1. INTRODUCTION

Fish exposure to different endocrine disrupting compounds (EDCs), both chemical and phytochemical, can occur through contaminated waste in surface water and coastal areas. In particular, phytoestrogens (flavonoids and non-flavonoids) originating from various anthropogenic activities (e.g. livestock, agriculture, aquaculture), with variable levels of isoflavones, metabolites and other phytochemicals, have been recorded in aquatic environments ranging from nanomolars to 0.20 mg l\(^{-1}\) (Kiparissis et al. 2001, Spengler et al. 2001, Rocha et al. 2013, Rearick et al. 2014). In terms of hormonal modulation and endocrine disruption in fish, it has been widely noted that all stages of development and life history, from embryo to adulthood, constitutively express the majority of mRNA transcripts related to hatching, reproduction and growth. All these crucial processes are modulated through oestrogen: apoptotic, proliferative and epigenetic signaling pathways, among others. It has also been reported that all fish life stages are sensitive and respond differently to both chemical and phytochemical EDCs (Pelissero et al. 1991, Kiparissis et al. 2003, Segner et al. 2003, Sassi-Messai et al. 2009, Schiller et al. 2013, Sarasquete et al. 2017, 2018a,b, 2019).

Phytochemical isoflavones act as selective oestrogen receptor modulators (SERMs), that is, as ligands (agonists or antagonists) of oestrogen receptors (ERs). This function depends on the coactivators and
corepressors present in the cells due to cross-talks between ERs and other transcription factors, and varies according to the basal levels of endogenous oestrogens (An et al. 2001, Setchell et al. 2005, Lecomte et al. 2017, Xiao et al. 2018). Unlike mammals, many species of fish harbor 3 ER genes. In zebrafish, the genes of these oestrogen receptors are designated ER1 (ERα), ER2b (ERβ1) and ER2a (ERβ2). Basal constitutive levels of ERs (ERα and β) are expressed at all life history stages of fish and show highly variable levels of transcription in both reproductive and non-reproductive organs. In addition, ERs have different binding profiles than natural and exogenous ligands, which vary according to development and life history stage. ERs also show selective and differential distribution in the organs and tissues of adult fish and also differ between males and females, as has been described extensively in zebrafish and other species (Bardet et al. 2002, Menuet et al. 2002, 2004, Chandrasekar et al. 2010, Lu et al. 2017).

Isoflavones (genistein, daidzein) are unique, in that these phytochemicals have a higher and more selective binding affinity for ERβ than for ERα. Most of the isoflavonoid phytoestrogens, which act as SERMs, also show a higher binding affinity for ERβ compared to many chemical endocrine disruptors and in contrast to the endogenous ligand (17β-oestradiol), which in mammals shows a very high binding affinity for both receptors (ERα and ERβ). In fish, in contrast to many higher vertebrates, a lower binding affinity of ERs (ERα and β) for the natural ligand, oestradiol, has also been demonstrated (Kuiper et al. 1998, Miyahara et al. 2003, Setchell et al. 2005, Patisaul & Adewale 2009, Nelson & Habibi 2013). Interestingly, in a recent in vitro study in zebrafish, genistein exhibited higher binding affinity for ERα than for ERβ1 or β2 (Sassi-Messai et al. 2009). In a different study with medaka Oryzias latipes and zebrafish embryos exposed to different doses of genistein, different and variable expression patterns were described in the transcription levels of the ERα and ERβ receptors which also varied by species and with the phytoestrogen dose (Schiller et al. 2013). Due to the highly conserved sequences and very high structural homologies (DNA-binding domain [DBD] and ligand binding domain [LBD]) between ERs in humans (ERα, β) and zebrafish (ERα, β1, β2) (Menuet et al. 2002, Chandrasekar et al. 2010, Schaaf 2017, Asnake et al. 2019), and also because of the higher expected affinity of genistein and daidzein for ERβ, in this work we analyzed the levels of expression of ERβ transcripts (β2 + β1 as a whole) in zebrafish, as has been done in studies with other fish species (Legler et al. 2000, Arukwe et al. 2008, Reyhanian et al. 2014, Sarasquete et al. 2017, 2018a, 2019).

In zebrafish and other fish species, reduced fertilisation, decreased hatching rates and reproductive imbalances have been described as the most common endocrine disorders (and/or the most toxic effects) that can be induced by endogenous factors (e.g. sex hormones) and by isoflavonoid phytoestrogens (Bennetou-Pelissero et al. 2001, Kiparissis et al. 2003, Kim et al. 2009, Sassi-Messai et al. 2009, Schiller et al. 2013, Sarasquete et al. 2018a). Hatching enzymes (HEs) are composed of the astacin–protease domain, and these zinc-dependent metalloproteases possess the ability to digest the envelope of the egg or chorion. While HEs (e.g. choriolysins) digest the egg's protein envelope to facilitate hatching, alveoline, which is a metalloprotease released from cortical alveoli of early vitellogenic oocytes, induces reassembly of chorionegens to harden the egg envelope at the time of fertilization. Some metalloproteases that are present in gametes are essential to ensure reproductive success, by facilitating the passage and penetration of sperm during fertilisation (Yamagami, 1996, Sano et al. 2008, Kawaguchi et al. 2012, Shibata et al. 2012, Nagasawa et al. 2015, Chakraborty et al. 2017).

Phytoestrogenic isoflavones have also been associated with the developmental processes of proliferation, apoptosis and sexual differentiation, as well as with chromatin remodelling, DNA repair and epigenetic mechanisms (Patisaul & Adewale 2009, Lecomte et al. 2017, Xiao et al. 2018). In mammals, the bromodomain testis-specific gene (BRDT) is an essential factor during spermatogenesis and for male fertility. It is widely known that BRDT proteins bind to hyper-acetylated chromatin during sperm elongation, and that these bromodomains recruit the machinery to remove acetylated histones which are replaced by proteasomes. In addition, bromodomain and extraterminal domain (BET) proteins can also maintain this structural role and participate in chromatin remodelling during oogenesis (Paillisson et al. 2007, Berkovits & Wolgemuth 2013, Bao & Bedford 2016, Taniguchi 2016). In fish, the role of the BRDT gene and proteins, which play a key role in hatching, development and differentiation, as well as embryogenesis and gametogenesis of males and females, is much less known and appears to be much more complex and variable than in other vertebrates (Úbeda-Manzanares et al. 2016, Sarasquete et al. 2018a). This complexity is potentially due to different strategies and reproductive behaviors, environmental and hormonal sensi-

The zebrafish *Danio rerio* is a small tropical freshwater cyprinid and an experimental model widely accepted in different fields of biomedical and animal research. The existence of a syntenic relationship between the human and zebrafish genomes (www. ensembl.org, *D. rerio*, genome release Zv9genomes) has been demonstrated, and both genomes can be compared (Mayden et al. 2007, Howe et al. 2013). In addition, due to its well-known embryogenesis and larval development, easy reproduction, sexual dimorphism, high fertility and short life cycle, all stages of zebrafish development and life history have been used in research disciplines such as embryogenesis, larval development, sexual differentiation, reproduction, environmental control, endocrine disruption and ecotoxicology (Selman et al. 1993, Kimmel et al. 1995, Segner 2009, Santos et al. 2017, Wang et al. 2019, and many others).

In the present study, our objective was to provide substantial evidence of the suitability of using a double transcriptional and cellular approach to evaluate the effects of 2 phytoestrogenic isoflavones. This research is mainly focussed on oestrogenic and choriolytic signals and transcriptional coregulation pathways. For this purpose, basal expression patterns and transcriptional variations of 3 molecular signals (ERβ [= β1 + β2, as a whole], HE1 and BRDT) were analysed in different organs and tissues (gonads, head portion, skeletal musculature and digestive system) of male and female zebrafish exposed to a non-toxic concentration of soya isoflavones (daidzein and genistein). Furthermore, in both gonads the histomorphological and histochemical basal patterns and possible cellular alterations induced by both phytochemicals were examined.

2. MATERIALS AND METHODS

2.1. Biological samples

Adult male and female zebrafish *Danio rerio* (genotype AB/Tübingen) were maintained at the Institute of Marine Sciences of Andalusia (ICMAN-CSIC, Puerto Real, Spain). The facilities of ICMAN-CSIC are in compliance with the European Convention for the Protection of Animals used for Experimental and Scientific Purposes, and the fish were approved for experimentation by the Ministry of Agriculture and Fisheries (REGA-ES110280000311) in accordance with current EU (Directive 2010/63/EU) and Spanish legislation. The experimental procedure (projects AGL2014-52906-R; RTI2018-0939-B-100) was approved by the Spanish National Research Council (CSIC) Ethics Committee, and by the Spanish Competent Authority, Junta de Andalucía (no. 09-7-15-278, RD53/2013; 832/2019).

2.2. Toxicity tests

Genistein (C15H10O5, LC Laboratories) and daidzein (C15H10O4, LC Laboratories) were dissolved in ethanol to make up 20 and 5 mM stock solutions, respectively, and then kept in the dark at 4°C until use. Mixed stock of 3−4 mo old male and female zebrafish *D. rerio* (AB) were kept in an automatic flow-through system supplied with recirculating filtered water at 27−28°C under 14 h light:10 h dark photoperiod cycle, keeping the physical–chemical parameters stable (Zebrafish Housing System from Aquatic Habitats B26U-2-A). For experimental trials, fish (n = 60; 1:1 sex ratio) with a standard total length between 3.5 and 4.0 cm were randomly distributed in 2.5 l glass tanks (n = 6 group−1) and fed daily with commercial pellets. Duplicates of controls (with and without ethanol) and triplicates of the experimental groups were used and analysed. For exposure tests to both isoflavones, a nominal concentration of 10 mg l−1 was chosen, taking into account toxicity parameters and non-harmful effects or responses that were calculated in zebrafish embryos and early larvae (Sarasquete et al. 2018a). During the 2 wk experimental period, under semi-static tests, 50% of the water was renewed daily with the corresponding proportion of freshly prepared stock solutions of both isoflavones. At the end of the experimental tests (15 d), all fish were anesthetised with an overdose of phenoxyethanol, and samples of the different organs and tissues were taken for molecular analysis (gonads, head, digestive system, skeletal musculature) and for histological–histochemical studies (ovary, testis), following similar procedures and standardised technical protocols conducted in recent research on zebrafish and other fish species (Sarasquete et al. 2017, 2018a, 2019).

2.3. Nucleic acid extraction and quantification of mRNA expression levels

For molecular analysis, triplicate samples were preserved into RNAlater (Sigma-Aldrich) for 24 h at
4°C and stored at −80°C. Total RNA was isolated from different organs and tissues (gonads, head, digestive system and musculature) using a RNeasy® Micro kit (Qiagen). Genomic DNA was removed via DNase digestion at 37°C for 30 min. Total RNA quality was verified (Bioanalyzer 2100; Agilent Technologies) and concentration was assessed by spectrophotometry ($A_{260}/A_{280}$ nm). For cDNA synthesis, total RNA (500 ng) was utilised for reverse transcription using a qScript™ cDNA Synthesis kit (BioRad). Real-time analysis was performed (Mastecycler ep gradient S Realplex2). Each reaction was carried out in triplicate with a mixture containing each of the specific primer pair (Table 1), cDNA (~10 ng), and 5 µl iTaq™ Universal SYBR® Green SuperMix (BioRad). Relative gene quantification was performed using the method of Pfaffl (2001), and results were normalised to elongation factor 1 alpha 1 (eEf1a1) and 18S rRNA, as recently described by Sarasquete et al. (2018a, 2019).

### 2.4. Statistical analysis

Temporal differences for each of the gene expression patterns and both treatments were tested by 1-way ANOVA performed after a base 10 logarithmic transformation to meet requirements for the parametric test. Normality was checked using Shapiro-Wilk’s test and homogeneity of variances with the Levene’s test. Tukey’s post hoc test was used to identify significantly different groups. Differences were considered statistically significant at $p < 0.05$. Statistical analyses of data from qPCR were performed using SPSS v.23.0.0.0 software (IBM).

### 2.5. Histological and histochemical approaches

Samples of both gonads were fixed in 4% v/v neutral formaldehyde (pH: 7.2) in sodium phosphate buffer (0.1 M) and embedded in paraffin wax. In the serial sections (5–6 µm), histomorphological staining techniques were used, such as Haematoxylin-eosin and Haematoxylin-VOF (Light green, Orange G, Acid Fuchsin) and Type III-G.S (Sarasquete & Gutierrez 2005). To analyse the cellular distribution of carbohydrates and proteins, several histochemical techniques were performed: PAS and diastase-PAS (neutral-glycoconjugates, glycoproteins, glycogen); Alcian Blue at pH 2.5, 1, and 0.5 (carboxylated and sulphated glycoconjugates); Mercury-Bromophenol Blue/Hg-BFB (general proteins); Ninhydrin-Schiff (proteins rich in lysine, NH$_2$); Ferric-Ferricyanide/ Thioglycollate reduction (cysteine and cystine; -SH and -S-S groups); 1,2-Naftoquinone-4-sulphonate sodium/NQS (arginine); P-dimethylaminobenzaldehyde/ DMAB-nitrite (tryptophan); and Hg-sulphate-sulphuric acid-sodium nitrate/Millon reaction (tyrosine). For the characterization of different glucidic residues bound to the glycoconjugates, several peroxidase-HPR-conjugated lectins were used, extracted from *Canavalia ensiformis* (ConA; Man/Glc), *Ulex europaeus* (UEA-I; Fuc), *Triticum vulgaris* (WGA; βGlcNAc >> NeuNAc/sialic acids/NANA), *Glycine max* (SBA; α/β GalNAc) and *Sambucus nigra*.
Lectin concentrations ranged between 15 and 30 µg ml−1, and specific sugar inhibitors were also used. The peroxidase activity was visualised with 3,3′-diaminobenzidine tetra hydrochloride/DAB and hydrogen peroxide (0.05%). All techniques were performed according to Pearse (1985), and following previously standardised techniques and protocols (Sarasquete et al. 2002). All reagents were purchased from Sigma-Aldrich.

3. RESULTS

ERβ, HE1 and BRDT were expressed in variable and differential ways in both male and female zebrafish gonads as well as in the different body regions studied. The highest levels of constitutive basal expression of ERβ were recorded in the testis >> digestive system > ovary = musculature = head portion (Fig. 1A). In parallel, HE1 transcripts showed the following basal expression patterns: testis >>> ovary > skeletal musculature > head = digestive system (Fig. 1B). Furthermore, the basal levels of the BRDT transcripts were differentially expressed as follows: ovary >> testis > head = musculature > digestive system (Fig. 1C).

The isoflavone genistein down-regulated the transcript levels of ERβ in the ovary and digestive system, and up-regulated ERβ levels in the skeletal musculature. Daidzein increased ERβ expression levels in the head portion and skeletal musculature. Nevertheless, no significant differences were observed in the basal expression levels of ERβ in the testes due to the effect of both isoflavones. On the other hand, baseline transcription levels of HE, which were expressed at a higher level in the testes than in the ovaries, were up-regulated in both gonads due to genistein exposure. Daidzein only increased the levels of HE1 expression in skeletal musculature. This isoflavone increased transcription levels of BRDT in the testes, and

![Fig. 1. Relative expression levels of (A) oestrogen receptor transcript (ERβ), (B) hatching enzyme (HE1) and (C) bromodomain testis-specific gene (BRDT) in controls and male and female zebrafish exposed to both isoflavones. Bars: means ± SE (n = 6 × 3) of the relative expression level of the gene for controls, and treatment with 10 mg l−1 daidzein and 10 mg l−1 genistein. Asterisks: significant differences between treatment group controls. Lowercase letters: significant differences between the mean values of each organ for the same treatment (1: control; 2: daidzein; 3: genistein).]
these bromodomains were down-regulated in the digestive system. Finally, genistein up-regulated the expression levels of BRDT in the head portion, and these BRDT transcripts were down-regulated in the digestive system (Fig. 1).

As illustrated briefly in Fig. 2, the male and female zebrafish gonads were in advanced stages of gametogenesis. Thus, in the ovaries the presence of pre-vitellogenic basophilic oocytes and acidoephilic oocytes was observed in the different stages of vitellogenesis, as well as the presence of some residual post-ovulatory follicles. In the testes, all the cells of the spermatogenic line were also present, distributed along the seminiferous tubules and contained cysts of spermatogonia, spermatocytes and spermatids in the testicular epithelia, as well as groups of spermatozoa released in the lumen of the tubules and seminiferous ducts. In the ovary, the cortical alveoli, which appeared at the periphery of the early vitellogenic oocytes, contained a moderate amount of carboxylated glycoconjugates and abundant neutral glycoproteins. As vitellogenesis progressed, acidoephilic yolk-granules and the characteristic zona radiata were evident, and both structures contained abundant proteins rich in all amino acids studied along with neutral glycoproteins. The yolk granules showed a remarkable staining affinity for all the lectins studied (ConA, UEA, WGA, SBA, SNA), which indicates great richness in glycoconjugates containing different sugar residues, such as mannose/glucose, fucose, N-acetyl galactosamine/glucosamine and sialic acids, whereas cortical alveoli and the zona radiata showed specific higher affinity for the WGA lectin (β-GlcNAc; NeuNAc). In the testes, most spermatogenic cells contained proteins rich in all amino acids studied, with marked increases in proteins rich in basic amino acids (i.e.arginine and lysine) and cysteine, especially in the spermatozoa. Most sugar residues bound to glycoconjugates increased, especially in spermatids and spermatozoa, as summarised in Table 2.

Finally, no histomorphological–histochemical modifications were observed in either the male or female gonads after exposure to both soya isoflavones. In addition, over the course of the 2 wk experiment, no significant variations were detected in the biometric parameters of the 3–4 mo old male and female zebrafish exposed to both phytoestrogens (average weights between 0.38 ± 0.03 and 0.40 ± 0.02 g) compared to the controls groups, and no mortalities were recorded.

4. DISCUSSION

The present study showed variable and differential basal expression patterns in the 3 genes studied (ERβ, HE1, and BRDT) according to sex and the different organs and tissues of the zebrafish that were analysed. Interestingly, basal constitutive transcript patterns for the 3 genes were quite unexpected in both gonads. In fact, for both transcript patterns (ERβ and HE1), higher basal expression levels were recorded in the testes than in the ovaries, while basal expression levels of BRDT were higher in the ovaries than in the testes. Similarly, exposure to the same non-toxic dose of both soya isoflavones (genistein and daidzein, at 10 mg l−1 for 2 wk) induced differential and variable transcriptional responses depending on the organs and tissues analysed (e.g. gonads, head, digestive system and skeletal musculature) and according to sex and the type of isoflavone provided. Most genes involved in reproduction, hatching, development and epigenetic processes (e.g. oestrogens and receptors, choriogenins, HEs, apoptotic and epigenetic markers, bromodomains) express constitutively at high or moderate levels in the
gonads and during the early stages of development and life, as has been reported in different species of fish including zebrafish (Segner et al. 2003, Chandrasekar et al. 2010, Úbeda-Manzanaro et al. 2016, Sarasquete et al. 2018a, González-Rojo et al. 2019). All these findings corroborate the maternal origin of mRNA transcripts and their transfer to the progeny, since the oocyte is the starting point of a new generation. As reviewed by Arukwe & Goksøyr (2003), most of the machinery for DNA and protein synthesis necessary for the developing embryo is performed autonomously by fertilised oocytes.

In the present study, as expected, ERs (e.g. ERβ) were expressed constitutively and differentially in most organs and tissues involved in reproductive and non-reproductive functions, displaying higher transcript levels for ERβ in the testes than in the ovaries. Several authors have also pointed out that the 3 ERs (α, β1, β2) overlap and are expressed mainly in the brain, pituitary, liver and gonads of zebrafish, displaying much lower basal expression levels for the ERs in the ovaries than in the testes. Furthermore, zebrafish males and females show a differential and opposite binding affinity for the different endogenous and exogenous ligands, indicating a sexually divergent mechanism and sex-specific control of ERs (Menuet et al. 2002, 2004, Chandrasekar et al. 2010, Lu et al. 2017, Schaaf 2017). Interestingly, some remarkable differences were observed in zebrafish compared to other species of fish, such as the hermaphrodite seabream Sparus aurata, which showed higher basal expression levels of ERβ in the ovaries and for ERα in the testes. In addition, very high basal expression levels have also been described for the 3 ERs (ERα, βs) in seabream testes, which corroborates the crucial role of oestrogens and their receptors for reproduction in both sexes. In parallel, in zebrafish and other fish species, moderately high basal expres-

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Table 2. Histochemical characteristics in gonads of zebrafish controls and exposed to isoflavones. Semi-quantitative data determined by 4 independent observers, showing different staining intensities and affinities to biological compounds, from negative (−) to weak, moderate and strongly positive (from 1+ to 3+). CA: cortical alveoli; YG: yolk granules; ZR: zona radiata; Spg: spermatogonia; Spc: spermatocyte; Spd: spermatid; Spz: spermatozoid.
sion levels for the ERs (i.e. ERα, βs) have been recorded in many other non-reproductive organs and tissues, such as the digestive system, muscles, gills, heart, kidney and eyes (Socorro et al. 2000, Pinto et al. 2006, Chandrasekar et al. 2010, Nelson & Habibi 2013). As has already been extensively reported, the baseline expression patterns of ERβ, which are constitutively expressed at a much higher level in the testes than in the ovaries of zebrafish and other fish species, allows corroboration of the obvious function of ERs (ERα, βs) during gametogenesis, and their involvement in the reproductive functions of females and males. In this context, it is appropriate to remember that oestrogens, which are commonly known as ‘female hormones’ and are studied more and better known during oogenesis, are also key actors in spermatogenesis. In fact, through genomic and non-genomic pathways via ERs (ERα, βs), including the pathway to the transmembrane G-protein coupled oestrogen receptors (GPERs), oestrogen receptors participate in reproductive functions. In particular, through the genomic signaling pathway, ERs play a fundamental role in the development, differentiation and maturation of female and male germ cells, as well as in the survival of spermatozoa. Recently, it has also been pointed out that GPER = GPR30 (i.e. a non-genomic pathway) is not necessary for the determination of sex, development or function of the ovary or fertility in female zebrafish (Leal et al. 2009, Schulz et al. 2010, Hao et al. 2013, Nelson & Habibi 2013, Crowder et al. 2018). Accordingly, the current results and the numerous previous studies allow corroboration of the generalised effects, integrating functions and even autoregulation of ERs (i.e. ERβs vs. ERα) in both gonads, as well as in different organs and somatic tissues.

Similar to oestrogens, soya isoflavones (i.e. genistein and daidzein), as characteristic phytooestrogens (= 17β-oestradiol) or better known as SERMs, can modulate (or interrupt) the gonadal development and reproductive cycle of both sexes (Pelissero et al. 1991, Kiparissis et al. 2003, Patisaul & Adewale 2009, Lecomte et al. 2017, Xiao et al. 2018). In the present study, baseline ERβ expression levels decreased significantly in the ovary (but not in the testes) and in the digestive system (including the liver) due to the action of genistein (at 10 mg l⁻¹). Conversely, ERβ expression levels were not altered by the effect of daidzein in either of the gonads. This isoflavone increased ERβ transcription levels in the head (i.e. brain and pituitary) and in skeletal musculature. In other recent studies, it was reported that both soya isoflavones (at 5 mg l⁻¹) increased ERβ expression levels in fertilised embryos and newly hatched zebrafish larvae. Nevertheless, during the larval development of Senegalese sole, baseline ERβ expression levels were modulated in variable ways due to exposure to both isoflavones, showing a greater induction pattern with daidzein (at 20 mg l⁻¹) that genistein at lower concentrations (3 and 10 mg l⁻¹; Sarasquete et al. 2017, 2018a, 2019). In a different study, in male and female zebrafish exposed to chemical endocrine disruptors (e.g. diethylstilbestrol and 4-nonylphenol), different transcriptional responses were described for the 3 ERs (ERα, ERβ1, β2) in the brain and in both gonads, depending on sex and the circulating basal levels of the natural ligand 17β-oestradiol (Chandrasekar et al. 2010). Recently, it was noted that ER transcript levels (ERα, βs) were not altered in zebrafish males when exposed to bisphenol-A. However, the expression levels of ERs coupled to the transmembrane G protein (GPER-1) were up-regulated by this plasticizer EDC (estrogen-dendrimer conjugate), which induced apoptotic and epigenetic effects in zebrafish testis (González-Rojo et al. 2019). Taken together, the current results and all the previous findings allow us to corroborate the differential and variable oestrogenic (and anti-) responses that can be induced by different endocrine disruptors, including phytoestrogenic isoflavones, through binding to ERs, as demonstrated in the gonads and other brain and somatic organs and tissues. In particular, a decrease in ERβ expression levels was observed in the ovary, but not in the testis, due to the action of genistein. In contrast, daidzein did not induce significant transcriptional effects (through ERβ), in either of the gonads, which is indicative of differential oestrogenic (anti-) effects of both genistein and daidzein in males and females, since a similar dose of each isoflavone was tested, and the male and female zebrafish were in a similar or parallel state of gonadal maturation.

On the other hand, the highest basal expression levels of HE1 were, surprisingly, expressed in the testes, with only moderate expression levels displayed in the ovaries. The lowest levels of HE1 were recorded in skeletal musculature > head = digestive system. Chakraborty et al. (2017) reported that in Japanese anchovy Engraulis japonicus, some isoforms of the HEs (i.e. AcHEs), which are considered to be ovastacin homologs, predominated both in the ovary and testis, and these isoforms showed no expression biased by sex, suggesting a conserved role in other areas unrelated to sex. In addition, some of these metalloproteases play the role of proteolytic agents by digesting the egg-envelope to facilitate...
penetration of spermatozoa and by promoting interaction between both gametes; other metalloproteases which are within the ovary, such as the alveolin peptidase, do not break the egg-envelope, but are instead involved in its hardening upon fertilisation (Yamagami 1996, Sano et al. 2008, Kawaguchi et al. 2012, Shibata et al. 2012, Nagasawa et al. 2015). Interestingly, in zebrafish, HE1 expression levels were up-regulated by exposure to genistein (at 10 mg l⁻¹) in both male and female gonads, while daidzein only increased HE1 expression levels in skeletal musculature. In a recent study, HE1 reached the highest levels of expression in zebrafish embryos (at 24 h post-fertilization, hpf), with a decreasing transcriptional pattern from this embryonic phase to hatching (at 72–96 hpf). In addition, expression levels of HE1 increased with exposure to daidzein (at 20 mg l⁻¹, LOTEC — Low Observable Toxic Effect Concentration — value, at 72 hpf), and this isoflavone accelerated the hatching process in zebrafish, while expression levels of HE1 were not modified by exposure to genistein at 5 mg l⁻¹ (Sarasquete et al. 2018a).

In the present study, by inducing increases in HE1 expression levels, genistein (at 10 mg l⁻¹) could activate or accelerate degradation of the egg or chorion envelope during the maturation of oocytes and/or increase the process of atresia (for example, autophagy or apoptosis), as described in other species due to increases in HEs (Donato et al. 2003, Chakraborty et al. 2017). Taken together, despite certain increases in the expression levels of HE1 and parallel decreases in ERβ levels (which are 2 signals oppositely modulated in the ovary [up−down regulations]) with exposure to genistein, these imbalances did not modify histomorphological or histochemical patterns of the vitelline structures, nor did they influence the maturation of the oocytes. Similarly, the expression levels of HE1 also increased in the testes with exposure to genistein, but neither histomorphological—histochemical alterations nor effects on the development and maturation of spermatogenic cells were observed.

It is widely known that the BRDT protein has 2 N-terminal bromodomains that bind to the tails of acetylated histones, and thus both bromodomain genes and proteins are involved in epigenetic processes, both constitutive and induced (Bao & Bedford 2016, Taniguchi 2016). In zebrafish, basal expression levels of BRDT were higher in the ovaries than in the testes, and moderate or low levels of expression were recorded in other organs and tissues (i.e. head, musculature, digestive system). In another recent study, higher levels of BRDT expression were recorded in fertilised eggs of zebrafish, showing lower expression levels during embryogenesis and hatching of larvae (Sarasquete et al. 2018a). In other fish species, such as the protandrous hermaphrodite Sparus aurata, basal levels of the BRDT transcripts were higher in the mature testes than in the ovaries, and moderately high levels of expression were also recorded in other organs and tissues (i.e. brain, pituitary, musculature, heart, kidney, spleen, intestine, gills), while in the gonochoric sea bass Dicentrarchus labrax, similar and elevated expression levels of BRDT were registered in both gonads, and very low levels of expression were recorded in the other organs and tissues. In addition, a constitutive expression of BRDT has been described in both spermatocytes and spermatids, which could be related to the role of BRDT in the termination of meiosis-I as well as in genome condensation or as a transcriptional regulator during these spermatogenetic states (Úbeda-Manzanaro et al. 2016). Interestingly, Kurtz et al. (2009) reported that histone acetylation (e.g. H4) occurs during fish spermatogenesis. In the gonochoric species D. labrax, the replacement of histones by protamines was observed during spermiogenesis, while in the hermaphrodite S. aurata there was no displacement of these basic nuclear proteins. The inter- and intra-specific differences indicated in BRDT expression levels in different species of fish (including zebrafish in the present study; i.e. higher levels of BRDT in the ovaries than in the testes) suggest a different role of BRDT protein during both spermatogenesis and oogenesis, as a transcriptional regulator and/or for chromatin remodeling and compaction depending on the species, as reported in mammals and fish (Taniguchi 2016, Úbeda-Manzanaro et al. 2016). In the present study, in the zebrafish testes, basal expression levels of BRDT transcripts increased significantly with exposure to daidzein (at 10 mg l⁻¹), but not with exposure to the same dose of genistein. In the ovary, no significant transcriptional changes were recorded due to the effect of either isoflavone. In another recent study, baseline expression levels of BRDT were up-regulated by genistein (at 5 mg l⁻¹, LOEC value, at 96 hpf), but BRDT transcript levels were not altered by exposure to daidzein (up to 20 mg l⁻¹), which is known to be less toxic than genistein, as reported in embryos and larvae of zebrafish at LC50 (lethal concentration 50 %) values of 67.63 mg l⁻¹ (daidzein) and 4.41 mg l⁻¹ (genistein) (Sarasquete et al. 2018a). As previously noted, the BRDT gene and the family of BETs are mainly involved in the modulation of gene transcription through epigenetic mechanisms, between bromo-
domains and acetylated histones (rich in lysine), regulating genome reprogramming during spermato-ogenesis and oogenesis, embryogenesis, sex-differentiation and maintenance of lineage commitment (Taniguchi 2016, Úbeda-Manzanaro et al. 2016). Accordingly, in zebrafish, some of the basic and crucial events that participate in certain structural aspects of gametogenesis, and that depend on the regulation and modulation of the bromodomain gene and proteins, would be enhanced by the effect of the phytoestrogenic daidzein, at least at the transcriptional level as demonstrated in mature testicles (i.e. up-regulation of BRDT). Nevertheless, baseline BRDT expression levels were not altered in the ovaries with exposure to both isoflavones. Therefore, despite the induced transcriptional imbalances (up−down regulations) in the expression levels of the studied signals (ERβ, HE1 and BRDT) in both gonads and in the other organs and tissues, no cellular alterations were observed. In this context, the histomorphological and histochemical characteristics of both gonads (controls and exposed to both isoflavones) are in agreement with numerous previous descriptions and findings. In particular, these gonadal characteristics highlight in the ovary the continuous development and asynchronous growth of germ cells at different stages of oogenesis, which is very characteristic of this species; that is, there is a permanent presence of oogonia, basophilic previtellogenic cells and oocytes in different stages of vitellogenesis and maturation, as well as the presence of post-ovulatory residual follicles and atretic oocytes. In the testis, these characteristics highlight the continuous and synchronous development of cysts of the spermatogenetic cell line, numerous from spermatogonia to spermatozoa, as has been described extensively in zebrafish (Selman et al. 1993, Leal et al. 2009, Gallo & Costantini 2012, Schulz et al. 2010, Deshpande & Pancharatna 2013, Lu et al. 2017).

In conclusion, the soya isoflavones genistein and daidzein differentially modulated the basal expression patterns of genes in the gonads of male and female zebrafish. Genistein resulted in a decrease in ERβ in the ovary but an increase in expression levels of HE1 in both gonads, whereas daidzein only increased the transcript levels of BRDT in the testes. These responses could suggest a greater anti-oestrogenic transcriptional effect (through ERβ) in the gonad of females induced by genistein, which also provokes a higher transcriptional choriolytic effect in both gonads. In addition, increased BRDT transcript levels in the testes caused by the action of daidzein could suggest a greater activating effect in spermatogenesis, chromatin remodelling and/or epigenetic processes. Altogether, these transcriptional responses (up−down regulations) must also be complemented and analysed at hormonal and physiological levels, since no histomorphological−histochemical changes were observed in the zebrafish ovary, as recorded through the pattern of distribution of carbohydrates, proteins and the presence of sugar residues in the content of glycoconjugates, which are detected in cortical alveoli, yolk granules and zona radiata of vitellogenic and mature oocytes. Similarly, no appreciable histomorphological or histochemical changes were observed in the spermatogenetic cells (from spermatogonia to spermatozoa). All these transcriptional and cellular responses, as well as changes at the protein, hormonal and enzymatic levels, should be studied in greater depth throughout the reproductive cycle of females and males, as well as during the process of sexual differentiation of zebrafish exposed to both phytoestrogenic isoflavones.

Acknowledgements. This study was financed by Spanish Projects (MINECO-Feder, AGL2014-52906R, RTI2018-0939- B-100). Grant contract-CSIC (M.U.M). Our thanks to Mrs. M. Carmen Carmona, Mr Juan A. Canales and Mr. J. Antonio Miquel for the excellent technical assistance.

LITERATURE CITED


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