Title: G protein-coupled estrogen receptor regulates heart rate and heart valve thickness in zebrafish

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Abstract: Estrogens act by binding to estrogen receptors alpha and beta (ERα, ERβ), ligand-dependent transcription factors that play crucial roles in sex differentiation, tumor growth and cardiovascular physiology. Estrogens also activate the G protein-coupled estrogen receptor (GPER), however the function of GPER in vivo is less well understood. Here we find that GPER is required for normal heart rate in zebrafish embryos and for normal valve thickness in zebrafish adults. Acute exposure to estrogens increased heart rate in wildtype and in ERα and ERβ mutant embryos but not in GPER mutants. Nuclear estrogen receptor signaling remained normal in GPER mutant embryos. However, GPER mutant embryos exhibited reduced basal heart rate while heart rate was normal in ERα and ERβ mutants. We detected gper transcript in discrete regions of the brain but not in the heart. In the brain, we observed gper expression in cells lacking nuclear estrogen receptor activity, suggesting that GPER acts in the brain to regulate heart rate independently of nuclear estrogen receptor signaling. Additionally, blood flow in embryos has been shown to influence heart valve maturation, suggesting the hypothesis that reduced heart rate during embryonic and juvenile development disrupts heart valve maturation. Consistent with this hypothesis, we find that adult GPER mutants have thinner heart valves than wildtype. Our results demonstrate that estradiol plays a previously unappreciated role in the acute modulation of heart rate during zebrafish embryonic development and that GPER functions as an autonomous estrogen receptor in vivo to regulate basal heart rate and heart valve thickness.
INTRODUCTION

Zebrafish are an established model for human cardiovascular development and function (1) with conserved estrogen signaling (2-4). While studying the function of ERα (esr1) in zebrafish embryonic heart valves (5, 6), we serendipitously observed that estrogen receptor modulators caused acute changes in heart rate. Estrogens bind two classes of receptors: nuclear hormone receptors (ERα, ERβ) that are ligand-dependent transcription factors (7), and the G protein-coupled estrogen receptor (GPER, also known as GPR30), an integral membrane protein (8, 9). It has been difficult to tease apart to what degree ERα and/or ERβ are involved in regulating GPER function in vivo. The observations that ERα can directly activate G proteins in cultured cells (10-13) and that GPER coimmunoprecipitated with ERα in tumor cells (14) has been used to argue that either GPER is dispensable for estrogen-dependent signaling or that GPER mediates interactions between ERα and G proteins (15). Studies using GPER-deficient mice implicate GPER in ventricular hypertrophy (16), regulation of blood pressure and vascular tone (17, 18) and atherosclerosis progression (19), but whether nuclear ER signaling is required for GPER function in these contexts is unknown. Additionally, these studies examined GPER function in adult animals, while the role of GPER during embryonic development is not well understood. Here we use zebrafish embryos, an established model of human development, to reveal a new function for GPER during cardiovascular development.

Estrogen signaling often differs between males and females. However, zebrafish embryos and larvae are bipotential hermaphrodites that have not begun to sexually differentiate before approximately 10 days post fertilization (dpf) (20), meaning that estrogen levels are uniform between age-matched embryos. Additionally, zebrafish embryos develop outside of the mother and not within a confined space, such as the uterus. Therefore, zebrafish embryos are not subject
to local estrogen concentration gradients, as has been reported to occur in rodents depending upon their position *in utero* and their proximity to embryos of the same or opposite sex (21, 22). These developmental traits make zebrafish a powerful model to study how sex hormone signaling influences the formation and function of non-gonadal tissues. Using complementary genetic and pharmacologic approaches, we sought to characterize how estradiol regulates heart rate and determine to what extent each estrogen receptor mediates estradiol-dependent changes in heart rate in zebrafish embryos.

RESULTS

We exposed 49 hour post fertilization (hpf) embryos to 17β-estradiol (estradiol) and assayed heart rate following one hour exposure. We found that estradiol exposure caused an approximately 20% increase in heart rate (Fig. 1, mean difference in heart rate between estradiol and vehicle exposed embryos was 26.51 ± 3.36 (standard error) beats per minute (bpm)). Exposure to progesterone, a structurally similar steroid sex hormone, had no effect on heart rate (Fig. 1, mean difference in heart rate 2.31 ± 6.54), suggesting that the effects on heart rate were specific to estrogens.

To explore whether heart rate was influenced by nuclear estrogen receptor or GPER signaling pathways, we employed a pharmacological approach. We exposed embryos to ICI182,780 (fulvestrant), a well-characterized ERα and ERβ antagonist (23) that also acts as a GPER agonist (8). Following one-hour exposure to ICI182,780, heart rate was significantly increased (Fig. 1, mean difference in heart rate 29.81 ± 4.75 bpm). This effect was blocked by co-administration of G36, a specific GPER antagonist (24) (Fig. 1, mean difference in heart rate 4.10 ± 6.23 bpm), suggesting that estradiol increases heart rate via GPER. We also exposed embryos to G1, a specific GPER agonist with no detectable agonist activity against nuclear estrogen receptors (25), and found that heart rate increased significantly (Fig. 1, mean difference in heart rate 40.98
± 6.35 bpm). Together, our pharmacological results suggest that GPER regulates heart rate acutely.

To definitively test the hypothesis that estradiol regulates heart rate via GPER, we generated GPER mutant embryos, exposed them to estrogen receptor modulators and assayed heart rate. Using CRISPR-Cas technology (26), we generated embryos with a 131 basepair deletion in the gper open reading frame (Fig. 2A, B). Embryos were viable and grossly normal, allowing us to measure heart rate (Fig. 2C, D). We exposed homozygous maternal zygotic gper mutant embryos (MZgper/-) to estradiol or to ICI182,780 and found no increase in heart rate compared to embryos exposed to vehicle (Fig. 2E). Our results demonstrate that estradiol increases heart rate in a GPER-dependent manner. Note that zygotic gper mutants exhibited increased heart rate in response to estradiol (Fig. S1, mean difference in heart rate 29.11 ± 3.56 bpm), indicating that GPER is maternally deposited into oocytes and expressed in embryos. This is consistent with previously published results that detected gper transcript in zebrafish embryos at 1 hour post fertilization, suggesting the presence of maternally loaded gper mRNA (27).

To test whether endogenous estrogens regulate heart rate during embryonic development, we examined basal heart rate in GPER mutant embryos reared in untreated water, reasoning that if heart rate was reduced, then that would suggest that endogenous estradiol regulates heart rate via GPER. We compared heart rate in wildtype versus MZgper/- embryos at 50 hpf and found that MZgper/- embryos had reduced heart rate compared to wildtype (Fig. 2F, mean difference in heart rate between wildtype and mutant -30.80 ± 7.07 bpm). These results demonstrate that GPER is required for normal basal heart rate in embryos and strongly suggest that endogenous estrogens influence heart rate via GPER.

Whether GPER acts as an autonomous estrogen receptor in vivo is controversial. Previous reports suggest that GPER activity might require interaction with nuclear estrogen receptors at the membrane or that estrogens activate GPER indirectly, by binding to nuclear receptors in the cytosol that then
activate downstream proteins, including GPER (15, 28). To determine whether nuclear estrogen receptors influence heart rate, we generated zebrafish with loss-of-function mutations in each nuclear estrogen receptor gene: esr1 (ERα), esr2a (ERβ1) and esr2b (ERβ2) (Fig. S2-S4). All mutant embryos were viable and grossly normal, allowing us to measure heart rate (Fig. S2-S4). To test whether estradiol increases heart rate via nuclear estrogen receptors, we exposed 49 hpf esr1−/−, esr2a−/− and esr2b−/− embryos to estradiol or vehicle for one hour and assayed heart rate. Following estradiol exposure, heart rate was increased in all mutants compared to vehicle control (Fig. 3A, mean difference in heart rate between estradiol and vehicle 25.04 ± 5.68 bpm for esr1−/−, 37.23 ± 7.66 bpm for esr2a−/−, 32.48 ± 1.92 bpm for esr2b−/−), similar to what we observed when wildtype embryos were exposed to estradiol (Fig. 1). These results demonstrate that nuclear estrogen receptors are not necessary for estradiol-dependent increase in heart rate.

To test whether endogenous estrogens regulate heart rate via nuclear estrogen receptors, we bred heterozygous fish to generate embryos homozygous for mutations in either esr1, esr2a or esr2b genes and assayed heart rate in 50 hpf embryos. We observed no significant difference in basal heart rate between homozygotes, heterozygotes or wild type siblings within the same clutch (Fig. 3B, mean difference in heart rate between homozygote and wildtype -4.34 ± 1.37 for esr1, -0.46 ± 3.75 for esr2a, -2.37 ± 3.26 for esr2b; between heterozygote and wildtype -3.34 ± 1.02 for esr1, -0.91 ± 1.53 for esr2a, 0.63 ± 1.66 for esr2b). These results demonstrate that nuclear estrogen receptors are not required for the establishment of normal basal heart rate in embryos.

It is possible that the mutations generated in each nuclear estrogen receptor gene do not cause loss of functional estrogen receptor proteins. To exclude this possibility and show that esr mutants exhibit loss of functional ER proteins, we generated esr mutants on the Tg(5xERE:GFP)c262/c262 transgenic background, where green fluorescent protein (GFP) expression occurs in cells with activated nuclear estrogen receptors (5) (referred to as 5xERE:GFP). Previous studies using whole mount in situ hybridization demonstrated that esr1
is expressed in embryonic heart valves while esr2b is expressed in the liver (6), therefore we hypothesized that mutants would fail to upregulate GFP in tissues where the relevant receptor is normally expressed. We exposed 2-3 day post fertilization (dpf) 5xERE:GFP, 5xERE:GFP;esr1<sup>−/−</sup>, 5xERE:GFP;esr2a<sup>−/−</sup> and 5xERE:GFP;esr2b<sup>−/−</sup> embryos to 100 ng/ml 17β-estradiol overnight and assayed fluorescence. Consistent with esr gene expression patterns, 5xERE:GFP;esr1<sup>−/−</sup> larvae exhibited fluorescence in the liver but not in the heart (Fig. S2), whereas 5xERE:GFP;esr2b<sup>−/−</sup> larvae exhibited fluorescence in the heart but not in the liver (Fig. S4). esr2a transcript was not detected at these embryonic and larval stages (6) and, as expected, we saw no change in fluorescence between 5xERE:GFP and 5xERE:GFP;esr2a<sup>−/−</sup> (Fig. S3). We conclude that the zebrafish nuclear estrogen receptor mutants lack estrogen receptor function.

Deleterious mutations can induce genetic compensation (29), however results from the 5xERE:GFP esr mutants suggest that compensatory expression of esr genes is not occurring. For example, it is possible that in the esr1 mutant there is compensatory upregulation of esr2a and/or esr2b that masks a heart rate phenotype. If esr2a or esr2b were upregulated in esr1 mutants, then we would expect to see fluorescence in the heart in 5xERE:GFP;esr1<sup>−/−</sup> embryos. Instead, we observed no fluorescence in the hearts of 5xERE:GFP;esr1<sup>−/−</sup> embryos (Fig. S2). Similarly, we observed no ectopic fluorescence in 5xERE:GFP;esr2b<sup>−/−</sup> embryos (Fig. S4), suggesting that esr genes are not compensating for one another in the multiple zebrafish esr mutants.

To further test whether nuclear estrogen receptor signaling is influenced by GPER, we generated gper mutants on the 5xERE:GFP transgenic background and asked whether estradiol exposure reduced nuclear estrogen receptor activity in mutants compared to wildtype. Following overnight exposure to estradiol, 3 dpf 5xERE:GFP and 5xERE:GFP;MZgper<sup>−/−</sup> larvae exhibited similar fluorescence (Fig. S5). These results demonstrate that nuclear estrogen receptor transcriptional activity does not require GPER and support the hypothesis that GPER acts as an autonomous estrogen receptor in vivo.
Heart rate can be modulated by cardiomyocytes in the heart, or by cells in the central nervous system, which directly innervates the heart to modulate heart rate and also regulates the release of humoral factors, such as thyroid hormone, that bind to receptors in cardiomyocytes and regulate heart rate (30). To determine whether GPER regulates heart rate tissue autonomously, we performed whole mount in situ hybridization to test whether gper transcripts are expressed in 50 hpf zebrafish embryo hearts. We did not detect transcript in the heart or in the vasculature. In contrast, we detected gper mRNA in three discrete anatomic areas of the brain: the preoptic and olfactory areas and in the ventral hypothalamus (Fig. 4A-C). Thus, gper localization is consistent with the hypothesis that GPER acts in the brain, and not through cells in the heart, to regulate heart rate.

Genetic evidence using esr mutants suggests that GPER acts independently of nuclear estrogen receptors to regulate heart rate (Fig. 3). To further test the hypothesis that GPER acts as an autonomous estrogen receptor in vivo, we asked whether GPER and nuclear estrogen receptors are expressed in the same cells in the brain, reasoning that if GPER and nuclear estrogen receptors fail to colocalize, this would support the idea that GPER acts as an autonomous estrogen receptor in vivo. We took 5xERE:GFP embryos at 1 day post fertilization and exposed them overnight to 100 ng/ml estradiol. When the embryos reached 48 hpf, we used two color fluorescent in situ hybridization to detect gfp and gper transcripts simultaneously. Since all three nuclear estrogen receptor genes activate the 5xERE:GFP transgene, detecting gfp allows us to monitor activity of all three estrogen receptors using a single RNA probe. In the olfactory and preoptic areas, we found no colocalization between gfp and gper (Fig. 4D, E). In the ventral hypothalamus, we found a cluster of cells at the midline expressing gper but not gfp. Surrounding this region of gper-positive cells was a bilaterally symmetric ‘U’-shaped labeling pattern of cells expressing both gper and gfp (Fig. 4F). Thus, GPER and nuclear estrogen receptors are expressed in unique and overlapping cells in the brain, supporting the hypothesis that GPER can act independently of nuclear estrogen receptors in vivo.
Because MZgper/- fish are viable, we asked whether loss of GPER and reduced basal heart rate in embryos might have an effect on adult heart morphology. We dissected hearts from 8-9 month old adult wildtype and MZgper/- zebrafish. Female mutants had a mean 47% reduction in atrioventricular (AV) valve width compared to wildtype (Fig. 5E; MZgper/- 33.5 ± 6.35 μm (mean ± standard deviation), n=10; wildtype 62.49 ± 18.53, n=9, p=0.0002 Student’s t test), while male mutants had a mean 40% reduction in AV valve width compared to wildtype (Fig. 5F; MZgper/- 29.39 ± 6.32, n=7; wildtype 49.14 ± 9.68, n=5, p=0.0016 Student’s t test). We also analyzed the single thickest part of AV valve leaflets in each heart and found that mutant valves were on average 45% thinner at their thickest point than wildtype valves from fish of the same sex (female MZgper/- 42.93 ± 10.38 vs wildtype 77.97 ± 24.32 μm, p=0.0006 Student’s t test; male MZgper/- 38.17 ± 11.62 vs wildtype 67.19 ± 24.77, p=0.02 Student’s t test). These results are consistent with previous observations that eliminating blood flow in embryos impairs heart valve formation (31). Together, these findings demonstrate a correlation between reduced blood flow in embryos and reduced valve thickness in adults and suggest that GPER regulation of heart rate is important for proper heart valve maturation.

DISCUSSION

Our results support the hypothesis that GPER acts as an autonomous estrogen receptor in vivo. Previous reports using cultured cells demonstrated that fluorescently labeled or isotopic estradiol specifically binds membranes from cells expressing GPER (8, 9). Additionally, estradiol exposure increased cyclic AMP and calcium levels in HEK293 and COS7 cells in a GPER-dependent manner (8, 9), while estradiol exposure increased phosphoinositide 3-kinase activity in SKBR3 breast cancer cell line in a GPER-dependent manner (8). However, because these studies utilized cells that either express artificially high levels of GPER or are tumorigenic, the findings do not address whether GPER acts as an estrogen receptor in vivo under normal physiologic conditions. Our genetic and
pharmacologic results strongly suggest that GPER is an estrogen receptor in vivo. If estradiol was binding to ERα or ERβ, and these receptors activated GPER, then we would expect to see no increase in heart rate in esr1, esr2a or esr2b mutants following exposure to estradiol. Instead, all esr mutants responded normally to estradiol exposure (Fig. 3), suggesting that ER and GPER signaling pathways are distinct in this context. Consistent with these results, we found gper transcript expressed in cells in the brain that lack nuclear estrogen receptor activity (Fig. 4), further supporting the hypothesis that GPER responds to estrogens independently of nuclear estrogen receptors in vivo.

A study in mouse hearts also supports the idea that GPER acts as an autonomous estrogen receptor in vivo. In adults, estradiol administration reduces cardiac damage following ischemia (32-34). To identify the receptor required for estradiol's protective effects, Kabir and colleagues subjected hearts from male mutant mice lacking either GPER, ERα or ERβ to ischemia-reperfusion injury in the presence of estradiol or vehicle. Estradiol treatment protected wildtype and ERα and ERβ mutant mice from injury, but had no effect on GPER mutant mice (35), demonstrating that estradiol exerts its protective effects via GPER, independently of ERα or ERβ. The extent to which ER and GPER signaling pathways interact likely depends on cell type, developmental stage, sex and/or pathology. Studying the influence of estrogens on heart rate in zebrafish embryos is a powerful in vivo system where GPER activity is dissociated from classical nuclear estrogen receptor signaling.

Between 2 and 5 dpf, zebrafish heart rate normally increases (36, 37). Our results support the hypothesis that endogenous estradiol regulates this increase in heart rate. The finding that GPER mutant embryos have lower basal heart rate compared to wildtype embryos implicates endogenous estradiol. Additionally, a recent HPLC analysis of endogenous estradiol concentration in zebrafish embryos found that estradiol concentrations increased from 137 pg/embryo at 48 hpf to 170 pg/embryo at 72 hpf (38). Taken together, these results support the hypothesis that endogenous estradiol regulates heart rate in zebrafish embryos.
and larvae. The source of embryonic estradiol, whether synthesized by the embryo or maternally deposited in the yolk, is not known.

Blood flow in embryos is a critical determinant of cardiovascular development (39). Zebrafish embryos in which blood flow is drastically reduced or eliminated exhibited a valve dysgenesis phenotype (31). MZgper-/- embryos exhibit a ~20% reduced heart rate compared to wildtype, yet are viable, allowing us to examine the adult heart following reduced basal heart rate in embryos. Based on the heart rate phenotype in MZgper-/- embryos and the detection of gper transcripts in the brain but not the heart, we propose that the reduced heart rate in GPER mutant embryos causes a reduction in blood flow that prevents heart valves from maturing properly and manifests in adults as thinner atrioventricular valves compared to wildtype. While gper transcript is expressed in the adult zebrafish brain (2), it is not known whether gper is expressed in the juvenile or adult zebrafish heart, where it could directly influence valve cell division and/or valve cell survival separately from influencing heart rate and blood flow.

There are several mechanisms by which GPER activity in the brain could regulate heart rate. Neurons that express GPER could be part of a chain of neurons that ultimately terminates in motor neurons that innervate the heart and regulate contractions. It is not known whether the zebrafish heart is innervated by 50 hpf and thus it is unclear whether GPER is influencing sympathetic or parasympathetic neuron activity. Parasympathetic neurons terminate at the heart and activate muscarinic acetylcholine receptors to reduce heart rate. In contrast, sympathetic neurons terminate at the heart and activate beta-adrenergic receptors to increase heart rate. GPER could be activating sympathetic activity or reducing parasympathetic activity. However, there is conflicting evidence as to whether autonomic control of the heart occurs in two-day old zebrafish embryos. Milan and colleagues reported that propranolol, a beta-adrenergic receptor antagonist, reduced heart rate in 2 dpf embryos, while isoproterenol, a beta-adrenergic receptor agonist, and atropine, a muscarinic acetylcholine receptor antagonist, both increased heart rate (40). These results suggest that at 2 dpf,
the zebrafish embryo heart can respond to sympathetic and parasympathetic activity. In contrast, Schwerte and colleagues reported that atropine and propranolol did not influence heart rate in embryos younger than 5 dpf, while zebrafish failed to respond to adrenergic stimulation before 4 dpf (41). Future studies are required to determine when functional autonomic innervation of the zebrafish heart initially occurs.

Even if the autonomic nervous system does not regulate heart rate in 2 dpf zebrafish embryos, GPER can still function in the brain to regulate heart rate. Previous work suggests that GPER influences neurotransmitter release (42). Therefore, GPER activity could trigger neuronal activity that leads to systemic release of humoral factors, such as thyroid hormone, known to regulate heart rate (30). Expression of gper transcript in the ventral hypothalamus / pituitary (Figure 4) is consistent with this hypothesis.

While our results illuminate GPER signaling in the context of embryonic heart rate, it is not clear to what extent GPER influences heart rate at later stages of development. At larval, juvenile and adult stages it is impossible to assess heart rate without immobilizing or anesthetizing zebrafish, manipulations that themselves influence heart rate. In adult mice with mutations in GPER, there was no significant difference in basal heart rate between mutant and wildtype mice of either sex (16, 17, 43). It is possible that GPER regulates heart rate in embryos but not in adults. Additionally, heart rate in GPER mutant mice was assayed using general anesthesia, which is known to depress heart rate compared to conscious mice (44). Anesthesia may mask the effect of GPER on basal heart that we observe in conscious animals. We also cannot exclude the possibility that the effects of GPER on heart rate are specific for zebrafish.

In summary, this study identified a role for GPER in the regulation of embryonic heart rate. The zebrafish estrogen receptor mutants we developed enable experiments to rapidly and conclusively identify the causative estrogen receptor associated with any estrogen signaling phenotype, as demonstrated with the estradiol-dependent increase in heart rate reported here. This has significant implications for studies of estrogenic environmental endocrine
disruptors, which are frequently tested on zebrafish to identify effects on embryonic development, organ formation and function (45). Zebrafish estrogen receptor mutants can now be used to determine whether such effects are specific for estrogen receptors and to identify the precise receptor target. Our results also establish a need to consider the impact on cardiac function when considering the toxicity of estrogenic environmental endocrine disruptors.
Figure 1. Estradiol and GPER agonists increased heart rate in zebrafish embryos. Wildtype embryos were incubated in water containing vehicle (0.1% DMSO), estradiol (3.67 μM, ER/GPER agonist), progesterone (1 μM), ICI (10 μM ICI182,780, ER antagonist/GPER agonist), G1 (1 μM, GPER agonist), G36 (1 μM, GPER antagonist) or two chemicals in combination at 49 hours post fertilization and heart rates were measured 1 hour following treatment. ***, p<0.0001 compared to vehicle, ANOVA with Dunnett’s test. Each black circle represents the mean heart rate from a single clutch of embryos (3-16 embryos per clutch). Clutches in the same treatment group were assayed on different days. Horizontal blue lines are the mean of each treatment.
Figure 2. Abnormal heart rate in gper mutant zebrafish. (A) Genomic DNA of gper<sup>uab102</sup> zebrafish contains a 131 basepair deletion in the gper coding region between CRISPR guide RNA targets 1 and 2, resulting in a premature stop codon in the GPER protein. Red dashes indicate DNA deletions, mutated amino acids are shown in red. (B) Genomic DNA was harvested from individual embryos, gper was PCR amplified and separated on an agarose gel to identify deletion mutations. (C-D) 3 day post fertilization wildtype and maternal zygotic gper<sup>uab102</sup> homozygous larvae (MZgper-/-) exhibit similar gross morphology. Images are lateral views, anterior to the left, dorsal to the top. Scale bar, 500 μm. (E) Neither estradiol (ER/GPER agonist, 3.67 μM) nor ICI182,780 (ER antagonist/GPER agonist, 10 μM) changed heart rate significantly compared to vehicle (0.1% DMSO) in MZgper-/-, two-way ANOVA followed by F test, p=0.27. (F) MZgper-/- exhibited lower basal heart rate than age-matched wildtype embryos. *, p<0.05 compared to wildtype, paired t test. Each black circle represents the mean heart rate from a single clutch of embryos (≥ 7 embryos per clutch). Clutches in the same treatment group or genotype were assayed on different days. Horizontal blue lines are the mean of each treatment.
Figure 3. Normal heart rate in nuclear estrogen receptor mutants. (A) Homozygous mutant embryos at 49 hour post fertilization were incubated in water containing estradiol (ER/GPER agonist, 3.67 μM) or vehicle (0.1% DMSO) and heart rate was measured 1 hour post treatment. Estradiol increased heart rate compared to vehicle in zebrafish with homozygous mutations in ERα (esr1 -/-), ERβ1 (esr2a -/-), ERβ2 (esr2b -/-). *p<0.05 compared to vehicle within genotype, paired t-test. (B) Basal heart rate was measured at 50 hours post fertilization in embryos reared in untreated water. Heart rate was not significantly different in homozygous mutant (-/-) embryos compared to heterozygous (-/+) and wildtype (+/+) siblings for each esr mutant, two-way ANOVA. Each black circle represents the mean heart rate from a single clutch of embryos (4-8 embryos per clutch). Clutches in the same treatment group or genotype were assayed on different days. Horizontal blue lines are the mean of each treatment.
Figure S1. Zygotic gper mutant embryos are sensitive to estradiol. Zygotic homozygous gper mutant embryos were incubated in water containing estradiol (ER/GPER agonist, 3.67 μM) or vehicle control (0.1% DMSO) at 49 hours post fertilization and heart rates were measured 1 hour post treatment. *, p<0.05 compared to vehicle, paired t test. Each black circle represents the mean heart rate from a single clutch of embryos (≥ 6 embryos per clutch). Clutches in the same treatment group were assayed on different days. Horizontal blue lines are the mean of each treatment.
Figure S2. Generation and validation of esr1 mutant zebrafish. (A) Genomic DNA of esr1mutant zebrafish contains a 4 basepair deletion in the esr1 coding region, resulting in a premature stop codon in the Esr1 (ERα) protein. Nucleotide deletions are shown as red dashes, amino acid mutations are in red. Map indicates site of frameshift mutation and premature stop codon (AF-1, activating function 1 domain; DBD, DNA binding domain; LBD, ligand binding domain; AF-2, activating function 2 domain). (B) High resolution melting curve analysis was used to distinguish mutants from wildtype. Curves represents DNA amplified from a wildtype AB (black) or esr1mutant zebrafish (cyan). (C) Strategy for validating zebrafish estrogen receptor mutants using transgenic 5xERE:GFP zebrafish. Mutants were generated on a transgenic background where estrogen receptor (ER) transcriptional activity is marked by green fluorescent protein (GFP) expression. Following exposure to estradiol, loss-of-function mutants should exhibit reduced fluorescence in cells expressing esr1. (D-K) 2-day post fertilization embryos were exposed to 367 nM (100 ng/mL) estradiol, live fluorescent images (D, F, H, J) and corresponding brightfield images (E, G, I, K) were taken at 3 d. 5xERE:GFPesr1mutant homozygous larvae (esr1−/−) exhibit normal morphology, but lack fluorescence in heart valves, whereas heterozygotes (esr1+/−) exhibit fluorescent heart valves. High magnification images of the heart are shown in H-K. Arrows indicate heart valves, arrow head indicates liver. Images are lateral views, anterior to the left, dorsal to the top. Scale bars, 500 μm (D-G), 100 μm (H-K).
Figure S3. Generation of esr2a mutant zebrafish. (A) Genomic DNA of esr2a<sup>uab134</sup> zebrafish contains a 2 basepair deletion (red dashes) in the esr2a coding region, resulting in a premature stop codon in the Esr2a (ERβ1) protein. Amino acid mutations are in red. Map indicates frameshift mutation and premature stop codon in the Esr2a protein. AF-1, activating function 1 domain; DBD, DNA binding domain; LBD, ligand binding domain; AF-2, activating function 2 domain. (B) High resolution melting curve analysis was used to distinguish mutants from wildtype. Curves represent DNA amplified from a wildtype AB (black) or esr2a<sup>uab134</sup> mutant zebrafish (cyan). (C-F) 5xERE:GFP<sup>262</sup> and 5xERE:GFP<sup>262</sup> (esr2a<sup>-/-</sup>) 3-day post fertilization (d) larvae were exposed to 367 nM (100 ng/mL) estradiol. Live fluorescent images (C, E) and corresponding brightfield images (D, F) were captured at 4 d. esr2a<sup>-/-</sup> larvae exhibit normal morphology and fluorescence, consistent with data demonstrating that esr2a is not expressed during these developmental stages. Arrows indicate heart valves, arrow head indicates liver. Images are lateral views, anterior to the left, dorsal to the top. Scale bar, 500 μm

| Amino acid sequence | Predicted ESR2A protein mutation |
|----------------------|---------------------------------|
| wildtype             | VGGHILSPIFNSSSPESLPVEMPFICPSPTDLGKAFSTLHFSYGLLHITYE | 554 amino acids frameshift at 31, premature stop at 115 |
| mutant               | VGGHLPSQTLISASGESPLHIALHRFPRLQNSALLQERASAG          |

| Frameshift | Premature stop |
|------------|----------------|
| C          | D              |
| E          | F              |
esr2b -/- E2 100 ng/mL

Predicted ESR2B protein mutation

**Amino acid sequence**

| A/B  | C     | D     | E     | F     |
|------|-------|-------|-------|-------|
| AF-1 | DBD   | hinge | LBD   | AF-2  |
|      |       |       |       | frameshift |

**Mutant**

MSSSPGPAPVLDSSKADRGASPALLPRLYASPLGMDNQTVCIPSPYVE ACQDYSFPVSGEFHNLTPVSSAVLGFHRPPFYSEILVFLSPTIL WPLSALPASAGLQRNTFTQRLGGQDGTRAQEPQFC

**Wildtype**

MSSSPGPAPVLDSSKADRGASPALLPRLYASPLGMDNQTVCIPSPYVE ACQDYSFPVSGEFHNLTPVSSAVLGFHRPPFYSEILVFLSPTIL WPLSALPASAGLQRNTFTQRLGGQDGTRAQEPQFC

593 amino acids

frameshift at 99, premature stop at 133

Figure S4. Generation and validation of esr2b mutant zebrafish. (A) Genomic DNA of esr2b<sup>mut127</sup> zebrafish contains a 5 basepair deletion (red) in the esr2b coding region, resulting in a premature stop codon in the Esr2b (ERβ2) protein. Amino acid mutations are in red. Map indicates frameshift mutation and premature stop codon in the Esr2b protein. AF-1, activating function 1 domain; DBD, DNA binding domain; LBD, ligand binding domain; AF-2, activating function 2 domain. (B) High resolution melting curve analysis was used to distinguish mutants from wildtype. Curves represents DNA amplified from a wildtype AB (black) or esr2b<sup>mut127</sup> mutant zebrafish (cyan). (C-F) 5xERE::GFP<sup>esr2b<sup>mut127</sup></sup>;esr2b<sup>-/-</sup> 3-day post fertilization (d) larvae were exposed to 367 nM (100 ng/mL) estradiol. Live fluorescent images (C, E) and corresponding brightfield images (D, F) were captured at 4 d. 5xERE::GFP<sup>esr2b<sup>mut127</sup></sup>;esr2b<sup>-/-</sup> homozygous larvae (esr2b -/-) exhibit normal morphology, but lack fluorescence in the liver. Arrows indicate heart valves, arrow head indicates liver. Images are lateral views, anterior to the left, dorsal to the top. Scale bar = 100 μm.
Figure 4. *gper* expression in the brain. (A-C) Whole mount colorimetric in situ hybridization was performed on wildtype embryos at 50 hours post fertilization (hpf). (A) *amhc* (alpha-myosin heavy chain) antisense RNA labels atrial myocardial cells in the heart (boxed). (B, C) *gper* antisense RNA labels a bilaterally symmetric cluster of cells in the olfactory area (white arrowheads) and preoptic area (black arrowhead) and a medial cluster of cells in the ventral hypothalamus (arrows). No label was detected in the heart. Lateral views with anterior to the left (A,B), ventral view with anterior to the top (C), scale bars = 100 μm. (D-F) Double fluorescent in situ hybridization performed on 48 hpf *Tg(5xERE:GFP)c262* embryos following overnight exposure to 100 ng/ml estradiol. *gfp* marks cells with active nuclear estrogen receptors. Confocal images of selected Z-slices (0.975 μm) show that *gper* is expressed in the olfactory area (D) and preoptic area (E) in cells lacking *gfp* (D'', E'', scale bars = 50 μm). In the ventral hypothalamus (F), *gper* is expressed in a medial cluster of cells lacking *gfp* (arrows, F, F''), whereas *gper* is expressed together with *gfp* more laterally (arrowheads, F'', scale bar = 10 μm). In merged images, *gper* is magenta, *gfp* is green and areas of colocalization are white. Dorsal views, anterior to the top.
Figure S5. Nuclear estrogen receptor transcriptional activity is normal in gper mutant zebrafish. (A-H) Maternal zygotic gper<sup>−/−</sup> homozygous larvae on the S<sub>vERE-GFP<sup>+</sup></sub> transgenic background (MZgper<sup>−/−</sup>) were exposed to 387 nM (100 ng/mL) estradiol at 2-days post fertilization (2 dpf). Fluorescence (A, C, E, G) and corresponding brightfield images (B, D, F, H) were taken at 3 dpf. Fluorescence in the heart valves (arrows) and liver (arrow heads) is similar between MZgper<sup>−/−</sup> and wildtype larvae. C, D, G, H. High magnification images of heart. Images are lateral views, anterior to the left, dorsal to the top. Scale bars, 500 μm (C-F), 100 μm (G-J).
Figure 5. GPER mutants exhibit decreased atrioventricular (AV) valve width in adulthood. (A-L) Representative images of MZgper -/- and wildtype (wt) hearts. (A-D) 5 μm coronal sections through the heart, stained with H&E. Atrium (A), ventricle (V), and bulbus arteriosus (BA) indicated with black arrows, AV valve leaflets are indicated with red arrows. Black boxes show the portion of the image that is digitally enlarged in A’-D’ to highlight AV valve morphology. Scale bars in A-D represent 200μm; scale bars in A’-D’ represent 50μm. (E,F) Graphs comparing average and maximum AV valve width between MZgper -/- and wildtype hearts in females (E) and males (F). Width was measured at the widest part of the valve over multiple coronal sections from the middle of the heart, perpendicular to the long axis of the valve, and corrected for standard length of each fish. Each circle represents measurements from a single fish, horizontal black lines are the mean width from each genotype. *p < 0.05; **p < 0.01; ***p < 0.001, unpaired Student’s t test.
Materials and Methods

Zebrafish

Zebrafish were raised at 28.5°C on a 14-h light, 10-h dark cycle in the UAB Zebrafish Research Facility in an Aquaneering recirculating water system (Aquaneering, Inc., San Diego, CA). Wildtype zebrafish were AB strain (46) and all mutant and transgenic lines were generated on the AB strain. To visualize nuclear estrogen receptor activity, transgenic line Tg(5xERE:GFP)c262/c262 was used for all studies unless otherwise mentioned (5). All procedures were approved by the UAB Institutional Animal Care and Use Committee.

Embryo collection

Embryos were collected during 10 minute intervals to ensure precise developmental timing within a group. Embryos were placed in Petri dishes containing E3B (60X E3B: 17.2g NaCl, 0.76g KCl, 2.9g CaCl₂·2H₂O, 2.39g MgSO₄ dissolved in 1 liter Milli-Q water; diluted to 1X in 9 liter Milli-Q water plus 100 μL 0.02% methylene blue) and placed in an incubator at 28.5°C on a 14-h light, 10-h dark cycle. At 24 hours post fertilization (hpf), embryos were incubated in E3B containing 200 μM 1-phenyl 2-thiourea (PTU) to inhibit pigment production (46). Between 24 and 48 hpf, embryos were manually dechorionated and randomly divided into control and experimental treatment groups (10 to 30 embryos per treatment group) in 60mm Petri dishes and kept at 28.5°C until 49 hpf.

Embryo treatments

At 49 hpf, embryos were incubated in E3B with estrogen receptor modulator(s) at 28.5°C for 1 hour. Estrogen receptor modulator treatments consisted of: 3.67 μM E2 (17β-estradiol, Sigma E8875; purity ≥ 98%), 10 μM ICI182,780 (fulvestrant, Sigma I4409; purity >98%), 1 μM G1 (Azano, AZ0001301; purity ≥ 98%), 1 μM G36 (Azano, AZ-0001303; purity ≥ 98%), 1 μM progesterone (Sigma P0130; purity ≥ 99%) or vehicle (0.1% dimethylsulfoxide (DMSO), Fisher D128-500; purity ≥ 99.9%). All chemical stocks were made in 100% DMSO at 1000x and diluted in E3B embryo media to final concentration at the time of treatment. For rescue experiments (ICI182,780 + G36), final DMSO concentration was 0.2%. There was no difference in heart rate between embryos incubated in 0.1% or 0.2% DMSO (not shown). All vehicle controls shown in figures are 0.1% DMSO.

Measurement of heart rates

All embryos were reared at 28.5°C and heart rate was measured at room temperature. Following one hour incubation in treatment compounds at 28.5°C, heart rate (beats per minute, bpm) was calculated by counting the number of heart beats in fifteen seconds and multiplying that number by four. Prior to measurements, each dish was removed from the incubator and placed under the microscope light for 4 minutes at room temperature, allowing embryos to acclimate to the light and eliminate any effect of the startle response. Control groups were counted first and last to ensure that the overall heart rate did not increase during the duration of counting due to natural increases in heart rate during development. All heart rates were measured on a Zeiss Stemi 2000.
dissecting microscope with a halogen transmitted light base (Carl Zeiss Microimaging, Thornwood, NJ).

**Generation of guide RNA and Cas9 mRNA**

Plasmids pT7-gRNA and pT3T3-nCas9n were obtained from Addgene (numbers 46759, 46757) (26). pT7-gRNA was digested simultaneously with BsmBI, BglII and Sall for one hour at 37 °C followed by one hour at 55 °C. To generate esr2a, esr2b and gper gRNAs, oligonucleotides containing target site sequences (see table below) were synthesized by Invitrogen. Oligos were hybridized to each other using NEBuffer3 restriction enzyme buffer (New England Biolabs) to generate double stranded target DNA and annealed into digested pT7-gRNA using Quick T4 DNA Ligase (New England Biolabs) as previously described (26). Guide RNAs were synthesized using the MegaShortScript T7 Kit (Life Technologies) using the relevant modified pT7-gRNA vector linearized with BamHI as a template. Guide RNA was purified using the RNA clean & concentrator kit (Zymo Research). To generate esr1 guide RNA, target-specific oligonucleotides containing the SP6 (5'-ATTTAGGTGACA)TA promoter sequence, a 20 base target site without the PAM, and a complementary region were annealed to a constant oligonucleotide encoding the reverse-complement of the tracrRNA tail as described (47). This oligo was used as a template for in vitro transcription using the MegaShortScript Sp6 Kit (LifeTechnologies). To generate Cas9 mRNA, the pT3TS-nCas9n plasmid was linearized with XbaI and transcribed using the mMessage mMACHINE T3 kit (LifeTechnologies) and purified using RNA clean & concentrator kit (Zymo Research). RNA concentration was quantified using a Nanodrop spectrophotometer (Nanodrop ND-1000, ThermoFisher).

Target site sequences for gper, esr1, esr2b and esr2a oligonucleotides:

| Gene      | CRISPR target (PAM in red) | Oligo 1          | Oligo 2          |
|-----------|----------------------------|------------------|------------------|
| esr1      | GTCCTCTCAGCAGGCA GCCGTTGG | ATTTAGGTGACA TA TA | GTTTTAGAGCTAGAA ATAGCAAG |
| esr2a     | GGAGAGGATGGTTGA AGATGGG  | TAGGAGAGGATGAG TTGAAGAT | AAACATCTTCAACTC ATCCTCT |
| esr2b     | GCAGGGCCAGTGCA AGTGAGG   | TAGGCGGGGAGTGC AGAGGTG | AAACACTCTCTGCAA CTGCCCG |
| gper target 1 | GGCTGTGGCAGATCTT ATTCTGG | TAGGCTGTGGCAGA TCTTATTC | AAACGAATAAGATCT GCCACAG |
| gper target 2 | GGAAAAGGAATGTGT GTACAGG  | TAGGAAAAGGAAT GTGTAC | AAACGTACACCATT TCTTTT |

**Embryo injections**
One-cell-stage embryos were injected using glass needles pulled on a Sutter Instruments Fleming/Brown Micropipette Puller, model P-97 and a regulated air-pressure micro-injector (Harvard Apparatus, NY, PL1–90). Each embryo was injected with a 1 nl solution of 150 ng/µl of Cas9 mRNA, 50 ng/µl of gRNA and 0.1% phenol red. Mixtures were injected into the yolk of each embryo. Injected embryos were raised to adulthood and crossed to wildtype fish (either AB or Tg5xERE:GFPc262) to generate F1 embryos. F1 offspring with heritable mutations were sequenced to identify loss of function mutations.

**Genomic DNA isolation**

Individual embryos or tail biopsies from individual adults were placed in 100 µL ELB (10 mM Tris pH 8.3, 50 mM KCl, 0.3% Tween 20) with 1 µL proteinase K (800 U/ml, NEB) in 96 well plates, one sample per well. Samples were incubated at 55°C for 2 hours (embryos) or 8 hours (tail clips) to extract genomic DNA. To inactivate Proteinase K, plates were incubated at 98°C for 10 minutes and stored at -20°C.

**High resolution melt curve analysis**

PCR and melting curve analysis was performed as described (48). PCR reactions contained 1 µl of LC Green Plus Melting Dye (BioFire Diagnostics), 1 µl of Ex Taq Buffer, 0.8 µl of dNTP Mixture (2.5 mM each), 1 µl of each primer (5 µM), 0.05 µl of Ex Taq (Takara Bio Inc), 1 µl of genomic DNA, and water up to 10 µl. PCR was performed in a Bio-Rad C1000 Touch thermal cycler, using black/white 96 well plates (Bio-Rad HSP9665). PCR reaction protocol was 98°C for 1 min, then 34 cycles of 98°C for 10 sec, 60°C for 20 sec, and 72°C for 20 sec, followed by 72°C for 1 min. After the final step, the plate was heated to 95°C for 20 sec and then rapidly cooled to 4°C. Melting curves were generated with either a LightScanner HR 96 (Idaho Technology) over a 70–95°C range and analyzed with LightScanner Instrument and Analysis Software (V. 2.0.0.1331, Idaho Technology, Inc, Salt Lake City, UT), or with a Bio-Rad CFX96 Real-Time System over a 70–95°C range and analyzed with Bio-Rad CFX Manager 3.1 software.

**Live imaging**

Live zebrafish embryos and larvae were visualized using a Nikon MULTIZOOM AZ100 equipped with epi-fluorescence and an Andor Clara digital camera unless otherwise noted. To validate mutants with 5xERE reporter activity, larvae were treated overnight with 100 ng/mL estradiol beginning at 2-3 dpf. Following overnight treatment, larvae were washed in E3B, anesthetized with 0.04% tricaine and imaged in Petri dish containing E3B. For Fig. S1 H-K, larvae were mounted in bridged coverslips in E3B with 0.04% tricaine (46). Images were captured on a Zeiss Axio Observer.Z1 fluorescent microscope equipped with an Axio HRm camera and Zen Blue 2011 software (Carl Zeiss Microscopy, Oberkochen, Germany). Adjustments, cropping and layout were performed using Photoshop CS6 and InDesign CS6 (Adobe Systems Inc., San Jose, CA).

**RNA in situ hybridization**

For synthesis of RNA probes, full-length gper open reading frame was amplified by PCR from genomic DNA extracted from 3 dpf larvae (gper is a single exon gene and therefore the open reading frame sequence is identical in genomic and cDNA) using primers 5’-ATGGAGGAGCAGACTACCAATGTG-3’ and 5’-
CTACACCTCAGACTCAGCTCTGACAG-3’ and TA cloned into pCR2.1 vector (Invitrogen). *amhc* and *gfp* probes were used as described (5, 49). All clones were verified by sequencing. Digoxigenin-labeled antisense RNA and FITC-labeled antisense RNA were transcribed using T7 and T3 polymerase, respectively, as previously described (5). Colorimetric whole-mount *in situ* hybridization was performed on zebrafish embryos and larvae as described previously, using 5% dextran in the hybridization buffer (50, 51). Following colorimetric *in situ* hybridization, embryos were sequentially cleared in glycerol (25%, 50%, 75% in phosphate buffered saline), mounted in 4% low-melting temperature agarose, and imaged using a Zeiss Axio Observer.Z1 microscope with Zeiss Axio MRC5 camera and Zen Blue 2011 software. Fluorescent *in situ* hybridization (FISH) was performed as previously described (51) with the following modifications: After rehydration, Proteinase K treatment was extended to 35 minutes. Following hybridization, embryos were washed in 2xSSC prior to being placed in PBT. Embryos were blocked in 2% Roche blocking reagent in 100 mM Maleic acid, 150 mM NaCl, pH 7.5 (52). For double labeling, following development of anti-DIG-POD antibody, reaction was inactivated in 100 mM glycine pH 2 for 10 minutes then incubated in anti-FITC antibody. Following florescent *in situ* hybridization, embryos were cleared in 50% glycerol, mounted on a bridged coverslip and imaged using a Nikon A1/R scanning confocal microscope with Nikon Advanced Elements software.

**Histology on adult zebrafish hearts**

Adult wild-type (*n*=15) and MZgper -/- (*n*=17) zebrafish at 8-9 months of age were used for heart dissections. Zebrafish were anesthetized in 0.2 mg/mL tricaine, measured with a digital caliper to obtain standard length (SL) (53), and decapitated dorsal to the pectoral fin, then hearts were dissected with forceps in phosphate buffered saline (PBS). Hearts were examined for structural integrity and fixed in 1 mL of 10% formaldehyde in PBS at 4°C for 16-20 hours overnight. Whole hearts were washed three times in PBS. To confirm heart integrity following dissection and fixation, whole hearts were imaged on a Nikon SMZ1500 stereomicroscope equipped with a Nikon DS-Qi1MC camera. Intact, properly dissected hearts were embedded in 10-15 μL of Histogel (Thermo Scientific) on 0.8 μm AA Millipore filter paper and oriented with atrium and ventricle in the same horizontal plane to ensure downstream collection of coronal sections. Histogel-embedded hearts were placed in tissue cassettes (Fisher Scientific #22-272420) and allowed to set on ice for 5 minutes. Excess Histogel was trimmed without disturbing the heart. Cassettes were closed and stored in 70% ethanol until further processing. Hearts were processed routinely into paraffin, embedded, sectioned at 5 μm and stained with hematoxylin and eosin (H&E). Bright-field images of sections were obtained using a Zeiss Axio Observer.Z1 microscope with a Zeiss Axio MRC5 camera and 20x objective (NA 0.8). Tiled images were captured and fused using the stitching algorithm of Zeiss ZEN 2 blue edition software. Cross-sectional length measurements were obtained by taking the mean of 6 sections from each heart (on average) determined to be in the middle of the heart by AV valve visibility. In each section, the thickness of the largest valve leaflet was measured perpendicular to the long axis of the valve. Zebrafish growth and size vary within groups of similarly aged adult fish (53). Therefore, measurements for each sex were normalized to standard length (SL), an established measure of postembryonic zebrafish development (53), by dividing the individual measurements for each fish into the mean SL for that sex. Comparisons between genotypes for each sex were made with two-tailed, unpaired Student’s *t* test. Statistical significance was accepted at a *P* value of <0.05.
Experimental design and data analysis

Hart rate assays were conducted in separate experiments. Each experiment included comparing groups (treated vs untreated or mutant vs wildtype) using at least 3 embryos per group with all embryos from the same clutch. All experiments were replicated for at least 3 times (n≥3) using different clutches on different days. This is essentially a complete block design with clutch/day as block. Mean heart rate of individual embryos from a clutch was used for comparing treatment groups (or mutant groups) within experiments using two-way ANOVA controlling for clutch/day effect. The overall treatment effect (or the genotype effect in some experiments) was tested using F test. If it was significant, Dunnett’s test was then used to compare each treatment group with the vehicle group or mutant group with the wildtype group. For some special individual pairs of comparisons, paired t test was used. Significance level is 0.05. All the analyses were conducted using R (version 3.0.2). Graphs were produced using GraphPad Prism 7.0a software.

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