptide of 22 amino acids. The predicted molecular weight of the mature peptide was 2365.87 Da and the isoelectric point was 7.70. The estimated half-life for Hep-1 (0.8 hours in mammalian reticulocytes *in vitro*, 10 min in yeast or 10

hours in *Escherichia coli*) was shorter than the half-life for Hep-2 (30 hours in mammalian reticulocytes *in vitro*, >20 hours in yeast or >10 hours in *E. coli*).

3.2. Patterns of nucleotide and amino acid variability in the sequence of both hepcidins

The comparison of the 25 sequences of the previously described Hepcidin-1 revealed only a single nucleotide polymorphism (SNP) in the mature peptide at the nucleotide position 241 of ORF: 5/25 of the sequences presented an A and 20/25 a T which coded for a different amino acid in the position 81 (Asn or Tyr, respectively). Hepcidin-2 however, presented a higher polymorphism showing several nucleotide substitutions. The presence of these changes resulted more prevalent in the mature peptide region, affecting in some cases the presence of the fifth cysteine (Figure 1). The amino acid sequences of turbot Hep-2 were submitted to GenBank under accession numbers JQ219831-9.

The alignment between both hepcidin consensus sequences revealed numerous differences in the amino acid sequences, but the eight Cys residues were relatively well conserved with some exception as it was mentioned above (Figure 2). The identity and similarity scores between both hepcidins were 51.1% and 66.67%, respectively. Interestingly, only Hep-1 presents the hypothetical iron regulatory sequence Q-S/I-H-L/I-S/A-L in the N-terminal region of the mature peptide.

3.3. Genomic organization

The hepcidin-2 genomic structure was composed by 3 exons and 2 introns. Intron 1 was 85 bp in length whereas intron 2 was 132 bp in length. The exon 1 contained the signal peptide region and 12 nucleotides belonging to the prodomain, the exon 2 was entirely composed by a prodomain sequence and the exon 3 contained 24 nucleotides of the prodomain and the mature peptide nucleotide sequence (Figure 3). Turbot Hep-2 gene organization followed the same pattern of hepcidin genes from other vertebrates. Interestingly, partially spliced Hep-1 transcripts retaining the first intron were identified, but not in Hep-2. The genomic sequence of turbot Hep-2 was submitted to GenBank under accession number JQ219840.

3.4. Phylogenetic analysis

Protein alignment with other hepcidin sequences from fish species and other vertebrates showed that hepcidins are relatively well conserved among the vertebrates, specially the

eight cysteine residues in the mature peptide (Figure 4A). The alignment revealed that hepcidin sequence similarity among fish species is higher than between fish and the other vertebrates. Moreover, notable variations in the mature peptide length are present among species and between hepcidin types of the same organism, as occurs in

Paralichthys olivaceus and *Xenopus tropicalis*, as well as in *S. maximus*.

Phylogenetic analysis placed the fish hepcidins in a separated cluster from the other vertebrates (Figure 4B). *Danio rerio* and *Puntius sarana* are cyprinids, and the hepcidins from these species were grouped in a branch. The salmonid proteins appeared also forming a separate group within the cluster of fish. Hepcidins from the perciforms *Sparus aurata*, *Siniperca chuatsi*, *Dicentrarchus labrax* and *Morone chrysops* also grouped together. *Scophthalmus maximus* Hep-1 was placed in the same cluster of *Paralichthys olivaceus* Hep-JF2, the other flatfish used in the analysis; *Scophthalmus maximus* Hep-2 was grouped within the same branch of *Paralichthys olivaceus* Hep-JF1. Between turbot Hep-1 and Japanese flounder Hep-JF2 the identity was 79.1% and the similarity was 83.5%, and the values between turbot Hep-2 and Japanese flounder Hep-JF1 were 58.3% and 74.1%, respectively.

3.5. Constitutive expression of both Hepcidins

Hepcidin-1 and Hepcidin-2 mRNA were detected in all tissues tested in healthy turbot. Fold-change for each tissue was referenced to the brain (1 fold), the organ presenting the lower expression level for both transcripts. Hep-1 was predominantly expressed in the liver (about 150 fold-change), a moderate level was detected in kidney, intestine and spleen (7-11 fold) and the lowest expression in gill, muscle plus skin, heart and brain (<2 fold) (Figure 5A). Hep-2 was also predominantly detected in liver (about 2255 fold-change); an elevated presence of transcripts was also observed in intestine (about 440 fold) and kidney (about 100 fold); the other tissues presented a lower fold-change of 20 fold) (Figure 5B). The relative rates of turbot hepcidin types expressed in the different tissues showed a higher presence of Hep-2 mRNA in most of the tissues tested, with the exception of muscle/skin and brain. The highest proportion of Hep-2 with regard to Hep-1 was observed in intestine and the larger proportion of Hepcidin-1 compared with Hep-2 was obtained in brain (Figure 5C).

3.6. Hepcidin expression after bacterial or viral infections and iron overload.

Regarding hepcidin expression after immunostimulation using the fish-pathogenic bacteria *Aeromonas salmonicida* subsp. *salmonicida*, Hep-1 increased its expression significantly in head kidney at 3 h post-infection (p.i.) (2.5 fold), and reached the larger fold-change at 24 h p.i. (35 fold). In liver, there were not significant differences compared to PBS-injected control at 3 h p.i., but the level of Hep-1 mRNA exhibited an increase at 8 h after bacterial challenge (2.6 fold), and the level remained more elevated at 24 and 72 h p.i. (about 5 fold) (Figure 6A). Hepcidin-2 expression was significantly inhibited in head kidney at 3 h p.i. (fold-change value of 0.2), but at 8 h p.i. the fold-change reached an increase of 10 fold and at 24 h after bacterial stimulation the level increased until 700 fold with regard to the control group. In liver, a significant up-regulation of Hep-2 was observed at 24 h post-injection (about 2.8 fold) and the level remained in the same value at 72 h. (Figure 6B). Interestingly, although the constitutive expression of hepcidin was predominant in liver, its induction after *Aeromonas salmonicida* subsp. *salmonicida* challenge was more pronounced in head kidney. Moreover, in head kidney the highest induction was for Hep-2 and in liver was reached for Hep-1.

Hep-1 expression was drastically reduced after viral stimulation with VHSV at 8 h p.i. in head kidney (fold-change value of 0.15) but at 72 h the expression level was significantly up-regulated (2.6 fold). In liver this increase occurs earlier, at 24 h after VHSV infection with the same fold-change value (2.6 fold) and at 72 h the increase with regard to the MEM + 2% FBS + P/S-injected control group was 3.5 fold (Figure 7A). As occurs with Hep-1, Hep-2 transcripts were highly down-regulated in head kidney at 8 h after viral stimulation (fold-change value of 0.01) but it was strongly up-regulated at 72 h (more than 42 fold). In liver a significant inhibition was also detected after VHSV injection at 8 h (about 0.7 fold-change value), but an important increase of Hep-2 mRNA was observed after 72 h (Figure 7B). The highest increase of hepcidin expression also occurred for Hep-2 in head kidney.

The administration of Iron-dextran (0.1 mg/fish) to juvenile turbot for generating an iron overload also affected the expression of hepcidins. Hep-1 transcripts were inhibited in head kidney and liver at 3 h after stimulation. In head kidney an up-regulation was detected at 72 h (3.1 fold) and in liver at 24 h (4.6 fold), remaining high values until 72 h (7 fold) (Figure 8A). A significant initial inhibition of Hep-2 level was also detected in head kidney at 3 h after iron administration with regard to PBS-injected control. No significant differences were found at subsequent sampling points. During the

experimental time course, iron overloading did not affect Hep-2 mRNA level in liver (Figure 8B). Moreover, hepcidin expression was Iron-dextran dose dependent at 24 h after administration. A significant up-regulation was detected for Hep-1 with the lowest iron dose (0.1 µg/fish) in head kidney (about 1.8 fold), but this increase disappeared with the intermediate dose (0.1 mg/fish) and a notable inhibition at concentrations of 0.5 mg/fish was observed. In liver the response was opposite, exhibiting significant elevation of Hep-1 expression with higher iron doses (Figure 9A). Regarding Hep-2, head kidney showed an inhibition in mRNA level, and this reduction was more pronounced using more concentrated solution (Figure 9B). The same tendency was observed in liver, but differences were non-significant.

4. Discussion

Mammalian hepcidins are multifunctional molecules involved in numerous immune processes but, on the other hand, present a pivotal role in iron metabolisms [10-18]. A new hepcidin-like antimicrobial molecule from the flatfish *Scophthalmus maximus* was characterized in this study. Turbot Hep-2 presents several differences in the sequence and length of the three regions (signal peptide, prodomain and mature peptide) with regard to the previously described turbot Hep-1. The identity score between both turbot hepcidins suggests that there are at least two different types of hepcidin in turbot. Genetic duplication and further mutation of AMPs has been suggested as a molecular strategy in the evolution of these molecules [42]. There are evidences for positive Darwinian selection on antimicrobial peptides [43], and more specifically in hepcidin-like variants of pleuronectiform and perciform [44]. The prevalence of mutations is usually most elevated in the mature peptide region compared to the signal sequence and prodomain [45]. Turbot Hep-2 presents amino acid substitutions in the three regions conforming the prepropeptide, but the prevalence of SNPs is more pronounced in the mature peptide, that is, the active region. This genetic variation due to accelerated evolutionary rates might possibly be directed by pathogens when the host is exposed to new environments [44,46]. As it was proposed for the complement component C3 in teleost fish, the presence of different isoforms would allow these animals to expand their innate immune recognition capabilities [47]. It could be interesting to investigate whether changes in the mature peptide region generate diversification or specialization in the functions of turbot hepcidins and affect their ability for bacterial inhibition.

In a similar way to other vertebrates, Hep-2 genomic organization is composed by three exons and two introns. As occurs in the majority of fish hepcidins, the first intron is larger than the second intron [19,20,22,28,48-54] with some exception [55]. Moreover, introns of turbot Hep-2 (85 and 132 bp) are shorter than introns of turbot Hep-1 (114 and 172 bp). A curious phenomenon of truncated hepcidin transcripts retaining the first intron was observed in some turbot Hep-1 transcripts. This event was also reported previously in rockbream hepcidins [52], where a significant increase of these partial spliced transcripts occurred after bacterial challenge or iron overload. In our work, we also cloned the turbot hepcidin from turbot previously stimulated with *A. salmonicida*. The resultant forms presented a stop codon in the intron and, as a result, these transcripts may not be translated into functional hepcidin. Cho *et al.* [52] proposed that the massive induction of transcripts causes a reduction in the specificity of the splicing process in immature transcripts, but further analysis will be necessary for determining the incomplete processing of these mRNA.

A phylogenetic analysis of hepcidins in vertebrate species revealed a strong conservation of the eight cysteine residues in the mature peptide region, conforming four disulphide bonds characteristic of hepcidins. Interestingly, Japanese flounder Hep-JF1 only presents six cysteines. Moreover, one mutation detected in turbot Hep-2 can affect the presence of the fifth cysteine. As was reported by Nemeth *et al.* [56] for human hepcidin, altered disulfide-bonding pattern allow a nearly full activity of the peptide *in vitro*. It is possible that in the case of turbot Hep-2 functionality also remains. Fish hepcidins formed a separated cluster from the other vertebrate hepcidins and within the fish group the similarities were higher among species belonging to the same order. The identities and similarities between turbot Hep-1 and Japanese flounder Hep-JF2 and between turbot Hep-2 and Japanese flounder Hep-JF1 were higher than the identity and similarity between both turbot hepcidins. These results suggest that Hep-1 and Hep-2 from turbot could have different functions.

As expected, both turbot hepcidins were abundantly expressed in liver. Our results for Hep-1 distribution differ in some tissues with those obtained by Chen *et al.* [28] using RT-PCR which could be due to differences in the age of the animals (adult fish or juveniles) or to the used technology (real-time quantitative PCR is more sensitive than the conventional PCR used by these investigators). The relative proportion of hepcidin types showed a predominant presence of Hep-2 in all tissues tested with the exception

of muscle plus skin and brain. This tissue-specific pattern of hepcidin mRNA in healthy juvenile turbot is in agreement with what it has been reported for other fish hepcidins [20-22,52,57].

Turbot hepcidins were significantly induced after bacterial challenge with *Aeromonas salmonicida*. Despite the highest presence of hepcidin transcripts in liver, the fold increase relative to control fish was more pronounced in head kidney for Hep-1 and Hep-2. Our findings contrast with the results obtained by Chen *et al.* [28] which observed a significantly up-regulation of turbot Hep-1 expression at 24, 48, 72 and 96 h after infection with *Listonella anguillarum* in liver and spleen but not in head kidney. Another experimental assay revealed a significant up-regulation of hepcidin transcripts in liver and spleen but not in head kidney after *Listonella anguillarum* infection in gilthead seabream [53]. Interestingly, Japanese flounder hepcidins were differentially affected by lipopolysaccharide (LPS) injection; Hep-JF1 expression was induced in liver but not in kidney and Hep-JF2 expression was up-modulated in kidney but not in liver [20]. Bo *et al.* [26] observed an increase in both *Oryzias melastigma* hepcidins (OM-hep1 and OM-hep2) in liver and spleen after *Vibrio parahaemolyticus* administration and, as in our case, the response was faster in the case of type 1 hepcidin. Induction of hepcidin expression after bacterial or LPS administration could vary as a function of the type of hepcidin in species presenting more than one form. There is scarce information about the response of fish hepcidin peptides to viral infections and the studies are usually based in the effect of Polyriboinosinic polyribocytidylic acid (poly I:C) injection, which is a synthetic nucleic acid widely used in the study of the immune response to double-stranded RNA virus [21,55,58]. These reports have shown that fish hepcidins can be induced by poly I:C. Cho *et al.* [52] analyzed the expression of hepcidin peptides in rockbream after iridovirus infection and detected an up-regulation of hepcidins in liver, intestine, kidney and spleen at 10 days post-challenge and the increase was especially high in liver. Hepcidin up-regulation was also reported in gilthead seabream after VHSV inoculation at 4 and 72 h post-injection in liver, head kidney and spleen [53]. Our analysis about the modulation of turbot hepcidins after VHSV challenge revealed an initial inhibition of Hep-1 and Hep-2 expression in head kidney and liver, but at 72 h after infection an up-regulation of both hepcidins was detected. This initial down-regulation of hepcidin levels could be related to increased levels of Tumor necrosis factor (TNF)- α after VHSV infection at 3, 8 and

24 h but not at 72 h post-challenge (data not shown). It has been found that TNF inhibits the release of iron from macrophages inducing hypoferremia and consequently hepcidin levels may be affected [59,60], but more research is needed to clarify this issue. Moreover, *in vitro* analysis will be necessary in order to determine if hepcidin peptides can inhibit viral replication, as was reported for marine medaka Om-hep1 against White Spot Syndrome Virus (WSSV) [61].

On the other hand, hepcidin is the main negative regulator in iron homeostasis in mammals [10-12]. Nicolas *et al.* [11] concluded in their study using upstream stimulatory factor-2 (USF2) gene knock-out mice that hepcidin was a negative regulator of iron uptake at the intestinal level and of the iron release from macrophages. When the level of iron in the organisms is high, the expression of hepcidin is elevated. Hepcidin acts by direct binding to its cellular receptor ferroportin, the unique known iron transporter on the cell membrane, and this receptor is internalized and degraded at lysosomal level [62]. As consequence, the iron absorption from enterocytes to plasma and its exportation from macrophages get canceled. Moreover, hepcidin reduces the iron available for pathogen proliferation since bacteria require iron as a growth-essential nutrient [63] and an efficient viral replication needs an iron-replete host [64]. The role of fish hepcidins in iron homeostasis remains still unclear. There are evidences about the conservation of the dual role of hepcidins in fish. Rodrigues *et al.* [51] detected an up-regulation of sea bass hepcidin level in liver at 4 days after experimental treatment using Iron-dextran. Huang *et al.* [21] studying the effect of an iron overload in the expression of the three different hepcidins from tilapia have suggested that these different types have their own functions in different organs for regulating iron metabolisms, since each hepcidin responds to the iron administration in different tissues. With regard to Japanese flounder, gene expression of Hep-JF1 in liver was down-regulated during experimental iron overloading (at 1, 2 or 3 weeks after treatment), whereas Hep-JF2 expression was not affected in liver but its expression in kidney increased after injection of iron-dextran [20]. Our results showed a possible role of Hep-1 in iron homeostasis in head kidney and liver, since a significant increase in the expression level was detected in both tissues. However, the gene expression of Hep-2 was not up-regulated during the experimental assay. On the other hand, a significant initial inhibition was detected for Hep-1 in head kidney (3 h) and liver (3 and 8 h) and for Hep-2 in head kidney (3 h). The reason for this initial down-regulation of hepcidin

transcripts is unknown and further studies are necessary in order to understand this process. Moreover, increasing doses of Iron-dextran significantly reduced the Hep-1 and Hep-2 expression in head kidney, but the Hep-1 levels in liver were progressively increased using higher doses of Iron-dextran.

The overall results of this paper seem to suggest that Hep-1 is more involved in body iron regulation, whereas the role of Hep-2 would be a more specialized one in the immune defense. Furthermore, the higher variability in the amino acid sequence of Hep-2 may be an evolutionary mechanism for the recognition of a diverse range of microbes and the longer half-life with regard to Hep-1 could favor the elimination of pathogens. Other evidence supporting this hypothesis is the presence of the Q-S/I-H-L/I-S/A-L sequence in the N-terminal region of the mature peptide. As was proposed by Robertson *et al* [23] the existence of these amino acids could be related with an iron-regulatory function. There are three hepcidins in *Oreochromis mossambicus* and one of those hepcidins (TH2-3) possesses the characteristic sequence and the other two peptides lack these amino acids; interestingly, tilapia hepcidin TH2-3 seems to be the peptide implicated in iron regulation in liver [21]. *Paralichthys olivaceus* possesses two types of hepcidin, and Hep-JF2 contains a similar N-terminal sequence, H-I-S-H-I-S-M. As was reported for tilapia TH2-3, only Japanese flounder Hep-JF2 was up-regulated in head kidney after iron overloading [20], Turbot Hep-1 contains the typical N-terminal sequences and, as occurs with their homolog in *P. olivaceus*, their expression was up-regulated after Iron-dextran administration, whereas the level of Hep-2 (hepcidin lacking this sequence) remained low.

In conclusion, a novel hepcidin-like antimicrobial peptide was characterized in the flatfish *Scophthalmus maximus*, presenting numerous differences in the sequence with the previously described hepcidin. Our results on the expression of both hepcidins suggest that Hep-1 and Hep-2 may have different functions, since their transcriptional modulation was different, the variability in the sequence and the half-life was higher in Hep-2 compared to Hep-1 and the presence of the Q-S/I-H-L/I-S/A-L sequence in the N-terminal region may be a marker to identify those most involved hepcidin in iron metabolism function. Hence, the dual role of mammalian hepcidins could be divided between both peptides in turbot. Further studies will be needed in order to elucidate this question.

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Figure legends

Figure 1. ORF sequence of turbot Hepcidin-2. The amino acid sequence corresponding to the predicted signal peptide is boxed, the prodomain region is in cursive letters and the mature peptide is shaded in grey colour. Polymorphic nucleotide and amino acid positions are marked in bold. Intron positions and lengths are indicated by vertical lines.

Figure 2. Alignment showing the identities between both turbot hepcidins. Identical amino acids are shaded in grey color. Hashes indicate the positions of the eight conserved Cys residues.

Figure 3. An exon-intron diagram showing the organization of turbot Hepcidin-2 gene.

Figure 4. (A) Amino acid alignment of turbot hepcidins with hepcidin peptides from other vertebrates. The signal peptides are underlined, the mature peptides are shaded in grey colour and the eight conserved Cys are indicated with hashes. (B) Phylogenetic tree of hepcidins from different vertebrates based on the complete amino acid sequence. The tree was constructed using the neighbour-joining method and gaps were completely deleted. Numbers indicate the percentage of bootstrapping after 1,000 replications.

Figure 5. (A) Hep-1 and (B) Hep-2 constitutive expression in various tissues of juvenile turbot. Fold-change units were calculated by dividing the normalized expression values for each tissue by the normalized expression values obtained in the organ presenting the lower expression level for each hepcidin (brain in both cases). (C) Relative proportions of turbot hepcidin types in different tissues calculated taking into account the normalized values of each hepcidin in the same tissue.

Figure 6. Expression of Hep-1 (A) and Hep-2 (B) in head kidney and liver at 3, 8, 24 and 72 h after *Aeromonas salmonicida* subsp. *salmonicida* challenge (black bars). Asterisks above the bars indicate significant differences between each sampling point and the corresponding control (PBS-injected groups) (white bars), and asterisks over lines show statistical significant differences across the different sampling points ($p < 0.05$).

Figure 7. Expression of Hep-1 (A) and Hep-2 (B) in head kidney and liver at 3, 8, 24 and 72 h after Viral Haemorrhagic Septicaemia virus infection (black bars). Asterisks above the bars indicate significant differences between each sampling point and the

corresponding control (MEM + 2% FBS + P/S-injected groups) (white bars), and asterisks over lines show statistical significant differences across the different sampling points ($p < 0.05$).

Figure 8. Expression of Hep-1 (A) and Hep-2 (B) in head kidney and liver at 3, 8, 24 and 72 h after Iron-dextran administration (black bars). Asterisks above the bars indicate significant differences between each sampling point and the corresponding control (PBS-injected groups) (white bars), and asterisks over lines show statistical significant differences across the different sampling points ($p < 0.05$).

Figure 9. Expression of Hep-1 (A) and Hep-2 (B) in head kidney and liver at 24 h after administration of three different Iron-dextran doses (black bars). Asterisks above the bars indicate statistical significant differences between each sampling point and the corresponding control (PBS-injected groups) (white bars) ($p < 0.05$).

Tables

Table 1. Primer sequences used in the ORF amplification and expression analysis.

	Gene	Primer	Sequence (5'-3')
ORF amplification	Hepcidin-1	Hepcidin Comp F1	CTCAAAATGAAGGCATTTCAG
		Hepcidin Comp R1	GAATCCTCAGAACTTGCAGC
	Hepcidin-2	AMP Comp F1	ATGAAGACTCTCACCGTTGCAG
		AMP Comp R1	CATGGCTGTTGGAGCAGGAATTC
q-PCR	Hepcidin-1	Hepc.-F3	CGAGTCACATCAGGCAGAAG
		Hepc.-R4	TCCTCAGAACTTGCAGCAGA
	Hepcidin-2	AMP prec2 F	ATGAAGACTCTCACCGTTGC
		AMP prec2 R	TTCTGTCTGTTACTCGGCATC
	EF1 alpha	T1-F2	GGAGGCCAGCTCAAAGATGG
		T1-R2	ACAGTTCCAATACCGCCGATT

A novel hepcidin-like antimicrobial peptide (Hep-2) was characterized in turbot
A. salmonicida and VHSV challenge affect the expression of both turbot hepcidins
Iron overload only induces the expression of Hep-1
A differential role of turbot hepcidins is suggested in this paper

Figure 1
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1 ATGAAGACTCTCACCGTTGCAGTCGCAGTGGCCGTCGTGCTCGCCTTC
1 M K T L T V A V A V A V V L A F
49 ATTTGGAT**K**CAGGAGAGCGCTGCCACATTCCACGGG|GCACAACAGCCG 85 bp
17 I W **M/I** Q E S A A T F H G A Q Q P
97 GAGGAGGCGGTGAGCAAT**R**AGGATCCAGCTGCTGATCCTCAGGAGACA
33 E E A V S N **K/E** D P A A D P Q E T
145 CCGGTGGACTCGTGGATG|ATGCCGAGTAACAGACAGAAGCGCGGCATG 132 bp
49 P V D S W M M P S N R Q K R G M
193 AAGTGCAAGTTTTGCTGCAACTGC**Y**GCAAC**W**T**K**AATGGCTGCGGCR**T**G
65 K C K F C C N C **C/R** N **F/I/L/M** N G C G M/V
241 TGCTGCGATTTCTGA
81 C C D F -

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A

B

Figure 8
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A

B

Figure 9
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