Thyroid hormone actions on male reproductive system of teleost fish

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Abstract

Thyroid hormones (THs) play important roles in the regulation of many biological processes of vertebrates, such as growth, metabolism, morphogenesis and reproduction. An increasing number of studies have been focused on the involvement of THs in the male reproductive system of vertebrates, in particular of fish. Therefore, this mini-review aims to summarize the main findings on THs role in male reproductive system of fish, focusing on sex differentiation, testicular development and spermatogenesis. The existing data in the literature have demonstrated that THs exert their roles at the different levels of the hypothalamic-pituitary-gonadal (HPG) axis. In general a positive correlation has been shown between THs and fish reproductive status; where THs are associated with testicular development, growth and maturation. Recently, the molecular mechanisms underlying the role of THs in spermatogenesis have been unraveled in zebrafish testis. THs promote germ cell proliferation and differentiation by increasing a stimulatory growth factor of spermatogenesis produced by Sertoli cells. In addition, THs enhanced the gonadotropin-induced androgen release in zebrafish testis. Next to their functions in the adult testis, THs are involved in the gonadal sex differentiation through modulating sex-related gene expression, and testicular development via regulation of Sertoli cell proliferation. In conclusion, this mini-review showed that THs modulate the male reproductive system during the different life stages of fish. The physiological and molecular mechanisms showed a link between the thyroid and reproduction, suggesting a possibly co-evolution and interdependence of these two systems.

1. Introduction

Thyroid hormones (THs) have been known to be involved in many biological processes, such as growth, morphogenesis, skin pigmentation, osmoregulation, and reproduction (Blanton and Specker, 2007). An increasing number of studies has shown that THs exert an important role in male reproductive system of vertebrates. In the past two decades, experimental and clinical studies have shown the involvement of THs in the testicular development and function in mammals (Wagner et al., 2008). Similar functions were reported in other vertebrates, such as fish, which is the focus of this mini-review. The interaction between THs and fish reproductive system was reviewed before by Cyr and Eales (1996), Blanton and Specker (2007), Habibi et al. (2012) and Flood et al. (2013). While there are significant variations in the nature of interaction between THs and reproduction in different fish species, it would appear that THs may exert stimulatory and/or inhibitory actions by different mechanisms (Habibi et al., 2012). In this context, this mini-review will address some of the direct and indirect actions of THs in male reproductive system of teleost fish, highlighting the main findings on sex differentiation, testicular development and adult tests.

Abbreviations: TH, Thyroid hormone; THs, Thyroid hormones; T3, 3,5,3′-triiodothyronine; HPT, Hypothalamic-pituitary-thyroid axis; HPG, Hypothalamic-pituitary-gonadal axis; CRH, Corticotropin-releasing hormone; CRF, corticotropin-releasing factor; GnhH, Gonadotropin inhibitory hormone; GnRH, Gonadotropin releasing hormone; DIOs, Deiodinases; T4, THs in the testicular development and function in mammals (Wagner et al., 2013). While there are significant variations in the nature of interaction between THs and reproduction in different fish species, it would appear that THs may exert stimulatory and/or inhibitory actions by different mechanisms (Habibi et al., 2012). In this context, this mini-review will address some of the direct and indirect actions of THs in male reproductive system of teleost fish, highlighting the main findings on sex differentiation, testicular development and adult tests.

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Thyroid function is activated by the hypothalamic-pituitary-thyroid axis (HPT), which has been reviewed in many vertebrates, including fish (Fig. 1A,B) (Brown and Cai, 2007; Blanton and Specker 2007; and Carr and Patiño, 2011). A particular characteristic of the HPT in fish is the non-involvement of thyrotrop releasing hormone (TRH) as occurs in mammals (Larsen et al., 1998). On the other hand, the pituitary secretion of TSH (Thyroid-stimulating hormone) seems to be dependent on corticotropin-releasing hormone (CRH, or known as corticotropin-releasing factor – CRF) (Fig. 1A,B) (Larsen et al., 1998), similarly as observed in amphibians, reptiles and birds (De Groef et al., 2003; Castañeda Cortés et al., 2014). The secreted TSH stimulates the synthesis and release of THs by thyroid follicles. In fish, the thyroid follicles are not encapsulated in an organ, but they are dispersed among the afferent branchial arterioles of the ventral region of pharynx (Cyr and Eales, 1996; Carr and Patiño, 2011). The l-Thyroxine (T4) is the main TH secreted under normal circumstances, while T3, known as the active hormone, is mostly produced by extrathyroidal sources from T4 conversion (Basset et al., 2003). The conversion of T4 into T3 is catalyzed by two distinct selenoenzymes, type 1 and type 2 deiodinases (DIO1 and DIO2, respectively) (Fig. 1A; Supplemental Fig. 1) (for review see Orozco et al., 2012).

The active form (T3) regulates nuclear gene expression by binding to the TRs in the regulatory regions of target genes, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003). In anamniotes (fish and amphibians), TR metabolises T3 and T4 into T2 and T1 respectively, activating or repressing their transcription (Basset et al., 2003).
terminal nerve salmon GnRH mRNA in sexually immature males (Fig. 1B) (Parhar et al., 2000). This fact is consistent with the hypothesis that TH, by suppressing terminal nerve GnRH expression, promotes inhibition of sexual maturation in tilapia males (Parhar et al., 2000).

More recently, studies have been focused on GnIH (Gonadotropin inhibitory hormone), which is a newly discovered hypothalamic neuropeptide that inhibits the pituitary gonadotropin synthesis and release (Tsutsui et al., 2018). Interestingly, it has been shown that mammalian GnIH neurons express TRα and TRβ (Tsutsui et al., 2018). In addition, female mice with thyroid dysfunction showed higher levels of GnIH, and presented lower circulating levels of gonadotropins and estradiol, which might be associated with the delayed puberty (Tsutsui et al., 2018). These data suggest a possible role of THs on regulating the GnIH synthesis and release in vertebrates (Fig. 1B).

3. Fish reproduction

The relationship between thyroid function and reproduction has been explored for many years in different species. In fish, it is established that reproductive strategies vary greatly among the species (Eales, 2006), and it has become clear that the nature of interaction between TH and reproduction is species-specific (Habibi et al., 2012). Thyroidal function usually increases during early gonadal development in fish, and it is maintained or enhanced during the period of reproduction, and commonly decreased during or after spawning (Cyr and Eales, 1996). Similar function is seen in reptiles, birds and mammals, where TH administration induced testicular growth in thryoidectomized or short photoperiod-induced males (Haldar-Misra and Thapliyal, 1981; Yoshimura et al., 2003; Freeman et al., 2007). Temporal relationship between thyroidal and reproductive status has been examined in some seasonal species of fish, in particular, non-salmonids and salmonids (for review see Cyr and Eales, 1996). In non-salmonids, a general positive correlation between thyroid activity and reproductive status has been seen among the species. On the other hand, thyroid function in Salmonidae is high in the early stages of gonadal maturation but decreases as vitellogenesis and testis development proceed. However, T3 plasma levels are higher in males than females just prior to spawning and may even increase (for review see Cyr and Eales, 1996). Table 1 summarizes selected data from the literature about the relationship between thyroid function and reproduction in non-mammalian and mammalian vertebrates. Therefore, the summarized data in the literature suggest that the basic function for THs is associated with gonadal maturation in fish as well as in other vertebrates.

4. Sex differentiation

THs have considerable influence on the sexual ontogeny of male vertebrates, through direct interactions with genes involved in sex determination and gonadal development along the HPG axis (Flood et al., 2013). The role of THs in the establishment of gonadal sex phenotype has been previously examined in amphibians and reptiles. For example, chronic treatment with TH synthesis inhibitors (perchlorate or thiouracil) during larval development resulted in a feminization effect in Xenopus laevis (Goleman et al., 2002; Hayes, 1997). In another study, premetamorphic tadpoles (Nieuwkoop and Faber stage 52–54) were exposed to exogenous T3 for 48 h and resulted in a concentration-dependent decrease of estrogen receptor beta isoform (ERβ) expression at concentrations of 5 and 50 nM T3. The expression of androgen-related genes also changed and significantly increased following treatment with T3 in relation to the control. Treatment with T3 also led to changes in ar (androgen receptor, 1.4–1.8 fold), srdα2 (steroid 5α-reductase 1; 1.5–1.9 fold) and srdβ2 (steroid 5α-reductase 2; 1.8–1.9 fold) in tropical clawed frogs (Duarte-Guterman and Trudeau, 2011). In reptiles, hyperthyroid treatment decreased cyp19α1 transcripts (gonadal aromatase), and consequently estradiol levels, inducing therefore more male offspring in a TSD (Temperature-dependent sex determination) turtle (Trachemys scripta) (Sun et al., 2016).

In fish and amphibians, an evidence that THs are involved in gonadal sex differentiation is the different expression of deiodinases and thr between males and females, as reported in striped parrotfish, Scarus iseri (Johnson and Lema, 2011) and tropical clawed frog Silurana tropicalis (Duarte-Guterman and Trudeau, 2011). In addition, An et al. (2010) observed a decrease in gonadal thrα expression in protandrous

Table 1

<table>
<thead>
<tr>
<th>Species</th>
<th>Treatment</th>
<th>Effects</th>
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<tbody>
<tr>
<td>Teleosts</td>
<td>Stimulated thyroid function</td>
<td>Coincides with sexual maturation</td>
</tr>
<tr>
<td></td>
<td>Elevated TH plasma levels</td>
<td>Occurs during gonadal maturation</td>
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<tr>
<td></td>
<td>Decreased TH plasma levels</td>
<td>Observed at spawning</td>
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<td></td>
<td>Goitrogen exposure</td>
<td>Associated with low egg production; low gonadal steroid levels in blood</td>
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<td></td>
<td>Presence of goiters in salmon from Great Lakes</td>
<td></td>
</tr>
<tr>
<td>Amphibians</td>
<td>Exogenous TH</td>
<td>No effect or mild enhancement of gonadal maturation</td>
</tr>
<tr>
<td></td>
<td>Surgical or chemical thyroidectomy of frogs</td>
<td>Prevents ovulation</td>
</tr>
<tr>
<td></td>
<td>Treatment with thyroxine of thyroidectomized frogs</td>
<td>Enhances action of estrogen on vitellogenesis</td>
</tr>
<tr>
<td></td>
<td>Treatment of anuran liver with TH</td>
<td>Not associated with reproductive activities</td>
</tr>
<tr>
<td></td>
<td>Thyroid function</td>
<td>Prevents testicular development</td>
</tr>
<tr>
<td></td>
<td>Goitrogen treatment of tadpoles (anurans)</td>
<td>Coincides with gonadal growth in males and females</td>
</tr>
<tr>
<td></td>
<td>Elevated thyroxine in the blood of salamanders</td>
<td></td>
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<tr>
<td>Reptiles</td>
<td>Thyroxine treatment</td>
<td>Atrophy of steroid secreting cells in testes</td>
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<tr>
<td></td>
<td>Decreased plasma levels and thyroid function</td>
<td>Associated with peak reproductive and high androgen levels</td>
</tr>
<tr>
<td></td>
<td>Elevated thyroxine levels in blood</td>
<td>Decreased androgen levels in males</td>
</tr>
<tr>
<td>Birds</td>
<td>TH treatment</td>
<td>Induced testicular growth of thyroidectomized</td>
</tr>
<tr>
<td></td>
<td>Thyroidectomized males</td>
<td>Testicular regression</td>
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<tr>
<td>Mammals</td>
<td>Exogenous T3</td>
<td>Induced gonadal growth in short-day hamsters</td>
</tr>
<tr>
<td></td>
<td>Exogenous T3</td>
<td>Delayed testicular regression in short photoperiod-induced males</td>
</tr>
</tbody>
</table>

Adapted from Norris (2006).

* TH, Thyroid Hormones.
black porgy (*Acanthopagrus schlegeli*) following male-to-female sex reversal. During zebrafish metamorphosis and early gonadal development, treatment with perchlorate (a well-known thyroid synthesis inhibitor), significantly skewed the sex ratio toward females (Fig. 2A) (Mukhi et al., 2007). However, the co-treatment with exogenous T4 not only blocked the feminizing effect of perchlorate but also overcompensated by skewing the ratio toward males. In another study, Sharma and Patiño (2013) showed that either perchlorate or methimazole (an antithyroid drug) induced feminization in zebrafish larvae at 45 dpf (days post-fertilization). Interestingly, the feminizing effect was still present in the perchlorate treatment, but not anymore in methimazole treatment at 60 dpf, suggesting that the effect disappeared as fish regained the euthyroid conditions. The conclusion is that methimazole does not cause male-to-female sex reversal but simply delays the ovary-to-testis transformation in presumptive males of this species (Fig. 2C). It is worth emphasizing that studies using antithyroid drugs can provide a powerful tool for understanding TH function but only with an appropriate control due to the extrathyroidal effects these drugs can have. A rigorous study requires a follow up to determine if those effects can be nullified or reversed by simultaneous administration of sufficient TH to regain euthyroid condition.

Interestingly, and more recently, Sharma et al. (2016) evaluated the molecular mechanisms involved in T4-induced masculinization in zebrafish larvae. These authors demonstrated that T4 treatment decreases cyp19a1 (aromatase) and erb (estrogen receptor beta) and increasing amh (anti-Müllerian hormone). T4 increases the ratio of females. (Fig. 2B).

5. Testicular development

In rodents, THs affect Sertoli cell proliferation and differentiation (Fig. 3A), as well as the onset of adult Leydig cell formation and function during neonatal-prepubertal period (Teerds et al., 1998; Ariyaratne et al., 2000). Induced-hypothyroidism in the neonatal rat impaired testicular growth, germ cell maturation, seminiferous tubule formation and other differentiation processes in the testis (França et al., 1995; Maran et al., 2002). Induced-neonatal hypothyroidism has also been shown to increase Sertoli cell proliferative window, which consequently increased Sertoli number and the Sertoli cell support capacity for germ cells, reflecting directly on the elevated spermatogenic yield and testis size/weight. On the other hand, Sertoli cell differentiation is delayed in the hypothyroidism model, increasing their proliferative window. In consequence, there is an elevated number of adult Sertoli cell which leads to higher spermatogenesis and testis size/weight. SC: Sertoli cell; SZ: spermatozoa. (B) Effects of hypothyroidism on testicular development in zebrafish. PTU-induced hypothyroidism increased Sertoli cell proliferation and Sertoli cell capacity reflecting directly on higher number of germ cells per cyst. In this model, higher sperm production and testis weight are seen. SC: Sertoli cell; SZ: spermatozoa.

![Fig. 2. Effects of thyroid hormones during sex differentiation of fish. (A) Treatment with perchlorate induces feminization in early gonadal development, skewing the sex ratio toward females. (B) Treatment with T4 induces masculinization in early gonadal development by reducing the expression of cyp19a1 (aromatase) and erb (estrogen receptor beta) and increasing amh (anti-Müllerian hormone). T4 increases the ratio of females. (C) Methimazole treatment delays masculinization from females during the early developmental stage of zebrafish. Sex ratio is 1:1.

![Fig. 3. (A) Effects of hypothyroidism and hyperthyroidism on testicular development in rodents. The induced-hyperthyroidism decreased the Sertoli cell proliferative window, reflecting directly on the final number of adult Sertoli cell, and consequently in the lower spermatogenic yield and testis size/weight. On the other hand, Sertoli cell differentiation is delayed in the hypothyroidism model, increasing their proliferative window. In consequence, there is an elevated number of adult Sertoli cell which leads to higher spermatogenesis and testis size/weight. SC: Sertoli cell; SZ: spermatozoa. (B) Effects of hypothyroidism on testicular development in fish. PTU-induced hypothyroidism increased Sertoli cell proliferation and Sertoli cell capacity reflecting directly on higher number of germ cells per cyst. In this model, higher sperm production and testis weight are seen. SC: Sertoli cell; SZ: spermatozoa.

early cessation of Sertoli cell proliferation, and a decrease in testis size and sperm production in rats (Fig. 3A) (van Haaster et al., 1993; Cooke et al., 1994a,b; Mendis-Handagma and Ariyaratne, 2005).

There are few papers describing the effects of hypothyroid state on
testicular development in fish (see review in Schulz et al., 2010; França et al., 2015). According to Matta et al. (2002), Nile tilapia (Oreochromis niloticus) juveniles (−0.95 g body weight and 3.5 cm of length) treated with PTU (propylthiouracil, anti-thyroid drug similar to methimazole) doubled the size of testis which is similar to the effects observed in rats, suggesting that mechanisms controlling testicular development by THs may be conserved among vertebrates. In hypothryoid Nile tilapia, Sertoli cells remained immature and were able to proliferate for longer period during testis development, resulting in a higher number of Sertoli cell (Matta et al., 2002) (Fig. 3B). The increased number of Sertoli cell per cyst reflected positively on Sertoli cell capacity and efficiency and consequently in the germ cell number in adult tilapia (Fig. 3B) (Matta et al., 2002; França et al., 2015). Sharma and Patiño (2013) reported that goitrogen (e.g. perchlorate) exposure inhibited pubertal development in male zebrafish, as clearly seen at 60 dpf. In the same work, all control and T4-treated males were mature (> 75% of them showed sperm in their testes), while those that were goitrogen-treated were still juvenile and less than 50% had sperm. In contrast, Timmermans et al. (1997) observed no significant increase in body weight and testis diameter or number of spermatogonia following treatment with T4 in two strains of juvenile common carp (Cyprinus carpio).

6. Adult testis

Although there is a general agreement that THs are important regulators of the testicular development prior to and during puberty in mammals, the role of these hormones is not very clear in the adult testis (Wagner et al., 2009). Considering that most of the experimental studies to date are focused on TH effects on the developing testes, only limited data are available on its role in vertebrate spermatogenesis (Wagner et al., 2009; Morais et al., 2013; Safian et al., 2016).

Altered thyroid status, natural or experimentally induced, is frequently associated with some sort of sexual dysfunction and/or morphological testicular alteration in mammals (Faraone-Menella et al., 2008). Similar studies in reptiles showed that thyroidectomy impaired spermatogenesis of adult garden lizards (Calotes versicolor), while T4 treatment nullified the thyroidectomy-induced testicular atrophy (Haldar-Miara and Thapliyal, 1981). In adult catfish (at early rerudescence/early preparatory phase), treatment with thiourea (a thyroid disrupter) reduced androgen plasma and tissue levels after thyroid depletion, leading to a male reproductive system disruption (Swapna et al., 2006). Interestingly, thiourea treatment withdrawal for 21 days induced a partial recovery of androgen levels, which indicates that thyroid plays a role in maintaining testis functions via androgen stimulation (Swapna et al., 2006). In this context, it has been observed that androgens could also regulate thyroid function. Testosterone propionate injected intramuscularly in immature rainbow trout (Salmo gairdneri Richardson) stimulates, directly or indirectly, the thyroid function, increasing T3 plasma levels (Hunt and Eales, 1979).

To study the effects of THs on fish spermatogenesis, a different experimental setting was employed, where African catfish (Clarias gariepinus) received T4 (100 ppm) for 21 days (Jacob et al., 2005). In this study, T4 depleted the number of spermatozoa/spermatids during the preparatory and pre-spawning phases (Jacob et al., 2005). Interestingly, the T4 overdose effects were more pronounced during the preparatory phase rather than pre-spawning, suggesting that altered TH status might have significant stage-dependent effects on the testis.

More recently, Morais et al. (2013) studied the effects of THs on zebrafish spermatogenesis using an ex vivo approach. These authors showed that T3 increased both mitotic index of type A undifferentiated spermatagonia (Aund) and Sertoli cell in zebrafish testis. In the follow up experiments, Safian et al. (2016) demonstrated that T3 not only promoted the formation of new cysts with type Aund spermatagonia, but also stimulated the accumulation of Adif (type A differentiated spermatagonia) by increasing their proliferation while reducing development into type B spermatagonia in zebrafish (Fig. 4).

To unravel the molecular mechanisms of THs on zebrafish spermatogenesis, Morais et al. (2013) first showed the localization of thrα and thrβ in Sertoli and Leydig cells, respectively. Similar data have been found in mammals, where TRs are localized in Sertoli cells, although these receptors were also found in mammalian germ cells (Buzzaard et al., 2000; Singh et al., 2011). Moreover, it has been demonstrated that T3 increased a stimulatory growth factor of spermatogenesis, named Igf3 (Insulin-like growth factor 3), which in turn promotes spermatogonial differentiation and proliferation towards meiosis. Igf3 is produced by Sertoli cells. T3 increases the transcript levels of two igfbp (insulin-like growth factor binding protein): igfbp1a and igfbp3. Fsh decreases RNAm levels of igfbp1a. The differential expression of the igfbps reflects directly on the Igf3 availability. Elevating Igf3 bioactivity by blocking Igfbps induced by T3 results in a decrease growth of spermatogonial differentiation, leading to a depletion of type Aund spermatagonia in zebrafish.

Next to their roles in Sertoli cells, THs also regulate Leydig cell functions in zebrafish testis (Fig. 4) (Morais et al., 2013). It is well known that Leydig cells express Fsh receptor, and Fsh is a potent steroidogenic hormone in zebrafish testis (Nóbrega et al., 2015). In this context, the Fsh steroidogenic capacity is increased in the presence of T3 in zebrafish testis (Morais et al., 2013). The observed increase in androgen secretion by zebrafish testis was accompanied by higher levels of ar and cyp17a1 (17α-hydroxylase/1720 lyase/1720 desmolase) mRNA levels, suggesting that Fsh effects are enhanced in the presence of T3 (Figs. 1 and 4).
In another study evaluating the action of THs in adult testis, de Montgolfier et al. (2009) examined testicular connexin 43 (cx43) mRNA in brook trout (Salvelinus fontinalis) and demonstrated a clear dose-response relationship between T3 and cx43, suggesting a possible regulation of this connexin by T3. These authors also investigated seasonal changes in the basal expression of several connexins and demonstrated higher levels of testicular cx43 in maturing fish, suggesting that changes in the connexins expression may precede the onset of spermatogenesis in this species.

7. Summary

In summary, this mini-review highlighted the main findings on THs role in male reproductive system of fish, focusing on sex differentiation, testicular development and spermatogenesis. The existing data in the literature have been demonstrated that THs exert their roles at the different levels of the HPG axis. In general, it has been shown a positive correlation between THs and fish reproductive status, where THs are frequently associated with testicular development, growth and maturation. More recently, the molecular mechanisms underlying the role of THs in spermatogenesis have been elucidated in the zebrafish testis model. In this model, ex vivo studies showed that T3 increases the Sertoli cell production of Igf3, which is a stimulatory growth factor of spermatogonial proliferation. Based on the evidences from the literature, it can be concluded that THs modulate the male reproductive system during the different life stages of fish. The physiological and molecular mechanisms showed a link between the thyroid and reproduction, suggesting a possibly co-evolution of these two systems. However, how crucial these systems are to each other remains a topic for future investigation.

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References


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