**Differential Expression of Duplicate Insulin-like Growth Factor-1 Receptors (igf1rs) in Medaka Gonads**

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**Abstract:** Insulin-like growth factor-1 receptors (igf1rs) play important roles in regulating development, differentiation, and proliferation in diverse organisms. In the present study, subtypes of medaka igf1r, igf1ra, and igf1rb were isolated and characterized. RT-PCR results showed that igf1ra and igf1rb mRNA were expressed in all tissues and throughout embryogenesis. Using real-time PCR, the differential expression of igf1ra and igf1rb mRNA during folliculogenesis was observed. The results of in situ hybridization (ISH) revealed that both of them were expressed in ovarian follicles at different stages, and igf1rb was also expressed in theca cells and granulosa cells. In the testis, both igf1ra and igf1rb mRNA were highly expressed in sperm, while igf1rb mRNA was also obviously detected in spermagonia. In addition, igf1ra mRNA was also present in Leydig cells in contrast to the distribution of igf1rb mRNA in Sertoli cells. Collectively, we demonstrated that differential igf1rs RNA expression identifies medaka meiotic germ cells and somatic cells of both sexes. These findings highlight the importance of the igf system in the development of fish gonads.

**Keywords:** igf1rs; gene expression; medaka; ovary; testis

1. Introduction

The igf signaling system is a growth factor complex containing ligands, receptors, and binding proteins, which exists in all vertebrates [1]. The role of the igf signaling system in growth regulation has been well established, and its important roles in gonadal development has been gradually explored [2]. Igf1 mRNA is expressed in spermatagonia and spermatocytes as well as Leydig cells in Nile tilapia (*Oreochromis niloticus*) [3]. Igf2 could influence germ cell proliferation in the testis of *Hypostomus garmani* [4]. Furthermore, the gonad-specific igf3 can regulate spermatogenesis and reproduction in teleosts, such as Nile tilapia and zebrafish (*Danio rerio*) [5,6]. Igf2bp3 deletion leads to abnormal germ plasm assembly and a reduction in the number of germ cells in zebrafish [7]. Furthermore, igf1r is also expressed in the testis and ovary, which indicates the important roles of igf1r in gonadal development and gametogenesis [8,9].

Igf1r is a cell surface receptor that belongs to the tyrosine kinase receptor superfam-ily, which is expressed in diverse tissues of organisms [10]. In mammals, there is only one igf1r. Igf1r has been detected in sperm of humans, and its levels is positively correlated with sperm concentration [11]. Besides, igf1r is also expressed in ovaries of alpaca (*Vicugna pacos*), including follicles, granulosa, and theca interna cells [12]. In the cultured testicular cells, igf1r is highly expressed in Sertoli cells but also in spermatogonia and primary spermatocytes [13].
Due to whole genome duplication, fish have two subtypes of \textit{igf1r}, \textit{igf1ra} and \textit{igf1rb} \cite{14}, and both of them play important biological functions. \textit{Igf1ra} and \textit{igf1rb} mRNA are highly expressed in gonads during vitellogenesis and spermatogonia proliferation in \textit{Pampus argenteus} \cite{15}. In adult zebrafish, \textit{igf1ra} and \textit{igf1rb} have distinct expression patterns, and the relative abundance of \textit{igf1ra} and \textit{igf1rb} is different in tissues \cite{16}. It has been shown that inhibition of the \textit{igf} signal pathway by knocking down \textit{igf1rb} in the embryo of zebrafish can result in mis-migration and apoptosis of primordial germ cells (PGCs) \cite{17}. Therefore, more attention has been focused on their cellular localization and biological activity in fish gonads. However, the role of \textit{igf1r} in reproduction and gonadal development has rarely been studied, especially in fish.

\textit{Medaka} (\textit{Oryzias latipes}) is a good model and has been widely used in investigating developmental biology \cite{18} and stem cell biology \cite{19,20}. In addition, the primordial germ cells specification \cite{21}, migration \cite{22}, and sex-determination mechanism \cite{23,24} have been systematically explored. Previously, we have investigated that \textit{igf1} is present in meiotic germ cells and somatic cells of both sexes in medaka \cite{25}. \textit{Igf2} is associated with self-renewal of the embryonic stem cell \cite{26}. \textit{Igf3} expression occurs both in germ cells and somatic cells in the ovary \cite{27}. In the present study, to figure out the distribution and differences between \textit{igf1ra} and \textit{igf1rb}, the temporal and spatial expression patterns of their mRNAs in gonads, developing embryos, and follicles at different developmental stages were investigated. The expression patterns between \textit{igf1ra} and \textit{igf1rb} mRNAs in the gonads were also carried out by in situ hybridization studies. Our findings highlight the potential roles of the \textit{igf} system in the reproduction and development of medaka and teleosts.

2. Materials and Methods

2.1. Fish and Embryos

Animal experiments were conducted strictly following the requirements of the Committee for Laboratory Animal Research at Shanghai Ocean University. Medaka was maintained in glass tanks with a water temperature of 26 °C, and an automatic photoperiod of 14 h light/10 h dark cycle was set. The developmental stage of medaka embryos has been previously described \cite{28}.

2.2. Isolation of Ovarian Follicles

The developing stages of the ovary were determined based on the original definition as described previously \cite{25}. The ovaries were dissected from the anesthetized female medaka. As follicles have different stages, the same developmental stages were manually collected together and used for subsequent experiments: primary growth (stage I, below 0.1 mm in diameter), pre-vitellogenic (stage II, about 0.30 mm), early vitellogenic (stage III, about 0.40 mm), mid-vitellogenic (stage IV, about 0.50 mm), and fully-grown (stage V, about 0.65 mm).

2.3. RNA Extraction

Total RNA of adult tissues and embryos at different developmental stages of medaka was extracted by a TRIzol reagent (Invitrogen, Carlsbad, CA, USA). The NanoDrop 2000 Spectrophotometer (Thermo Fisher Scientific, Waltham, MA, USA) was used to detect the RNA quality and quantity. Furthermore, the integrity of RNA was verified by 1% agarose gel stained with nucleic acid dyes. Then, the cDNA was synthesized according to M-MLV reverse transcriptase (Takara, Shiga, Kusatsu, Japan) with an oligo(dT)\textsubscript{18} primer.

2.4. Cloning and Sequence Analysis of Medaka \textit{Igf1r}

By searching the NCBI Gene database (https://www.ncbi.nlm.nih.gov/gene/ accessed on 1 March 2020), two computational predicted cDNAs encoding medaka \textit{igf1ra} (Gene ID: 101173298) and \textit{igf1rb} (Gene ID: 101163560) were obtained, respectively. To verify the accuracy of the two sequences, the Open Reading Frame (ORF) of \textit{igf1ra} and \textit{igf1rb} was cloned and sequenced. Afterwards, the \textit{igf1ra} and \textit{igf1rb} putative proteins were aligned.
with the igf1r orthologs from other examined organisms by using Vector NTI Advance 11.5 software (Thermo Fisher Scientific). The phylogenetic tree was based on the MEGA X program with the neighbor-joining (NJ) method [29]. All the primers used in the present study are listed in Table 1.

Table 1. Sequences of primers used in the present study.

| Primer       | Sequence (5’ to 3’ Direction) | Purpose         |
|--------------|--------------------------------|-----------------|
| igf1ra-F     | ATGGACATCCGAAGAC               | RT-PCR          |
| igf1ra-R     | TCGTCTGATACGGCCAGTGGTTG       | RT-PCR          |
| igf1rb-F     | TGCTTCGGGAAATGGGCTCC          | RT-PCR          |
| igf1rb-R     | TAGATGTCAAGTCCCTTTTGAT        | RT-PCR          |
| igf1ra-F1    | ATGGGACGCTGACCTTGTGTG         | CDS cloning     |
| igf1ra-R1    | TCGTCTGAGTGTACATCTC           | CDS cloning     |
| igf1ra-F2    | ATGTACACTCCAGAGAAGGG          | CDS cloning     |
| igf1ra-R2    | TGCTTCGGGAAATGGGCGGG          | CDS cloning     |
| igf1ra-F1    | ATGGGACGCTGACCTTGTGTG         | CDS cloning     |
| igf1ra-R1    | TCGTCTGAGTGTACATCTC           | CDS cloning     |
| igf1ra-F2    | ATGTACACTCCAGAGAAGGG          | CDS cloning     |
| igf1ra-R2    | TGCTTCGGGAAATGGGCGGG          | CDS cloning     |
| Actin-F      | TTCAACAGCTGCAGAGACT           | Internal control|
| Actin-R      | CCTCAAATCCAGAAGACT           | Internal control|

2.5. RT-PCR and Real-Time PCR Analysis

The expression of igf1r isoforms was detected by RT-PCR amplification using the primers in Table 1, and β-actin was used for calibration. PCR was performed for 35 cycles, and the reaction system contained 1 µL of cDNA template, 12.5 µL of 5 U/µL Premix Taq (EX Taq version) (Takara), and 0.5 µL of 10 mM each of the forward and reverse primers, and deionized water was added to replenish the total volume to 25 µL. The reaction procedures were as follows: 95 °C for 10 s, annealing at 58 °C for 10 s, and extension at 72 °C for 1 min. The PCR products were then detected on a 1.5% agarose gel stained with nucleic acid dyes and analyzed by a bio-imaging system (Bio-Rad, Hercules, CA, USA).

Real-time PCR (qPCR) was performed under the CFX96™ Real-Time System (Bio-Rad) using the SYBR Green PCR Master Mix Kit (Takara). Using β-actin as calibration, the relative abundance of igf1ra and igf1rb mRNA was determined using the 2−ΔΔct, as described previously [30]. The data were presented as the mean ± SEM (n = 3). Statistical analyses were evaluated using a one-way ANOVA (p < 0.05) in GraphPad Prism 7 software.

2.6. RNA In Situ Hybridization

Sections in situ hybridization (SISH) were carried out as described previously [31]. Briefly, the gonads were fixed in 4% paraformaldehyde and then dehydrated gradiently with 20% and 30% sucrose. Next, the gonads were soaked with an embedding agent, Optimal Cutting Temperature (O.C.T., Sakura, Torrance, CA, USA), and subjected for sections with a freezing microtome (Leica, Wetzlar, Germany). To synthesize probes, the partial cDNA sequences of igf1ra and igf1rb obtained by PCR were inserted into the pGEM-T vector and sequenced for verification. The plasmids were then linearized with an appropriate restriction enzyme for the synthesis of probes by using the DIG or FITC RNA Labelling Kit (Roche, Basel, Switzerland). RNA of SISH was stained with BCIP/NBT and Fluorescent in situ hybridization (FISH) was carried out by using the (TSA™) Plus Fluorescence Systems according to the product manual (Life Technologies, Carlsbad, CA, USA). The nucleus was stained with DAPI.
2.7. Microscopic Observation

Microscopy was performed as described [32]. In brief, microscopic observation and micrographs of gonadal sections were taken on a Nikon Ds-Ri2 camera (Nikon, Tokyo, Japan).

3. Results

3.1. Cloning and Sequence Analysis of Medaka Igf1r

PCR was employed to amplify the sequences of *igf1r* derived from tissues and embryo samples. The *igf1ra* ORF was obtained by TA cloning, which was 4197 nt and encoded 1398 amino acid residues (GenBank accession no. BK061359) (Figure S1). However, the *igf1rb* ORF was 4224 nt for 1407 amino acid residues (GenBank accession no. BK061360) (Figure S2). The IGF1Rs’ alignment showed the intracellular protein tyrosine kinase domain on the β-subunit of Igf1r (Figure 1). Igf1ra protein and Igf1rb protein of medaka are highly similar to those of zebrafish, with 74% and 71% identity, respectively, according to the Igf1rs multiple sequence alignment (Figure S3). Besides, the phylogenetic tree showed that compared with medaka Igf1ra, zebrafish Igf1rb is much closer to medaka Igf1rb (Figure 2). Such a divergence was also observed for Takifugu rubripes and Salmo salar (data not shown). Furthermore, although both Igf1ra and Igf1rb existed in different regions of different chromosomes in other species, they all showed strong chromosome synteny (Figure S4).

3.2. RT-PCR Analysis of Igf1r RNA Expression

The results of the RT-PCR showed that *igf1ra* and *igf1rb* were expressed in all adult tissues and embryos. Evidently, the expression of *igf1ra* and *igf1rb* was lower in the kidney, liver, and gut, in comparison with the eye, brain, ovary, and testis (Figure 3A). Besides, these two genes were similarly expressed during embryogenesis (Figure 3B). Furthermore, qPCR was performed to further analyze the expression profiles of the two *igf1r* types at different stages of folliculogenesis. The level of *igf1ra* increased from PG (stage I) to PV (stage II) and then decreased slowly, and it was hardly expressed in FG (stage V) (Figure 3G). The expression of *igf1rb* increased from the PG stage, peaked in EV (stage III), and weakened afterward (Figure 3H). These data indicate that *igf1r* was dynamically and differentially expressed during folliculogenesis in medaka.

![Figure 1.](image) **Figure 1.** Multiple sequence alignment of medaka Igf1rs. The alignment sequences show the intracellular protein tyrosine kinase domain. Conserved regions between species are highlighted. The length and percentage identity values of Igf1rs homologs are given at the end of the alignment.
Figure 2. Phylogenetic tree of Igf1r. The insulin receptor (InsR) served as the out-group. Bootstrap values are given, and the bar indicates number of substitutions per site. Accession numbers are after the organism. Igf1rs from different species are clustered together, indicating that generation of igf1r and insr took place in early vertebrate evolution.

Figure 3. Expression of igf1ra RNA and igf1rb RNA. (A and B) RT-PCR analysis of medaka igf1ra and igf1rb in adult tissues (A) and developing embryos (B). (C–F) Ovarian and testicular cryosections using antisense igf1ra and igf1rb probes and the signals were visualized by chromogenic staining. (G,H) qPCR results of igf1ra and igf1rb at different stages of follicles. I–V, stages of oocytes; sg, spermatogonia; sc, spermatocytes; st, spermatids; sm, sperm; gc, granulosa cells; tc, theca cells; se, Sertoli cells; le, Leydig cells. Scale bars, 100 µm.
3.3. Gonadal Expression of Igf1ra RNA and Igf1rb RNA by ISH

To better understand the subcellular distributions of igf1r, a chromogenic SISH was performed on cryosections. In the ovary, igf1ra RNA and igf1rb RNA were found in the cytoplasm of oocytes from stages I–IV (Figure 3C,D). Furthermore, igf1rb was also found in theca cells and granulosa cells, while igf1ra was absent (Figure 3D). In the testis, the igf1ra mRNA was highly expressed in sperm at the later stage of spermatogenesis (Figure 3E). Remarkably, igf1rb mRNA was detected in spermatogonia and sperm and was weakly detected in spermatocytes as well as spermatids, which was significantly different from the expression of igf1ra mRNA (Figure 3F). Furthermore, a positive signal for igf1ra was also found around Leydig cells, whereas the signal of igf1rb existed around Sertoli cells (Figure 3E,F). On the contrary, both of the sense probes, as controls, got no signal above the background (data not shown).

3.4. Ovarian Differential Expression of Igf1ra and Igf1rb RNAs by Fluorescence ISH

To further accurately identify the RNA expression of igf1r, the co-distribution of igf1rb and vasa was carried out by FISH. Vasa is a well-studied gene in medaka and other species, which is restrictively expressed in the germ cells of both sexes [33]. In the ovary, the vasa signal was expressed obviously in pre-meiotic oocytes and was decreased with the process of oogenesis (Figure 4A). Conversely, the igf1rb signal was easily found in oocytes from stages I–IV (Figure 4B). Notably, the igf1rb positive signal was also obvious in the granulosa cells and theca cells at later folliculogenesis (Figure 4C,D).

Figure 4. Expression of igf1rb RNA and vasa RNA in the ovary. FISH on ovarian cryosections using antisense RNA probes and the signals were visualized by fluorescence staining. The vasa RNA was stained in red, and the igf1rb was stained in green. Nuclei were stained in blue with DAPI. (A,B) Different stages of oocytes (I–V), granulosa cells and theca cells were indicated by sticks. (C,D) The merges of vasa with igf1rb and vasa with igf1rb and DAPI. I–V, stages of oocytes; gc, granulosa cells; tc, theca cells. Scale bars, 25 µm.
We compared the RNA expression of *igf1ra* with *igf1rb* by dual-color FISH. Conforming to their observations by chromogenic staining, both *igf1ra* and *igf1rb* RNAs were expressed in oocytes from stages I to IV (Figure 5A,B). Furthermore, *igf1rb* was detected abundantly in somatic cells, including granulosa cells, as well as theca cells at later developmental oocytes, while *igf1ra* was not detected in these somatic cells (Figure 5C,D).

Figure 5. Expression of *igf1ra* RNA and *igf1rb* RNA in the ovary. FISH on ovarian cryosections using antisense RNA probes and the signals were visualized by fluorescence staining. The *igf1ra* RNA was stained in red, and the *igf1rb* was stained in green. Nuclei were stained in blue with DAPI. (A, B) Different stages of oocytes (I–V), granulosa cells and theca cells were indicated by sticks. (C, D) Merges of *igf1ra* with *igf1rb* and *igf1ra* with *igf1rb* and DAPI. I–V, stages of oocytes; gc, granulosa cells; tc, theca cells. Scale bars, 25 µm.

3.5. Testicular Differential Expression of *Igf1ra* and *Igf1rb* RNAs by Fluorescence ISH

In the next step, we compared the RNA expression of *igf1rb* with *vasa* by dual color FISH. In the testis, the *vasa* signal was strong in spermatogonia, and with the progress of spermatogenesis, the intensity of the *vasa* signal was decreased until it disappeared in sperm (Figure 6A). In contrast, the *igf1rb* mRNA was abundantly expressed in the spermatogonia and sperm, and it was relatively low in spermatocytes as well as in spermatids (Figure 6B). Surprisingly, a positive signal for *igf1rb* was also detected between the cyst and the cyst of the testis structure (Figure 6C,D), which is generally thought to be the location of Sertoli cells [34].
Figure 6. Expression of *vasa* RNA and *igf1rb* RNA in the testis. After hybridization with antisense *vasa* and *igf1rb* RNA probes, the signals were visualized by fluorescence staining. Nuclei were stained blue by DAPI. (A–D) Merges of *vasa* with DAPI, *igf1rb* with DAPI, *vasa* with *igf1rb*, and *vasa* with *igf1rb* and DAPI. Sertoli cells were indicated by arrows. *Vasa* and *igf1rb* showed significantly different expression patterns. The *vasa* signal peaked in spermatogonia and then gradually decreased until it disappeared in sperm. Conversely, the *igf1rb* was obviously detected in spermatogonia, sperm, and Sertoli cells. sg, spermatogonia; sc, spermatocytes; st, spermatids; sm, sperm; se, Sertoli cells. Scale bars, 25 \( \mu \)m.

Then, a dual-color FISH was performed to precisely compare the expression of *igf1ra* and *igf1rb* RNA. Results indicated that *igf1ra* mRNA was richly expressed in sperm, but it was not in other stages of spermatogenic cells (Figure 7A,C). In contrast, *igf1rb* mRNA was expressed in almost all stages of spermatogenesis, especially spermatogonia and sperm. (Figure 7B–E). Furthermore, *igf1ra* mRNA and *igf1rb* mRNA were also found in somatic cells, with *igf1ra* expressed in the intertubular space where the Leydig cells usually located, while *igf1rb* mRNA delineated the germinal cysts called Sertoli cells (Figure 7A–F).
Figure 7. Expression of igf1ra RNA and igf1rb RNA in the testis. Adult testicular cryosections were hybridized to antisense RNA probes and the signals were visualized by fluorescence staining. Nuclei were stained blue by DAPI. (A,B) Lower magnification view showing different stages of spermatogenesis. Sertoli cells and Leydig cells were indicated by arrows. (C,D) Merges of igf1ra with igf1rb and igf1ra with igf1rb and DAPI. (E,F) Larger Magnification of panels D (white frame), highlighting the different cells. The igf1ra signal exists in sperm and Leydig cells. Notably, the igf1rb signal exists in spermatogonia, sperm, as well as Sertoli cells. se, Sertoli cells; le, Leydig cells; sm, sperm; sc, spermatocytes; sg, spermatogonia. Scale bars, 25 µm.

4. Discussion

In contrast to a single igf1r in mammals, many fish species have two igf1r paralogous isoforms, igf1ra and igf1rb, such as zebrafish [16], gilthead seabream (Sparus aurata) [9], and Epinephelus coioides [35]. According to the present study, medaka also has two igf1r subtypes. A sequence comparison and phylogenetic tree of the medaka two Igf1rs protein sequences with other vertebrates indicated that the receptors were highly conservative in the process of vertebrate evolution. The structure of the medaka igf1rs is most similar to that of zebrafish igf1rs [16]. Not only is their sequence identity of medaka igf1rs and zebrafish igf1rs greater than 60%, but structural motifs are nearly conserved in the two medaka igf1rs, such as the ATP-binding site, ligand-binding region, tyrosine kinase domain, autophosphorylation site, and IRS-I docking site. Igf1r is composed of extracellular α-subunits.
and transmembrane-spanning β-subunits, which contain cytoplasmic tyrosine kinase activity [36]. Activated igf1r phosphorylates specific reaction components, including IRS-1, IRS-2, and SRC homology collagen, and it regulates downstream responses through the phosphatidylinositol 3-kinase (PI3K)/AKT and mitogen-activated protein kinase (MAPK) pathways [1,37].

Both igf1r mRNAs were extensively expressed in different tissues in the adult medaka, which was consistent with reports from other teleosts [38]. The rich expression of two igf1r mRNAs in the eye, brain, and gonad generally agreed with the previous reports of igf1 mRNA expression levels in these tissues of medaka [25]. Similar to Paralichthys olivaceus [39], the igf1r involvement in reproduction and development was consistent with the functions of the igf system, such as igf1 promoted spermatogenesis, oocyte maturation, and steroidogenesis [40,41], and igf3 maintained the differentiation of ovary [42]. Furthermore, the expression of igf1rs mRNAs in the embryos at different developmental stages was demonstrated. Our present study further supports the potential functions of the igf system in fish.

In adult medaka, igf1ra and igf1rb mRNAs were detected in the oocytes at different stages; however, the expression levels were significantly different. During folliculogenesis, the level of igf1ra mRNA in mature follicles was extremely low in comparison with igf1rb, suggesting that igf1ra may play a minor role in mature follicles, while igf1rb plays a dominant role. Furthermore, through co-localization with the germ cell marker gene vasa [43], it was found that igf1rb RNA was highly expressed in many somatic cells, including theca cells and granulosa cells at later folliculogenesis, while igf1ra was absent. Similarly, igf1r was also found in theca cells and granulosa cells from the ovary of Oncorhynchus kisutch [44], gilthead seabream [9], alpaca [12], and mice [45]. Based on these findings, it indicates that the two igf1rs of medaka share some similar biological functions, which means that they play the same roles in many body activities. Meanwhile, it predicts that both of them also have their own unique roles in promoting individual growth and gonadal development. Igf1rb may also be involved in hormone production in the ovary of medaka according to previous studies that theca cells and granulosa cells regulate steroid hormone production [44].

In this study, the two subtypes of igf1r were differentially expressed in the ovary as well as in the testis. Through the co-localization of vasa and igf1rb, as well as igf1ra and igf1rb, we found that both types of igf1r were expressed in sperm. In addition, igf1rb was also expressed in spermatogonia. Furthermore, it seemed that igf1ra preferred to express in Leydig cells, whereas igf1rb preferred to express in Sertoli cells, suggesting that the two isoforms may have different functions during spermatogenesis. These results agreed with the reported presence of igf1 receptors in somatic cells of zebrafish [46], rainbow trout (Oncorhynchus mykiss) [13], and gilthead seabream [9]. In recent research on zebrafish, it has been found that igf1rb was expressed in spermatogonia and could mediate igf3 to activate the β-catenin-dependent signal pathway to regulate spermatogenesis [46]. Moreover, it was shown that igf1 and igf2 interact with igf1r, causing the receptor activation and regulating organism growth, development, and reproduction [36]. Therefore, we speculate that the two subtypes of igf1r are essential for spermatogenesis, while igf1rb is also involved in the growth and proliferation of spermatogonia in the early stage. Overall, igf1rs play critical roles in the development of fish gonads, which is worthy of further research and provides a reference for other fish. Besides, the igf system is conserved in diverse species, and the interaction among ligands, receptors, and binding proteins of the system ensures that various life activities are carried out on the normal track, so the follow-up study of their co-localization is necessary for the further investigation of the igf system.

In summary, we demonstrated the differential expression of two igf1r subtypes in the adult gonads and embryos of medaka. The distinct expression patterns of the two subtypes of igf1r indicate that they play different roles in gonadal and embryonic development. Overall, the present study provides conclusive evidence for the potential roles of igf1r in gonadal development and gametogenesis in fish, as well as a reference for further research.
**Supplementary Materials:** The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/life12060859/s1. Figure S1: Nucleotide and deduced amino acid sequence of medaka igf1ra; Figure S2: Nucleotide and deduced amino acid sequence of medaka igf1rb; Figure S3: Multiple sequence alignment of IGF1 receptors; Figure S4: Chromosome synteny diagrams for Igf1rs.

**Author Contributions:** Conceptualization, Experimental designing, and Project administration: M.L.; W.W.; Y.Z. (Yuli Zhao); C.Y.; Methodology: W.W.; Y.Z. (Yefei Zhu); C.Y.; Supervision: M.L.; W.W.; Writing—original draft: W.W.; Y.Z. (Yuli Zhao); Writing—review and editing: W.W.; M.L.; W.Z. All authors have read and agreed to the published version of the manuscript.

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**Institutional Review Board Statement:** The study was conducted according to the guidelines of the Declaration of Helsinki, and it was approved by the Shanghai Ocean University Animal Care and Use Committee, with approval number SHOU-2021-118.

**Informed Consent Statement:** Informed consent was obtained from all subjects involved in the study.

**Data Availability Statement:** Not applicable.

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**Abbreviations**

Igf, insulin-like growth factor; RT-PCR, reverse transcription-polymerase chain reaction; ISH, in situ hybridization; FISH, fluorescence in situ hybridization; P13K, phosphoinositide 3 kinase; aa, amino acid residues; nt, nucleotide.

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