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1 Variations of salinity during reproduction and development affect

- 2 ontogenetic trajectories in a coastal amphibian
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30 Abstract

Although coastal ecosystems are naturally submitted to temporal variations of salinity, 31 salinization has been increasing over time threatening coastal biodiversity. Species that 32 exploit such habitats can thus be exposed to brackish water at different life stages. 33 However, the impacts of variations of salinity on wildlife remain poorly understood. 34 This is particularly true for coastal amphibians, due to the strong dependency of early 35 life stages (embryos and larvae) on aquatic environments. In order to investigate the 36 effect of salinity during egg laying and embryonic and larval development of coastal 37 amphibians, we used a full-factorial design to expose reproductive adults, eggs and 38 39 larvae of coastal spined toads (*Bufo spinosus*) to fresh (0 g.l⁻¹) or brackish water (4 g.l⁻¹). At egg laying, we evaluated parental investment in reproduction. During embryonic 40 and larval development, we assessed effects on survival, development, and growth. 41 We highlighted strong effects of environmental salinity on reproduction (reduced egg 42 laying time, marginally reduced egg size, and reduced investment in reproduction). 43 Responses to salinity were highly dependent on the developmental stages of exposure 44 (stronger effects when individuals were exposed during embryonic development). 45 These effects carried over when exposure occurred at egg laying or during embryonic 46 development, highlighting the importance of the environmental conditions during 47 48 early life on ontogenetic trajectories. We also highlighted partial compensation when individuals were transferred back to freshwater. Whether the magnitude of these 49 responses can allow coastal biodiversity to overcome the observed detrimental effects 50 of salinization remain to be assessed. 51

- 52 Key-words: Carry-over effects, Embryonic development, Freshwater, Recovery,
- 53 Reproductive investment, Tadpoles

54 **1. INTRODUCTION**

Salinization of freshwater bodies is strongly increasing over time (Williams 2001). 55 Coastal wetlands are particularly subjected to salinization because of salt water 56 intrusions (Knighton et al. 1991; Visschers et al. 2022), reductions in freshwater flow 57 resulting from dams and reservoirs, droughts and increased abstraction (Reid et al. 58 59 2019), rising sea levels (Vermeer and Rahmstorf 2009; Church and White 2011; Bakker et al. 2017), and the predicted increase in frequency and intensity of marine surges 60 (Nicholls et al. 1999; McLean et al. 2001; Knutson et al. 2010; Dettinger 2011; Trenberth 61 62 et al. 2015; IPCC 2022). Due to their particular position between terrestrial and oceanic 63 systems, coastal ecosystems are also naturally submitted to variations of salinity (Xue 64 et al. 2013), and are thus known to experience temporal and spatial heterogeneity of salinity (Estévez et al. 2019; Ranjbar and Ehteshami 2019; Fu et al. 2021). Importantly, 65 salinity can increase through time, but can also decrease when precipitations increase 66 (Lorrain-Soligon et al. 2023c). However, the impacts of these short term variations of 67 salinity on wildlife remain poorly understood. 68

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Salinization is expected to strongly affect coastal biodiversity (Amores et al. 2013; Debue et al. 2022). Indeed, most organisms need to maintain their internal homeostasis and rely on physiological compensatory mechanisms to regulate water and ion fluxes (Schultz and McCormick 2012; Evans and Kültz 2020). Most species have specific ranges of tolerance, and crossing these limits of tolerance activates mechanisms that aim at regulating ionic and hydric fluxes (Evans 2009). These mechanisms include modulation of drinking and urination rates, regulation of the permeability of

membranes to water, remodeling of cells and tissues, and increased expression and 77 activity of ion transporters (Evans 2009; Kültz 2015; Rivera-Ingraham and Lignot 2017). 78 However, these mechanisms are often metabolically costly and the associated 79 energetic expenditures can affect energetic allocation to other functions (Herbert et al. 80 2015; Rivera-Ingraham and Lignot 2017). As such, salinity can influence critical 81 functions such as foraging (Chuang et al. 2022), activity (Leite et al. 2022; 82 Moniruzzaman et al. 2022), anti-predator responses (Hoover et al. 2013) and growth 83 (Liu et al. 2022). In addition, elevated salinity can also increase the occurrence of 84 malformations (Hieu et al. 2021), potentially leading to increased mortality (Cañedo-85 Argüelles et al. 2013; Moniruzzaman et al. 2022; Woodley et al. 2023). In adults, 86 increased salinity can reduce egg production and fecundity (Pinder et al. 2005; 87 Froneman 2023; Woodley et al. 2023), egg size (Lorrain-Soligon et al. 2023a), or the age 88 at first reproduction (Woodley et al. 2023). In addition, increasing salinity is known to 89 reduce sperm performance, motility and velocity (Wilder and Welch 2014; Byrne et al. 90 2015, 2022; Green et al. 2021). Ultimately, salinization can alter community structure 91 (Hart et al. 2003; Anufriieva and Shadrin 2018), across all trophic levels (Hintz and 92 Relyea 2019). 93

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This is especially true for coastal amphibians. Indeed, amphibians are particularly sensitive to salinity due to their highly permeable skin used for gas exchanges, and for ions and water transport (Shoemaker and Nagy 1984; Martin and Nagy 1997), and to their relatively low osmoregulatory abilities (Katz 1989). Early life stages (i.e. eggs and larvae) are particularly susceptible to environmental salinity (Albecker and McCoy 100 2017) because of their strong dependency on aquatic environment compared to adults (Wells 2007) and to their comparatively lower osmoregulatory abilities (Karraker and 101 102 Gibbs 2011). Increasing salinity from various sources can lead to reduced thyroid hormones levels (Gomez-Mestre et al. 2004) and to changes in osmoregulatory 103 hormones such as corticosterone and aldosterone (Tornabene et al. 2021b, 2022) in 104 105 larval amphibians. Ultimately, this lead to decreased embryonic and larval survival (Christy and Dickman 2002; Chinathamby et al. 2006; Dougherty and Smith 2006; 106 Sanzo and Hecnar 2006; Karraker et al. 2008; Bernabò et al. 2013; Jones et al. 2015), and 107 to increased malformations (Gosner and Black 1957; Sanzo and Hecnar 2006; 108 Haramura 2007; Karraker 2007; Hopkins et al. 2013; Brady 2013; Tornabene et al. 109 2021a), time to hatching (Haramura 2016; Tornabene et al. 2021a), hatching success 110 (Tornabene et al. 2021a), larval duration (Christy and Dickman 2002; Gomez-Mestre 111 and Tejedo 2003; Lukens and Wilcoxen 2020; Tornabene et al. 2021a), and to reduced 112 growth and thus to induce smaller body size (Wu and Kam 2009; Bernabò et al. 2013; 113 Wood and Welch 2015; Haramura 2016; Lukens and Wilcoxen 2020; Tornabene et al. 114 2021a, b). 115

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Despite their sensitivity, amphibians are known to be very plastic in behavioral and morphological traits (Hoverman and Relyea 2008), and their developmental traits can vary in response to environmental cues (Chivers et al. 2001; Warkentin 2011). For instance, tadpoles can modulate their growth rates according to the quality of environmental conditions (Werner 1986; Vonesh and Warkentin 2006). Additionally, it has been suggested that the influence of a given environmental factor on

development and growth of tadpoles is highly dependent on the stage and condition 123 during which they experience such factor (Denver et al. 2002). This can be highly 124 125 relevant for changes in salinity. For example, Fejervarya limnocharis and Litoria ewingii tadpoles released from salinity stress at early larval stages reached a size at 126 metamorphosis similar to that of tadpoles maintained in low salinity throughout 127 development, indicating compensatory growth (Squires et al. 2010; Wu et al. 2012). 128 Yet, the efficiency of such compensatory mechanisms may vary if individuals are 129 exposed earlier or later during larval development (Hsu et al. 2018). Additionally, 130 whether parents can influence the tolerance of their offspring according to the salinity 131 in which they are laying eggs is unknown. 132

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In order to investigate the effect of salinity during egg-laying and embryonic and larval 134 development of coastal amphibians, we used a full-factorial experimental design to 135 expose reproductive adults, eggs and larvae of coastal spined toads (Bufo spinosus, a 136 species which coastal individuals are supposedly adapted to exposure to salinity) to 137 fresh (0 g.l⁻¹) or brackish water (4 g.l⁻¹, Figure 1). More specifically, reproductive pairs 138 (amplexus) were exposed to fresh or brackish water prior to egg-laying (Figure 1). 139 Following egg laying, we exposed eggs to either fresh or brackish water until hatching 140 (Figure 1). Following hatching, we exposed larvae to either fresh or brackish water 141 (Figure 1). At egg laying, we evaluated fecundity and parental investment in 142 reproduction. During both embryonic and larval development, we assessed effects of 143 each experimental treatment on survival (hatching rate, larval survival), development 144 (duration of embryonic and larval development until Gosner stage 30 [Gosner 1960], 145

malformation), and growth (body length). We predicted that (1) Brackish water would 146 impact reproductive investment and egg size, and alter fecundity because salinity can 147 impact sperm motility (Byrne et al. 2015, 2022). (2) Salt exposure would decrease 148 embryonic survival, larval survival and growth (Albecker and McCoy 2017; Lukens 149 and Wilcoxen 2020; Tornabene et al. 2021a) as well as malformations (Gosner and Black 150 1957; Sanzo and Hecnar 2006; Haramura 2007; Karraker 2007; Hopkins et al. 2013; 151 Brady 2013; Tornabene et al. 2021a), regardless of the stage of exposure. These effects 152 should be alleviated in individuals transferred to freshwater, because of the resilience 153 capacities of amphibians to salt exposure (Squires et al. 2010; Wu et al. 2012; Hsu et al. 154 2018; Lorrain-Soligon et al. 2022). 155

156 **2. METHODS**

157 2.1. Study species

Spined toads (*Bufo spinosus*) are a common amphibian species found in western Europe (Speybroeck et al. 2018; Meek 2022). Breeding occurs in late winter, during which adults converge to reproductive ponds where they mate (amplexus) and lay their eggs constituted of 2 eggs strings which can be constituted of up to 10000 eggs (Miaud and Muratet 2018). Egg laying, embryonic and larval developments all occur in ponds.

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164 2.2. Captures and measurements

Amplectant pairs were captured on one coastal pond (salinity 3.1 g.l⁻¹ during captures, conductimeter YSI Professional Plus) situated in the Réserve Naturelle Nationale du Marais d'Yves (location: 46.04013, -1.05363). This site can display seasonal variations of salinity ranging from 0.5 g.l⁻¹ to 6 g.l⁻¹, as what is found in nearby ponds. Captures occurred at night (between 20 pm and 4 am) on 21/02/2023 and 22/02/2023. Twenty one amplectant pairs were captured by hand, placed in a transport box (14*16*9cm), and brought to the laboratory immediately after field sessions.

At the laboratory, individuals were kept in a thermally controlled room with temperature set at 17 °C and controlled photoperiod (12 h dark-12 h light), temperature and photoperiod being set for the totality of the experiment. Pairs were transitorily (for measurements only) separated and all individuals were measured (snout-vent length, SVL) using a caliper (\pm 1 mm) and weighed using an electronic balance (\pm 0.1g). Once measurements were completed, separated amplectant individuals were shortly reunited and pairs re-formed systematically as soon as thepartners were brought back into contact.

180

181 **2.3. Experimental procedures**

Amplectant pairs, embryos and larvae were exposed to either freshwater or brackish 182 183 water (sea salt, NaCl), following a full factorial design (Figure 1). At the end of the experiment, larval individuals were thus exposed to one of the eight following 184 185 treatments: (1) FFF: clutch obtained in freshwater, embryonic development in 186 freshwater, larval development in freshwater; (2) FSF: clutch obtained in freshwater, embryonic development in brackish water, larval development in freshwater; (3) FFS: 187 clutch obtained in freshwater, embryonic development in freshwater, larval 188 development in brackish water; (4) FSS: clutch obtained in freshwater, embryonic 189 development in brackish water, larval development in brackish water; (5) SFF: clutch 190 191 obtained in brackish water, embryonic development in freshwater, larval development in freshwater; (6) SSF: clutch obtained in brackish water, embryonic 192 development in brackish water, larval development in freshwater; (7) SFS: clutch 193 194 obtained in brackish water, embryonic development in freshwater, larval development in brackish water; (8) SSS: clutch obtained in brackish water, embryonic 195 development in brackish water, larval development in brackish water. Water of these 196 197 treatments was changed every seven days.

Tadpoles were fed ad libitum with thawed frozen organic spinach. Individuals were checked twice a day for their specific developmental stage, for mortality and for abnormalities (which were checked visually).

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203 2.3.1. Egg laying (n=21)

Once the amplectant pairs were reunited, they were placed in a plastic container 204 (35*55*26 cm) containing either freshwater (~20L, salinity: 0.3 g.l-1, 10 amplectant pairs) 205 206 or brackish water (~20L, salinity: 4.0 g.l-1, 11 amplectant pairs) as well as branches for laying support. The two groups did not differ in size (freshwater: females= 74.7 mm ± 207 208 1.61 SE, males= 64.1 mm ± 1.00 SE; brackish water: females= 75.9 mm ± 2.08 SE, males= 62.7 mm ± 1.22 SE; differences between females in the two groups: Estimate=1.209, 209 SE=2.604, t_{1.19}=0.464, p-value=0.648; differences between males in the two groups: 210 Estimate=1.373, SE=1.601, t_{1.19}=0.857, p-value=0.402). Amplectant pairs were left in 211 212 these tanks until egg laying. Individuals were checked every two hours to compute time to egg laying (measured from their placement in a plastic container to the 213 completion of egg laying). Adult individuals were then released at their site of capture. 214 Adults were thus kept under laboratory conditions for a maximum of seven days 215 216 (maximal time to egg laying 151 hours), and the water of their treatment was left unchanged during this duration. They were unfed during this period, as adults 217 anurans usually don't fed during the reproduction. Twelve pieces of each egg strings, 218 219 containing 30 eggs each (i.e. a total of 360 eggs for each egg string) were selected 220 randomly and kept for our experiment, and the remaining eggs (i.e. 100-4500) were 221 released at the site of origin of their parents.

223 2.3.2. Embryonic development (n=232)

The two treatments were applied in 2 L individual tanks (13*18*18cm, n=232, see table 224 1) containing one of the two treatment (freshwater: 0 g.l⁻¹ salinity: 0.3 ± 0.0 g.l⁻¹; 225 226 brackish water : 4 g.l⁻¹ salinity 4.0 ± 0.06 g.l⁻¹). Water was changed once a week. Each 227 clutch was represented by 6 segments of 30 eggs in each of the experimental treatments. To prevent osmotic shock, segments for which the clutch was obtained in 228 229 another salinity than which their embryonic development occur (i.e. clutch that were obtained at 0 g.l-1 but for which embryonic development will occur at 4 g.l-1 and clutch 230 that were obtained at 4 g.l-1 but for which embryonic development will occur at 0 g.l-231 232 ¹) were gradually exposed to the salinity of the treatment, with an increase or a decrease of 1 g.l-1 a day (Hsu et al. 2018). This way, segments were exposed to their 233 final treatment over 4 days. Each segment were kept in the laboratory until tadpoles 234 235 hatched and reached a free feeding stage (Gosner stage 25, hereafter GS25) (Gomez-Mestre and Tejedo 2003). 236

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238 2.3.3. *Larval development (n=232)*

Upon hatching (GS25), we selected 1 individual per segment (randomly, among alive and not malformed tadpoles of each segment), and these individuals were once again distributed in two treatments: freshwater (0 g.l⁻¹ salinity: 0.3 ± 0.0 g.l⁻¹), and brackish water (4 g.l⁻¹ salinity 4.0 ± 0.06 g.l⁻¹). For each clutch, three segments and thus three tadpoles were thus exposed to each treatment (FFF, FFS, FSF, FSS, SFF, SFS, SSF, SSS). In some segments, no individuals survived until GS25. In these cases, individuals were replaced by other individuals from the same treatments but originating from different clutches in order to keep a total of 232 tadpoles that were followed during larval development (see table 1 for the count of tadpoles in each treatment, and Appendix A for the contribution of each clutch). All other tadpoles were released at the site where their parents were captured.

Similarly to embryos, when individuals were transferred to a different treatment (i.e. tadpoles for which embryonic development occur at 0 g.l⁻¹ but larval development will occur at 4 g.l⁻¹ and tadpoles for which embryonic development occur at 4 g.l⁻¹ but larval development will occur at 0 g.l⁻¹), tadpoles were gradually exposed to the salinity of the treatment, with an increase or a decrease of 1 g.l⁻¹ a day to prevent osmotic shock (Hsu et al. 2018).

Larval development was monitored until G30 (Gosner 1960), in order to have approximatively the same time considered for embryonic development and larval development, and because GS30 is a pivotal stage when somatic growth decreases and significant morphological changes occur Cheron et al. (2021). At the end of this experimental procedure, individuals were released at the site where their parents were captured.

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263 2.4. Measurements

264 2.4.1. *Reproductive effort (n=21)*

265 Once egg laying was completed, individuals were again weighed to calculate 266 variations in body mass. Additionally, each clutch was weighed using an electronic 267 balance (± 0.1g). Moreover, in order to assess fecundity, each egg string was placed in

a container $(35 \times 20 \times 25 \text{ cm})$ containing 2 cm of dechlorinated tap water and a scale 268 (graph paper). A picture was taken from above in order to measure the total length of 269 270 the egg string using ImageJ software (Schneider et al. 2012). For each clutch, we randomly selected six 10-cm segments and individually counted the number of eggs 271 within each segment. The mean number of eggs per 10-cm segment was calculated and 272 used to assess fecundity (number of eggs) for each clutch based on the length of the 273 egg strings. Finally, on a subsample of 100 randomly selected eggs in each egg strings, 274 we measured egg diameter. 275

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277 2.4.2. Embryonic development (n=232)

At GS 25, hatching success (proportion of eggs that hatched, on the segment of 30 eggs) and deformation rates (proportion of larvae that hatched but were malformed, on the total number of hatched individuals) were calculated. Additionally, for each segments, we described general activity of all hatchlings as being, on a scale of 1 to 5, the proportion of live individuals that were active; 1=0-20%, 2=20-40%, 3=40-60%, 4=60-80%, 5=80-100%) over a 1-min period.

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285 2.4.3. *Larval development (n=232)*

At GS25 and GS30, we measured total length of individuals. Each tadpole was put into a Petri dish with the water from its own tank, and photographed. Individuals were than measured using the software ImageJ (Schneider et al. 2012). We also measured activity following a scale ranging from 1 to 5 (from 1 if the individual is amorphous, to 5 if the individual move energetically, following Lorrain-Soligon et al. [2022]). The
tadpoles were not stimulated to record these observations.

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293

294 2.5. Statistical analyses

295 We computed Linear Models (LMs) or Linear Mixed Models (LMMs) and Generalized Linear Mixed Models (GLMMs) using the lme4 (Bates et al. 2015) and lmerTest 296 297 (Kunzetsova et al. 2017) packages. For all test computed, models accuracy was tested using the check_model function from the performance package (Lüdecke et al. 2020). 298 When models did not fit the data, the response variable was log+1 transformed. 299 300 Binomials models accuracy was also tested by checking overdispersion. Significance 301 level was set at 0.05, differences at a p-value level <0.1 were considered marginal. For 302 all comparisons studied, mean \pm SE of each group are given in Appendix B.

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304 2.5.1. Reproductive effort

We evaluated the effect of salinity (clutches laid in freshwater [0 g.l-1] or in brackish 305 water [4 g.l-1]) on clutch length, clutch size (number of eggs), clutch mass, egg 306 307 diameter, laying time, variation of body mass (for females and males). For these analyses, we set linear models (LMs), with salinity as an explanatory variable. We also 308 tested for the relationships between egg diameter and clutch size in each salinity by 309 setting a LM with egg diameter as a dependent variable, and clutch size, salinity and 310 their interaction as explanatory variables. Finally, we tested for the effect of the 311 312 interaction between female mass and salinity on clutch size and clutch mass, using LMs. For these models, best variables were retained using a top-down selection (usingp-values), and only the retained variables are presented.

315

316 2.5.2. Embryonic development

We evaluated the effect of salinity during spawning (clutch laid in freshwater [0 g.l⁻¹] or in brackish water [4 g.l⁻¹]), and during embryonic development (individuals developing in freshwater [0 g.l⁻¹] or brackish water [4 g.l⁻¹]).

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We computed these effects on time to hatching (log+1 transformed) and individuals 321 322 total size with Linear Mixed models (LMMs), and on hatching success, malformation rate, general activity and tadpole activity (both rated on a scale from 1 to 5)using 323 binomial Generalized Linear Mixed Models (GLMMS). For all these models, we 324 included salinity during spawning, salinity during embryonic development and their 325 interaction as explanatory variables, and the clutch as a random effect (as multiple 326 individuals originated from the same clutch). Best variables were retained using a top-327 down selection (using p-values), and only the retained variables are presented. 328

329

330 2.5.3. Larval development

We evaluated the effect of salinity at different stages of exposure. We computed the effect of salinity during spawning (clutches laid in freshwater [0 g.l⁻¹] or in brackish water [4 g.l⁻¹]), salinity during embryonic development (eggs developing in freshwater [0 g.l⁻¹] or brackish water [4 g.l⁻¹]), and salinity during larval development (individuals

developing in freshwater [0 g.l⁻¹] or brackish water [4 g.l⁻¹]). We computed these effects 335 on development duration between S25 and S30, growth rate between GS25 and GS30, 336 and total size (log+1 transformed) at GS30 with LMMs, and on mortality and activity 337 (rated on a scale from 1 to 5) using binomial GLMMs. For all these models, we 338 included salinity during spawning, salinity during embryonic development, salinity 339 during larval development and their interaction as explanatory variables, and the 340 clutch as a random effect (as multiple individuals originated from the same clutch). 341 Best variables were retained using a top-down selection (using p-values), and only the 342 retained variables are presented. At GS30, only 2 individuals were malformed (in FFS 343 and FSF) and thus statistics were not computed. All data analysis were performed 344 345 using R 3.6.3 (R Core Team 2020) and Rstudio v1.1.419.

346 **3. RESULTS**

347 3.1. Reproductive effort

Clutches produced in freshwater or in brackish water were similar for length 348 (Estimate=70.05, SE= 263.58, t_{1.19}=0.27, p-value=0.793) and number of eggs 349 (Estimate=182.60, SE= 405.40, t_{1.19}=0.45, p-value=0.658). However, clutches produced 350 in freshwater were heavier (Estimate=28.21, SE= 13.07, $t_{1.19}$ =2.16, p-value=0.044), and 351 352 their eggs were marginally larger (Estimate=0.11, SE= 0.06, $t_{1.19}$ =1.80, p-value=0.088, Figure 2A). Laying time was shorter for individuals that laid eggs in brackish water 353 compared to those that laid eggs in freshwater (Estimate=-31.77, SE= 14.84, $t_{1.19}$ =-2.14, 354 p-value=0.046, Figure 2B). 355

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Egg diameter was related to the interaction between clutch size and salinity (Estimate<0.001, SE<0.001, $t_{3.17}$ =2.65, p-value=0.017, Figure 2C). Egg diameter was not influenced by clutch size in clutches laid in freshwater (Estimate<0.001, SE<0.001, $t_{1.8}$ =1.21, p-value=0.260, Figure 2C), but increased with increasing clutch size in clutches laid in brackish water (Estimate<0.001, SE<0.001, $t_{1.9}$ =2.40, p-value=0.040, Figure 2C).

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Interestingly, females spawning at 0 g.l⁻¹ (-13.56 % ± 2.23 SE) and 4 g.l⁻¹ (-12.21 % ± 1.36 SE) lost similar mass during egg laying (Estimate=1.35, SE=2.56, $t_{1.19}$ =0.53, pvalue=0.604). Males reproducing at 0 g.l⁻¹ lost mass (-1.63 % ± 2.19 SE) while males reproducing at 4 g.l⁻¹ gained mass (3.79% ± 1.27 SE) during egg laying (Estimate=5.41, SE=2.47, $t_{1.19}$ =2.19, p-value=0.041).

Clutch size increased with female mass (Estimate=40.876, SE=10.303, $t_{1.19}$ =3.967, pvalue<0.001). When testing for the interaction between females body mass and salinity, clutch mass increased with female body mass (Estimate=1.211, SE=0.362, $t_{1.19}$ =3.347, p-value=0.004) and was higher in the freshwater treatment compared to the brackish one (Estimate=25.192, SE=10.579, $t_{1.19}$ =2.831, p-value=0.029).

375

376 3.2. Embryonic development

Hatching success was higher for clutches produced in freshwater (Estimate=0.480, SE=0.194, z=2.649, p-value=0.014), and higher in embryos that developed in freshwater (Estimate=0.972, SE=0.064, z=15.103, p-value<0.001). There was an interaction between the salinity during spawning and the salinity during embryonic development (Estimate=-1.493, SE=0.141, z=-10.580, p-value<0.001, Figure 3A, Appendix C), hatching success being higher for individuals originating from a clutch laid in freshwater, and that developed in freshwater.

Malformation rate was higher in embryos which developed in brackish water (Estimate=0.893, SE=0.114, z=7.840, p-value<0.001), but not related to salinity during spawning (Estimate=0.086, SE=0.242, z=0.357, p-value=0.721). There was an interaction between the salinity during spawning and the salinity during embryonic development (Estimate=-1.059, SE=0.301, z=-3.524, p-value<0.001, Figure 3B, Appendix C), malformation rate being higher for individuals originating from a clutch laid in freshwater, and that developed in brackish water.

Time to hatching was higher in embryos which developed in brackish water (Estimate=0.107, SE=0.013, $t_{1.221}$ =8.387, p-value<0.001), but not related to spawning in brackish water (Estimate=0.004, SE=0.025, t_{1.27}=0.168, p-value=0.868). There was an
interaction between the salinity during spawning and the salinity during embryonic
development (longer time to hatching for individuals that were laid in freshwater but
developed in brackish water, Estimate=-0.065, SE=0.021, t_{1.218}=-3.104, p-value=0.002,
Appendix C).

General activity (Estimate=1.164, SE=0.173, z=6.735, p-value<0.001), tadpole activity
(Estimate=1.141, SE=0.117, z=9.706, p-value<0.001) and total length (Estimate=-2.157,
SE=0.0.139, t_{1.228}=-15.540, p-value<0.001, Figure 3C) were all influenced by salinity
experienced during embryonic development solely, and were higher for embryos that
developed in freshwater.

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404

405 3.3. Larval development

Mortality at GS30 was influenced by salinity to which individuals were exposed
during embryonic development (higher for individuals which embryonic
development occurred in brackish water [FSF, FSS, SSF and SSS], Estimate=3.405,
SE=0.005, z=676.3, p-value<0.001, Figure 4A) and during larval development (higher
for larvae developing in brackish water [FFS, FSS, SFS and SSS], Estimate=1.421,
SE=0.005, z=282.5, p-value<0.001, Figure 4A), but not their interaction.

Development duration between S25 and S30 varied with salinity during embryonic
development (longer for individuals which embryonic development occurred in
brackish water [particularly FSS, SSF and SSS], Estimate=1.409, SE=0.635, t_{1.101}=2.218,

p-value=0.029, Figure 4B), and there was an interaction between salinity during
embryonic development and salinity during larval development (Estimate=3.307,
SE=1.095, t_{1.91}=3.020, p-value=0.003, Figure 4B, Appendix D), as well as between
salinity during spawning and salinity during larval development (Estimate=-2.914,
SE=0.768, t_{1.94}=-3.793, p-value<0.001, Figure 4B, Appendix D).

Growth rate between GS25 and GS30 was only influenced by salinity during embryonic development, and was higher for individuals which embryonic development occurred in brackish water (particularly FSF for individuals laid in freshwater, and SSF and SSS for individuals laid in brackishwater, Estimate=0.081, SE=0.036, t_{1.110}=2.240, p-value=0.027, Figure 4C).

Total length was influenced by salinity experienced during embryonic development solely, decreasing for individuals which embryonic development occurred in brackish water (Estimate=-0.033, SE=0.012, t_{1.110}=-2.653, p-value=0.009, Figure 4D). Conversely, activity (Estimate=-0.350, SE=0.147, z=-2.387, p-value=0.017) was influenced by salinity during larval development solely, decreasing for larvae that developed in brackish water (particularly FFS, FFS and SFS).

431 **4. DISCUSSION**

Our study demonstrated that environmental salinity can negatively affect reproduction, as well as embryonic and larval development in a coastal anuran. Importantly, our full-factorial design allowed us to highlight that early exposure to brackish water did not induce increased tolerance during later exposure (i.e., tadpoles spawned in brackish water did not perform better in brackish water). However, individuals returned to freshwater after being exposed to brackish water expressed partial recovery, indicating resilience to salinity stress.

439

440 4.1. *Reproduction*

441 Clutch size did not differ between freshwater and brackish water treatment, and clutch size increased with female size and female mass loss for both treatments, with no 442 difference between the two treatments. Such results further suggest that clutch size 443 444 might depend on energetic investment that occurred prior to egg laying (Sinervo and 445 DeNardo 1996; Nilsson and Svensson 1997; Lorrain-Soligon et al. 2023a). Interestingly, 446 we found that salinity influenced some reproductive parameters in adults, suggesting 447 a rapid modulation of reproductive effort based on the salinity to which reproductive 448 adults were exposed. Indeed, reproductive adults exposed to brackish water laid their eggs more rapidly than those kept in fresh water. Two opposing hypotheses can 449 explain such result. First, females may lay their eggs more rapidly in brackish water 450 in order to evade an osmotically constraining environment. Conversely, because adult 451 452 individuals originated from salt-exposed coastal environment, they may consider brackish water as a favourable environment if they are locally adapted to such 453

454 conditions and if there are less competitors in this environment (Hopkins and Brodie 2015; Lorrain-Soligon et al. 2023b); but delay laying in freshwater because they 455 456 consider this environment as suboptimal. This later hypothesis seems relatively unlikely as our first hypothesis is further supported by the fact that, in brackish water, 457 eggs were smaller but not fewer. This result suggests that females produced eggs of 458 comparatively lower quality in brackish water, as eggs produced in the freshwater 459 treatment are marginally larger, and larger eggs often have higher hatching success 460 and offspring fitness (Xu et al. 2019; Renoirt et al. 2022). The number of undeveloped 461 eggs (presumably unfertilized given the negative effect of salinity on sperm, (Wilder 462 and Welch 2014; Green et al. 2021)) was also higher in brackish water. 463

464

Interestingly, despite the marginal effect of salinity on egg size, females from both 465 treatments lost a similar amount of mass. In combination with smaller eggs, this result 466 could be linked to a relatively larger investment in egg jellies (which are critical to egg 467 development and protection, Bonnell & Chandler, 1996; Yurewicz et al., 1975) in 468 females exposed to brackish water, a hypothesis that is not supported by clutch mass 469 470 which was higher for females exposed to freshwater. However, investigations focusing on investment in egg protection (egg jellies) in response to environmental salinity 471 remain too scarce. More likely, the fact that females exposed to brackish water have 472 lost a similar amount of mass than their counterpart exposed to freshwater but laid 473 smaller eggs might be linked to direct costs of osmoregulation in adult amphibians 474 exposed to brackish water (Lorrain-Soligon et al. 2022). These variations should be 475

476 investigated through the evaluation of metabolic rates in reproducing adults exposed477 to different salinities.

478

The fact that clutch mass was higher in the freshwater treatment is likely linked to 479 480 water absorption due to osmotic exchanges with the environment (Venturino et al. 481 2003; Karraker and Gibbs 2011; López-Alcaide and Macip-Ríos 2011), or due to higher 482 investment in glycoproteins surrounding the eggs (i.e. egg jellies, see above). More 483 importantly, we found a positive correlation between egg size and clutch size for clutches laid in brackish water but not for clutches laid in freshwater, while fecundity 484 was expected to negatively trade off with egg size (Lasne et al., 2018; Smith & Fretwell, 485 1974). Based on the fractional egg hypothesis (Ricklefs 1968; Nussbaum 1981; Ford and 486 Seigel 2010), the negative relation between fecundity and egg size is expected because 487 species with large clutch sizes and small eggs (typical of Bufonid toads, Wells, 2007) 488 should add additional offspring should extra energy become available (Ricklefs 1968; 489 Nussbaum 1981). In our study, we found that increased clutch size is concomitant 490 with increased egg size in clutches laid in brackish water suggesting that complex 491 492 interactions between prior energetic investment in reproduction and current spawning conditions govern reproductive effort. 493

494

495 Lastly, reproducing males have lost mass in freshwater, but gained mass in brackish
496 water. Such result could be linked to reduced ejaculate volume, with males in
497 freshwater producing more sperm than their counterparts exposed to brackish water.

This hypothesis seems corroborated by the reduced proportion of fertilized eggs in the brackish treatment. However, we did not evaluate sperm volume in each treatment, which might need to be investigated. However, this gain of mass in males can also be linked to the fact that they can be osmotically imbalanced (Hall et al. 2020; Tornabene et al. 2021b).

503

504 Overall, the effects of salinity we found on reproductive adults and clutches clearly affect the investment in reproduction, which suggest that reproductive pond selection 505 in heterogeneous environment may allow to circumvent such environmental 506 constraint (Albecker and McCoy 2017). However, if coastal salinization continues to 507 increase (Herbert et al. 2015; Singh 2021; Cunillera-Montcusí et al. 2022) and 508 individuals are constrained to reproduce in brackish water, this could affect the quality 509 of subsequent offspring development, and thereby negatively influence coastal 510 511 population persistence.

512

513 4.2. Embryonic development

We show that embryonic development duration increased for individuals that were exposed to 4 g.l⁻¹ salinity during embryonic development, but only when they were spawned in freshwater. Increased development time in response to salinity has been described in amphibians (see Christy and Dickman 2002; Sanzo and Hecnar 2006; Wu and Kam 2009; Wijethunga et al. 2016; Lukens and Wilcoxen 2020), an effect which can be due to increasing energy expenditure in osmotically stressful environments. At

hatching, individuals in the 4 g.l-1 treatment were smaller (irrespective of the salinity 520 in which they were spawned) (Wu and Kam 2009; Wood and Welch 2015; Haramura 521 522 2016; Lukens and Wilcoxen 2020; Tornabene et al. 2021a), which is probably linked to higher costs of osmoregulation (Gomez-Mestre et al. 2004). Individuals in the 4 g.l-1 523 524 treatment also presented lower activity. Interestingly, hatching success was always lower for individuals originating from a clutch that was spawned in brackish water 525 irrespective of the salinity in which the remaining embryonic development occurred. 526 Additionally, individuals that were laid in freshwater but developed in brackish water 527 had a higher malformation rate compared to individuals in any other treatments. As 528 individuals developing in the SF and SS groups (that were laid in brackish water) 529 presented a higher mortality rate, this could indicate selective mortality affecting 530 individuals susceptible to salinity, while more tolerant - surviving - individuals 531 displayed lower malformation rates. This hypothesis is further supported by the fact 532 that higher malformation rates were found in individuals spawned in freshwater that 533 developed in brackish water (Gosner and Black 1957; Hopkins et al. 2013). Taken 534 together, these results indicate long lasting effects of the salinity in which the clutch 535 was produced (and thus long-lasting effect of very early exposure to salinity during 536 537 embryonic development). Our results thus emphasize that embryos are sensitive to change in salinity during their development, which can occur through changes of 538 temperatures and precipitations in coastal environments (Lorrain-Soligon et al. 2023c), 539 540 which may carry-over during later larval development and potentially during post-541 metamorphic stages (Traversari 2021; Dahrouge and Rittenhouse 2022), affecting 542 overall amphibian populations.

543

544 4.3. Larval development

Overall, development duration was longer for individuals that were spawned in 545 brackish water, indicating long lasting, carry-over effects of the salinity in which adults 546 547 lay eggs, and accumulating effects during larval development (see Tornabene et al. 2021a). Individuals that experienced salinity during either embryonic or larval 548 549 development (FSF, FFS, SSF and SFS) or during both stages (FSS and SSS) expressed 550 higher mortality (see also Squires et al. 2010; Wu et al. 2012; Albecker and McCoy 2017; Hsu et al. 2018). Additionally, individuals that developed in brackish water during 551 552 embryonic development were smaller (see also Squires et al. 2010), while there was no 553 effects of salinity experienced during larval development on total length. These 554 responses remain consistent regardless of the salinity in which clutches were spawned. 555 This further suggest that salinity experienced during embryonic development is more detrimental than salinity experienced during larval development, which is in line with 556 the idea that early developmental stages (embryos) are more sensitive to salinity than 557 later developmental stages (larvae) (Albecker and McCoy 2017). This is likely due to 558 the development of internal gills (the main organs responsible for ion and water 559 balance in tadpoles, Uchiyama and Yoshizawa 1992) which improve the 560 osmoregulation ability at this developmental stage. However, in the context of 561 accumulating effects, larvae can be found to be more sensitive than embryos 562 (Tornabene et al. 2021a), which highlight that this effect might be context dependent. 563

The detrimental effects of salinity on growth could be linked to changes in activity we found. Indeed, larvae exposed to higher salinity have been shown to express reduced activity and foraging behaviors (Hall et al. 2017; Tornabene et al. 2021a), lower responses to stimuli (Sanzo and Hecnar 2006; Karraker 2007), lower swimming performance (Haramura 2016), erratic movements (Tornabene et al. 2021a), and reduced speed and distance (Denoël et al. 2010).

571

572

Importantly, if individuals exposed to freshwater during larval development after 573 being exposed to brackish water during their embryonic development (FSF and SSF), 574 were smaller compared to those that were exposed to freshwater during most of their 575 development (FFF and SFF), they expressed a higher growth rate. This result suggest 576 577 partial compensatory responses, rather than full compensation or over-compensation (Ali et al. 2003). Compensatory growth may act as an adaptive mechanism because it 578 increases the chance of an organism attaining a critical size at a given developmental 579 stage and (i.e., metamorphosis, Wilbur and Collins 1973), which can ultimately 580 influence fitness (Dmitriew 2011). It is a likely mechanisms in response to osmotic 581 stress in coastal organisms (Hsu et al. 2018), allowing to overcome the negative effects 582 of salinity by expressing catch-up growth when the salt stress ceases. Interestingly, 583 584 some individuals were able to express compensatory growth, but not all, which may 585 indicate differences in genotype or bacterium composition between individuals that 586 may lead to local adaptation (Albecker et al. 2019, 2021; Gabriel et al. 2021).

Importantly, recent research suggests that, although compensatory growth can 587 produce rapid benefits, it is also associated with delayed costs during later stages of 588 589 development and/or adult life (Metcalfe and Monaghan 2001). Such costs include increased risk of predation, decreased resistance to starvation, decreased protein 590 maintenance and increased muscle lesions, reduction in locomotor performance, 591 592 reduced age at sexual maturation and higher telomere attrition rate (Metcalfe and Monaghan 2001). Indeed, accelerated growth is known to induce accumulated cellular 593 damages which ultimately may impair fitness later in life (Mangel and Munch 2005), 594 which could thus affect adults amphibians living in coastal environments 595

596

597 5. CONCLUSION

Reproduction, embryonic and larval development in amphibians are all affected by 598 environmental salinity. Salinity in which individuals were spawned or salinity 599 600 experienced during embryonic development can also carry-over during subsequent 601 developmental stages, highlighting the importance of the environmental conditions during early development on ontogenetic trajectories. Our study focused on a salt-602 exposed - presumably salt tolerant -coastal anuran, highlighting that temporal and 603 spatial variations of salinity might outperform local adaptation, potentially 604 605 threatening coastal populations if salinity happens to increase. Importantly, we also 606 highlighted partial compensation when individuals were transferred to freshwater, 607 which can be a useful response in coastal environments. This partial compensation 608 (which will still depend on the response of each life stage to these fluctuating levels of 609 salinity) might be helpful for individuals in order to express resilience when

freshwater become available, and might help to set-up conservation measures. Indeed, our results might indicate that, following a massive salinization, individuals may be transferred to less impacted adjacent ponds, which might induce survival of the populations given individuals' resilience. This might be applicable by practitioners following a brutal marine submersion, focusing preferentially on embryos, that are shown to be more sensitive. Whether such approach reveals cost-effective in response to ongoing global changes remain to be considered.

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References

620	Albecker MA, Belden LK, McCoy MW (2019) Comparative analysis of anuran
621	amphibian skin microbiomes across inland and coastal wetlands. Microb Ecol
622	78:348–360. https://doi.org/10.1007/s00248-018-1295-9
623	Albecker MA, McCoy MW (2017) Adaptive responses to salinity stress across
624	multiple life stages in anuran amphibians. Front Zool 14:1–16.
625	https://doi.org/10.1186/s12983-017-0222-0
626	Albecker MA, Stuckert AMM, Balakrishnan CN, McCoy MW (2021) Molecular
627	mechanisms of local adaptation for salt-tolerance in a treefrog. Molecular
628	Ecology 30:2065–2086. https://doi.org/10.1111/mec.15867
629 630 631	Ali M, Nicieza A, Wootton RJ (2003) Compensatory growth in fishes: a response to growth depression. Fish and fisheries 4:147–190. https://doi.org/10.1046/j.1467-2979.2003.00120.x
632	Amores MJ, Verones F, Raptis C, et al (2013) Biodiversity impacts from salinity
633	increase in a coastal wetland. Environmental science & technology 47:6384–
634	6392. https://doi.org/10.1021/es3045423
635	Anufriieva E, Shadrin N (2018) Diversity of fauna in Crimean hypersaline water
636	bodies. Journal of Siberian Federal University Biology 11:294–305.
637	https://doi.org/10.17516/1997-1389-0073
638	Bakker AM, Wong TE, Ruckert KL, Keller K (2017) Sea-level projections representing
639	the deeply uncertain contribution of the West Antarctic ice sheet. Scientific
640	Reports 7:1–7. https://doi.org/10.1038/s41598-017-04134-5
641	Bates D, Mächler M, Bolker B, Walker S (2015) Fitting linear mixed-effects models
642	using lme4. Journal of Statistical Software 67:1–48.
643	https://doi.org/10.18637/jss.v067.i01
644	Bernabò I, Bonacci A, Coscarelli F, et al (2013) Effects of salinity stress on <i>Bufo</i>
645	<i>balearicus</i> and <i>Bufo bufo</i> tadpoles: Tolerance, morphological gill alterations and
646	Na+/K+-ATPase localization. Aquatic toxicology (Amsterdam, Netherlands)
647	132-133C:119–133. https://doi.org/10.1016/j.aquatox.2013.01.019
648	Bonnell BS, Chandler DE (1996) Egg jelly layers of <i>Xenopus laevis</i> are unique in
649	ultrastructure and sugar distribution. Molecular Reproduction and
650	Development: Incorporating Gamete Research 44:212–220.
651	https://doi.org/10.1002/(SICI)1098-2795(199606)44:2<212::AID-
652	MRD10>3.0.CO;2-4
653	Brady SP (2013) Microgeographic maladaptive performance and deme depression in

response to roads and runoff. PeerJ 1:e163. https://doi.org/10.7717/peerj.163

655	Byrne PG, Anastas ZM, Silla AJ (2022) A test for plasticity in sperm motility
656	activation in response to osmotic environment in an anuran amphibian.
657	Ecology and Evolution 12:e9387. https://doi.org/10.1002/ece3.9387
658	Byrne PG, Dunne C, Munn AJ, Silla AJ (2015) Environmental osmolality influences
659	sperm motility activation in an anuran amphibian. Journal of Evolutionary
660	Biology 28:521–534. https://doi.org/10.1111/jeb.12584
661 662 663	Cañedo-Argüelles M, Kefford BJ, Piscart C, et al (2013) Salinisation of rivers: An urgent ecological issue. Environmental Pollution 173:157–167. https://doi.org/10.1016/j.envpol.2012.10.011
664	Cheron M, Raoelison L, Kato A, et al (2021) Ontogenetic changes in activity,
665	locomotion and behavioural complexity in tadpoles. Biological Journal of the
666	Linnean Society 134:165–176. https://doi.org/10.1093/biolinnean/blab077
667 668 669	Chinathamby K, Reina RD, Bailey PCE, Lees BK (2006) Effects of salinity on the survival, growth and development of tadpoles of the brown tree frog, <i>Litoria ewingii</i> . Aust J Zool 54:97–105. https://doi.org/10.1071/ZO06006
670	Chivers DP, Kiesecker JM, Marco A, et al (2001) Predator-induced life history
671	changes in amphibians: egg predation induces hatching. Oikos 92:135–142.
672	https://doi.org/10.1034/j.1600-0706.2001.920116.x
673	Christy M, Dickman C (2002) Effects of salinity on tadpoles of the Green and Golden
674	Bell Frog (<i>Litoria aurea</i>). Amphibia-Reptilia 23:1–11.
675	https://doi.org/10.1163/156853802320877582
676	Chuang M, Cheng Y-J, Andersen D, et al (2022) Increasing salinity stress decreases
677	the thermal tolerance of amphibian tadpoles in coastal areas of Taiwan.
678	Scientific Reports 12:9014. https://doi.org/10.1038/s41598-022-12837-7
679	Church JA, White NJ (2011) Sea-level rise from the late 19th to the early 21st century.
680	Surv Geophys 32:585–602. https://doi.org/10.1007/s10712-011-9119-1
681	Cunillera-Montcusí D, Beklioğlu M, Cañedo-Argüelles M, et al (2022) Freshwater
682	salinisation: a research agenda for a saltier world. Trends in Ecology &
683	Evolution 37:440–453. https://doi.org/10.1016/j.tree.2021.12.005
684	Dahrouge NC, Rittenhouse TAG (2022) Variable temperature regimes and wetland
685	salinity reduce performance of juvenile wood frogs. Oecologia 199:1021–1033.
686	https://doi.org/10.1007/s00442-022-05243-3
687	Debue M, Ouédraogo D-Y, Sordello R, Reyjol Y (2022) Impacts of coastal realignment
688	on biodiversity. A systematic review and meta-analysis. Basic and Applied
689	Ecology 60:48–62. https://doi.org/10.1016/j.baae.2022.01.009

690	Denoël M, Bichot M, Ficetola GF, et al (2010) Cumulative effects of road de-icing salt
691	on amphibian behavior. Aquatic Toxicology 99:275–280.
692	https://doi.org/10.1016/j.aquatox.2010.05.007
693	Denver RJ, Glennemeier KA, Boorse GC (2002) Endocrinology of complex life cycles:
694	amphibians. In: Hormones, brain and behavior. Elsevier, pp 469–XI
695	Dettinger M (2011) Climate change, atmospheric rivers, and floods in California - A
696	multimodel analysis of storm frequency and magnitude changes. JAWRA
697	Journal of the American Water Resources Association 47:514–523.
698	https://doi.org/10.1111/j.1752-1688.2011.00546.x
699	Dmitriew CM (2011) The evolution of growth trajectories: what limits growth rate?
700	Biological Reviews 86:97–116. https://doi.org/10.1111/j.1469-
701	185X.2010.00136.x
702	Dougherty C, Smith G (2006) Acute effects of road de-icers on the tadpoles of three
703	anurans. Appl Herpetol 3:87–93.
704	https://doi.org/10.1163/157075406776984266
705	Estévez E, Rodríguez-Castillo T, González-Ferreras AM, et al (2019) Drivers of spatio-
706	temporal patterns of salinity in Spanish rivers: a nationwide assessment.
707	Philosophical Transactions of the Royal Society B 374:20180022.
708	https://doi.org/10.1098/rstb.2018.0022?rss=1
709	Evans DH (2009) Osmotic and ionic regulation: cells and animals. CRC Press.
710 711 712	Evans TG, Kültz D (2020) The cellular stress response in fish exposed to salinity fluctuations. Journal of Experimental Zoology Part A: Ecological and Integrative Physiology 333:421–435. https://doi.org/10.1002/jez.2350
713	Ford NB, Seigel RA (2010) An experimental test of the fractional egg size hypothesis.
714	Herpetologica 66:451–455. https://doi.org/10.1655/09-057.1
715	Froneman PW (2023) The effect of salinity on the egg production rate of the sac-
716	spawning calanoid copepod, <i>Pseudodiaptomus hessei</i> , in a temporarily
717	open/closed Southern African estuary. Diversity 15:263.
718	https://doi.org/10.3390/d15020263
719	Fu Z, Wu F, Zhang Z, et al (2021) Sea surface salinity estimation and spatial-temporal
720	heterogeneity analysis in the gulf of Mexico. Remote Sensing 13:881.
721	https://doi.org/10.3390/rs13050881
722 723 724 725	Gabriel A, Costa S, Henriques I, Lopes I (2021) Effects of long-term exposure to increased salinity on the amphibian skin bacterium <i>Erwinia toletana</i> . Arch Environ Contam Toxicol 80:779–788. https://doi.org/10.1007/s00244-021-00845-z

726 727 728	Gomez-Mestre I, Tejedo M (2003) Local adaptation of an anuran amphibian to osmotically stressful environments. Evolution 57:1889–1899. https://doi.org/10.1111/j.0014-3820.2003.tb00596.x
729 730 731 732	Gomez-Mestre I, Tejedo M, Ramayo E, Estepa J (2004) Developmental alterations and osmoregulatory physiology of a larval anuran under osmotic stress. Physiological and Biochemical Zoology 77:267–274. https://doi.org/10.1086/378143
733 734	Gosner KL (1960) A simplified table for staging anuran embryos and larvae with notes on identification. Herpetologica 16:183–190
735 736	Gosner KL, Black IH (1957) The effects of acidity on the development and hatching of New Jersey frogs. Ecology 38:256–262. https://doi.org/10.2307/1931684
737 738 739	Green L, Niemax J, Herrmann J-P, et al (2021) Sperm performance limits the reproduction of an invasive fish in novel salinities. Diversity and Distributions 27:1091–1105. https://doi.org/10.1111/ddi.13258
740 741 742 743	Hall EM, Brady SP, Mattheus NM, et al (2017) Physiological consequences of exposure to salinized roadside ponds on wood frog larvae and adults. Biological Conservation 209:98–106. https://doi.org/10.1016/j.biocon.2017.02.013
744 745 746	Hall EM, Brunner JL, Hutzenbiler B, Crespi EJ (2020) Salinity stress increases the severity of ranavirus epidemics in amphibian populations. Proc R Soc B 287:20200062. https://doi.org/10.1098/rspb.2020.0062
747 748 749	Haramura T (2016) Hatching plasticity in response to salinity levels in a rhacophorid frog inhabiting a coastal area. J Zool 299:125–131. https://doi.org/10.1111/jzo.12323
750 751 752	Haramura T (2007) Salinity tolerance of eggs of <i>Buergeria japonica</i> (Amphibia, Anura) inhabiting coastal areas. Zoological Science 24:820–823. https://doi.org/10.2108/zsj.24.820
753 754	Hart BT, Lake PS, Webb JA, Grace MR (2003) Ecological risk to aquatic systems from salinity increases. Aust J Bot 51:689–702. https://doi.org/10.1071/bt02111
755 756 757	Herbert ER, Boon P, Burgin AJ, et al (2015) A global perspective on wetland salinization: ecological consequences of a growing threat to freshwater wetlands. Ecosphere 6:1–43. https://doi.org/10.1890/ES14-00534.1
758 759 760 761	 Hieu DQ, Hang BTB, Huong DTT, et al (2021) Salinity affects growth performance, physiology, immune responses and temperature resistance in striped catfish (<i>Pangasianodon hypophthalmus</i>) during its early life stages. Fish Physiol Biochem 47:1995–2013. https://doi.org/10.1007/s10695-021-01021-9

762 Hintz WD, Relyea RA (2019) A review of the species, community, and ecosystem impacts of road salt salinisation in fresh waters. Freshwater biology 64:1081-763 1097. https://doi.org/10.1111/fwb.13286 764 765 Hoover Z, Ferrari MC, Chivers DP (2013) The effects of sub-lethal salinity concentrations on the anti-predator responses of fathead minnows. 766 Chemosphere 90:1047-1052. 767 https://doi.org/10.1016/j.chemosphere.2012.08.051 768 Hopkins G, French S, Brodie III E (2013) Increased frequency and severity of 769 770 developmental deformities in rough-skinned newt (Taricha granulosa) embryos exposed to road deicing salts (NaCl & MgCl2). Environmental pollution 771 772 (Barking, Essex: 1987) 173C:264-269. https://doi.org/10.1016/j.envpol.2012.10.002 773 774 Hopkins GR, Brodie ED (2015) Occurrence of amphibians in saline habitats: A review 775 and evolutionary perspective. Herpetological Monographs 29:1-27. 776 https://doi.org/10.1655/HERPMONOGRAPHS-D-14-00006 Hoverman JT, Relyea RA (2008) Temporal environmental variation and phenotypic 777 plasticity: a mechanism underlying priority effects. Oikos 117:23-32. 778 https://doi.org/10.1111/j.2007.0030-1299.15969.x 779 Hsu W-T, Wu C-S, Hatch KA, et al (2018) Full compensation of growth in salt-780 tolerant tadpoles after release from salinity stress. Journal of Zoology 304:141-781 149. https://doi.org/10.1111/jzo.12508 782 IPCC (2022) Climate change 2022: Impacts, adaptation, and vulnerability. 783 Contribution of Working Group II to the Sixth Assessment Report of the 784 Intergovernmental Panel on Climate Change. [H.-O. Pörtner, D.C. Roberts, M. 785 Tignor, E.S. Poloczanska, K. Mintenbeck, A. Alegría, M. Craig, S. Langsdorf, S. 786 Löschke, V. Möller, A. Okem, B. Rama (Eds.)], Cambridge Univ. Press. 787 Cambridge Univ. Press. 788 Jones B, Snodgrass JW, Ownby DR (2015) Relative toxicity of NaCl and road deicing 789 salt to developing amphibians. Copeia 103:72-77. 790 https://doi.org/10.1643/CP-13-082 791 Karraker NE (2007) Are embryonic and larval green frogs (Rana clamitans) insensitive 792 to road deicing salt? Herpetol Conserv Biol 2:35-41 793 Karraker NE, Gibbs JP (2011) Road deicing salt irreversibly disrupts osmoregulation 794 of salamander egg clutches. Environmental Pollution 159:833-835. 795 796 https://doi.org/10.1016/j.envpol.2010.11.019 797 Karraker NE, Gibbs JP, Vonesh JR (2008) Impacts of road deicing salt on the 798 demography of vernal pool-breeding amphibians. Ecological Applications 799 18:724–734. https://doi.org/10.1890/07-1644.1

800	Katz U (1989) Strategies of adaptation to osmotic stress in anuran Amphibia under
801	salt and burrowing conditions. Comparative Biochemistry and Physiology
802	Part A: Physiology 93:499–503. https://doi.org/10.1016/0300-9629(89)90001-7
803	Knighton AD, Mills K, Woodroffe CD (1991) Tidal-creek extension and saltwater
804	intrusion in northern Australia. Geology 19:831–834.
805	https://doi.org/10.1130/0091-7613(1991)019<0831:TCEASI>2.3.CO;2
806	Knutson TR, McBride JL, Chan J, et al (2010) Tropical cyclones and climate change.
807	Nature geoscience 3:157–163. https://doi.org/10.1002/wcc.371
808	Kültz D (2015) Physiological mechanisms used by fish to cope with salinity stress.
809	The Journal of Experimental Biology 218:1907–1914.
810	https://doi.org/10.1242/jeb.118695
811	Kunzetsova A, Brockhoff P, Christensen R (2017) lmerTest package: Tests in linear
812	mixed effect models. Journal of Statistical Software 82:1–26.
813	https://doi.org/10.18637/jss.v082.i13
814	Lasne E, Leblanc CA-L, Gillet C (2018) Egg size versus number of offspring trade-off:
815	Female age rather than size matters in a domesticated Arctic charr population.
816	Evolutionary Biology 45:105–112. https://doi.org/10.1007/s11692-017-9433-8
817	Leite T, Branco P, Ferreira MT, Santos JM (2022) Activity, boldness and schooling in
818	freshwater fish are affected by river salinization. Science of The Total
819	Environment 819:153046. https://doi.org/10.1016/j.scitotenv.2022.153046
820 821 822	Liu P, Zhao B, Zhang J, et al (2022) Responses of survival, growth, and feeding of the invasive Golden Apple Snail (<i>Pomacea canaliculata</i>) to salinity stress. Freshwater Science 41:000–000. https://doi.org/10.1086/721026
823 824 825	López-Alcaide S, Macip-Ríos R (2011) Effects of climate change in amphibians and reptiles. In: Biodiversity loss in a changing planet, Grillo, O., Y. G., Venora, Eds.; InTech: Rijeka,. InTech Rijeka, pp 163–184
826	Lorrain-Soligon L, Bichet C, Robin F, Brischoux F (2022) From the field to the lab:
827	Physiological and behavioural consequences of environmental salinity in a
828	coastal frog. Frontiers in Physiology 13:919165.
829	https://doi.org/10.3389/fphys.2022.919165
830	Lorrain-Soligon L, Périsse L, Robin F, et al (2023a) The costs of living on the coast:
831	Reduction in body size and size-specific reproductive output in coastal
832	populations of a widespread amphibian. Functional Ecology 37:2703–2717.
833	https://doi.org/10.1111/1365-2435.14413
834	Lorrain-Soligon L, Robin F, Barbraud C, Brischoux F (2023b) Some like it salty:
835	Spatio-temporal dynamics of salinity differentially affect anurans and
836	caudates in coastal wetlands. Freshwater Biology 68:1279–1292.
837	https://doi.org/10.1111/fwb.14103

838 Lorrain-Soligon L, Robin F, Bertin X, et al (2023c) Long-term trends of salinity in coastal wetlands: Effects of climate, extreme weather events, and sea water 839 level. Environmental Research 237:116937. 840 https://doi.org/10.1016/j.envres.2023.116937 841 Lüdecke D, Makowski D, Waggoner P, Patil I (2020) Performance: assessment of 842 regression models performance. Journal of Open Source Software 6:3139. 843 844 https://doi.org/10.21105/joss.03139 Lukens E, Wilcoxen TE (2020) Effects of elevated salinity on Cuban treefrog 845 Osteopilus septontrionalis aldosterone levels, growth, and development. Marine 846 847 and Freshwater Behaviour and Physiology 53:99-111. https://doi.org/10.1080/10236244.2020.1772062 848 849 Mangel M, Munch SB (2005) A life-history perspective on short-and long-term consequences of compensatory growth. The American Naturalist 166:E155-850 E176. https://doi.org/10.1086/444439 851 Martin KL, Nagy KA (1997) Water balance and the physiology of the amphibian to 852 amniote transition. In: Amniote Origins. Elsevier, pp 399–423 853 McLean RF, Tsyban A, Burkett V, et al (2001) Coastal zones and marine ecosystems. 854 Climate change 343–379 855 Meek R (2022) Long-term changes in four populations of the spiny toad, Bufo 856 spinosus, in Western France; Data from road mortalities. Conservation 2:248-857 261. https://doi.org/10.3390/conservation2020017 858 Metcalfe NB, Monaghan P (2001) Compensation for a bad start: grow now, pay later? 859 Trends in ecology & evolution 16:254–260. https://doi.org/10.1016/S0169-860 5347(01)02124-3 861 Miaud C, Muratet J (2018) Identifier les oeufs et les larves des amphibiens de France. 862 863 Editions Quae Moniruzzaman M, Mukherjee M, Kumar S, Chakraborty SB (2022) Effects of salinity 864 stress on antioxidant status and inflammatory responses in females of a "Near 865 Threatened" economically important fish species Notopterus chitala: a 866 mechanistic approach. Environ Sci Pollut Res 29:75031-75042. 867 https://doi.org/10.1007/s11356-022-21142-9 868 Nicholls R, Hoozemans F, Marchand M (1999) Increasing flood risk and wetland 869 losses due to global sea-level rise: regional and global analyses. Global 870 Environmental Change 9:S69-S87. https://doi.org/10.1016/S0959-871 3780(99)00019-9 872 Nilsson J-Åke, Svensson E (1997) The cost of reproduction: a new link between 873 current reproductive effort and future reproductive success. Proceedings of 874

875	the Royal Society of London Series B: Biological Sciences 263:711–714.
876	https://doi.org/10.1098/rspb.1996.0106
877	Nussbaum RA (1981) Seasonal shifts in clutch size and egg size in the side blotched
878	lizard <i>Uta stansburiana</i> Baird and Girard. Oecologia 49:8–13.
879	https://doi.org/10.1007/BF00376891
880 881 882	Pechenik JA (2006) Larval experience and latent effects – metamorphosis is not a new beginning. Integrative and Comparative Biology 46:323–333. https://doi.org/10.1093/icb/icj028
883	Pinder AM, Halse SA, McRae JM, Shiel RJ (2005) Occurrence of aquatic invertebrates
884	of the wheatbelt region of Western Australia in relation to salinity.
885	Hydrobiologia 543:1–24. https://doi.org/10.1007/s10750-004-5712-3
886	R Core Team (2020) R: A language and environment for statistical computing. R
887	Foundation for Statistical Computing. Austria: Vienna. https://www.R-
888	project.org/
889 890 891	Ranjbar A, Ehteshami M (2019) Spatio-temporal mapping of salinity in the heterogeneous coastal aquifer. Appl Water Sci 9:1–14. https://doi.org/10.1007/s13201-019-0908-x
892	Reid AJ, Carlson AK, Creed IF, et al (2019) Emerging threats and persistent
893	conservation challenges for freshwater biodiversity. Biological Reviews
894	94:849–873. https://doi.org/10.1111/brv.12480
895 896 897 898	Renoirt M, Angelier F, Cheron M, Brischoux F (2022) What are the contributions of maternal and paternal traits to fecundity and offspring development? A case study in an amphibian species, the spined toad <i>Bufo spinosus</i> . Current Zoology zoac072. https://doi.org/10.1093/cz/zoac072/6694174
899 900 901	Ricklefs RE (1968) On the limitation of brood size in passerine birds by the ability of adults to nourish their young. Proceedings of the National Academy of Sciences 61:847–851. https://doi.org/10.1073/pnas.61.3.847
902	Rivera-Ingraham GA, Lignot J-H (2017) Osmoregulation, bioenergetics and oxidative
903	stress in coastal marine invertebrates: raising the questions for future research.
904	J Exp Biol 220:1749–1760. https://doi.org/10.1242/jeb.135624
905 906 907	Sanzo D, Hecnar SJ (2006) Effects of road de-icing salt (NaCl) on larval wood frogs (<i>Rana sylvatica</i>). Environmental Pollution 140:247–256. https://doi.org/10.1016/j.envpol.2005.07.013
908 909 910	Schneider CA, Rasband WS, Eliceiri KW (2012) NIH Image to ImageJ: 25 years of image analysis. Nature methods 9:671–675. https://doi.org/10.1038/nmeth.2089

911 912	Schultz ET, McCormick SD (2012) Euryhalinity in an evolutionary context. Fish physiology 32:477–533. https://doi.org/10.1016/B978-0-12-396951-4.00010-4
913	Shoemaker VH, Nagy KA (1984) Osmoregulation in the Galápagos marine iguana,
914	<i>Amblyrhynchus cristatus</i> . Physiological zoology 57:291–300.
915	https://doi.org/10.1086/physzool.57.3.30163717
916 917 918	Sinervo B, DeNardo DF (1996) Costs of reproduction in the wild: path analysis of natural selection and experimental tests of causation. Evolution 50:1299–1313. https://doi.org/10.1111/j.1558-5646.1996.tb02370.x
919	Singh A (2021) Soil salinization management for sustainable development: A review.
920	Journal of Environmental Management 277:111383.
921	https://doi.org/10.1016/j.jenvman.2020.111383
922 923 924	Smith CC, Fretwell SD (1974) The optimal balance between size and number of offspring. The American Naturalist 108:499–506. https://doi.org/10.1086/282929
925	Speybroeck J, Beukema W, Bok B, Van Der Voort J (2018) Guide Delachaux des
926	amphibiens & reptiles de France et d'Europe. Delachaux et Nieslé
927 928 929	Squires ZE, Bailey PCE, Reina RD, Wong BBBM (2010) Compensatory growth in tadpoles after transient salinity stress. Mar Freshwater Res 61:219–222. https://doi.org/10.1071/MF09123
930	Tornabene BJ, Breuner CW, Hossack BR (2021a) Comparative effects of energy-
931	related saline wastewaters and sodium chloride on hatching, survival, and
932	fitness-associated traits of two amphibian species. Environmental Toxicology
933	and Chemistry 40:3137–3147. https://doi.org/10.1002/etc.5193
934	Tornabene BJ, Crespi EJ, Breuner CW, Hossack BR (2022) Testing whether adrenal
935	steroids mediate phenotypic and physiologic effects of elevated salinity on
936	larval tiger salamanders. Integrative Zoology 18:27–44.
937	https://doi.org/10.1111/1749-4877.12669
938	Tornabene BJ, Hossack BR, Crespi EJ, Breuner CW (2021b) Corticosterone mediates a
939	growth-survival tradeoff for an amphibian exposed to increased salinity.
940	Journal of Experimental Zoology Part A: Ecological and Integrative
941	Physiology 335:703–715. https://doi.org/10.1002/jez.2535
942 943 944	Traversari BA (2021) Carry-over effects of elevated salinity and temperature on wood frog (<i>Rana sylvatica</i>) survival, growth, and behavior. PhD Thesis, Washington State University
945	Trenberth KE, Fasullo JT, Shepherd TG (2015) Attribution of climate extreme events.
946	Nature Clim Change 5:725-730. https://doi.org/10.1038/nclimate2657

947 948 949	Uchiyama M, Yoshizawa H (1992) Salinity tolerance and structure of external and internal gills in tadpoles of the crab-eating frog, <i>Rana cancrivora</i> . Cell Tissue Res 267:35–44. https://doi.org/10.1007/BF00318689
950	Venturino A, Rosenbaum E, Caballero A, et al (2003) Biomarkers of effect in toads
951	and frogs. Biomarkers : biochemical indicators of exposure, response, and
952	susceptibility to chemicals 8:167–86.
953	https://doi.org/10.1080/1354700031000120116
954	Vermeer M, Rahmstorf S (2009) Global sea level linked to global temperature.
955	Proceedings of the National Academy of Sciences 106:21527–21532.
956	https://doi.org/10.1073/pnas.0907765106
957	Visschers LLB, Santos CD, Franco AMA (2022) Accelerated migration of mangroves
958	indicate large-scale saltwater intrusion in Amazon coastal wetlands. Science of
959	The Total Environment 836:155679.
960	https://doi.org/10.1016/j.scitotenv.2022.155679
961 962 963	Vonesh JR, Warkentin KM (2006) Opposite shifts in size at metamorphosis in response to larval and metamorph predators. Ecology 87:556–562. https://doi.org/10.1890/05-0930
964	Warkentin KM (2011) Plasticity of hatching in amphibians: evolution, trade-offs, cues
965	and mechanisms. Integrative and comparative biology 51:111–127.
966	https://doi.org/10.1093/icb/icr046
967	Wells KD (2007) The Ecology and Behavior of Amphibians, University of Chicago
968	Press. University of Chicago Press
969 970 971	Werner EE (1986) Amphibian metamorphosis: growth rate, predation risk, and the optimal size at transformation. The American Naturalist 128:319–341. https://doi.org/10.1086/284565
972 973 974 975	Wijethunga U, Greenlees M, Shine R (2016) Living up to its name? The effect of salinity on development, growth, and phenotype of the "marine" toad (<i>Rhinella marina</i>). J Comp Physiol B 186:205–213. https://doi.org/10.1007/s00360-015-0944-2
976	Wilbur HM, Collins JP (1973) Ecological aspects of amphibian metamorphosis:
977	nonnormal distributions of competitive ability reflect selection for facultative
978	metamorphosis. Science 182:1305–1314
979 980 981	Wilder AE, Welch AM (2014) Effects of salinity and pesticide on sperm activity and oviposition site selection in green Treefrogs, <i>Hyla cinerea</i> . Copeia 2014:659–667. https://doi.org/10.1643/CE-14-053
982	Williams WD (2001) Anthropogenic salinisation of inland waters. In: Melack JM,
983	Jellison R, Herbst DB (eds) Saline Lakes. Springer Netherlands, Dordrecht, pp
984	329–337

985 986 987 988	Wood L, Welch AM (2015) Assessment of interactive effects of elevated salinity and three pesticides on life history and behavior of southern toad (<i>Anaxyrus</i> <i>terrestris</i>) tadpoles. Environmental Toxicology and Chemistry 34:667–676. https://doi.org/10.1002/etc.2861
989 990 991	Woodley A, Hintz LL, Wilmoth B, Hintz WD (2023) Impacts of water hardness and road deicing salt on zooplankton survival and reproduction. Sci Rep 13:2975. https://doi.org/10.1038/s41598-023-30116-x
992 993 994	Wu C-S, Gomez-Mestre I, Kam Y-C (2012) Irreversibility of a bad start: early exposure to osmotic stress limits growth and adaptive developmental plasticity. Oecologia 169:15–22. https://doi.org/10.1007/s00442-011-2170-2
995 996 997	Wu C-S, Kam Y-C (2009) Effects of salinity on the survival, growth, development, and metamorphosis of <i>Fejervarya limnocharis</i> tadpoles living in brackish water. Zoological science 26:476–82. https://doi.org/10.2108/zsj.26.476
998 999 1000	Xu F, Yang W, Li Y (2019) Enlarged egg size increases offspring fitness of a frog species on the zhoushan archipelago of china. Sci Rep 9:11653. https://doi.org/10.1038/s41598-019-48147-8
1001 1002 1003	Xue Z, He R, Fennel K, et al (2013) Modeling ocean circulation and biogeochemical variability in the Gulf of Mexico. Biogeosciences 10:7219–7234. https://doi.org/10.5194/bg-10-7219-2013
1004 1005 1006	Yurewicz EC, Oliphant G, Hedrick JL (1975) Macromolecular composition of <i>Xenopus laevis</i> egg jelly coat. Biochemistry 14:3101–3107. https://doi.org/10.1021/bi00685a010
1007	

1008	Statements	& Dec	larations
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- 1009 Ethical approval
- 1010 This work was approved by the French authorities under permits
- 1011 DREAL/2020D/8041 (animals capture) and APAFIS #33592-2021102610033982
- 1012 (animals husbandry).
- 1013
- 1014 Consent to participate
- 1015 Not applicable.
- 1016
- 1017 Consent to publish
- 1018 Not applicable.
- 1019

1020 Authors contribution

- 1021 LLS and FB have conceptualized the study. LLS, FR and MJ participated to field
- 1022 prospection. LLS and TB participated to data collection. LLS analyzed the data. LLS
- 1023 and FB wrote the initial draft. All authors have reviewed and edited the manuscript
- 1024 and approved the final version.
- 1025

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- 1031

1032 **Competing interest**

1033 The authors declare that they have no conflict of interest.

1034

1035 Data archiving statement

1036 Data supporting the findings of this study are made available as a supplementary

1037 material.

1038

1040 Caption to figures

Figure 1 – Experimental design used to assess the effects of salinity during different stages of development in *Bufo spinosus*. Colours represent salinity treatments (light blue = fresh water $[0 \text{ g.}]^{-1}$, dark blue = brackish water $[4 \text{ g.}]^{-1}$).

1044

Figure 2 – Effects of salinity during spawning on reproductive parameters. (A) Egg
diameter, (B) clutch size (number of eggs), (C) relationships between clutch size and
egg diameter, for clutches produced in freshwater (0 g.l⁻¹) or brackish water (4 g.l⁻¹).
Data are presented as mean ± SE. See Figure 1 for sample size.

1049

Figure 3 – Effects of salinity during spawning and embryonic development on different aspects of embryonic development assessed upon hatching (Gosner stage 25). (A) Hatching success, (B) malformation rate, and (C) total length (mm). FF: spawning in freshwater [0 g.l⁻¹], embryonic development in freshwater; FS: spawning in freshwater, embryo in brackish water [4 g.l⁻¹]; SF: spawning in brackish water, embryo in freshwater; SS: spawning in brackish water, embryo in brackish water. Data are presented as mean ± SE. See Figure 1 for sample size.

1057

Figure 4 - Effects of salinity during spawning, embryonic development and larval 1058 development on different parameters of larval development measured at Gosner stage 1059 1060 30. (A) Mortality rate, (B) duration between GS25 and GS30, (C) growth between GS25 and GS30, and (D) total length (mm). FFF: spawning in freshwater [0 g.l-1], embryonic 1061 development in freshwater, larval development in freshwater; FSF: spawning in 1062 freshwater, embryo in brackish water [4 g.l-1], larvae in freshwater; FFS: spawning in 1063 freshwater, embryo in freshwater, larvae in brackish water; FSS: spawning in 1064 freshwater, embryo in brackish water, larvae in brackish water; SFF: spawning in 1065 brackish water, embryo in freshwater, larvae in freshwater; SSF: spawning in brackish 1066 1067 water, embryo in brackish water, larvae in freshwater; SFS: spawning in brackish water, embryo in freshwater, larvae in brackish water; SSS: spawning in brackish 1068 water, embryo in brackish water, larvae in brackish water. Data are presented as mean 1069 ± SE. Treatments and sample size are illustrated in Figure 1. 1070

1071

1073 Figure 1



1075 Figure 2





1077 Figure 3



1081 Caption to tables

Embryonic treatment	Number of individuals in the treatment	Surviving individuals at GS25	Larval treatment	Number of individuals in the treatment	Surviving individuals at GS25	Surviving individuals at GS30
	66	66	FFF	36	36	30
11	00	00	FFS	30	30	15
FS	102	70	FSF	53	37	14
15	102	70	FSS	49	33	2
SE	66	65	SFF	33	33	22
51	00	individuals at GS25 66 70 65 31	SFS	33	32	18
22	83	21	SSF	46	20	7
33	55 65 51	SSS	37	11	2	

1082 Table 1 – Number of surviving individuals in each treatment at each life stage

1084 Caption to Appendices

1085 Appendix A - Contributions of clutches to each treatment (number of tadpoles1086 originating from each clutch in each treatment)

Appendix B - Mean ± SE values for all studied variables and treatments during egg
laying, embryonic development and larval development.

1089 Appendix C – Post-hoc analyses for the differences found at Gosner stage 25 (GS25) 1090 with the interaction between salinity during spawning (clutches laid in freshwater [0 1091 g.l⁻¹] or brackish water [4 g.l⁻¹]) and salinity during embryonic development 1092 (freshwater [0 g.l⁻¹] or brackish water [4 g.l⁻¹]), for hatching success, malformation rate, 1093 and time to hatching.

1094

1095 Appendix D – Post-hoc analyses for the differences found for duration between Gosner 1096 stage 25 (GS25) and Gosner stage 30 (GS30) with the interaction between salinity 1097 during embryonic development (freshwater $[0 \text{ g.l}^{-1}]$ or brackish water $[4 \text{ g.l}^{-1}]$) and 1098 salinity during larval development (freshwater $[0 \text{ g.l}^{-1}]$ or brackish water $[4 \text{ g.l}^{-1}]$), and 1099 the interaction between salinity during spawning (clutches laid in freshwater $[0 \text{ g.l}^{-1}]$ 100 or brackish water $[4 \text{ g.l}^{-1}]$) and salinity during larval development (freshwater $[0 \text{ g.l}^{-1}]$ 1101 or brackish water $[4 \text{ g.l}^{-1}]$).

1103 Appendix A

	Number of		
Treatment	Clutch	tadpoles	
	F01	3	
	F02	9	
	F03	3	
	F04	3	
	F05	3	
FFF	F06	3	
	F07	3	
	F08	3	
	F09	3	
	F10	3	
	F01	3	
	F02	3	
	F03	3	
	F04	3	
	F05	3	
FFS	F06	3	
	F07	3	
	F08	3	
	F09	3	
	F10	3	
	F01	3	
	F02	1	
	F05	3	
FSF	F07	21	
	F08	5	
	F09	4	
	F01	5	
	F02	1	
	F03	3	
FSS	F06	1	
	F07	16	
	F08	5	
	F09	2	
	S01	3	
	S02	3	
	S03	3	
	S04	3	
	S05	3	
SFF	S06	3	
	S07	3	
	S08	3	
	S09	3	
	S10	3	

	S11	3
	S01	3
	S02	3
	S03	3
	S04	3
	S05	3
SFS	S06	3
	S07	2
	S08	3
	S09	3
	S10	3
	S11	3
	S05	1
SSF	S10	8
	S11	11
	S02	2
	S06	3
SSS	S07	1
	S10	4
	S11	1

1106 Appendix B

Period	Parameter	Group	Mean	SE
	Longth (cm)	F	1608.2097	185.0155628
	Length (Chi)	S	1678.259455	186.6813434
	Number of	F	2427.04	358.3985591
	eggs	S	2609.6	208.7763819
Egg laying		F	92.82	10.71980307
	iviass (g)	S	64.60909091	7.777607187
	Eggs diameter	F	1.76688	0.039757389
	(mm)	S	1.660572727	0.043287962
	Laying time (h)	F	79.1	13.10847902
		S	47.33272727	7.662623464
	Females loss in	F	-13.55765323	2.230276924
	mass (%)	S	-12.20855975	1.364961958
	Males loss in	F	-1.627788792	2.18901737
	mass (%)	S	3.788011248	1.265685304
		FF	0.68	0.039014277
	Hatching	FS	0.284444444	0.043299898
	success	SF	0.439393939	0.032995599
		SS	0.037373737	0.018445671
		FF	0.252273761	0.018871836
	Malformation rate	FS	0.609116088	0.026959964
		SF	0.253329084	0.031584961
		SS	0.269217687	0.04932672
		FF	17.73333333	0.150101261
Embryonic	Time to hatching (days)	FS	18.81428571	0.223183124
development		SF	17.81538462	0.191014885
		SS	19.06451613	0.270147247
		FF	8.839393939	0.144306052
	Total lenght	FS	7.127058824	0.075127201
	(mm)	SF	8.619230769	0.122971289
		SS	6.529354839	0.188660779
	General activity (scale on 1 to 5)	FF	3.766666667	0.140376386
		FS	1.071428571	0.031346695
		SF	3.415384615	0.163924683
		SS	1.214285714	0.076478688
		FFF	0.142857143	0.059172634
	Mortality	FFS	0.5	0.092847669
		FSF	0.735849057	0.073173841
		FSS	0.959183673	0.034800726
Larval		SFF	0.333333333	0.083333333
development		SFS	0.4375	0.089098306
		SSF	0.847826087	0.081204698
		SSS	0.945945946	0.069119496
		FFF	10.87096774	1.811827957
		FFS	10.86666667	1.983972819

Duration between S25 and S30 (days)	FSF	10.07142857	1.655732658
	FSS	13.5	2.350048355
	SFF	12.90909091	2.247184286
	SFS	10.31578947	1.823591173
	SSF	13.28571429	2.970776027
	SSS	13.5	4.070403152
	FFF	0.268393183	0.03012627
	FFS	0.22382476	0.015880458
Cuevuth vete	FSF	0.341870161	0.029992171
Growth rate	FSS	0.159178572	0.001028693
and S30	SFF	0.231039881	0.029893008
	SFS	0.182156128	0.015580051
	SSF	0.2555141	0.014131725
	SSS	0.286321429	0.028431838
	FFF	11.57616129	0.275041152
	FFS	11.493	0.148745132
Total lenght (mm)	FSF	10.57214286	0.18434435
	FSS	10.0815	0.029665049
	SFF	11.64254545	0.331575428
	SFS	10.86410526	0.119641317
	SSF	10.83157143	0.198921964
	SSS	10.4225	0.450066712
	FFF	3.774193548	0.170563757
Activity (scale of 1 to 5)	FFS	2.533333333	0.23771398
	FSF	3.642857143	0.189178582
	FSS	1.5	0.123091491
	SFF	3.681818182	0.164587583
	SFS	2.842105263	0.251820854
	SSF	4.285714286	0.109108945
	SSS	3	0

1109 Appendix C

Gosner

stage	Variable		Contrast	Estimate	SE	z/t	p-value
-		Embryonic					
		0	Spawning0-Spawning4	0.48	0.194	2.469	0.014
		Embryonic					
	Hatching success	4	Spawning0-Spawning4	1.97	0.227	8.697	<0.001
			Embryonic0-				
		Spawning 0	Embryonic4	0.972	0.064	15.103	<0.001
			Embryonic0-				
		Spawning 4	Embryonic4	2.466	0.126	19.625	<0.001
		Embryonic					
		0	Spawning0-Spawning4	0.087	0.242	0.357	0.721
		Embryonic					
	Malformation	4	Spawning0-Spawning4	1.146	0.361	3.169	0.002
S25	rate		Embryonic0-				
		Spawning 0	Embryonic4	-0.893	0.114	-7.84	<0.001
			Embryonic0-				
		Spawning 4	Embryonic4	0.166	0.278	0.598	0.55
		Embryonic				0.4.60	
		0	Spawning0-Spawning4	-0.004	0.025	-0.168	0.868
		Embryonic		0.004	0.000	2 4 4 2	0.040
		4	Spawning0-Spawning4	0.061	0.029	2.118	0.042
	Time to hatching	c · · ·	Embryonic0-	0.407	0.010	0.400	0.004
		Spawning 0	Embryonic4	-0.107	0.013	-8.438	<0.001
			Embryonic0-				
		Spawning 4	Embryonic4	-0.042	0.017	-2.544	0.012

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1112 Appendix D

			Estimate	SE	t value	p-value
	Embryonic0 - Embryonic4		-1.40912	0.63540	-2.2177	0.028809
	Spawning0 - Spawning4	-0.87456	0.81189	-1.0772	0.294830	
Duation	Larval0 - Larval4		-0.53475	0.55331	-0.9665	0.336348
	Embryonic0:Larval0 Embryonic0:Larval0 Embryonic0:Larval0	Embryonic4:Larval0	0.24447	0.58293	0.4194	0.675807
		Embryonic0:Larval4	1.11884	0.42624	2.6249	0.010097
		Embryonic4:Larval4	-1.94387	1.03537	-1.8775	0.063455
	Embryonic4:Larval0	Embryonic0:Larval4	0.87437	0.58972	1.4827	0.141317
between	Embryonic4:Larval0	Embryonic4:Larval4	-2.18834	1.01502	-2.1560	0.033720
S25 and S30	Embryonic0:Larval4	Embryonic4:Larval4	-3.06271	1.03309	-2.9646	0.003827
	Spawning0:Larval0 -	Spawning4:Larval0	-2.33170	0.84095	-2.7727	0.011038
	Spawning0:Larval0 -	Spawning0:Larval4	-1.99189	0.66563	-2.9925	0.003549
	Spawning0:Larval0 -	Spawning4:Larval4	-1.40932	0.96120	-1.4662	0.151424
	Spawning4:Larval0 -	Spawning0:Larval4	0.33981	1.00336	0.3387	0.736590
	Spawning4:Larval0 -	Spawning4:Larval4	0.92238	0.68144	1.3536	0.179157
	Spawning0:Larval4 -	Spawning4:Larval4	0.58257	0.95197	0.6120	0.544697