

Epigenetics and Probiotics Application in Fish Reproductive Performance

Subjects: Reproductive Biology

Contributor: Md Afsar Ahmed Sumon, Mohammad Habibur Rahman Molla, Israa J. Hakeem, Foysal Ahammad, Ramzi H. Amran, Mamdoh T. Jamal, Mohamed Hosny Gabr, Md. Shafiqul Islam, Md. Tariqul Alam, Christopher L. Brown, Eun-Woo Lee, Mohammed Moulay, Amer H. Asseri, F A Dain Md Opo, Ahad Amer Alsaiani, Md. Tawheed Hasan

Fish represent an excellent source of animal protein as well as a biomedical research model as a result of their evolutionary relatedness and similarity with the human genome. Commercial and ornamental fish culture has achieved popularity, but reproductive dysfunctions act as a limiting factor for quality fry production, interfering with the sustainability of the aquaculture industry. Fish reproduction is crucial for any species' existence, and reproductive performance can potentially be improved through applications of epigenetics and probiotics. Epigenetics is a highly sensitive molecular approach that includes chromatin structure and function alteration, DNA methylation, and modification of non-coding RNA molecules for the transfer of desired information from parents to offspring. DNA methyltransferase improves reproductive *cyp11a1*, *esr2b*, and *figla* gene expression and feminizes zebrafish (*Danio rerio*). Moreover, epigenetics also contributes to genome stability, environmental plasticity, and embryonic development. However, methylation of specific genes can negatively affect sperm quality, resulting in poor fertilization. Probiotic administration is able to induce responsiveness of incompetent follicles to maturation-inducing hormones and can change oocyte chemical composition during vitellogenic development. The positive role of probiotics on testicular cells is validated by upregulating the transcription levels of *leptin*, *bdnf*, and *dmrt1* genes facilitating the spermatogenesis.

Keywords: epigenetics ; probiotics ; reproductive dysfunctions ; gene transcription ; ornamental fish ; commercial fish

1. Introduction

Aquaculture has been known for millennia, when the trend of captive fish rearing began, and it is now playing a crucial role in solving the world food crisis, particularly in meeting the protein demand ^[1]. However, broodstock management, which includes the optimization of critical reproductive processes, such as nutrition, maturation, egg and sperm production, and spawning, remains a major obstacle to the advancement of the aquaculture industry. Captive broodstocks are most vulnerable to the disruption of reproductive activities by hormonal imbalance and unfavorable environmental parameters ^[2] resulting in reproductive dysfunction. Such reproductive impairments include no or poor-quality egg or sperm production, defective or weak spawn, less growth and high mortality of fry, and sometimes mortality of the brood. Reproductive dysfunctions can be triggered by direct or indirect variations of the gametes and endocrine system, captivity-induced stress, unsuitability of the spawning environment, and deficiencies in the nutritional profiles of feeds ^{[3][4]}. Some taxonomic groups of fishes are far more sensitive to such inhibitory influences than others. Reproductive dysfunction is common in *Anguilla* spp. (catadromous eels), *Seriola* spp. (greater amberjack and Japanese yellowtail), *Caranx ignobilis* (giant trevally), *Epinephelus* spp. (groupers) and *Thunnus* spp. (bluefin tuna) ^[5], *Anoplopoma fimbria* (Sablefish) ^[6], and *Oncorhynchus mykiss* (Rainbow trout) ^[2].

To avoid reproductive dysfunction, scientific management of broodstock animals ensures proper physiology, immunology, reproductive enzymes pathways activation, and transcription of specific reproductive genes. Many commercial farms seek to maintain healthy broodstocks by supplementation with probiotics to increase reproductive enzymes activities and gene transcription, but dysfunction remains fairly common during breeding seasons ^{[7][8]}. Moreover, epigenetic mechanisms include DNA methylation, histone modification, and noncoding RNA action underlying various processes have recently received considerable attention in aquaculture and offer some promise in the amelioration of breeding performance ^[9].

The term epigenetics literally means “above” or “on top of” genetics, and epigenetic traits refer to stable heritable phenotypes resulting from changes in the status of chromosomes without alteration of DNA sequences ^[10]. Epigenetic reprogramming underpins many developmental processes such as gametogenesis and embryogenesis, foundation of environment regulating events of fish sex differentiation, providing a linkage between phenotypic and metabolic changes during domestication ^[11]. In several studies, epigenetic inheritance was found to be more feasible in fish as compared with terrestrial animals ^{[12][13][14]}. Epigenetic information has the potential to contribute to lower disease prevalence and the potential eradication of the use of antibiotics in commercial aquaculture ^[15]. Sperm quality has been linked to DNA methylation in spermatozoa and in *Morone saxatilis* (striped bass) sperm DNA methylation has a positive relationship with male reproductive capacity ^[16]. Epigenetics influence fish sex determination and differentiation, facilitating interactions between these processes with other surroundings ^[17]. When it comes to valuable fish species, such as grouper, it is essential to know sex patterns, especially when their sex is changed. In aquaculture, a stable and predictable mating

system is critical for healthy fry production [9] and alterations in DNA methylation patterns triggered by higher temperatures can lead to more masculine traits in females [18][19]. Although epigenetics has huge potential, studies on economically important aquaculture species remain preliminary, and many unanswered questions remain [12][20]. An understanding of epigenetic mechanisms for commercial species could contribute to expansions and improvements in the economic viability of large-scale aquaculture.

Aquaculture technology has developed a positive view of probiotics application as an alternative to the application of synthetic antibiotics or chemicals [21][22][23]. Recognition of advantages of practical use of probiotics has grown in light of evidence of upregulation of fish growth, stress adaptation, immune modulation, and disease resistance [24][25]. Probiotics are defined as “live microorganisms administered at an appropriate concentration that exert beneficial effects on host health and immune parameters” [26]. Ghosh, et al. [27] initially reported probiotics capacity to restore viable nutrients in female live-bearing ornamental fishes such as *Poecilia reticulata* (Guppy), *P. sphenops* (molly), *Xiphophorus helleri* (Swordtail) and *X. maculatus* (Platyfish). A later investigation was also carried out in *X. helleri* [28], *Danio rerio* (Zebrafish) [29][30][31][32][33][34], *Carassius auratus* (Goldfish) [35], and *Fundulus heteroclitus* (Killifish) [36]. Limited probiotics studies were also conducted to improve reproductive performance in commercial fishes such as *Oreochromis niloticus* (Nile Tilapia) [37], *O. mykiss* [38], *Ompok pabda* (Butter catfish) [39], *Clarias gariepinus* (African catfish) [40] and *A. anguilla* (European eel) [41] on the subject of reproductive indicators such as fecundity (Fec), hatching rate (HR), gamete quality, gonadosomatic index (GSI), fertilization rate (FR), and survival rate (SR). Pathogenic infections often leading to mortality of brood or offspring may be unnoticed by hatchery managers. Synthetic antibiotics or chemical applications to control infection may lead to mass environmental bacterial killing, antibiotic deposition in fish body and the generation of antibiotic-resistant pathogens [42][43]. As a result, probiotics may be a favorable option for broodstocks and fry for the control of infections and for improved reproductive success [44]. The use of probiotics helps to stabilize and diversify the intestinal microbial community, leading to improved reproduction processes through activation of different hormones, enzymes, and genes transcription resulting better FR, HR, SR, and larval growth [7][38].

2. Fish Reproductive Dysfunctions

Globally, aquaculture is continually striving for the consistency of physiological integrity of fingerlings through standardized reproductive programmes, which are a key objective for sustainable aquaculture [45]. Brood fish rearing and their management strategies have been considered as vital for aquaculture production for the last three decades. Generally, many species of captive fishes do not perform normal reproduction (especially females), a phenomenon potentially caused by a lack of natural spawning stimuli, specifically failure of oocyte maturation [46][47]. Supplied protein, fatty acids, lipid, vitamins, especially E and C, and carotenoids also influence fish Fec, FR, HR, and larval development [48][49]. Moreover, presence of chemical fertilizers, antibiotics, hormones and industrial effluents, environmental degradation, and frequent fluctuation of temperature in nature and captivity suppress the immunity and beneficial microbial activity in broodstock fishes, making them more susceptible to infectious disease [27][50][51][52]. Hypothalamus, pituitary, and gonads form the hypothalamic-pituitary-gonadal (HPG) axis, which regulates reproductive function in most fishes [53]. The hypothalamus and the pituitary glands modulate the production of pituitary gonadotropins luteinizing hormone (LH) and follicle stimulating hormone (FSH), and gonadal sex steroids (SS) regulate vitellogenesis and oocyte maturation. FSH and LH have been confirmed to figure importantly in oocyte growth and maturation [54]. Gioacchini et al. [7] have categorized three types of female broodstocks reproductive dysfunctions based on the affected pattern of reproductive cycle whereas, Selvaraj, et al. [55] documented two modes of dysfunction of teleost fishes (**Figure 1**). Both groups of investigators associated the most disruptive reproductive dysfunctions occurred with the vitellogenic phase, as exemplified by failed vitellogenesis in *Anguilla* spp. and *Seriola* spp. [56] and *Mugil cephalus* (Grey Mullet) [57]. The second mode is categorized as the inability to reach final oocyte maturation to ovulation as seen in farmed white and striped bass [58][59][60] and in Cyprinidae [61]. The third set of dysfunctions leads to failures to spawn in the breeding season; sometimes *E. aeneus*, *M. saxatilis*, and *Dentex dentex* (common dentex) females may release eggs after ovulations without exhibiting characteristic breeding behavior [7][62]. Under some circumstances, cultured salmonids fail to complete the reproductive cycle and eggs in the abdominal cavity are reabsorbed over the following months [63]. These are examples of failures among female broodstocks that do not appear to be accompanied by reproductive problems that can be attributed to male fishes.

One common problem faced by cultured male fishes is low quantity or quality of milt production during spermiation [60]. The presence of both male and female germ cells in the testis tissue is referred to as an intersex state caused by endocrine disruption through chemical exposure and hormonal imbalance [64]. The histopathological assessment of gonadal tissue during the development stage allows the detection of intersex reproductive dysfunction in male Japanese medaka (*Oryzias latipes*) [65].

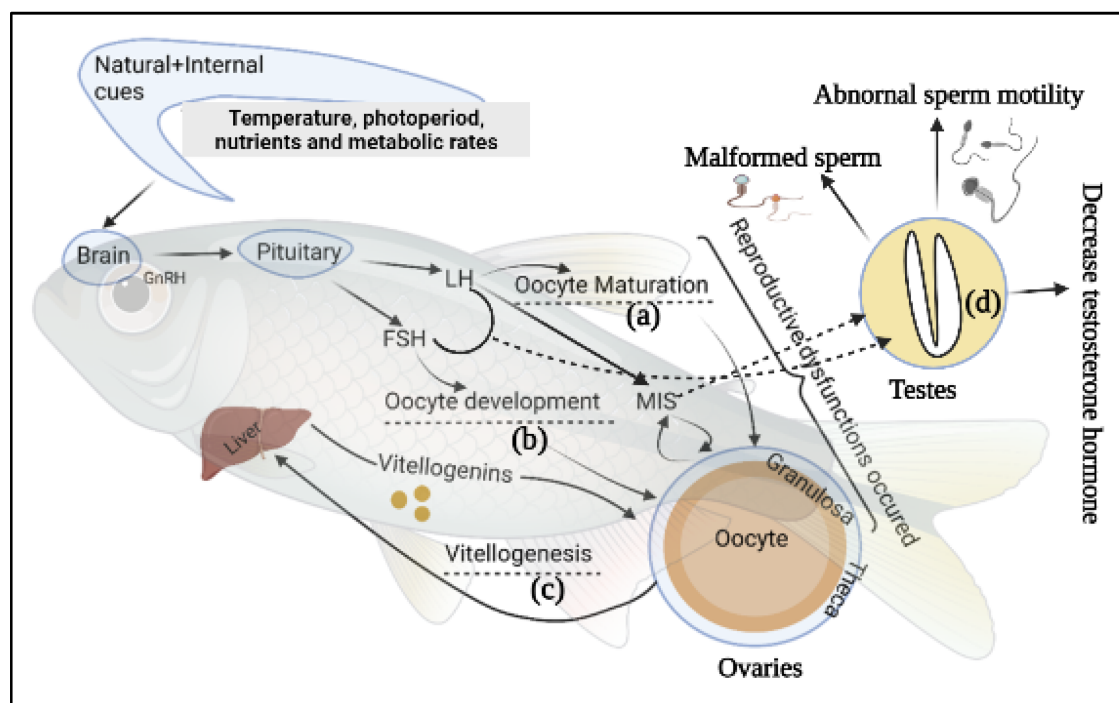


Figure 1. Schematic representation of the reproductive system and gonadal dysfunctions of fish. The broken line indicates the reproductive developmental stages. These dysfunctions may occur: (a) Lack of effectiveness of luteinizing hormone (LH) for oocyte maturation; (b) outreaching of follicle-stimulating hormone (FSH) for oocyte development; (c) lower or no production of vitellogenin from the liver; (d) malformation of sperm and its motility, lowered testosterone to the testis. GnRH: gonadotrophin releasing hormone, MIS: maturation-inducing steroid.

3. Epigenetics Mechanism and Modifications of Fish Reproductive Performance

Considerable attention is focused in aquaculture on brood rearing, breeding, feeding, sex control, and disease management. The study of epigenetic mechanisms underlying various molecular mechanisms has the potential to contribute to favorable hatchery productivity. Environmental cues can cause phenotypic changes in organisms via epigenetic mechanisms (**Figure 2**). Moreover, environmental temperature, dissolved oxygen, water, pH, pollutants, and other factors influenced epigenetic changes in fish gonad and sex-dependent behavior [11][66][67]. Epigenetic modifications include DNA methylation, histone modification, chromatin re-modelling, and the action of noncoding RNAs (ncRNAs), attracting interest for their practical potential in fish reproduction [12].

The structure and consequently the function of chromatin can be altered by epigenetic mechanisms, resulting in the modulation of patterns of gene expression [68]. DNA methylation is the most well-known and well-understood epigenetic mechanism, in response to variable environmental factors, such as photoperiod, toxins, temperature, nutrients, etc. [69]. This entails the addition or removal of a methyl group to DNA, which alters gene function and transcription level as well. The methyl group is covalently added to the 5-carbon location of the cytosine ring, resulting in 5-methylcytosine (5-mC), known colloquially as the “fifth base” of DNA. DNA methylation takes place at CpG doublets either within CpG islands (dinucleotide CG), intergenic regions, or the gene body [70]. It has been reported that CpG islands influence gene expression by regulating transcription factor binding with chromatin structure. De novo DNA methylation and its maintenance are carried out by a family of DNA methyltransferase enzymes (DNMTs) and during embryogenesis, DNMT3A and DNMT3B are responsible for de novo methylation [71]. DNA demethylation is also equally important and accompanied by DNA methylation which is necessary for epigenetic reprogramming of genes. Through the incorporation of histone variants and post-translational modification of histones, chromatin structure can be altered to enhance or repress transcription. These altered chromatin states can be inherited both mitotically and meiotically, suggesting that they may transmit epigenetic information to the next generation. Evidence suggests that certain modified histones are retained non-randomly during spermatogenesis in both mammals and zebrafish, and these marks are thought to play a role in transferring epigenetic information to embryos [72][73]. The ncRNAs, which are made up of small and long RNA molecules, can influence gene expression [74]. Investigations reveal that ncRNAs play important roles in genome stability, environmental plasticity, and embryonic development [75][76]. The majority of research on ncRNAs in fish and shellfish, including important aquaculture species, has been conducted in Atlantic salmon and rainbow trout [77][78]. Aside from developmental programming, broodstock holding/conditioning is an important consideration for the potential transmission of epigenetic information transfer from parents to offspring [79]. Importantly, epigenetic transmission can occur on both the maternal and paternal sides [80] and the study of transgenerational plasticity in fish has grown in popularity [81].

Recent investigations have analyzed the impact of epigenetics, specifically DNA methylation on breeding dynamics and productivity in finfish aquaculture. DNA methylation patterns have been found to change in response to temperature

increases, leading to the masculinization of genetically female *D. labrax* [82], *Cynoglossus semilaevis* (Halfsmooth tongue sole) [18][19]; a trait which can be passed to offspring. In another study, Campos, et al. [83] found that *Solea senegalensis* (Senegalese sole) larvae undergoing metamorphosis had higher methylation levels on the *myog* promoter in skeletal muscle when reared at lower temperatures (15 °C). Hatchery offspring grown in captivity acquired epigenetic alterations in sperm which may explain rapid genetic and phenotypic alterations in the hybrid fishes. The differential methylation in hatchery salmon displayed the presence of their TATA-binding protein (a transcription factor that binds specifically to a DNA sequence) during spermiogenesis and embryonic development [84][85][86]. Epigenetics and broodstock nutrition also influence sperm production and quality in the aquaculture industry. Diet-induced methylation influence freshwater and saltwater trout and higher salt-containing diets caused dramatic changes in global methylation patterns [87]. Moreover, hydrogen sulfide stimulates DNA methylation in environments that can be continued generationally through the germline even after withdrawal of this toxic chemical [14]. Hypoxia-induced reproductive impairment of gonadal development, low sperm count, and motility through different methylation in sperm genes are inherited by the next generation [88]. Transgenerational epigenetic alteration in spermatozoa of aquatic animals resulted in phenotypic variation and methylation of specific gene groups, with potential negative effects on sperm quality and fertilization in male striped bass [16].

Global and gene-specific methylation in spermatozoa significantly affect the fertilization performance of *C. carpio*, in which methylation at CpG sites markedly increased and decreased after 24 and 96 h of post stripping, respectively [89]. In Nile tilapia, the gonadal transcriptomic study demonstrated that all of the DNMTs were expressed in both male and female reproductive organs while specific DNMTs were more highly expressed in the testis. Incubation of gonads with DNMTs inhibitor showed downregulated DNMTs with increased expression of male and female sex determinant gene *dmrt1* and *cyp19a1a*, respectively [90].

It has also been revealed that *cyp19a1a* expression in Nile tilapia was critically controlled by environmental factors like temperature [91]. Similarly in zebrafish, DNMTs treatment feminizes the fish, stimulating long-term expression of key reproduction-related genes (e.g., *cyp11a1*, *esr2b* and *figla*) [92]. However, integrative transcriptome of Atlantic salmon testis revealed the involvement of differentially expressed micro (mi) RNA, thereby resulting in early puberty. This was the first study to link specific groups miRNA involvement in testis maturation in an inversely correlated relationship with targets [93]. Moreover, the endogenous non-coding RNAs (MicroRNAome) of sperm are affected by overexpression of growth hormone and consequently reduced sperm quality and fertilization potentiality of transgenic zebrafish [94]. Finally, the insufficiency of miR-202 compromised oogenesis or folliculogenesis, and significantly decreased the number of follicles [95].

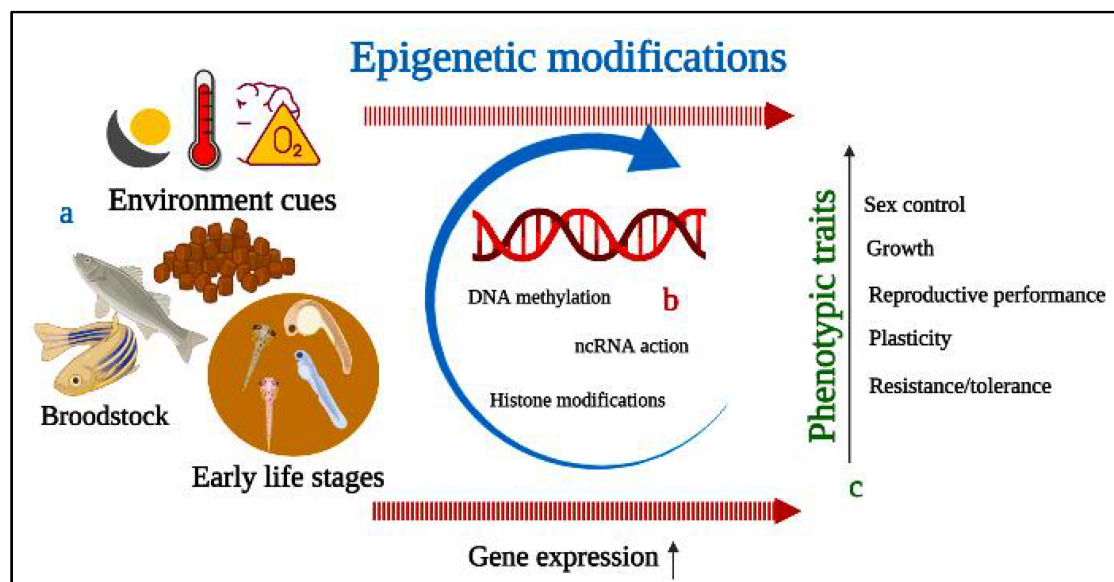


Figure 2. A diagram depicting the potential applications of epigenetics for modulation of reproductive performance. (a) Environmental cues can alter gene transcription via epigenetic mechanisms; (b) DNA methylation, histone modifications, and non-coding RNA action resulting in phenotypic changes; (c) outcomes of epigenetics in phenotypic traits from early to later life stage.

4. Influence of Probiotics on Reproductive Performance of Fish

Host-derived probiotic bacteria are being used increasingly in the aquaculture sector. The host-associated probiotics have not only improved the reproductivity of fish [96] but also helped to improve the endocrine and reproductive signaling of fish. Isolated *Enterococcus xiangfangensis*, *Citrobacter freundii*, *Pseudomonas aeruginosa*, *P. stutzeri*, and *B. subtilis* from fish resulted in growth, hematological, and reproductive performance upregulation among host fishes [97].

The action mechanism of probiotics on reproductive function is normally indicated by increasing egg production along with improvements in the vitellogenic follicles and GSI [98]. However, the lipidic and glucidic components elevated during pre-vitellogenesis result in oocytes maturation, modification of the secondary structure of protein, and impacts on hydration and phosphorylation [99]. Probiotic administration can induce the responsiveness of incompetent follicles (stage IIIa) to MIH in the maturation period and changes oocyte chemical composition, fostering the vitellogenic development [100]. Modifications of the electrophoretic pattern at maturation, and, to a lesser extent, at yolk protein levels have changed the ooplasm components [101]. In fish, probiotics inhibit apoptosis and increase the rate of follicular survival during the developmental stage. Furthermore, effects of probiotics also increase the production of sperm from the testis, with stimulatory actions on two components known as intertubular (or interstitial) and tubular sections [34]. Some probiotics have been shown to stimulate steroidogenic Leydig cells, blood/lymphatic vessels, macrophages and mast cells, neural, and connective tissue cells. This is the case although the somatic Sertoli cells and the germ cells are found at different stages of development and Sertoli cells function in the determination of spermatogenic capacity [102]. The close and continuous interaction with Sertoli cells may contribute to germ cell survival. Consequently, the positive role of probiotics in molecular parameters in testicular cells is validated by upregulating the transcription level of leptin, bdnf and dmrt1 genes facilitated the potentialities of spermatogenesis [41]. Moreover, increased transcription levels of activin, ar α , ar β , pr1, and fshr contribute to sperm quality improvement during spermatogenesis [41].

Probiotics are widely used as feed enrichment for farming aquatic organisms especially fish. The initial application of probiotics in aquaculture was for growth promoters and fish health. However, new areas of research, such as their effect on reproduction, maturation, and fecundity, have been found, though these require more comprehensive development [96]. Several studies have found that probiotics improve host microbial balance and thus improve health, disease resistance, growth performance, feed utilization, and reproductive performance [96][103][104]. Nutrient supplements for broodstock are critical for aquaculture success. Major nutrients such as lipid, protein, fatty acids, vitamins E and C, and carotenoids are essential for various reproduction processes such as fecundity, fertilization, hatching, and larval development [49]. In general, studies in fish show that decreased food availability or starvation causes gonad regression and a decrease in female spawning and egg production, whereas increased food availability and supplemented adequate nutrients promote growth and larger body sizes, causing earlier maturation and higher fecundity in some species [105]. However, the administration of different growth promoters (hormones, antibiotics, nutrient mixtures) can cause suppression of the beneficial microbial activity in the intestinal tract of the broodfish. In such conditions, probiotic supplements can be used to repair these deficiencies and improve the fish health conditions in breeding time.

Additionally, supplemented probiotics such as lactic acid bacteria improved feed utilization and reproductive function of fish [106]. The administration of probiotics through feed supplementation could regulate and modify the expression patterns of genes or hormones responsible for the regulation of fish reproduction [107]. Therefore, using probiotics as feed supplements can potentially improve reproductive function and activate reproductive genes to correct reproductive dysfunction.

4.1. Ornamental Fish

Ornamental fish have popularity for their unique colors and their importance is increasing not only for hobby purposes, but also for their use as animal models for scientific research. A diverse array of probiotics induces beneficial effects on ornamental fish reproductive performance, as presented in **Table 1**.

4.1.1. Male

Ornamental fish especially, zebrafish were used over the last 20 years for studying genetics and gonadal development [108][109]. Probiotics have been used in zebrafish trials to observe the transcription of the genes related to reproductive maturity and reproduction [110][111]. Transcription of KiSS1, KiSS2, and gnrh3 genes in the brain can trigger the reproductive system of male fish by the direct or indirect secretion of many hormones in response to probiotics, especially leptin [112]. Valcarce et al. [34] first reported *P. acidilactici* supplementation correlates with and upregulates male reproductive performance of zebrafishes. This probiotic triggered five genes, including brain-derived neurotrophic factor (bdnf), BCL2-interacting killer (bik), double-sex and mab-3 related transcription factor 1 (dmrt1), and FSH beta subunit, and leptin a (lepa) transcription, as an indication of good sperm quality (SQ) marker. These genes all produce positive effects on testicular cells, potentially improving reproductive performance [34][113]. The blended dietary administration *P. acidilactici* (0.2%) and nucleotide (0.5%) demonstrated positive effects on SQ, sperm motility (SM) and sperm density (SDn) in goldfish (*C. auratus*) [35]. Furthermore, *Lab. rhamnosus* and *Bifidobacterium longum* have been showing antioxidant and anti-inflammatory features to assist zebrafish SQ and male reproductive behavior [114]. These probiotics demonstrated a positive effect on SQ, SDn, total and progressive SM, and fast spermatozoa subpopulations.

4.1.2. Female

Probiotic *Lab. rhamnosus* IMC 501 has striking effects on the ovarian development of female zebrafish [71][31]. Moreover, in this fish, dietary administration of *L. rhamnosus* at 10⁶ CFU g⁻¹ improved Fec, GSI, and oocyte maturation (FD and FM), supporting reproductive performance by improving fecundity [29][101]177. This probiotic increases the transcription level of transforming growth factor b1 (tgfb1), growth differentiation factor9 (gdf9), and bone morphogenetic protein15 (bmp15) contributing to oocyte development of that fish. Carnevali, et al. [115] also monitored the long-term effects of the same

probiotic on zebrafish and reported dietary effectiveness on FD; ovulated oocytes quantification; embryo quality and larval growth performance. This probiotic stimulates sexual maturation of this species by improving the expression of aromatase cytochrome p 19 (*cyp19a*), vitellogenin (*vtg*), an isoform of the E2 receptor (*era*), LH receptor, 20-b hydroxysteroid dehydrogenase (*20b-hsd*), membrane progesterone receptors a and b, cyclin B, activinbA1, smad2, tgfb1, gdf9 and bmp15, which are responsible for regulating reproductive hormone secretion. Autophagy was observed during follicle development in the ovarian tissue and *L. rhamnosus* has a key role in follicle maturation as confirmed by focal plane array analysis [30]. Miccoli et al. [33] conducted an experiment with the same probiotic including similar dose, duration, and species as reported by Gioacchini et al. [30], and reported that probiotics promote embryonic development by changing both maternal and zygotic mRNA levels. Similarly, the use of *L. rhamnosus* CICC 6141 and *L. casei* BL23 probiotic effects on *D. rerio* were documented and later probiotic markedly improved Fec, OVr, HR, and FR [116].

Probiotic, *L. rhamnosus* IMC 501 treatment has profound effects on killifish (*Fundulus heteroclitus*) GSI, Fec, and embryo SR [36]. Dietary supplementation of probiotic positively improves egg and ovum diameter, absolute fecundity, and some other properties in goldfish [35]. Probiotic administration caused a substantial impact on reproductive performance in four live-bearing ornamental species: *Poecilia reticulata*, *P. sphenops*, *X. helleri*, and *X. maculatus* [27]. In these fishes, administration of *B. subtilis* for 1-year improved GSI, Fec, and fry production of spawning females. Additionally, probiotic could synthesize vitamin B1 and vitamin B12 that controlled the mortality or body deformities of fry. The live-bearing ornamental female swordtails displayed improved GSI, Fec, and fry production after taking commercial probiotic (PrimaLac) (see Table 1) as a feed additive for 182 days [28].

Table 1. Dietary supplemented probiotics effects on ornamental fish reproduction. Symbol: no change (→); increase (↑); decrease (↓) versus controls.

Supplemented Probiotics	Fish Species	Fish Number	Duration	Concentration	Effects on Fish	R
<i>Bacillus subtilis</i>	<i>Poecilia reticulata</i> (Guppy), <i>P. sphenops</i> (Valenciennes), <i>Xiphophorus helleri</i> (Swordtail fish) and <i>X. maculatus</i> (Platyfish)	60 virgin females of each species	365 days	5×10^7 – 5×10^8 CFU g ⁻¹ and 5×10^6 – 5×10^6 CFU g ⁻¹	EP Fec and GSI ↑; SR (fry) ↑; Fry death and deformities ↓	
<i>Lactobacillus rhamnosus</i> IMC 501	<i>Danio rerio</i> (Zebrafish)	10 females	10 days	10^6 CFU g ⁻¹	EP Fec, GSI, and Ovulation rate ↑; Oocyte maturation G and FD ↑; Oocyte maturation FD and FM ↑	
<i>Lab. rhamnosus</i> IMC 501	<i>D. rerio</i>	40 males and females	10 days	10^6 CFU g ⁻¹	Follicular survival ↑ and apoptosis ↓	
<i>Pediococcus acidilactici</i> (Bactocell®)	<i>D. rerio</i>	5 wild males	10 days	10^6 CFU g ⁻¹	Embryo development ↑; HR ↑	
<i>Lab. rhamnosus</i> CECT8361 and <i>Bifidobacterium longum</i> CECT7347	<i>D. rerio</i>	36 Males	21 days	10^9 CFU g ⁻¹	SP testicular cells ↑	
PrimaLac® (<i>Lab. acidophilus</i> , <i>Lab. casei</i> , <i>Enterococcus faecium</i> , <i>Bifidobacterium thermophilum</i>)	<i>X. helleri</i>	10 females and 3 males	182 days	0.04%, 0.09% and 0.14%	SP SQ, SDn, SM ↑	
<i>P. acidilactici</i>	<i>Carassius auratus</i> (Goldfish)	720 fishes	180 days	0.1, 0.2, and 0.3%	EP Fec and GSI ↑; SR (fry) ↑	
<i>Lab. rhamnosus</i> IMC 501	<i>Fundulus heteroclitus</i> (Killifish)	10 females and 10 males	8 days	10^6 CFU mL ⁻¹	EP GSI, HSI, AF, RF, ED, OD, FR, and HR ↑; SP SM, SD, SDn, and Stc ↑	
					EP GSI, Fec ↑ and HR →; SR (fry) ↑; GP L and W ↑	

AF: absolute fecundity; EP: egg parameters; ED: egg diameter; Fec: fecundity; FD: follicle development; FM: follicle maturation; FR: fertilization rate; G: growth; GP: growth parameters; GSI: gonadosomatic index; HR: hatching rate; HSI: hepatosomatic index; OD: ovum diameter; RF: relative fecundity; SR: survival rate; SP: sperm parameters; SM: sperm motility (%); SD: sperm duration; SDn: sperm density; Stc: spermatocriet (%); W: weight.

References

1. Liao, I.C.; Chao, N.-H. Aquaculture and food crisis: Opportunities and constraints. *Asia Pac. J. Clin. Nutr.* 2009, 18, 564–569.
2. Merrifield, D.L.; Ringo, E. *Aquaculture Nutrition: Gut Health, Probiotics and Prebiotics*; John Wiley & Sons: Hoboken, NJ, USA, 2014.
3. Akhtar, M.; Ciji, A.; Sarma, D.; Rajesh, M.; Kamalam, B.; Sharma, P.; Singh, A. Reproductive dysfunction in females of endangered golden mahseer (*Tor putitora*) in captivity. *Anim. Reprod. Sci.* 2017, 182, 95–103.
4. Mañanós, E.; Duncan, N.; Mylonas, C. Reproduction and control of ovulation, spermiation and spawning in cultured fish. In *Methods in Reproductive Aquaculture: Marine and Freshwater Species*; CRC Press, Taylor and Francis Group: Boca Raton, FL, USA, 2008; pp. 3–80.
5. Ottolenghi, F.; Silvestri, C.; Giordano, P.; Lovatelli, A.; New, M.B. *Capture-Based Aquaculture: The Fattening of Eels, Groupers, Tunas and Yellowtails*; FAO: Mexico City, Mexico, 2004.
6. Guzmán, J.; Luckenbach, A.; Goetz, F.W.; Fairgrieve, W.T.; Middleton, M.A.; Swanson, P. Reproductive dysfunction in cultured sablefish (*Anoplopoma fimbria*). *Bull. Fish. Res. Agency* 2015, 40, 111–119.
7. Gioacchini, G.; Giorgini, E.; Vaccari, L.; Carnevali, O. Can Probiotics Affect Reproductive Processes of Aquatic Animals? In *Aquaculture Nutrition: Gut Health, Probiotics and Prebiotics*; John Wiley & Sons, Ltd.: Hoboken, NJ, USA, 2014; p. 328–346.
8. Martínez Cruz, P.; Ibáñez, A.L.; Monroy Hermosillo, O.A.; Ramírez Saad, H.C. Use of probiotics in aquaculture. *Int. Sch. Res. Not.* 2012, 2012, 916845.
9. Granada, L.; Lemos, M.F.; Cabral, H.N.; Bossier, P.; Novais, S.C. Epigenetics in aquaculture—The last frontier. *Rev. Aquac.* 2018, 10, 994–1013.
10. Berger, S.L.; Kouzarides, T.; Shiekhattar, R.; Shilatifard, A. An operational definition of epigenetics. *Genes Dev.* 2009, 23, 781–783.
11. Herráez, M.P.; Lombó, M.; González-Rojo, S. The Role of Epigenetics in Fish Biology and Reproduction: An Insight into the Methods Applied to Aquaculture. In *Cellular and Molecular Approaches in Fish Biology*; Elsevier: Amsterdam, The Netherlands, 2022; pp. 69–104.
12. Gavary, M.R.; Roberts, S.B. Epigenetic considerations in aquaculture. *PeerJ.* 2017, 5, e4147.
13. Hanson, M.A.; Skinner, M.K. Developmental origins of epigenetic transgenerational inheritance. *Environ. Epigenet.* 2016, 2, dvw002.
14. Kelley, J.L.; Tobler, M.; Beck, D.; Sadler-Riggleman, I.; Quackenbush, C.R.; Rodriguez, L.A.; Skinner, M.K. Epigenetic inheritance of DNA methylation changes in fish living in hydrogen sulfide—Rich springs. *Proc. Natl. Acad. Sci. USA* 2021, 118, e2014929118.
15. González-Recio, O. Epigenetics: A new challenge in the post-genomic era of livestock. *Front. Genet.* 2012, 2, 106.
16. Woods, L.C., III; Li, Y.; Ding, Y.; Liu, J.; Reading, B.J.; Fuller, S.A.; Song, J. DNA methylation profiles correlated to striped bass sperm fertility. *BMC Genom.* 2018, 19, 244.
17. Renn, S.C.; Hurd, P.L. Epigenetic regulation and environmental sex determination in cichlid fishes. *Sex. Dev.* 2021, 15, 93–107.
18. Chen, S.; Zhang, G.; Shao, C.; Huang, Q.; Liu, G.; Zhang, P.; Song, W.; An, N.; Chalopin, D.; Volff, J.-N.; et al. Whole-genome sequence of a flatfish provides insights into ZW sex chromosome evolution and adaptation to a benthic lifestyle. *Nat. Genet.* 2014, 46, 253–260.
19. Shao, C.; Li, Q.; Chen, S.; Zhang, P.; Lian, J.; Hu, Q.; Sun, B.; Jin, L.; Liu, S.; Wang, Z.; et al. Epigenetic modification and inheritance in sexual reversal of fish. *Genome Res.* 2014, 24, 604–615.
20. Labbé, C.; Robles, V.; Herráez, M.P. Epigenetics in fish gametes and early embryo. *Aquaculture* 2017, 472, 93–106.
21. Hasan, K.N.; Banerjee, G. Recent studies on probiotics as beneficial mediator in aquaculture: A review. *J. Basic Appl. Zool.* 2020, 81, 53.
22. Hasan, M.T.; Jang, W.J.; Kim, H.; Lee, B.-J.; Kim, K.W.; Hur, S.W.; Lim, S.G.; Bai, S.C.; Kong, I.-S. Synergistic effects of dietary *Bacillus* sp. SJ-10 plus β -glucan oligosaccharides as a synbiotic on growth performance, innate immunity and streptococcal resistance in olive flounder (*Paralichthys olivaceus*). *Fish Shellfish Immunol.* 2018, 82, 544–553.
23. Jamal, M.T.; Sumon, A.A.; Pugazhendhi, A.; Al Harbi, M.; Hussain, A.; Haque, F. Use of probiotics in commercially important finfish aquaculture. *Int. J. Probiotics Prebiotics* 2020, 15, 7–21.

24. Hasan, M.T.; Jang, W.J.; Lee, B.-J.; Hur, S.W.; Lim, S.G.; Kim, K.W.; Han, H.-S.; Lee, E.-W.; Bai, S.C.; Kong, I.-S. Dietary Supplementation of *Bacillus* sp. SJ-10 and *Lactobacillus plantarum* KCCM 11322 Combinations enhance growth and cellular and humoral immunity in olive flounder (*Paralichthys olivaceus*). *Probiotics Antimicrob. Proteins* 2021, 13, 1277–1291.
25. Hasan, M.T.; Je Jang, W.; Lee, J.M.; Lee, B.-J.; Hur, S.W.; Gu Lim, S.; Kim, K.W.; Han, H.-S.; Kong, I.-S. Effects of immunostimulants, prebiotics, probiotics, synbiotics, and potentially immunoreactive feed additives on olive flounder (*Paralichthys olivaceus*): A review. *Rev. Fish. Sci. Aquac.* 2019, 27, 417–437.
26. FAO/WHO. Health and Nutritional Properties of Probiotics in Food including Powder Milk with Live Lactic Acid Bacteria. In Report of a Joint FAO/WHO Expert Consultation on Evaluation of Health and Nutritional Properties of Probiotics in Food Including Powder Milk with Live Lactic Acid Bacteria; FAO Food and Nutrition Paper—Series Number: 0254-4725[1014-2908; FAO/WHO: Cordoba, Argentina, 2001; p. 56.
27. Ghosh, S.; Sinha, A.; Sahu, C. Effect of probiotic on reproductive performance in female livebearing ornamental fish. *Aquac. Res.* 2007, 38, 518–526.
28. Abasali, H.; Mohamad, S. Effect of dietary supplementation with probiotic on reproductive performance of female livebearing ornamental fish. *Res. J. Anim. Sci.* 2010, 4, 103–107.
29. Gioacchini, G.; Bizzaro, D.; Giorgini, E.; Ferraris, P.; Sabbatini, S.; Carnevali, O. P-230 Oocytes maturation induction by *Lactobacillus rhamnosus* in *Danio rerio*: In vivo and in vitro studies. *Hum. Reprod.* 2010, 25, I205–I206.
30. Gioacchini, G.; Dalla Valle, L.; Benato, F.; Fimia, G.M.; Nardacci, R.; Ciccocanti, F.; Piacentini, M.; Borini, A.; Carnevali, O. Interplay between autophagy and apoptosis in the development of *Danio rerio* follicles and the effects of a probiotic. *Reprod. Fertil. Dev.* 2013, 25, 1115–1125.
31. Gioacchini, G.; Giorgini, E.; Ferraris, P.; Tosi, G.; Bizzaro, D.; Silvi, S. Could probiotics improve fecundity? *Danio rerio* as a case of study. *J. Biotechnol.* 2010, 150, 59–60.
32. Giorgini, E.; Conti, C.; Ferraris, P.; Sabbatini, S.; Tosi, G.; Rubini, C.; Vaccari, L.; Gioacchini, G.; Carnevali, O. Effects of *Lactobacillus rhamnosus* on zebrafish oocyte maturation: An FTIR imaging and biochemical analysis. *Anal. Bioanal. Chem.* 2010, 398, 3063–3072.
33. Miccoli, A.; Gioacchini, G.; Maradonna, F.; Benato, F.; Skobo, T.; Carnevali, O. Beneficial bacteria affect *Danio rerio* development by the modulation of maternal factors involved in autophagic, apoptotic and dorsalizing processes. *Cell. Physiol. Biochem.* 2015, 35, 1706–1718.
34. Valcarce, D.G.; Pardo, M.; Riesco, M.; Cruz, Z.; Robles, V. Effect of diet supplementation with a commercial probiotic containing *Pediococcus acidilactici* (Lindner, 1887) on the expression of five quality markers in zebrafish (*Danio rerio* (Hamilton, 1822)) testis. *J. Appl. Ichthyol.* 2015, 31, 18–21.
35. Mehdinejad, N.; Imanpour, M.R.; Jafari, V. Combined or individual effects of dietary probiotic, *Pediococcus acidilactici* and nucleotide on reproductive performance in goldfish (*Carassius auratus*). *Probiotics Antimicrob. Proteins* 2019, 11, 233–238.
36. Lombardo, F.; Gioacchini, G.; Carnevali, O. Probiotic-based nutritional effects on killifish reproduction. *Fish. Aquacult J. FAJ-33* 2011, 2011, FAJ-33.
37. Dias, D.d.C.; Furlaneto, F.d.P.B.; Sussel, F.R.; Tachibana, L.; Gonçalves, G.S.; Ishikawa, C.M.; Natori, M.M.; Ranzani-Paiva, M.J.T. Economic feasibility of probiotic use in the diet of Nile tilapia, *Oreochromis niloticus*, during the reproductive period. *Acta Scientiarum. Anim. Sci.* 2020, 42, e47960.
38. Akbari Nargesi, E.; Falahatkar, B.; Sajjadi, M.M. Dietary supplementation of probiotics and influence on feed efficiency, growth parameters and reproductive performance in female rainbow trout (*Oncorhynchus mykiss*) broodstock. *Aquac. Nutr.* 2020, 26, 98–108.
39. Rahman, M.L.; Akhter, S.; Mallik, M.K.M.; Rashid, I. Probiotic enrich dietary effect on the reproduction of butter catfish, *Ompok pabda* (Hamilton, 1872). *Int. J. Curr. Res. Life Sci.* 2018, 7, 866–873.
40. Ariole, C.N.; Okpokwasili, G.C. The effect of indigenous probiotics on egg hatchability and larval viability of *Clarias gariepinus*. *Rev. Ambiente Água* 2012, 7, 81–88.
41. Vílchez, M.C.; Santangeli, S.; Maradonna, F.; Gioacchini, G.; Verdenelli, C.; Gallego, V.; Peñaranda, D.S.; Tveiten, H.; Pérez, L.; Carnevali, O.; et al. Effect of the probiotic *Lactobacillus rhamnosus* on the expression of genes involved in European eel spermatogenesis. *Theriogenology* 2015, 84, 1321–1331.
42. Hasan, M.T.; Jang, W.J.; Lee, S.; Kim, K.W.; Lee, B.J.; Han, H.S.; Bai, S.C.; Kong, I.S. Effect of β -glucan oligosaccharides as a new prebiotic for dietary supplementation in olive flounder (*Paralichthys olivaceus*) aquaculture. *Aquac. Res.* 2018, 49, 1310–1319.
43. Hasan, M.T.; Jang, W.J.; Lee, B.-J.; Kim, K.W.; Hur, S.W.; Lim, S.G.; Bai, S.C.; Kong, I.-S. Heat-killed *Bacillus* sp. SJ-10 probiotic acts as a growth and humoral innate immunity response enhancer in olive flounder (*Paralichthys olivaceus*). *Fish Shellfish Immunol.* 2019, 88, 424–431.
44. Verschuere, L.; Rombaut, G.; Sorgeloos, P.; Verstraete, W. Probiotic bacteria as biological control agents in aquaculture. *Microbiol. Mol. Biol. Rev.* 2000, 64, 655–671.
45. Boyd, C.E.; D'Abramo, L.R.; Glencross, B.D.; Huyben, D.C.; Juarez, L.M.; Lockwood, G.S.; McNevin, A.A.; Tacon, A.G.; Teletchea, F.; Tomasso, J.R., Jr.; et al. Achieving sustainable aquaculture: Historical and current perspectives and f

- uture needs and challenges. *J. World Aquac. Soc.* 2020, 51, 578–633.
46. Mylonas, C.C.; Zohar, Y. Promoting oocyte maturation, ovulation and spawning in farmed fish. In *The Fish Oocyte*; Springer: Berlin/Heidelberg, Germany, 2007; pp. 437–474.
 47. Papadaki, M.; Peleteiro, J.B.; Alvarez-Blázquez, B.; Villanueva, J.L.R.; Linares, F.; Vilar, A.; Rial, E.P.; Lluch, N.; Fakriddis, I.; Sigelaki, I.; et al. Description of the annual reproductive cycle of wreckfish *Polyprion americanus* in captivity. *Fishes* 2018, 3, 43.
 48. Fernando, A.; Phang, V.; Chan, S. Diets and feeding regimes of poeciliid fishes in Singapore. *Asian Fish. Sci* 1991, 4, 99–107.
 49. Izquierdo, M.; Fernandez-Palacios, H.; Tacon, A. Effect of broodstock nutrition on reproductive performance of fish. *Aquaculture* 2001, 197, 25–42.
 50. Nayak, S.K. Role of gastrointestinal microbiota in fish. *Aquac. Res.* 2010, 41, 1553–1573.
 51. Sullam, K.E.; Essinger, S.D.; Lozupone, C.A.; O'Connor, M.P.; Rosen, G.L.; Knight, R.; Kilham, S.S.; Russell, J.A. Environmental and ecological factors that shape the gut bacterial communities of fish: A meta-analysis. *Mol. Ecol.* 2012, 21, 3363–3378.
 52. Wong, S.; Rawls, J.F. Intestinal microbiota composition in fishes is influenced by host ecology and environment. *Mol. Ecol.* 2012, 21, 3100–3102.
 53. Maruska, K.P.; Fernald, R.D. Social regulation of gene expression in the hypothalamic-pituitary-gonadal axis. *Physiology* 2011, 26, 412–423.
 54. Zhang, Z.; Zhu, B.; Ge, W. Genetic analysis of zebrafish gonadotropin (FSH and LH) functions by TALEN-mediated gene disruption. *Mol. Endocrinol.* 2015, 29, 76–98.
 55. Selvaraj, S.; Chidambaram, P.; Ezhilarasi, V.; Kumar, P.P.; Moses, T.; Antony, C.; Ahilan, B. A Review on the Reproductive Dysfunction in Farmed Finfish. *Annu. Res. Rev. Biol.* 2021, 36, 65–81.
 56. Abellan, E.; Basurco, B. Marine Finfish Species Diversification: Current Situation and Prospects in Mediterranean Aquaculture; CIHEAM-IAMZ: Zaragoza, Spain; FAO: Rome, Italy, 1999.
 57. Monbrison, D.D.; Tzchori, I.; Holland, M.C.; Zohar, Y.; Yaron, Z.; Elizur, A. Acceleration of gonadal development and spawning induction in the Mediterranean grey mullet, *Mugil cephalus*: Preliminary studies. *Isr. J. Aquac.* 1997, 49, 214–221.
 58. Mylonas, C.C.; Magnus, Y.; Klebanov, Y.; Gissis, A.; Zohar, Y. Reproductive biology and endocrine regulation of final oocyte maturation of captive white bass. *J. Fish. Biol.* 1997, 51, 234–250.
 59. Mylonas, C.C.; Woods, L., III; Zohar, Y. Cyto-histological examination of post-vitellogenesis and final oocyte maturation in captive-reared striped bass. *J. Fish. Biol.* 1997, 50, 34–49.
 60. Mylonas, C.C.; Fostier, A.; Zanuy, S. Broodstock management and hormonal manipulations of fish reproduction. *Gen. Comp. Endocrinol.* 2010, 165, 516–534.
 61. Podhorec, P.; Kouřil, J. Hypothalamic factors (GnRH and DA) and their utilization to elimination of reproductive dysfunction in Cyprinidae fish (a review). *Bull. VÚRH Vodňany* 2009, 45, 10–17.
 62. Zohar, Y.; Mylonas, C. Endocrine manipulations of spawning in cultured fish: From hormones to genes, Reproductive Biotechnology in Finfish Aquaculture. *Aquaculture* 2001, 197, 99–136.
 63. Bromage, N.; Jones, J.; Randall, C.; Thrush, M.; Davies, B.; Springate, J.; Duston, J.; Barker, G. Broodstock management, fecundity, egg quality and the timing of egg production in the rainbow trout (*Oncorhynchus mykiss*). *Aquaculture* 1992, 100, 141–166.
 64. Hutchinson, T.H.; Ankley, G.T.; Segner, H.; Tyler, C.R. Screening and testing for endocrine disruption in fish—Biomarkers as “signposts,” not “traffic lights,” in risk assessment. *Environ. Health Perspect.* 2006, 114, 106–114.
 65. Grim, K.C.; Wolfe, M.; Hawkins, W.; Johnson, R.; Wolf, J. Intersex in Japanese medaka (*Oryzias latipes*) used as negative controls in toxicologic bioassays: A review of 54 cases from 41 studies. *Environ. Toxicol. Chem.* 2007, 26, 1636–1643.
 66. Pierron, F.; Lorient, S.; Hérouin, D.; Daffe, G.; Etcheverria, B.; Cachot, J.; Morin, B.; Dufour, S.; Gonzalez, P. Transgenerational epigenetic sex determination: Environment experienced by female fish affects offspring sex ratio. *Environ. Pollut.* 2021, 277, 116864.
 67. Li, R.; Yang, L.; Han, J.; Zou, Y.; Wang, Y.; Feng, C.; Zhou, B. Early-life exposure to tris (1, 3-dichloro-2-propyl) phosphate caused multigenerational neurodevelopmental toxicity in zebrafish via altering maternal thyroid hormones transfer and epigenetic modifications. *Environ. Pollut.* 2021, 285, 117471.
 68. Stephens, K.E.; Miaszkowski, C.A.; Levine, J.D.; Pullinger, C.R.; Aouizerat, B.E. Epigenetic regulation and measurement of epigenetic changes. *Biol. Res. Nurs.* 2013, 15, 373–381.
 69. Breton-Larivière, M.; Elder, E.; McGraw, S. DNA methylation, environmental exposures and early embryo development. *Anim. Reprod.* 2019, 16, 465–474.
 70. Jin, B.; Li, Y.; Robertson, K.D. DNA methylation: Superior or subordinate in the epigenetic hierarchy? *Genes Cancer* 2011, 2, 607–617.

71. Moore, L.D.; Le, T.; Fan, G. DNA methylation and its basic function. *Neuropsychopharmacology* 2013, 38, 23–38.
72. Brykczynska, U.; Hisano, M.; Erkek, S.; Ramos, L.; Oakeley, E.J.; Roloff, T.C.; Beisel, C.; Schübeler, D.; Stadler, M.B.; Peters, A.H. Repressive and active histone methylation mark distinct promoters in human and mouse spermatozoa. *Nat. Struct. Mol. Biol.* 2010, 17, 679–687.
73. Wu, S.-F.; Zhang, H.; Cairns, B.R. Genes for embryo development are packaged in blocks of multivalent chromatin in zebrafish sperm. *Genome Res.* 2011, 21, 578–589.
74. Statello, L.; Guo, C.-J.; Chen, L.-L.; Huarte, M. Gene regulation by long non-coding RNAs and its biological functions. *Nat. Rev. Mol. Cell Biol.* 2021, 22, 96–118.
75. Bizuayehu, T.T.; Babiak, I. MicroRNA in teleost fish. *Genome Biol. Evol.* 2014, 6, 1911–1937.
76. Pauli, A.; Rinn, J.L.; Schier, A.F. Non-coding RNAs as regulators of embryogenesis. *Nat. Rev. Genet.* 2011, 12, 136–149.
77. Andreassen, R.; Worren, M.M.; Høyheim, B. Discovery and characterization of miRNA genes in Atlantic salmon (*Salmo salar*) by use of a deep sequencing approach. *BMC Genom.* 2013, 14, 482.
78. Juanchich, A.; Bardou, P.; Rué, O.; Gabillard, J.-C.; Gaspin, C.; Bobe, J.; Guiguen, Y. Characterization of an extensive rainbow trout miRNA transcriptome by next generation sequencing. *BMC Genom.* 2016, 17, 164.
79. Nilsson, E.E.; Skinner, M.K. Environmentally induced epigenetic transgenerational inheritance of reproductive disease. *Biol. Reprod.* 2015, 93, 145.
80. Heard, E.; Martienssen, R.A. Transgenerational epigenetic inheritance: Myths and mechanisms. *Cell* 2014, 157, 95–109.
81. Donelson, J.M.; Wong, M.; Booth, D.J.; Munday, P.L. Transgenerational plasticity of reproduction depends on rate of warming across generations. *Evol. Appl.* 2016, 9, 1072–1081.
82. Navarro-Martín, L.; Viñas, J.; Ribas, L.; Díaz, N.; Gutiérrez, A.; Di Croce, L.; Piferrer, F. DNA methylation of the gonadal aromatase (*cyp19a*) promoter is involved in temperature-dependent sex ratio shifts in the European sea bass. *PLoS Genet.* 2011, 7, e1002447.
83. Campos, C.; Valente, L.; Conceição, L.; Engrola, S.; Fernandes, J. Temperature affects methylation of the myogenin putative promoter, its expression and muscle cellularity in Senegalese sole larvae. *Epigenetics* 2013, 8, 389–397.
84. Akhtar, W.; Veenstra, G.J.C. TBP-related factors: A paradigm of diversity in transcription initiation. *Cell Biosci.* 2011, 1, 23.
85. Le Luyer, J.; Laporte, M.; Beacham, T.D.; Kaukinen, K.H.; Withler, R.E.; Leong, J.S.; Rondeau, E.B.; Koop, B.F.; Bernatchez, L. Parallel epigenetic modifications induced by hatchery rearing in a Pacific salmon. *Proc. Natl. Acad. Sci. USA* 2017, 114, 12964–12969.
86. Rodríguez Barreto, D.; García de Leaniz, C.; Verspoor, E.; Sobolewska, H.; Coulson, M.; Consuegra, S. DNA methylation on changes in the sperm of captive-reared fish: A route to epigenetic introgression in wild populations. *Mol. Biol. Evol.* 2019, 36, 2205–2211.
87. Morán, P.; Marco-Rius, F.; Megías, M.; Covelo-Soto, L.; Pérez-Figueroa, A. Environmental induced methylation changes associated with seawater adaptation in brown trout. *Aquaculture* 2013, 392, 77–83.
88. Wang, S.Y.; Lau, K.; Lai, K.-P.; Zhang, J.-W.; Tse, A.C.-K.; Li, J.-W.; Tong, Y.; Chan, T.-F.; Wong, C.K.-C.; Chiu, J.M.-Y.; et al. Hypoxia causes transgenerational impairments in reproduction of fish. *Nat. Commun.* 2016, 7, 12114.
89. Cheng, Y.; Vechtova, P.; Fussy, Z.; Sterba, J.; Linhartová, Z.; Rodina, M.; Tučková, V.; Gela, D.; Samarin, A.M.; Lebeda, I. Changes in phenotypes and DNA methylation of in vitro aging sperm in common carp *Cyprinus carpio*. *Int. J. Mol. Sci.* 2021, 22, 5925.
90. Wang, F.-L.; Yan, L.-X.; Shi, H.-J.; Liu, X.-Y.; Zheng, Q.-Y.; Sun, L.-N.; Wang, D.-S. Genome-wide identification, evolution of DNA methyltransferases and their expression during gonadal development in Nile tilapia. *Comp. Biochem. Physiol. Part B Biochem. Mol. Biol.* 2018, 226, 73–84.
91. Wang, Y.Y.; Sun, L.X.; Zhu, J.J.; Zhao, Y.; Wang, H.; Liu, H.J.; Ji, X.S. Epigenetic control of *cyp19a1a* expression is critical for high temperature induced Nile tilapia masculinization. *J. Therm. Biol.* 2017, 69, 76–84.
92. Ribas, L.; Vanezis, K.; Imués, M.A.; Piferrer, F. Treatment with a DNA methyltransferase inhibitor feminizes zebrafish and induces long-term expression changes in the gonads. *Epigenet. Chromatin* 2017, 10, 59.
93. Skafnesmo, K.O.; Edvardsen, R.B.; Furmanek, T.; Crespo, D.; Andersson, E.; Kleppe, L.; Taranger, G.L.; Bogerd, J.; Schulz, R.W.; Wargelius, A. Integrative testis transcriptome analysis reveals differentially expressed miRNAs and their mRNA targets during early puberty in Atlantic salmon. *BMC Genom.* 2017, 18, 801.
94. Domingues, W.B.; Silveira, T.L.; Nunes, L.S.; Blodorn, E.B.; Schneider, A.; Corcine, C.D.; Varela Junior, A.S.; Acosta, I.B.; Kütter, M.T.; Greif, G.; et al. GH Overexpression alters spermatogenic cells microRNAome profile in transgenic zebrafish. *Front. Genet.* 2021, 12, 1712.
95. Gay, S.; Bugeon, J.; Bouchareb, A.; Henry, L.; Delahaye, C.; Legeai, F.; Montfort, J.; Le Cam, A.; Siegel, A.; Bobe, J.; et al. MiR-202 controls female fecundity by regulating medaka oogenesis. *PLoS Genet.* 2018, 14, e1007593.
96. Aydin, F.; Şehriban, Ç.-Y. Effect of probiotics on reproductive performance of fish. *Nat. Eng. Sci.* 2019, 4, 153–162.

97. Salam, M.A.; Islam, M.; Paul, S.I.; Rahman, M.; Rahman, M.L.; Islam, F.; Rahman, A.; Shaha, D.C.; Alam, M.S.; Islam, T. Gut probiotic bacteria of *Barbonymus gonionotus* improve growth, hematological parameters and reproductive performances of the host. *Sci. Rep.* 2021, 11, 10692.
98. Dimitroglou, A.; Merrifield, D.L.; Carnevali, O.; Picchietti, S.; Avella, M.; Daniels, C.; Güroy, D.; Davies, S.J. Microbial manipulations to improve fish health and production—A Mediterranean perspective. *Fish Shellfish Immunol.* 2011, 30, 1–16.
99. Jayasankar, V.; Tomy, S.; Wilder, M.N. Insights on molecular mechanisms of ovarian development in decapod crustacea: Focus on vitellogenesis-stimulating factors and pathways. *Front. Endocrinol.* 2020, 11, 1790–1796.
100. Gioacchini, G.; Giorgini, E.; Merrifield, D.L.; Hardiman, G.; Borini, A.; Vaccari, L.; Carnevali, O. Probiotics can induce follicle maturational competence: The *Danio rerio* case. *Biol. Reprod.* 2012, 86, 65.
101. Gao, Y.; Liu, J.; Wang, X.; Liu, D. Genetic manipulation in zebrafish. *Sheng Wu Gong Cheng Xue Bao Chin. J. Biotechnol.* 2017, 33, 1674–1692.
102. Koulis, S.; Kramer, C.R.; Grier, H.J. Organization of the male gonad in a protogynous fish, *Thalassoma bifasciatum* (Teleostei: Labridae). *J. Morphol.* 2002, 254, 292–311.
103. Sumon, T.A.; Hussain, H.M.A.; Sumon, M.A.A.; Jang, W.J.; Guardiola, F.A.; Sharifuzzaman, S.M.; Brown, C.L.; Lee, E.W.; Kim, C.H.; Hasan, M.T. Functionality and prophylactic role of probiotics in shellfish aquaculture. *Aquac. Rep.* 2022, 25, 101220.
104. Sumon, M.A.A.; Sumon, S.T.A.; Hussain, M.A.; Lee, S.J.; Jang, W.J.; Sharifuzzaman, S.M.; Brown, C.L.; Lee, E.W.; Hasan, M.T. Single and multi-strain probiotics supplementation in commercially prominent finfish aquaculture: Review of the current knowledge. *J. Microbiol. Biotechnol.* 2022, 32, 681–698.
105. Volkoff, H.; London, S. Nutrition and reproduction in fish. *Encycl. Reprod.* 2018, 9, 743–748.
106. Giri, S.S.; Yun, S.; Jun, J.W.; Kim, H.J.; Kim, S.G.; Kang, J.W.; Kim, S.W.; Han, S.J.; Sukumaran, V.; Park, S.C. Therapeutic effect of intestinal autochthonous *Lactobacillus reuteri* P16 against waterborne lead toxicity in *Cyprinus carpio*. *Front. Immunol.* 2018, 9, 1824.
107. Gioacchini, F.; Lombardo, F.; Merrifield, D.; Silvi, S.; Cresci, A.; Avella, M.; Carnevali, O. Effects of probiotic on zebrafish reproduction. *J. Aquac. Res. Dev.* 2011, S1, 1–6.
108. Grunwald, D.J.; Eisen, J.S. Headwaters of the zebrafish—Emergence of a new model vertebrate. *Nat. Rev. Genet.* 2002, 3, 717–724.
109. Ye, M.; Chen, Y. Zebrafish as an emerging model to study gonad development. *Comput. Struct. Biotechnol. J.* 2020, 18, 2373–2380.
110. Scaramuzzi, R.J.; Campbell, B.K.; Downing, J.A.; Kendall, N.R.; Khalid, M.; Muñoz-Gutiérrez, M.; Somchit, A. A review of the effects of supplementary nutrition in the ewe on the concentrations of reproductive and metabolic hormones and the mechanisms that regulate folliculogenesis and ovulation rate. *Reprod. Nutr. Dev.* 2006, 46, 339–354.
111. Xia, Y.; Yu, E.; Lu, M.; Xie, J. Effects of probiotic supplementation on gut microbiota as well as metabolite profiles within Nile tilapia, *Oreochromis niloticus*. *Aquaculture* 2020, 527, 735428.
112. Ohga, H.; Selvaraj, S.; Matsuyama, M. The roles of kisspeptin system in the reproductive physiology of fish with special reference to chub mackerel studies as main axis. *Front. Endocrinol.* 2018, 9, 147.
113. Yu, G.; Zhang, D.; Liu, W.; Wang, J.; Liu, X.; Zhou, C.; Gui, J.; Xiao, W. Zebrafish androgen receptor is required for spermatogenesis and maintenance of ovarian function. *Oncotarget* 2018, 9, 24320–24334.
114. Valcarce, D.G.; Riesco, M.F.; Martínez-Vázquez, J.M.; Robles, V. Diet supplemented with antioxidant and anti-inflammatory probiotics improves sperm quality after only one spermatogenic cycle in zebrafish model. *Nutrients* 2019, 11, 843.
115. Carnevali, O.; Avella, M.; Gioacchini, G. Effects of probiotic administration on zebrafish development and reproduction. *Gen. Comp. Endocrinol.* 2013, 188, 297–302.
116. Qin, C.; Xu, L.; Yang, Y.; He, S.; Dai, Y.; Zhao, H.; Zhou, Z. Comparison of fecundity and offspring immunity in zebrafish fed *Lactobacillus rhamnosus* CICC 6141 and *Lactobacillus casei* BL23. *Reproduction* 2014, 147, 53–64.
117. Gioacchini, F.; Maradonna, F.; Lombardo, F.; Bizzaro, D.; Olivetto, I.; Carnevali, O. Increase of fecundity by probiotic administration in zebrafish (*Danio rerio*). *Reproduction* 2010, 140, 953–959.
118. Major, K.M.; DeCourten, B.M.; Li, J.; Britton, M.; Settles, M.L.; Mehinto, A.C.; Connon, R.E.; Brander, S.M. Early life exposure to environmentally relevant levels of endocrine disruptors drive multigenerational and transgenerational epigenetic changes in a fish model. *Front. Mar. Sci.* 2020, 7, 471.