

# <sup>1</sup> Estradiol promotes habituation learning <sup>2</sup> despite the suppressive effects of the <sup>3</sup> recognized Estrogen Receptors

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## <sup>9</sup> Abstract

<sup>10</sup> Habituating to the constant stimuli in the environment is a critical learning process conserved across species. We  
<sup>11</sup> use a larval zebrafish visual response to sudden darkness as a model for studying habituation learning, where  
<sup>12</sup> zebrafish reduce their responses to repeated stimulations. In this paradigm, treatment with estradiol strongly  
<sup>13</sup> increases learning rate, resulting in more strongly suppressed responses. We used knockout mutant lines for the  
<sup>14</sup> Estrogen Receptors (*esr1*, *esr2a*, *esr2b*, *gper1*) in an attempt to identify the receptor(s) mediating these effects. These  
<sup>15</sup> experiments failed to identify a necessary receptor (or combination of receptors). Surprisingly, *esr1*, *esr2a*, and *gper1*  
<sup>16</sup> mutants showed weak but consistent increases in habituation, indicating these receptors suppress habituation  
<sup>17</sup> learning. These experiments demonstrate that estradiol is a complex modulator of learning in our model, where the  
<sup>18</sup> learning-promoting effects are mediated by an unidentified estradiol target, and the classical Estrogen Receptors  
<sup>19</sup> act in competition to subtly suppress learning.

<sup>20</sup>

## <sup>21</sup> Introduction

<sup>22</sup> A primary function of the brain is to learn from experiences and adjust behavior in response. One aspect of learned  
<sup>23</sup> behaviour involves sharpening attention and behavioural resources toward salient cues by ignoring irrelevant  
<sup>24</sup> background stimuli. For instance, it may be critical to recognize the alarm calls of a nearby animal, whereas  
<sup>25</sup> continually registering the steady hum from distant traffic is far less important. The capacity to reduce responses  
<sup>26</sup> to repetitive, non-essential stimuli is known as habituation, which is considered the simplest form of learning and  
<sup>27</sup> memory (*Rankin et al., 2009*).

<sup>28</sup> We study a paradigm for long-term habituation where larval zebrafish reduce their responsiveness to sudden  
<sup>29</sup> pulses of whole-field darkness, or dark flashes (DFs) (*Wolman et al., 2011; Randlett et al., 2019; Lamiré et al., 2023*).  
<sup>30</sup> We recently reported that multiple hormonal signaling pathways show strong modulation of habituation learning  
<sup>31</sup> performance, including melatonin, progesterone, and estrogen (*Lamiré et al., 2023*). The ability of these signaling  
<sup>32</sup> pathways to modulate learning is consistent with previous results in other systems and paradigms (*Nilsson and*

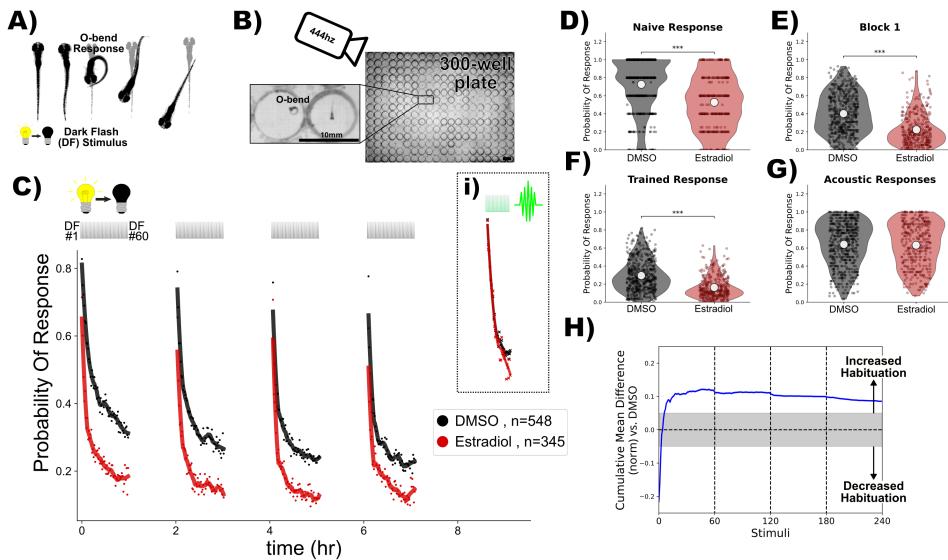
33 *Gustafsson, 2002; Naderi et al., 2020; Dillon et al., 2013; Rawashdeh et al., 2007; Jilg et al., 2019; El-Sherif et al.,*  
34 *2003; Barros et al., 2015*), and may be an important mechanism to shift learning and memory performance or  
35 strategies based on biological rhythms or external fluctuations like seasons, weather or the day/night cycle.

36 In this project we have focused on estrogen signaling. We identified multiple estradiol analogs which strongly  
37 increased habituation learning when added at 5-10 $\mu$ M doses in the water (ethinyl estradiol, estradiol valerate, and  
38 hexestrol, *Lamiré et al., 2023*). 17 $\beta$ -estradiol (here referred to as estradiol) is the most potent and biologically  
39 active form of estrogen, and is used in a variety of clinical contexts including contraception, hormone replacement  
40 therapy, and feminizing hormone therapy. Our discovery of a role for estradiol in promoting habituation learning  
41 is not surprising, as it has well-documented effects on other learning and memory processes. This has been most  
42 extensively characterized in the hippocampus, where estradiol promotes behavioural performance and the cellular/circuit hallmarks of hippocampal plasticity, including Long-term potentiation (LTP) and modulation of dendritic  
43 spine density (*Iqbal et al., 2024; Luine, 2014; Finney et al., 2020; Nilsson and Gustafsson, 2002*). While the role  
44 of estradiol in habituation is less well explored, it has previously been shown to increase memory retention for  
45 olfactory habituation in mice (*Dillon et al., 2013*), indicating it plays conserved roles in plasticity regulation across  
46 paradigms.

47 Estradiol signals via two classes of Estrogen Receptors (ERs): the ligand-activated transcription factors ER $\alpha$ ,  
48 and ER $\beta$ , and the seven-transmembrane G-protein coupled receptor Gper1. ER $\alpha/\beta$  are thought to mediate the  
49 long-term "genomic" effects of estrogens through transcriptional activation of target genes, and thus are typically  
50 termed nuclear ERs. Estradiol also has acute effects on physiology – often called "non-genomic", or "membrane-  
51 initiated". These acute effects are thought to be mediated largely by the G protein coupled receptor Gper1, which  
52 signals via multiple G-proteins, and potentially epidermal growth factor (EGF) receptor transactivation (*Prossnitz*  
53 *and Barton, 2023; Revankar et al., 2005; Filardo et al., 2000*). In this way, Gper1 signalling impacts multiple core  
54 second-messenger systems, including: adenylyl cyclase, ERK, PI3K-Akt, and nitric oxide synthase. There is evidence  
55 from receptor-specific pharmacology and genetic knockout experiments in mice for a role of all of these receptors  
56 in hippocampal plasticity (*Finney et al., 2020; Koitmäe et al., 2023; Briz et al., 2015*).

57 Pharmacological experiments in adult zebrafish indicate that nuclear ERs are involved in the consolidation of  
58 object recognition memory, while Gper1 is involved in the consolidation of object placement memory (*Naderi et al.,*  
59 *2020*). Therefore, both classes of receptors are good candidates for plasticity regulation in zebrafish. However, we  
60 are unaware of any previous studies using genetic knockout lines to test for the function of zebrafish ERs in learning  
61 or memory.

62 In this project we aimed to identify the relevant ER(s) mediating the effects of estradiol on habituation using  
63 genetic knockout alleles. Zebrafish have single gene encoding ER $\alpha$  (ER1, *esr1*) and Gper1 (*gper1*), and two homologs  
64 of ER $\beta$ : ER2a (*esr2a*) and ER2b (*esr2a*) (*Romano et al., 2017; Menuet et al., 2002*). We found that none of these  
65 mutants were insensitive to estradiol's effects, indicating that estradiol acts in this context via an alternative receptor  
66 or pathway. Surprisingly, our experiments found that mutants for *esr1*, *esr2a*, and *gper1* actually habituate more  
67 than their sibling controls. While the effect size is small and behavioural-genetic experiments can be variable, these  
68 data indicate that these ERs actually act to inhibit habituation learning, rather than mediating the habituation-  
69 promoting effects of estradiol that we observe pharmacologically.



**Figure 1. Estradiol increases habituation learning.**

A) In response to a dark flash (DF), larval zebrafish perform a large turning manoeuvre termed an "O-bend" response.

B) High-throughput setup for recording and quantifying responsiveness using a high-speed camera recording at 444hz observing larvae in 300-well plates. Scale bar = 10mm.

C) Treatment with estradiol (red) results in more rapid and profound decreases in the probability of response to DF stimuli during habituation training relative to DMSO vehicle controls (black). DF stimuli are delivered at 1-minute intervals, in 4 blocks of 60 stimuli, separated by 1hr of rest (from 0:00-7:00). 1.5 hours later a block of 30 vibration stimuli are delivered at 1-minute intervals (i). Each dot is the probability of response to one DF. Lines are smoothed in time with a Savitzky-Golay filter (window = 15 stimuli, order = 2).

D-G) Distributions responsiveness for different epochs of the experiment. Each dot is the per-fish average of the epoch. Statistical significance was calculated using Mann-Whitney U test, \*\*\* =  $p < 0.001$ . D) the naive response to the first 5 DF stimuli; E) the mean response to the remaining DF stimuli in the Block 1 (DFs 6:60); F) the trained response to the last 45 DFs in all four training blocks (DFs 16:60,76:120,136:180,196:240); G) the 30 vibration stimuli delivered with a tap from a solenoid on the 300-well plate platform.

H) Cumulative mean difference (CMD) plot quantifying relative habituation performance after estradiol treatment. These plots display the cumulative average differences in the mean response across larvae of the treatment group (estradiol) relative to the control group (DMSO). Difference from 0 reflect a divergence in the change in responsiveness across the 240 DF stimuli in the 4 training blocks, with positive values reflecting increased habituation. The widths of the line is a bootstrapped 99.5% confidence intervals. The gray boxed region reflects the expected non-significant effect size (Randlett et al., 2019).

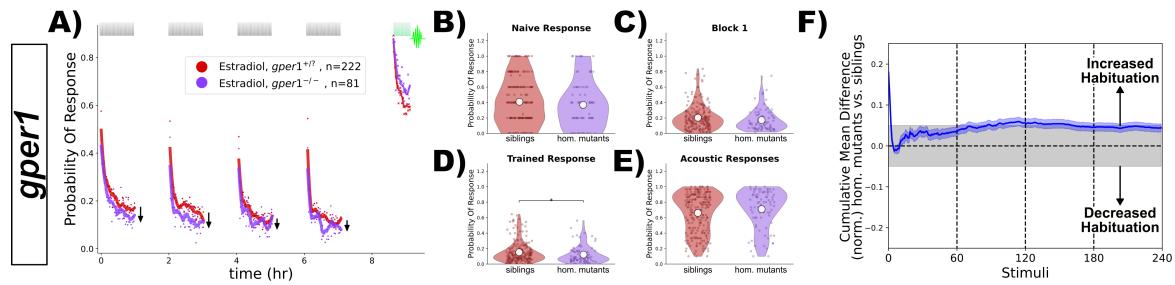
Treatment groups are: Estradiol = 10 $\mu$ M estradiol treatment ( $n = 345$  fish); DMSO = 0.1% DMSO vehicle controls ( $n = 548$  fish)

## Results

### Estradiol increases visual habituation learning

In response to a sudden global darkening stimulus, which we refer to as a dark flashe (DF), larval zebrafish execute an "O-bend" maneuver, characterized by a deep "O"-shaped bend and a high-amplitude turn (Burgess and Granato, 2007, Figure 1a). Habituation learning manifests as a progressive reduction in response to repeated stimuli, and this learning can be retained for seconds/minutes, or hours/days for short-, and long-term habituation, respectively (Rankin et al., 2009). We use high-speed cameras, machine-vision analysis, and 300-well plates to quantify habituation across large populations of larvae to identify molecular/genetic mechanisms of long-term habituation (Figure 1A,B, Randlett et al., 2019; Lamiré et al., 2023). When stimulated with DFs repeated at 1-minute intervals in blocks of 60 stimuli, larval zebrafish exhibit long-term habituation, reducing not only the probability of executing a response, but also modulating the latency and other kinematic aspects of the response (Randlett et al., 2019).

Our previous small-molecule screening experiments identified multiple synthetic Estrogen Receptor agonists as positive modulators of DF habituation learning, including ethinyl estradiol, estradiol valerate, and hexestrol (Lamiré et al., 2023). The major effect we observed was a stronger decrease in the probability of executing a O-bend response during the training/learning blocks. We have confirmed and extended these results using estradiol, which is the major natural estrogen in vertebrates.



**Figure 2. *gper1* mutants do not show habituation deficits after treatment with estradiol.**

**A)** Homozygous *gper1*<sup>-/-</sup> mutants ( $n = 81$  fish, purple) do not show impaired habituation relative to sibling controls (*gper1*<sup>+/+</sup>) and *gper1*<sup>+/+</sup> ( $n = 222$  fish, red). Rather, there is a slight suppression of responsiveness in the mutant group (arrows), indicating weakly increased habituation. Each dot is the probability of response to one stimulus. Lines are smoothed in time with a Savitzky-Golay filter (window = 15 stimuli, order = 2).

**B-E)** No significant differences are observed in the responsiveness distributions for the naive response to the first 5 DF stimuli (B), during the first training block (C), or the vibration response (E), while a subtle but statistically significant decrease in responsiveness is observed in the trained response (D). Statistical significance was calculated using Mann-Whitney U test, \* =  $p < 0.05$ .

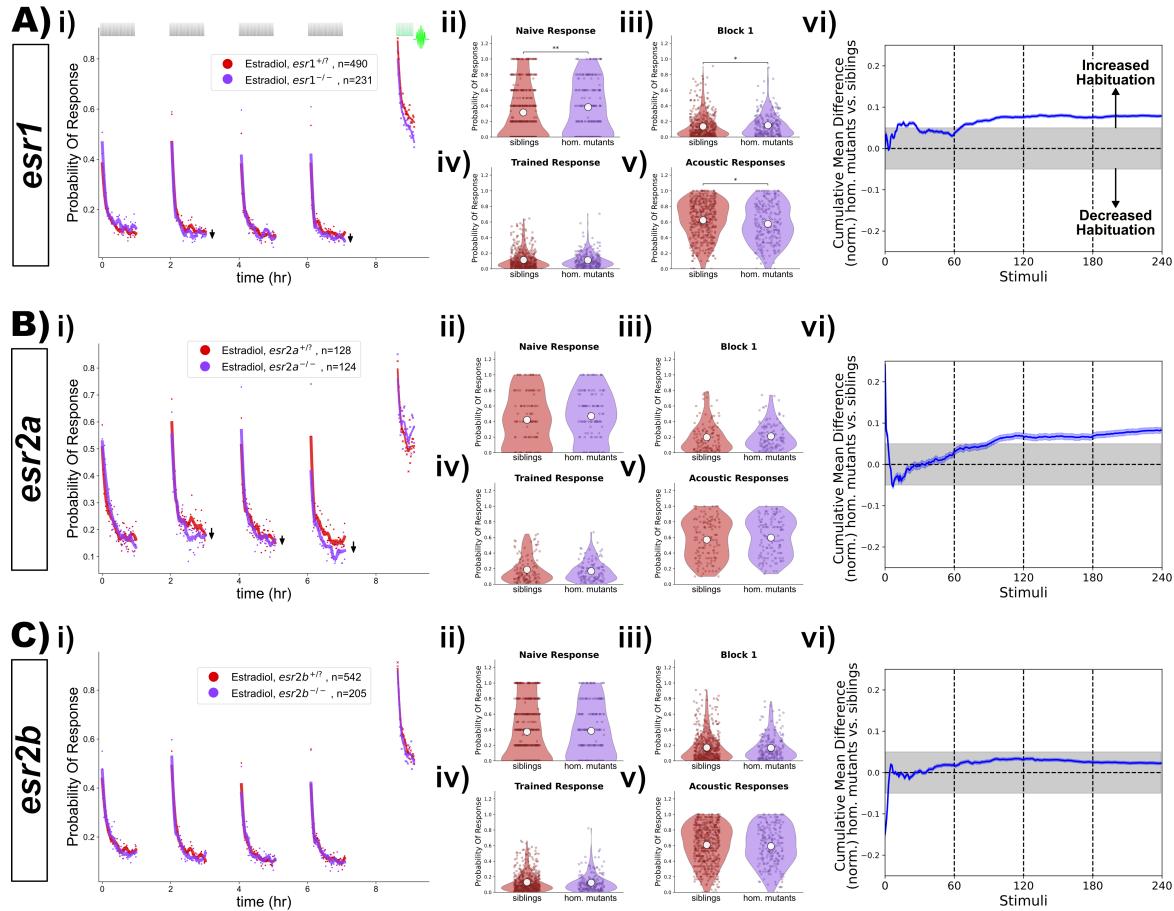
**F)** Cumulative mean difference (CMD) plot quantifying habituation performance of mutants relative to sibling controls, consistent with slightly increased habituation rate in mutant larvae.

An acute dose of 10 $\mu$ M estradiol potently increases habituation learning, which is observable when the response probability of the population of estradiol-treated larvae is compared with DMSO-treated vehicle controls (Figure 1C-H). Consistent with our previous experiments (Lamiré et al., 2023), there is a reduction in the naive responsiveness of the estradiol-treated larvae to the first DF stimuli (Figure 1D), but the major effect is observed during the training phase (Figure 1C,E,F), as is revealed by the consistent positive deviation in the cumulative mean difference (CMD) plots that are normalized to the naive response level in order to quantify response suppression indicative of habituation performance (Figure 1H, Randlett et al., 2019). Importantly, the responsiveness of the larvae to vibration stimuli delivered after the DF stimulation (Figure 1Ci), which elicit c-bend escape responses (Kimmel et al., 1974), is indistinguishable from controls (Figure 1G). From this we conclude that estradiol does not affect global arousal levels but rather has specific effects on habituation learning.

### Gper1 is dispensable for estradiol-promoted habituation learning

The effects of estradiol that we have observed occur very rapidly – larvae are only pretreated with estradiol for  $\approx 25\text{min}-1\text{hr}$  before the first DF. This is the time necessary to set the apparatus and begin the experiment. Since the nuclear hormone receptors are thought to primarily exert their effects via transcriptional alterations, this necessitates a delay in their signaling. For this reason, we first hypothesized that membrane-initiated signaling through Gper1 was the most likely mechanism.

To test this we used a knockout allele *gper1*<sup>uab102</sup> (Romano et al., 2017), and generated larvae from *gper1*<sup>uab102</sup> heterozygous or homo/heterozygous crosses to generate clutches of larvae of mixed genotypes. Larvae were treated with estradiol during habituation, and were subsequently genotyped. We reasoned that if *gper1* is required for the effect of estradiol on habituation, mutants would be insensitive to estradiol and habituate significantly less than sibling controls. Contrary to this hypothesis, we found that *gper1* mutants showed no deficits in habituation (Figure 2). Remarkably, rather than observing the anticipated inhibition of habituation, *gper1* mutants appeared to habituate slightly more than controls, with the responsiveness level slightly but consistently below the sibling controls across stimuli (Figure 2A). This is further supported by a weak but statistically significant decrease in the responsiveness of the larvae during the training period (Figure 2D), and a deviation towards positive values in the CMD plot (Figure 2F). From these experiments we conclude that Gper1 agonism does not promote habituation learning, but rather may act to suppress it.



**Figure 3. *esr1*, *esr2a* and *esr2b* mutants do not show habituation deficits after treatment with estradiol.**

- A) Homozygous *esr1*<sup>-/-</sup> mutants (n = 231 fish) do not show impaired habituation relative to sibling controls (*esr1*<sup>+/+</sup>) and *esr1*<sup>(+/-)</sup>, n = 490 fish.  
B) Homozygous *esr2a*<sup>-/-</sup> mutants (n = 214 fish) do not show impaired habituation relative to sibling controls (*esr2a*<sup>+/+</sup>) and *esr2a*<sup>(+/-)</sup>, n = 128 fish.  
C) Homozygous *esr2b*<sup>-/-</sup> mutants (n = 205 fish) do not show impaired habituation relative to sibling controls (*esr2b*<sup>+/+</sup>) and *esr2b*<sup>(+/-)</sup>, n = 542 fish.

i)-vi) For each lettered section:

i) Responsiveness to stimuli comparing homozygous mutants to sibling controls (heterozygous or wild-type). Each dot is the probability of response to one stimulus. Lines are smoothed in time with a Savitzky-Golay filter (window = 15 stimuli, order = 2). Suppression of responsiveness is indicated by arrows, potentially reflecting increased habituation.

ii-v) Distributions responsiveness for different epochs of the experiment. Each dot is the per-fish average of the epoch. Statistical significance was calculated using Mann-Whitney U test, \* = p < 0.05, \*\* = p < 0.01. ii) the naive response to the first 5 DF stimuli; iii) the mean response to the remaining DF stimuli in the Block 1 (DFs 6:60); iv) the trained response to the last 45 DFs in all four training blocks (DFs 16:60,76:120,136:180,196:240); v) the 30 vibration stimuli delivered with a tap from a solenoid on the 300-well plate platform.

vi) Cumulative mean difference (CMD) plot quantifying habituation performance of mutants relative to sibling controls

## 115 Nuclear ERs are dispensable for estradiol-promoted habituation learning

116 Since we found the *gper1* was unnecessary for the habituating-promoting effects of estradiol, we next focused on  
117 the three nuclear receptors in the zebrafish genome: *esr1*, *esr2a* and *esr2a*. Using the same strategy as for *gper1*, we  
118 analyzed previously established knockout mutants (*esr2a<sup>uab134</sup>*, *esr2a<sup>uab134</sup>*, and *esr2b<sup>uab127</sup>*), looking for a mutant  
119 with insensitivity to estradiol. However, we failed to identify any deficits in habituation (*Figure 3*). To our surprise,  
120 we again found that both *esr1* and *esr2a* mutants showed subtle increases in habituation (*Figure 3Avi,Bvi*), similar  
121 in magnitude to what we had seen for *gper1* mutants (*Figure 2F*). From these data we conclude that none of the  
122 nuclear ERs are required for the effects of estradiol on promoting habituation. As for Gper1, ER1 and ER2a show a  
123 weak inhibitory effect on habituation, indicating that they also act to suppress habituation learning.

## 124 Nuclear ERs are simultaneously dispensable for estradiol-promoted habituation learning

125 While our experiments demonstrated that *esr1*, *esr2a* and *esr2a* mutants remain sensitive to estradiol, it is plausible  
126 that they could act in a redundant fashion to mediate the effects of estradiol on habituation, perhaps via co-  
127 expression in a critical cell type. To test for this possibility, we generated combinations of mutants by crossing  
128 individual lines together (*Figure 4*). A likely scenario could be that the two ER $\beta$  paralogs, ER2a and ER2b, act  
129 redundantly. However, we found that double mutants for *esr2a;esr2b* did not show habituation deficits (*Figure 4A*).  
130 Neither did we observe suppression of habituation in double mutants for *esr1;esr2a* (*Figure 4B*), or *esr1;esr2b*  
131 (*Figure 4C*). Finally, we tested triple mutants for all three nuclear ERs (*esr1;esr2a;esr2b*), but again failed to identify  
132 suppressions in habituation (*Figure 4D*).

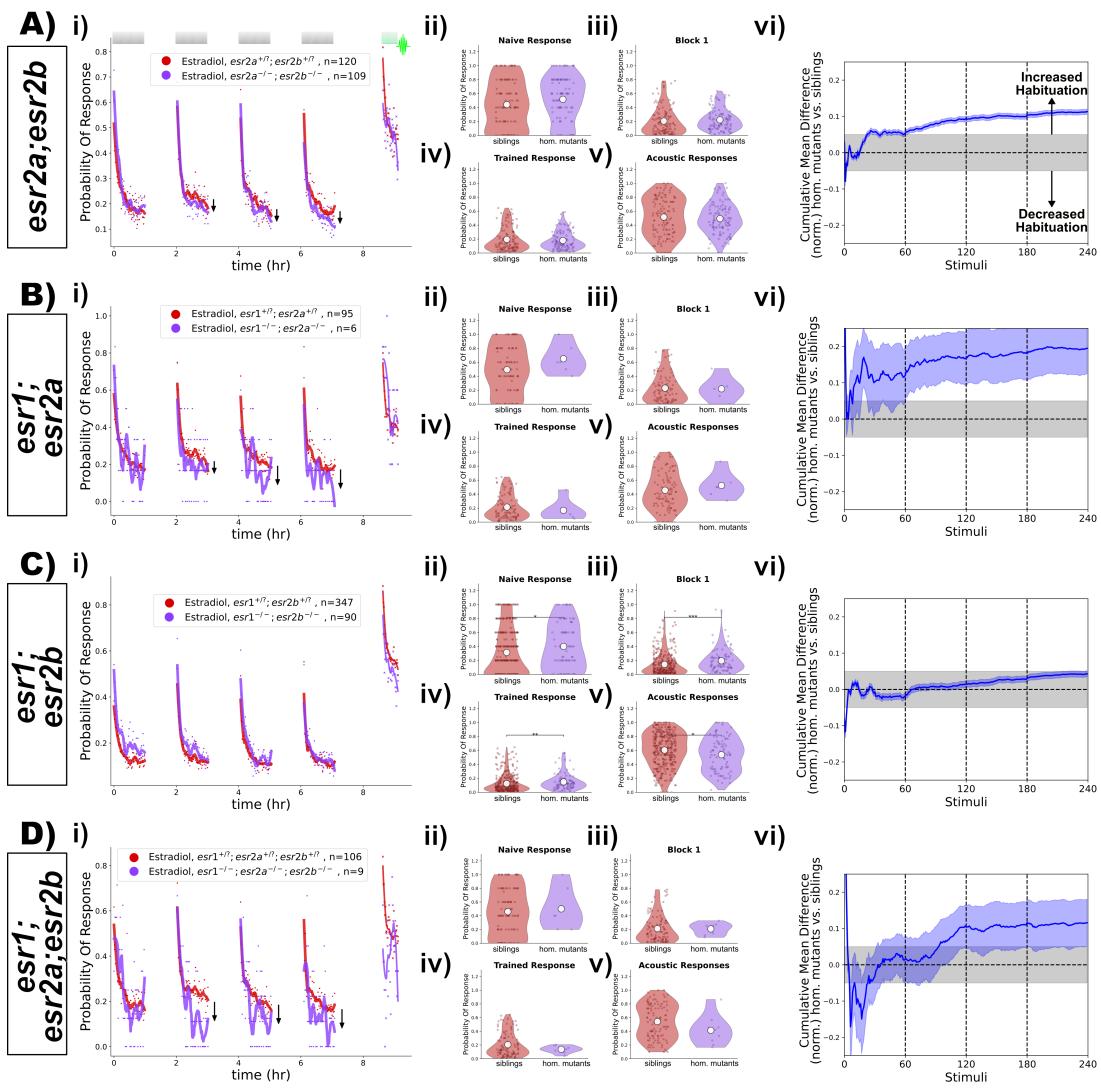
133 We note that we did see a statistically "significant" increase in DF responsiveness in the *esr1;esr2b* double mutants  
134 (*Figure 4Cii-iv*). This is attributable to a small increase in naive responsiveness, and did not result from  
135 habituation deficits according to our normalized CMD analysis (*Figure 4Cvi*). Considering that we did not observe  
136 any increased responsiveness in the triple mutants (*Figure 4D*), we conclude that the observed increased respon-  
137 siveness in the double mutants is likely a result of biological noise, that only rose to statistical "significance" due to  
138 the very large number of larvae tested in our high-throughput experiments.

139 Consistent with the paradoxical effect of increased habituation in *esr1* and *esr2a* single mutants (*Figure 3A,B*), we  
140 again observed that double and triple mutants containing these genes also showed a slight increase in habituation  
141 (with the exception of the *esr1;esr2b* double mutants). This adds further support to the model in which ER1 and  
142 ER2b act to suppress learning in this context, rather than promote it.

## 143 Gper1 and nuclear ERs are simultaneously dispensable for estradiol-promoted habitua- 144 tion learning

145 While it is unclear how the GPCR Gper1 might act redundantly with the nuclear ERs, we nevertheless decided to  
146 test this possibility by combining the *gper1* and the *esr1*, *esr2a*, and *esr2b* mutations (*Figure 5*). As with the previous  
147 iterations of this experiment, we did not find combinations of mutants with suppressed habituation (*Figure 5*).  
148 Consistent with our model of (*gper1*, *esr1* and *esr2a*) having inhibitory affects on learning, we again found that most  
149 of these combinations of mutants showed evidence of increased habituation (*Figure 5i,vi*). While the responsiveness  
150 distributions did not show significant differences (*Figure 5ii-iv*), the normalized CMD plots consistently showed  
151 positive deviations, which reflect increased habituation (*Figure 5vi*).

152 Despite the fact that we analyzed the behaviour of 1152 larvae and successfully genotyping all 4 ER genes  
153 in 373 individuals (after >4600 genotyping PCRs), we were only able to identify a single quadruple mutant larva  
154 lacking all known ERs (*Figure 5E*). This is likely simply due to the limitations of combinatorial Mendelian inheritance,  
155 and the fact that *esr1* and *esr2a* are linked on chromosome 20. While it is dubious to conclude much from an  
156 n = 1 experiment, we find it remarkable that this larva exhibits the strongest increased habituation of all of our  
157 experiments, with suppression of responses (*Figure 5Ei*), and strong positive deviation in the CMD plot (*Figure 5Ev*).



**Figure 4. Double and triple mutant combinations of *esr1*, *esr2a* and *esr2b* do not show habituation deficits after treatment with estradiol.**

A) Homozygous *esr2a<sup>(-/-)</sup>;esr2b<sup>(-/-)</sup>* double mutants (n = 109 fish) do not show impaired habituation relative to double heterozygous or homozygous sibling controls (<sup>(+/-)</sup>/<sup>(+/-)</sup>, n = 120 fish).

B) Homozygous *esr1<sup>(-/-)</sup>;esr2a<sup>(-/-)</sup>* double mutants (n = 6 fish) do not show impaired habituation relative to double heterozygous or homozygous sibling controls (<sup>(+/-)</sup>/<sup>(+/-)</sup>, n = 95 fish).

C) Homozygous *esr1<sup>(-/-)</sup>;esr2b<sup>(-/-)</sup>* double mutants (n = 90 fish) do not show impaired habituation relative to double heterozygous or homozygous sibling controls (<sup>(+/-)</sup>/<sup>(+/-)</sup>, n = 347 fish).

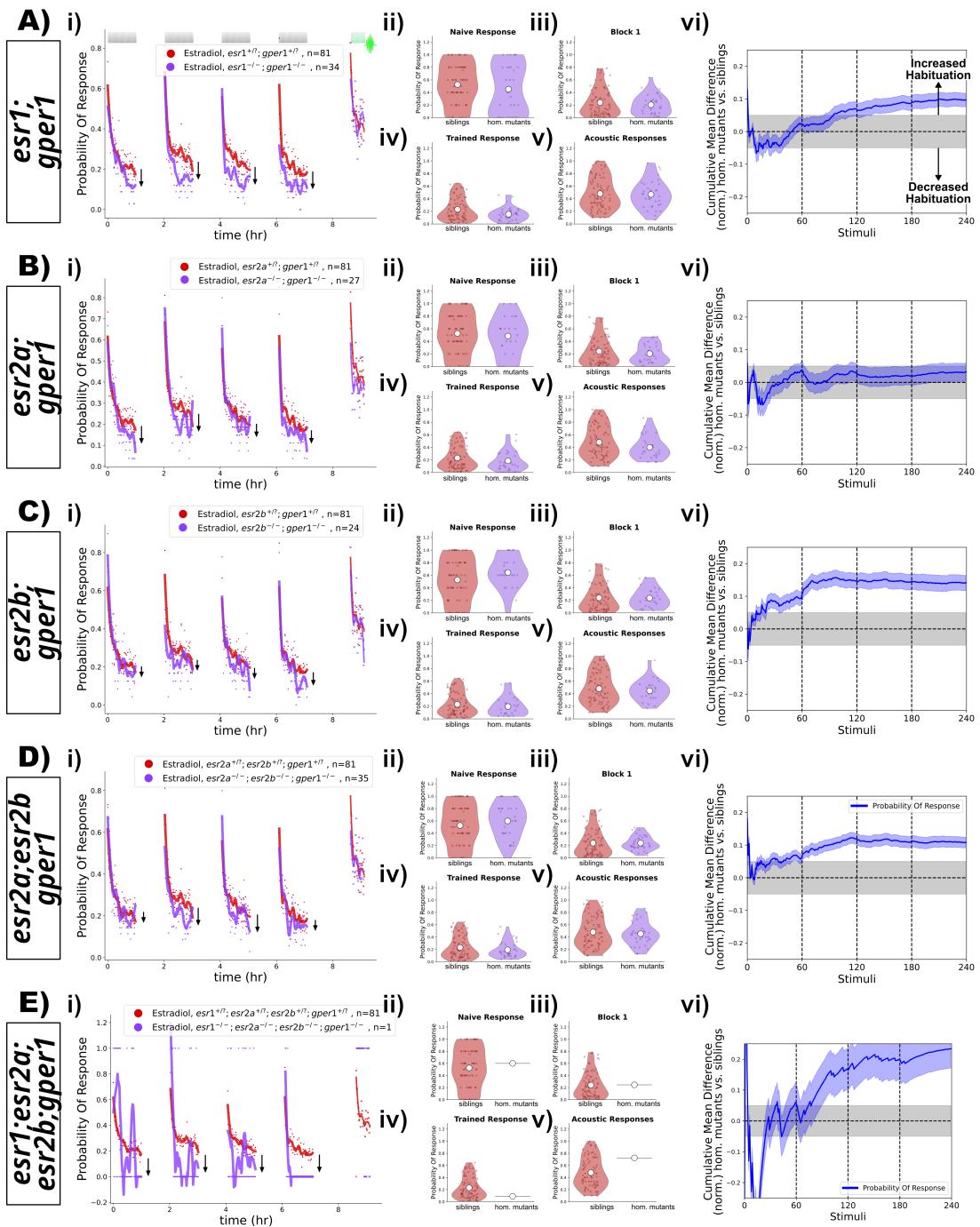
D) Homozygous *esr1<sup>(-/-)</sup>;esr2a<sup>(-/-)</sup>;esr2b<sup>(-/-)</sup>* triple mutants (n = 9 fish) do not show impaired habituation relative to triple heterozygous or homozygous sibling controls (<sup>(+/-)</sup>/<sup>(+/-)</sup>/<sup>(+/-)</sup>, n = 106 fish).

i)-vi) For each lettered section:

i) Responsiveness to stimuli comparing homozygous mutants to sibling controls (heterozygous or wild-type). Each dot is the probability of response to one stimulus. Lines are smoothed in time with a Savitzky-Golay filter (window = 15 stimuli, order = 2).

ii-v) Distributions responsiveness for different epochs of the experiment. Each dot is the per-fish average of the epoch. Statistical significance was calculated using Mann-Whitney U test, \* = p < 0.05, \*\* = p < 0.01. ii) the naive response to the first 5 DF stimuli; iii) the mean response to the remaining DF stimuli in the Block 1 (DFs 6:60); iv) the trained response to the last 45 DFs in all four training blocks (DFs 16:60,76:120,136:180,196:240); v) the 30 vibration stimuli delivered with a tap from a solenoid on the 300-well plate platform.

vi) Cumulative mean difference (CMD) plot quantifying habituation performance of mutants relative to sibling controls.



**Figure 5. Double, triple and quadruple mutant combinations of *esr1*, *esr2a*, *esr2b*, and *gper1* do not show habituation deficits after treatment with estradiol.**

**A)** Homozygous *esr1<sup>-/-</sup>;gper1<sup>-/-</sup>* double mutants ( $n = 34$  fish, purple) do not show impaired habituation relative to double heterozygous or homozygous sibling controls ( $n = 81$  fish, red). **B)** Homozygous *esr2a<sup>-/-</sup>;gper1<sup>-/-</sup>* double mutants ( $n = 27$  fish, purple) do not show impaired habituation relative to double heterozygous or homozygous sibling controls ( $n = 81$  fish, red). **C)** Homozygous *esr2b<sup>-/-</sup>;gper1<sup>-/-</sup>* double mutants ( $n = 24$  fish, purple) do not show impaired habituation relative to double heterozygous or homozygous sibling controls ( $n = 81$  fish, red). **D)** Homozygous *esr2a<sup>-/-</sup>;esr2b<sup>-/-</sup>;gper1<sup>-/-</sup>* triple mutants ( $n = 35$  fish, purple) do not show impaired habituation relative to heterozygous or homozygous sibling controls ( $n = 81$  fish, red). **E)** A single homozygous *esr1<sup>-/-</sup>;esr2a<sup>-/-</sup>;esr2b<sup>-/-</sup>;gper1<sup>-/-</sup>* quadruple mutant ( $n = 1$  fish, purple) does not appear to show impaired habituation relative to heterozygous or homozygous sibling controls ( $n = 81$  fish, red).

**i-vi)** For each lettered section: **i)** Responsiveness to stimuli comparing homozygous mutants to sibling controls (heterozygous or wild-type). Each dot is the probability of response to one stimulus. Lines are smoothed in time with a Savitzky-Golay filter (window = 15 stimuli, order = 2). **ii-v)** Distributions responsiveness for different epochs of the experiment. Each dot is the per-fish average of the epoch. Statistical significance was calculated using Mann-Whitney U test, \* =  $p < 0.05$ , \*\* =  $p < 0.01$ . **ii)** the naive response to the first 5 DF stimuli; **iii)** the mean response to the remaining DF stimuli in the Block 1 (DFs 6:60); **iv)** the trained response to the last 45 DFs in all four training blocks (DFs 16:60,76:120,136:180,196:240); **v)** the 30 vibration stimuli delivered with a tap from a solenoid on the 300-well plate platform. **vi)** Cumulative mean difference (CMD) plot quantifying habituation performance of mutants relative to sibling controls. As mutant larvae were all derived from the same experiments, the sibling control data (red) is the same in **A-E**.

# 158 Discussion

## 159 The promotion of habituation learning by estradiol is mediated by an unidentified target

160 Our experiments indicate that Gper1, ER1, ER2a and ER2b do not mediate the positive effects of estradiol on  
161 habituation learning. As this is fundamentally a negative result, it is difficult to conclusively demonstrate this  
162 beyond any doubt. One major caveat relates to the actual functional nature of the mutant alleles that we have  
163 used. These are all Cas9-generated small deletions resulting in frameshift mutations that lead to early stop codons,  
164 and are thus predicted null/knockout lines. Despite this genetic confidence, it is always possible that residual  
165 activity could still remain, perhaps via alternate splicing or alternate start codons. For this reason, we were originally  
166 unsure if we would publish these results at all. However, we recognize that this bias against publishing negative  
167 results is bad for science. Negative results from well-designed and executed experiments are of value for the  
168 community and making this knowledge public is our duty as responsible scientists (*Mlinarić et al., 2017*). These  
169 high-throughput neurogenetic experiments are also a massive amount of work – raising, testing and genotyping  
170 thousands of zebrafish larvae. This was undertaken primarily by technicians and Master's students, all of whom  
171 deserve recognition, irrespective of the perceived "impact" of the outcome of their experiments.

172 While the possibility of "residual activity" in our mutants is a clear limitation of our approach, we argue that this  
173 alternative interpretation is very unlikely. The *esr1<sup>uab118</sup>* and *esr2b<sup>uab127</sup>* alleles both exhibited a lack of estradiol  
174 responsiveness in other tissues (*Romano et al., 2017*), and *esr2b<sup>uab127</sup>* mutants are female sterile/subfertile (D.  
175 Gorelick, personal communication), indicating a non-functional receptor. Similarly, *gper1<sup>uab102</sup>* mutants show a lack  
176 of estradiol responsiveness in heart rate modulation (*Romano et al., 2017*). Interestingly this was only observed in  
177 maternal-zygotic mutants. While it seems unlikely that sufficient maternal mRNA/protein for Gper1 could persist  
178 in 5dpf larvae, we can formally rule this out with our current datasets. The *esr2a<sup>uab134</sup>* mutants have no previously  
179 published phenotype, and so we do not have an independent positive control for the nature of this allele. However,  
180 the best evidence we have against the "residual activity" hypothesis is that we actually found phenotypes in our  
181 assays for *esr1<sup>uab118</sup>*, *esr2a<sup>uab134</sup>*, and *gper1<sup>uab102</sup>* mutants. These phenotypes are just of the unexpected sign, where  
182 mutants show increased habituation (discussed below).

## 183 Potential mechanisms for estradiol-promoted habituation learning

184 We have concluded that the lack of habituation deficits in our mutants is due to the presence of an alternative re-  
185 ceptor or pathway that mediates the learning-promoting effects of estradiol. In support of this model, a very recent  
186 study of estradiol-induced thrombosis came to the same conclusion, namely that an unidentified target mediates  
187 this process in larval zebrafish (*Yu et al., 2024*). Importantly, these thrombosis experiments were performed on  
188 different knockout alleles which are full genetic deletions, in which "residual activity" is not plausible. Together with  
189 our observations, these data suggest that a novel estradiol target exists which has multiple critical functions (at  
190 least in zebrafish).

191 What might this unidentified target be? Various leads exist in the literature. One hypothesis posits the existence  
192 of an unidentified "Gq-mER" (Gq-coupled membrane estrogen receptor) (*Qiu et al., 2006; Vail and Roepke, 2019*),  
193 and therefore estradiol may signal via additional GPCRs beyond Gper1. Another possibility is an interaction between  
194 estradiol and other membrane receptors, for example: the Voltage-Gated Sodium Channel Nav1.2 (*Sula et al., 2021*;  
195 *Treviño and Gorelick, 2021*), transient receptor potential (TRP) channels (*Payrits et al., 2017; Ramírez-Barrantes  
196 et al., 2020*), or various other ion channels (*Kow and Pfaff, 2016*). The robust nature of our "non-canonical" but  
197 clearly estradiol-dependent phenotype, combined with the high-throughput nature of our behavioural assays, could  
198 be an ideal assay for future screening efforts to attempt to identify novel estradiol target(s).

## 199 **Multiple Estrogen Receptors act to suppress habituation learning.**

200 While we were surprised to find that the classical ERs do not promote habituation, we were shocked to find clear  
201 evidence of the opposite! We found that *esr1<sup>uab118</sup>*, *esr2a<sup>uab134</sup>*, and *gper1<sup>uab102</sup>* mutants habituate more than their  
202 sibling controls (**Figure 1**, **Figure 2**), consistent with a role for these ERs in inhibiting habituation. While effect sizes of  
203 these magnitudes border on those that are easily dismissible as "noise", they were not only observed in the single  
204 mutants experiments, but also in the double and triple mutant combinations of these alleles (**Figure 3-Figure 5**), pro-  
205 viding good evidence that they are biologically meaningful effects. In fact, these multi-mutants generally exhibited  
206 larger effect sizes, consistent with an additive interaction.

207 Untangling the mechanisms of ER1-, ER2a- and Gper1-dependent suppression of habituation will require con-  
208 siderable further work. The additive interaction we observed genetically indicates that the ERs act cooperatively to  
209 suppress habituation learning. All three receptors are expressed in the larval zebrafish brain (**Thisse and Thisse,**  
210 **2008; Romano et al., 2017**), but whether they are acting in the same or different cell types awaits characterization.  
211 The study of this inhibitory pathway may be challenging since it opposes the major learning-promoting effect of  
212 estradiol, and therefore may be more straightforward to study after the identification and deletion of the estradiol  
213 target that promotes habituation.

## 214 **Conclusion**

215 What began as a straightforward study to identify the receptor(s) that mediate the habituation-promoting effects  
216 of estradiol has instead led us to a surprising and paradoxical result; canonical ERs do regulate habituation, but  
217 are suppressive and act in opposition to the habituation-promoting effects of estradiol. This fits with the general  
218 theme of our studies of this habituation paradigm – we find increasing complexity and contradiction within this  
219 "simple" learning process the deeper we look. This began with our detailed observations of behaviour, leading us  
220 to conclude that habituation results from a distributed plasticity processes that adapts different aspects of behavior  
221 independently (**Randlett et al., 2019**). We believe that this property underlies our subsequent discoveries of phar-  
222 macological and genetic manipulations that can result in either specific changes in specific aspects of habituation  
223 (but not others), or even opposing effects, where a single manipulation can simultaneously increase and decrease  
224 habituation, depending on which component of behavior is measured (**Randlett et al., 2019; Lamiré et al., 2023**).  
225 This complexity appears to be a fundamental property of habituation (**McDiarmid et al., 2019**), and that the study of  
226 habituation will likely continue to surprise us, hopefully leading to unexpected insights into the nature of plasticity  
227 underling learning and memory.

# 228 **Materials and Methods**

## 229 **Animals**

230 All experiments were performed on larval zebrafish at 5 days post fertilization (dpf), raised at a density of ≈1  
231 larvae/mL of E3 media supplemented with 0.02% HEPES pH 7.2. Larvae were raised in a 14:10h light/dark cycle at  
232 28-29°C. Adult zebrafish were housed, cared for, and bred at the following zebrafish facilities: Plateau de Recherche  
233 Expérimentale en Criblage In Vivo (PRECI, SFR Biosciences, Lyon) and the Animalerie Zebrafish Rockefeller (AZR, SFR  
234 Santé Lyon Est, Lyon). Adult zebrafish used to generate larvae were housed in accordance with regulations of the  
235 PRECI and AZR facilities, which are regulated by an internal animal wellbeing committee, and were approved by  
236 the animal welfare committee (comité d'éthique en expérimentation animale de la Région Rhône-Alpes: CECCAPP,  
237 Agreement # C693870602). Behaviour experiments were performed at the 5dpf stage, and are thus not subject to  
238 ethical review, but these procedures do not harm the larvae.

239 Mutant lines were obtained from D. Gorelick's lab, and were of the following alleles:

240 *esr1<sup>uab118</sup>* is a 4bp deletion (ZDB-ALT-180420-2), yielding a predicted null frameshift/stop mutation, confirmed  
241 by a lack of estradiol responsiveness in the heart as assayed by *Tg(5xERF:GFP)<sup>c262</sup>* expression (**Romano et al., 2017**).

242     *esr2a<sup>uab134</sup>* is a 2bp deletion (ZDB-ALT-180420-3), yielding a predicted null frameshift/stop mutation (*Romano*  
243     *et al., 2017*)

244     *esr2b<sup>uab127</sup>* is a 4bp deletion (ZDB-ALT-180420-4), yielding a predicted null frameshift/stop mutation, confirmed  
245     by a lack of estradiol responsiveness in the liver as assayed by *Tg(5xERE:GFP)<sup>c262</sup>* expression (*Romano et al., 2017*).

246     *gper1<sup>uab102</sup>* is a 133bp deletion (ZDB-ALT-180420-1), yielding a predicted null frameshift/stop mutation, con-  
247     firmed by a lack of estradiol responsiveness in heart beating rate in maternal-zygotic mutants (*Romano et al., 2017*).

## 248     **Genotyping**

249     *esr1<sup>uab118</sup>* was genotyped by PCR using the forward/reverse primer pair:

250     GCTGGTCACCTTGAATGCTT/TGAGATGTGAGAGATGACTAGGA with a  $T_M$  of 58°C yielding a 381 bp PCR product that  
251     was digested with the restriction enzyme ApeKI. The mutant product is not digested, and the wild type has two  
252     bands at 177 and 204 bp.

253     *esr2a<sup>uab134</sup>* was genotypes by PCR using the forward/reverse primer pair:

254     CTTCAGCTGCAGGAAGTGGAAAGTCGGGCTTAGCGACTG with a  $T_M$  of 58°C yielding a 236 bp PCR product that was  
255     digested with the restriction enzyme MboI. The mutant product is not digested, and the wild type has two bands at  
256     180 and 56 bp

257     *esr2b<sup>uab127</sup>* was genotypes by PCR using the forward/reverse primer pair:

258     TGGGCCTGAGATGCAGTAGT/GTGTGTGCTTGGCCTC with a  $T_M$  of 60°C yielding a 431 bp PCR product that was  
259     digested with the restriction enzyme Mbil. The mutant product is digested into two bands of 150 and 281 bp and  
260     the wild type into 3 bands of 78, 150 and 198 bp.

261     *gper1<sup>uab102</sup>* was genotypes by PCR using the forward/reverse primer pair:

262     ATGGAGGAGCAGACTACCAATGTG/CCATCCAGATGAGGCTGCAA with a  $T_M$  of 60°C yielding a mutant product of 372bp  
263     and a wild type product of 505 bp.

## 264     **Pharmacology**

265     β-Estradiol (Sigma E2758, here referred to as "estradiol") was dissolved in dimethyl sulfoxide (DMSO) and stored  
266     at -20°C. Larvae were treated with estradiol immediately before the behavioural assay by pipetting 10-30uL of 10x  
267     solution directly into the behavioural wells, always with a final concentration of 0.1% DMSO in E3.

## 268     **Habituation behaviour testing**

269     Larval behavior was evaluated in 300-well plates using an updated version of the experimental setup previously  
270     described (*Randlett et al., 2019; Lamiré et al., 2023*). Briefly, 300-well plates were custom made using laser-cut  
271     acrylic sheets where each well measures 8 mm in diameter and 6 mm in depth, corresponding to an approximate  
272     water volume of 300 µL. These plates are suspended under a water bath held at 31°C, acting as a heated lid to  
273     minimize condensation and maintain a 29°C water temperature within each well. Behavioral recordings were made  
274     using a Mikrotron CXP-4 camera running at 444hz in conjunction with a Silicon Software frame grabber (Marathon  
275     ACX-QP, Basler), illuminated by IR LEDs (TSHF5410, digikey.com). Visual stimuli were presented using a rectangular  
276     array of 155 WS2813 RGB LEDs (144LED/M, aliexpress.com). For the DF stimulus, the LEDs were briefly switched off  
277     (1 s), then linearly returned to the original brightness over a 20 s interval. Vibration/Tap stimuli were administered  
278     using a solenoid (ROB-10391, Sparkfun). This behavioral paradigm was designed to be symmetrical: each 1 hr block  
279     of stimulation was followed by 1 hr of rest. During these rest periods, the camera was moves using a stepper motor  
280     controlled linear actuator (Hanpose HPV4, 500cm), which moved the camera between two plates, allowing us to  
281     screen up to 600 fish per experiment across two 300-well plates.

282     Control of the apparatus (RGB LEDs, solenoid, camera linear actuator) was implemented via a Raspberry Pi Pico  
283     microcontroller running CircuitPython (<https://circuitpython.org/>) (code : [code.py](#)) and custom Python software  
284     which handled the camera acquisition via the [Python wrapper of the Silicon Software Framegrabber SDK](#), triggered

285 stimuli via the Raspberry Pi Pico, and tracked the head and tail coordinates of the larvae across the 300-wells at a  
286 baseline framerate of between 20-30hz (code : [Run\\_BigRig2.py](#)). When a stimulus is delivered (DF or Vibration/Tap),  
287 a 1-second "Burst" video is recorded at the full frame rate as a Tiff file, from which the head and tail coordinates  
288 are subsequently tracked offline (code : [TrackBurst\\_BigRig.py](#)). Larval zebrafish tracking was done via background  
289 subtraction and morphological operations implemented using multiple open-source packages, including: OpenCV  
290 ([Bradski, 2000](#)), scikit-image ([Van der Walt et al., 2014](#)), NumPy ([Harris et al., 2020](#)), SciPy ([Virtanen et al., 2020](#)),  
291 and Numba ([Lam et al., 2015](#)).

## 292 Data analysis

293 Data was analyzed in Python using custom written analysis scripts (code : [Analyze\\_EsrHab.py](#)). Responses to DFs  
294 and vibration/taps were identified as movement events that had a cumulative tail bend angle greater than 3 radian  
295 and 1 radian, respectively. Data was analyzed using multiple open-source packages, including: NumPy ([Harris et al.,  
296 2020](#)), SciPy ([Virtanen et al., 2020](#)), Numba ([Lam et al., 2015](#)) and Pandas ([Wes McKinney, 2010](#)). Data was plotted  
297 using Matplotlib ([Hunter, 2007](#)) and seaborn ([Waskom, 2021](#)). Statistical "significance" between the distributions  
298 was tested using the Mann-Whitney U test implemented in Scipy ([Virtanen et al., 2020](#)).

299 The cumulative difference plots to assess changes in habituation performance for the treatments were calcu-  
300 lated as previously ([Randlett et al., 2019](#)), where we first calculated the average response across larvae for each  
301 group for each DF. This generated a mean vector for each group. These two vectors were normalized by dividing  
302 them by the naive response (mean response to the first 5 DFs), and then the treatment group was subtracted  
303 from the control group, yielding a "mean difference" vector between stimulus and controls at each flash. From  
304 this mean difference vector we calculated the cumulative mean distribution using Numpy's 'nancumsum' function  
305 divided by the number of stimuli experienced, or the index of the vector. To generate statistical confidence in  
306 these distributions, we bootstrapped 2000 replicates, and calculated the 99.5% confidence intervals using SciPy's  
307 'stats.norm.interval' function. The assumption of this analysis is that if the two groups are habituating similarly, then  
308 the "mean difference" vector will exhibit a noise distribution centered at a mean of 0, and thus the cumulative mean  
309 distribution would remain near 0. Treatments that affect habituation will show strong increasing or decreasing  
310 cumulative mean distributions, reflecting increased or decreased habituation performance throughout training,  
311 respectively. We use an empirically defined threshold of  $\pm 0.05$  as the statistically meaningful effect size in this  
312 analysis, as is reflected in the shaded gray regions in the plots ([Randlett et al., 2019](#)).

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## 316 Data Availability

317 Software and analysis code is available here: [https://github.com/owenrandlett/2025\\_HabEstrogen](https://github.com/owenrandlett/2025_HabEstrogen). All datasets  
318 used in these analyses are available here: [HabEstrogen\\_Datasets](#).

## 319 References

- 320 Barros LA, Tufik S, Andersen ML. The role of progesterone in memory: an overview of three decades. Neurosci Biobehav Rev.  
321 2015 Feb; 49:193–204.
- 322 Bradski G. The openCV library. Dr Dobb's Journal: Software Tools for the Professional Programmer. 2000; 25(11):120–123.

- 323 **Briz V**, Liu Y, Zhu G, Bi X, Baudry M. A novel form of synaptic plasticity in field CA3 of hippocampus requires GPER1 activation  
324 and BDNF release. *J Cell Biol.* 2015 Sep; 210(7):1225–1237.
- 325 **Burgess HA**, Granato M. Sensorimotor Gating in Larval Zebrafish. *The Journal of Neuroscience.* 2007 May; 27(18):4984–4994.
- 326 **Dillon TS**, Fox LC, Han C, Linster C.  $17\beta$ -estradiol enhances memory duration in the main olfactory bulb in CD-1 mice. *Behav  
327 Neurosci.* 2013 Dec; 127(6):923–931.
- 328 **El-Sherif Y**, Tesoriero J, Hogan MV, Wierszko A. Melatonin regulates neuronal plasticity in the hippocampus. *J Neurosci Res.*  
329 2003 May; 72(4):454–460.
- 330 **Filardo EJ**, Quinn JA, Bland KI, Frackelton AR Jr. Estrogen-induced activation of Erk-1 and Erk-2 requires the G protein-coupled  
331 receptor homolog, GPR30, and occurs via trans-activation of the epidermal growth factor receptor through release of HB-EGF.  
332 *Mol Endocrinol.* 2000 Oct; 14(10):1649–1660.
- 333 **Finney CA**, Shvetcov A, Westbrook RF, Jones NM, Morris MJ. The role of hippocampal estradiol in synaptic plasticity and memory:  
334 A systematic review. *Front Neuroendocrinol.* 2020 Jan; 56(100818):100818.
- 335 **Harris CR**, Millman KJ, van der Walt SJ, Gommers R, Virtanen P, Cournapeau D, Wieser E, Taylor J, Berg S, Smith NJ, Kern R, Picus M,  
336 Hoyer S, van Kerkwijk MH, Brett M, Haldane A, Del Río JF, Wiebe M, Peterson P, Gérard-Marchant P, et al. Array programming  
337 with NumPy. *Nature.* 2020 Sep; 585(7825):357–362.
- 338 **Hunter JD**. Matplotlib: A 2D graphics environment. *Computing in Science & Engineering.* 2007; 9(3):90–95.
- 339 **Iqbal J**, Huang GD, Xue YX, Yang M, Jia XJ. Role of estrogen in sex differences in memory, emotion and neuropsychiatric disorders.  
340 *Mol Biol Rep.* 2024 Mar; 51(1):415.
- 341 **Jilg A**, Bechstein P, Saade A, Dick M, Li TX, Tosini G, Rami A, Zemmar A, Stehle JH. Melatonin modulates daytime-dependent  
342 synaptic plasticity and learning efficiency. *J Pineal Res.* 2019 Apr; 66(3):e12553.
- 343 **Kimmel CB**, Patterson J, Kimmel RO. The development and behavioral characteristics of the startle response in the zebra fish.  
344 *Dev Psychobiol.* 1974 Jan; 7(1):47–60.
- 345 **Koitmäe A**, Karsten Y, Li X, Morellini F, Rune GM, Bender RA. GPER1 deficiency causes sex-specific dysregulation of hippocampal  
346 plasticity and cognitive function. *J Endocrinol.* 2023 Sep; 258(3).
- 347 **Kow LM**, Pfaff DW. Rapid estrogen actions on ion channels: A survey in search for mechanisms. *Steroids.* 2016 Jul; 111:46–53.
- 348 **Lam SK**, Pitrou A, Seibert S. Numba: A LLVM-based python jit compiler. In: *Proceedings of the Second Workshop on the LLVM  
349 Compiler Infrastructure in HPC;* 2015. p. 1–6.
- 350 **Lamiré LA**, Haesemeyer M, Engert F, Granato M, Randlett O. Functional and pharmacological analyses of visual habituation  
351 learning in larval zebrafish. *Elife.* 2023 Dec; 12.
- 352 **Luine VN**. Estradiol and cognitive function: past, present and future. *Horm Behav.* 2014 Sep; 66(4):602–618.
- 353 **McDiarmid TA**, Yu AJ, Rankin CH. Habituation is more than learning to ignore: Multiple mechanisms serve to facilitate shifts in  
354 behavioral strategy. *Bioessays.* 2019 Sep; 41(9):e1900077.
- 355 **Wes McKinney**. Data Structures for Statistical Computing in Python. In: Stéfan van der Walt, Jarrod Millman, editors. *Proceedings  
356 of the 9th Python in Science Conference;* 2010. p. 56 – 61. doi: 10.25080/Majora-92bf1922-00a.
- 357 **Menuet A**, Pellegrini E, Anglade I, Blaise O, Laudet V, Kah O, Pakdel F. Molecular characterization of three estrogen receptor forms  
358 in zebrafish: Binding characteristics, transactivation properties, and tissue Distributions1. *Biol Reprod.* 2002 Jun; 66(6):1881–  
359 1892.
- 360 **Mlinarić A**, Horvat M, Šupak Smolčić V. Dealing with the positive publication bias: Why you should really publish your negative  
361 results. *Biochem Med (Zagreb).* 2017 Oct; 27(3):030201.

- 362 Naderi M, Salahinejad A, Attaran A, Niyogi S, Chivers DP. Rapid effects of estradiol and its receptor agonists on object recognition  
363 and object placement in adult male zebrafish. *Behav Brain Res.* 2020 Apr; 384(112514):112514.
- 364 Nilsson S, Gustafsson JA. Biological role of estrogen and estrogen receptors. *Crit Rev Biochem Mol Biol.* 2002; 37(1):1–28.
- 365 Payrits M, Sághy É, Csekő K, Pohóczky K, Bölcskei K, Ernszt D, Barabás K, Szolcsányi J, Ábrahám IM, Helyes Z, Szőke É. Estradiol  
366 sensitizes the transient receptor potential vanilloid 1 receptor in pain responses. *Endocrinology.* 2017 Oct; 158(10):3249–3258.
- 367 Prossnitz ER, Barton M. The G protein-coupled oestrogen receptor GPER in health and disease: an update. *Nat Rev Endocrinol.*  
368 2023 Jul; 19(7):407–424.
- 369 Qiu J, Bosch MA, Tobias SC, Krust A, Graham SM, Murphy SJ, Korach KS, Chambon P, Scanlan TS, Rønnekleiv OK, Kelly MJ.  
370 A G-protein-coupled estrogen receptor is involved in hypothalamic control of energy homeostasis. *J Neurosci.* 2006 May;  
371 26(21):5649–5655.
- 372 Ramírez-Barrantes R, Carvajal-Zamorano K, Rodriguez B, Cordova C, Lozano C, Simon F, Díaz P, Muñoz P, Marchant I, Latorre  
373 R, Castillo K, Olivero P. TRPV1-estradiol stereospecific relationship underlies cell survival in oxidative cell death. *Front Physiol.*  
374 2020 May; 11:444.
- 375 Randlett O, Haesemeyer M, Forkin G, Shoenhard H, Schier AF, Engert F, Granato M. Distributed plasticity drives visual habituation  
376 learning in larval zebrafish. *Curr Biol.* 2019 Apr; 29(8):1337–1345.e4.
- 377 Rankin CH, Abrams T, Barry RJ, Bhatnagar S, Clayton DF, Colombo J, Coppola G, Geyer MA, Glanzman DL, Marsland S,  
378 McSweeney FK, Wilson DA, Wu CF, Thompson RF. Habituation revisited: an updated and revised description of the behavioral  
379 characteristics of habituation. *Neurobiol Learn Mem.* 2009 Sep; 92(2):135–138.
- 380 Rawashdeh O, de Borsetti NH, Roman G, Cahill GM. Melatonin suppresses nighttime memory formation in zebrafish. *Science.*  
381 2007 Nov; 318(5853):1144–1146.
- 382 Revankar CM, Cimino DF, Sklar LA, Arterburn JB, Prossnitz ER. A transmembrane intracellular estrogen receptor mediates rapid  
383 cell signaling. *Science.* 2005 Mar; 307(5715):1625–1630.
- 384 Romano SN, Edwards HE, Souder JP, Ryan KJ, Cui X, Gorelick DA. G protein-coupled estrogen receptor regulates embryonic heart  
385 rate in zebrafish. *PLoS Genet.* 2017 Oct; 13(10):e1007069.
- 386 Sula A, Hollingworth D, Ng LCT, Larmore M, DeCaen PG, Wallace BA. A tamoxifen receptor within a voltage-gated sodium channel.  
387 *Mol Cell.* 2021 Mar; 81(6):1160–1169.e5.
- 388 Thisse C, Thisse B. High-resolution *in situ* hybridization to whole-mount zebrafish embryos. *Nat Protoc.* 2008; 3(1):59–69.
- 389 Treviño LS, Gorelick DA. The interface of nuclear and membrane steroid signaling. *Endocrinology.* 2021 Aug; 162(8).
- 390 Vail G, Roepke TA. Membrane-initiated estrogen signaling via Gq-coupled GPCR in the central nervous system. *Steroids.* 2019  
391 Feb; 142:77–83.
- 392 Virtanen P, Gommers R, Oliphant TE, Haberland M, Reddy T, Cournapeau D, Burovski E, Peterson P, Weckesser W, Bright J,  
393 van der Walt SJ, Brett M, Wilson J, Millman KJ, Mayorov N, Nelson ARJ, Jones E, Kern R, Larson E, Carey CJ, et al. SciPy 1.0:  
394 fundamental algorithms for scientific computing in Python. *Nat Methods.* 2020 Mar; 17(3):261–272.
- 395 Van der Walt S, Schönberger JL, Nunez-Iglesias J, Boulogne F, Warner JD, Yager N, Gouillart E, Yu T. scikit-image: image processing  
396 in Python. *PeerJ.* 2014; 2:e453.
- 397 Waskom M. seaborn: statistical data visualization. *J Open Source Softw.* 2021 Apr; 6(60):3021.
- 398 Wolman MA, Jain RA, Liss L, Granato M. Chemical modulation of memory formation in larval zebrafish. *Proceedings of the  
399 National Academy of Sciences.* 2011 Sep; 108(37):15468–15473.
- 400 Yu X, Zhao QY, Yaman M, Emly SM, Lee JK, Su H, Ferguson AC, Nagaswami C, Chaturantabut S, Goessling W, Weisel JW, Auchus  
401 RJ, Shavit JA. Hormone-induced thrombosis is mediated through non-canonical fibrin(ogen) aggregation and a novel estrogen  
402 target in zebrafish. *bioRxiv.* 2024 Nov; .