"Heat waves" experienced during larval life have species-specific consequences on life-history traits and sexual development in anuran amphibians

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Abstract

Extreme temperatures during heat waves can induce mass-mortality events, but can also exert sublethal negative effects by compromising life-history traits and derailing sexual development. Ectothermic animals may, however, also benefit from increased temperatures via enhanced physiological performance and the suppression of cold-adapted pathogens. Therefore, it is crucial to address how the intensity and timing of naturally occurring or human-induced heat waves affect life-history traits and sexual development in amphibians, to predict future effects of climate change and to minimise risks arising from the application of elevated temperature in disease mitigation. We raised agile frog (*Rana dalmatina*) and common toad (*Bufo bufo*) tadpoles at 19 °C and exposed them to a simulated heat wave of 28 or 30 °C for six days during one of three ontogenetic periods (early, mid or late larval development). In agile frogs, exposure to 30 °C during early larval development increased mortality. Regardless of timing, all heat-treatments delayed metamorphosis, and exposure to 30 °C decreased body mass at metamorphosis. Furthermore, exposure to 30 °C during any period and to 28 °C late in development caused female-to-male sex reversal, skewing sex ratios strongly towards males. In common toads, high temperature only slightly decreased survival and did not influence phenotypic sex ratio, while it reduced metamorph mass and length of larval development. Juvenile body mass measured two months after metamorphosis was not adversely affected by temperature treatments in either species. Our results indicate that heat waves may have devastating effects on amphibian populations, and the severity of these negative consequences, and sensitivity can vary greatly between species and with the timing and intensity of heat. Finally, thermal treatments against cold-adapted pathogens have to be executed with caution,

52 taking into account the thermo-sensitivity of the species and the life stage of animals to be
53 treated.
54
55 Keywords:
56 *Batrachochytrium dendrobatidis*; Bufonidae; Chytridiomycosis; Heat stress; Ranidae; Thermal
57 tolerance

Introduction

Earth's wildlife and ecosystem face the sixth mass extinction event today due to anthropogenic environmental alterations, including extreme climatic conditions (Ceballos et al. 2015). Due to global climate change, heat waves occur with increasing frequency, intensity and duration (Gardner et al. 2016). Extreme temperatures during heat waves expose species to intensified physiological stress (Williams et al. 2016) and can even induce mass-mortality events (Welbergen et al. 2008, McKechnie and Wolf 2019). Warming climate with frequently reappearing heat waves can alter species distributions (Krockenberger et al. 2012, Stillman 2019), trigger shifts in the timing of the breeding season and directly affect breeding success in a taxonomically diverse range of species (Blaustein et al. 2001, Oswald et al. 2008, Truebano et al. 2018, Stillman 2019). These factors can generate profound changes in community structure and ecosystem functioning via the formation of interactions between species with previously non-overlapping spatial or temporal distributions (Williams et al. 2016) and the alteration of predator-prey and host-pathogen systems (Blaustein et al. 2010, Cohen et al. 2019, Stillman 2019, Carreira et al. 2020). Fluctuations in temperature affect ectotherms in particular because they lack the metabolic, physiological and anatomical mechanisms that would allow them to maintain constant body temperature, and, therefore, ectothermsare able to maintain high physiological performance only within a narrower environmental temperature range than are endotherms (Clarke and Pörtner 2010).

Amphibians are among the most threatened vertebrate groups, because 41 % of the species are endangered (IUCN 2021), and almost 50 % show population declines worldwide, mainly due to anthropogenic environmental change (Stuart et al. 2004, Wake and Vredenburg 2008, Hof et

al. 2011, Monastersky 2014, Campbell Grant et al. 2016). The growing incidence of meteorological extremes and rising temperatures resulting from global climate change and anthropogenic heat pollution (i.e. urban heat islands; Arnfield 2003, Brans et al. 2018) are major threats to amphibians. Their complex life cycle, usually including an aquatic stage, the unshelled eggs and a highly permeable integument make amphibians excessively sensitive to water availability. Also, though amphibian larvae generally exhibit a relatively high thermal tolerance (Ultsch et al. 1999, Sunday et al. 2011, but also see Harkey and Semlitsch 1988, Wallace and Wallace 2000, Bellakhal et al. 2014, Goldstein et al. 2017) temperatures as low as 30 °C experienced during the larval period can be detrimental to them. Heat can result in delayed metamorphosis (Goldstein et al. 2017), reduced body mass (Harkey and Semlitsch 1988, Phuge 2017, Lambert et al. 2018), disabled locomotor activity (Goldstein et al. 2017), sex reversal (Dournon et al. 1984, Wallace and Wallace 2000, Mikó et al. 2021) and biased sex ratios (Phuge 2017, Lambert et al. 2018, Ruiz-Garciá et al. 2021). Exposure of adult frogs to 30 °C or higher can increase stress hormone levels (Juráni et al. 1973, Narayan and Hero 2014) and enhance the processes that contribute to accelerated ageing (Burraco et al. 2020).

Emerging infectious diseases represent another serious threat to amphibians (Harvell et al. 2002, Pounds et al. 2006). Due to repeated introductions arising from human activities (Lips 2016, O'Hanlon et al. 2018), chytridiomycosis caused by the chytrid fungi *Batrachochytrium dendrobatidis* (*Bd*) and *Batrachochytrium salamandrivorans* (*Bsal*) (Van Rooij et al. 2015) has already led to the decline or extinction of several hundred species and continues to cause mass mortality events on five continents (Scheele et al. 2019). Since *Bsal* was only discovered eight years ago (Martel et al. 2013) and its known geographic distribution is much smaller (Spitzen-van der Sluijs et al. 2016), we focus here on the much better known and more widespread *Bd*.

104 The fungus infects keratinous epidermal layers of the skin with waterborne motile zoospores
105 (Berger et al. 1998), impairs its osmoregulatory function, which leads to shifts in electrolyte
106 balance that can ultimately result in cardiac asystolic death in metamorphosed anurans (Voyles et
107 al. 2009). Tadpoles exhibit keratinous elements only in their mouthparts; therefore they are less
108 susceptible to *Bd* infection than subsequent life stages (Marantelli et al. 2004, Blaustein et al.
109 2005). Nonetheless, it is often the early ontogeny (larval and metamorphic stages) when
110 individuals become infected, due to their aquatic lifestyle (Kilpatrick et al. 2010). The thermal
111 optimum of this cold-adapted fungus is between 18-24 °C, and its growth ceases above 27-28 °C
112 (Cohen et al. 2017, Voyles et al. 2017), while the vast majority of amphibian species can survive
113 temperatures above 30 °C (Ultsch et al. 1999, Sunday et al. 2011). Consequently, when and
114 wherever microclimatic conditions allow amphibians to sufficiently raise their body temperature
115 via thermoregulation, *Bd* infection prevalence and intensity are low (Richards-Zawacki 2010,
116 Forrest and Schlaepfer 2011, Becker et al. 2012), and mass mortalities typically only occur in
117 constantly cool environments (Berger et al. 2004, Woodhams and Alford 2005). Accordingly,
118 thermal treatment of amphibians with 28 °C and higher for a few days can be effectively applied
119 for *Bd*-disinfection of larval, juvenile or adult amphibians in captive populations (Woodhams et
120 al. 2003, Retallick and Miera 2007, Chatfield and Richards-Zawacki 2011, Geiger et al. 2011,
121 McMahon et al. 2014) and may also prove effective for fighting *Bd in situ* (Hettyey et al. 2019).
122 Based on the above information, heat waves may exert several opposing effects on
123 developing amphibians, which may be beneficial for combating *Bd* but harmful for other fitness-
124 related traits. Heat waves usually last for only a couple of days, and just a few days of heat
125 treatment can be sufficient for the suppression or even the complete clearance of cold-adapted
126 pathogens, such as *Bd* (Woodhams et al. 2003, Retallick and Miera 2007, McMahon et al. 2014).

However, we still know little about the developmental costs of brief periods of high temperatures for larval amphibians because most experiments that investigated the effects of heat on larval fitness exposed animals to heat chronically for several weeks, and the effects of shorter heat pulses are rarely tested (Mikó et al. 2021). Because the effects of high temperatures are likely to depend on the intensity, timing and duration of exposure, and may differ between species, studies focusing on these sources of variation are necessary to assess potential malign impacts of heat waves on amphibians and uncover hidden risks arising from thermal treatment of diseased animals.

In this study, we experimentally investigated the developmental effects of six-day long exposures to 28 and 30 °C during early, mid, and late larval development of two amphibian species. We assessed the effects of these experimental heat waves on the survival, growth, somatic and sexual development of agile frogs (*Rana dalmatina*; Bonaparte, 1840) and common toads (*Bufo bufo*; Linnaeus, 1758). These two species are common in Europe, they inhabit various types of water bodies, have different thermal optima (Morand et al. 1997) and represent two globally widespread families (Bufonidae and Ranidae). The temperatures we applied occur during heat waves in aquatic habitats of amphibian larvae in the temperate climate zone (Lambert et al. 2018, Lindauer et al. 2020) and are also recommended for thermal treatment of diseased amphibians (Chatfield and Richards-Zawacki 2011, McMahon et al. 2014, Cohen et al. 2017, Hettyey et al. 2019). Thus, our aim was twofold: to reveal developmental effects of heat waves that may occur in natural habitats, and to assess possible negative consequences of thermal treatment applied against cold-adapted pathogens.

Methods

Experimental procedures

In March 2019, we collected 50 eggs from each of 12 freshly laid egg clutches of the agile frog from three ponds in the Pilis-Visegrádi Mountains, Hungary (Katlan: 47.71110° N, 19.04570° E, Ilona-tó: 47.71326° N, 19.04050° E, and Apátkúti pisztrángos: 47.76656° N, 18.98121° E; four clutches from each population). We transported the eggs to the Experimental Station of the Plant Protection Institute in Julianna-major, Budapest, and placed each clutch (family hereafter) into a plastic container (24 × 16 ×13 cm) filled with 1.3 L continuously aerated reconstituted soft water (RSW; APHA et al. 1992, USEPA 2002). In the laboratory, we maintained 16.3 ± 0.3 °C (mean ± SD) and the lighting was adjusted weekly to outdoor conditions, starting with 12:12 h light:dark cycles in late March, which we gradually changed to 14:10 h by the end of April. In April, we collected 50 eggs from each of 12 freshly laid egg strings of the common toad from two ponds in the Pilis-Visegrádi Mountains (Apátkúti tározó: 47.77444° N, 18.98624° E, and Határréti tó: 47.64644° N, 18.90920° E) and one pond in Budapest (Hidegkúti horgásztó: 47.56942° N, 18.95509° E), i.e. four clutches from each population. We housed common toad eggs as described above for agile frogs.

Four days after hatching, when all individuals reached the free-swimming stage (development stage 25; according to Gosner 1960), we started the experiment by haphazardly selecting 36 healthy-looking larvae from each family (36 individuals × 12 families = 432 individuals per species). Tadpoles not used in the experiment were released at the site of their origin. We reared tadpoles individually in opaque plastic containers (18 × 13 × 12 cm) filled with 1 L RSW, arranged in a randomised block design, where each block contained members of one family. Air temperature in the laboratory was 20.1 ± 1.1 °C resulting in 19.0 ± 0.2 °C water

temperature in tadpole containers. We changed water in the tadpole rearing containers twice a week and fed tadpoles *ad libitum* with slightly boiled, chopped spinach.

We exposed tadpoles to 19 (unheated control), 28, or 30 °C water temperature for six days, starting 6, 12, or 18 days after hatching (Fig. 1). Thus, thermal treatments were applied during three ontogenetic periods: in early, mid, and late larval stages (hereafter 1^{st}, 2^{nd} and 3^{rd} larval period). This resulted in nine treatments with 48 replicates (4 individuals per family × 12 families) in each treatment for each species. In agile frogs, data from the 19 and 30 °C treatments presented here were also used (combined with data from additional treatment groups) for testing another *a-priori* study question, which we published elsewhere (Mikó et al. 2021). We performed thermal treatments in a separate room adjacent to the room where we reared tadpoles. Lighting conditions and room temperature were set to be identical in the two rooms. Immediately before starting thermal treatments, we performed a water change and topped up the RSW to reach a depth of 10 cm (1.7 L RSW in each container during treatment). We placed the containers in 80 × 60 × 12 cm trays filled with tap water to a depth of 8 cm (to avoid floating of the rearing containers), and started to heat the water in the trays to the treatment-specific temperature using thermostated aquarium heaters (Tetra HT 200 in 28 °C treatments and Tetra HT 300 in 30 °C treatments, Tetra GmbH, Melle, Germany). Thereby, water temperature increased gradually to the desired level in ca. two hours, allowing tadpoles to adapt. Opposite to heaters, we placed water pumps (Tetra WP 300) to ensure homogeneous water temperatures, resulting in < 0.5 °C difference among tadpole containers within trays. Overall, this resulted in 28.1 ± 0.4 and 30.0 ± 0.3 °C (mean ± SD) in heated tadpole containers in respective treatments (for details on temperature setting and validation, see the electronic Supporting Information; Fig S1, Table S1). Each tray hosted twelve containers, one from each family (Fig S2), resulting in

four trays in each thermal treatment at a time. During the treatment period, we changed water in the tadpole containers every other day with aerated RSW pre-heated to the treatment-specific temperature, and fed tadpoles with a reduced (ca. 1/3) amount of spinach to prevent water fouling and anoxia. Control individuals experienced the same handling and treatment conditions, except that their trays lacked heaters. At the end of the six-day long thermal treatment periods, we changed water with 1 L heated and aerated RSW, removed the containers from the trays and placed them back into their original position in the laboratory, allowing tadpoles to cool down gradually.

After the last thermal treatments, when tadpoles approached metamorphosis, we checked all rearing containers daily. When an individual started to metamorphose (emergence of forelimbs; development stage 42), we measured its body mass to the nearest 0.1 mg with an analytical balance (Ohaus Pioneer PA-114, Ohaus Europe Gmb, Nanikon, Switzerland), replaced its rearing water with 0.1 L fresh RSW, lifted one side of the container by ca. 2 cm to provide the metamorphs with both water and a dry surface, and covered the container with a transparent, perforated lid. When metamorphosis was completed (complete tail resorption; development stage 46), we placed the individual into a new, lidded container of the same size as before, equipped with wet paper towel lining and a piece of cardboard egg-holder as a shelter. Twice a week, we fed froglets *ad libitum* with small crickets (*Acheta domestica*, instar stage 1-2) sprinkled with a 3:1 mixture of Reptiland 76280 (Trixie Heimtierbedarf GmbH & Co. KG, Tarp, Germany) and Promotor 43 (Laboratorios Calier S.A., Barcelona, Spain) to provide the necessary vitamins, minerals and amino acids. Due to their smaller size, we fed toadlets with springtails (*Folsomia* sp.) in the first three weeks after metamorphosis, and switched to crickets afterwards. For each

individual we recorded the dates of starting metamorphosis, completion of tail resorption, and eventual mortality.

Between 6-8 (for agile frogs) or 9-12 (for common toads) weeks after metamorphosis (depending on species and development), when gonads became sufficiently differentiated and easy to observe (Ogielska and Kotusz 2004, Nemesházi et al. 2020), we measured body mass to the nearest 0.01 g and euthanized juvenile individuals in a water bath containing 6.6 g/L tricaine-methanesulfonate (MS-222) buffered to neutral pH with the same amount of Na_2HPO_4. We dissected the animals and examined the internal organs under an Olympus SZX12 stereomicroscope (Olympus Europa SE & Co. KG, Hamburg, Germany) at 16× magnification and assigned fat bodies into one of four ordinal categories based on their size: lacking, small, regular-sized, or large. We also categorised phenotypic sex as male (testes), female (ovaries) or uncertain (abnormally looking gonads). Because many animals' guts contained food remains, we cut out the entire digestive tract, measured its mass to the nearest 0.01 g, and subtracted it from the body mass of juveniles to obtain 'net body mass'. We removed both feet of euthanized agile frogs and stored them in 96 % ethanol until DNA analyses.

We extracted DNA from agile frog foot samples with Geneaid Genomic DNA Extraction Kit for animal tissue (Thermo Fisher Scientific, Waltham USA) following the manufacturer's protocol, except that digestion time was 2 hours. We used a recently developed molecular marker set for genetic sexing validated on agile frog populations in Hungary (Nemesházi et al. 2020). We first tested all froglets for the Rds3 marker (≥ 95 % sex linkage) applying high-resolution melting (HRM). We considered an individual to be concordant male or female if its Rds3 genotype was in accordance with its phenotypic sex. Individuals that appeared to be sex-reversed based on the Rds3 marker were also tested using PCR for Rds1 (≥ 89 % sex linkage). For a

detailed description of HRM and PCR methods, see Nemesházi et al. (2020). When both markers congruently suggested sex reversal, we considered the given individuals to be sex-reversed. In case of contradiction between the results of analyses based on Rds1 and Rds3, we considered genetic sex to be unknown (Table S2). We did not investigate sex reversal in common toads because phenotypic sex ratios suggested no treatment effects on sex (see Results).

Statistical analyses

We analysed the data of the two species separately. We assessed treatment effects on survival, length of larval development, body mass at metamorphosis, net body mass at dissection, size of fat bodies, and phenotypic sex ratio. For each dependent variable, we ran a model (see model specifications below) with temperature and treatment period as categorical fixed factors and their interaction, the difference between the mean temperature in each tadpole container and the nominal temperature of the given treatment (measured as described in the electronic Supporting Information) as a numeric covariate, and family nested in population as random factors. We tested the effect of temperature within each treatment period by calculating pre-planned linear contrasts (Ruxton and Beauchamp 2008), correcting the significance threshold for multiple testing using the false discovery rate (FDR) method (Pike 2011). All analyses were conducted in 'R' (version 3.6.2), with the 'emmeans' package for linear contrasts.

For the analysis of survival, we used Cox's proportional hazards model (R package 'coxme'). Individuals were divided into five ordered categories; 1: died during treatment, 2: died after treatment, but before the start of metamorphosis, 3: died during metamorphosis, 4: died after metamorphosis, but before dissection, 5: survived until dissection. Animals that died before the treatment (four agile frog and five common toad larvae) were excluded from survival

263 analyses. We entered the ordinal survival categories as the dependent variable and treated the
264 fifth survival category as censored observations.

265 To analyse variation in the length of larval development, body mass at metamorphosis and
266 net body mass at dissection, we used linear mixed-effects models (LMM; 'lme' function of the
267 'nlme' package), allowing the variances to differ among treatment groups ('varIdent' function)
268 because graphical model diagnostics indicated heterogeneous variances. In the analysis of net
269 body mass at dissection, we included age (number of days from finishing metamorphosis to
270 dissection) as a further covariate. In the case of agile frogs, we entered the log-transformed
271 values of the length of larval development to achieve normal distribution of model residuals. For
272 the analysis of fat-body size, we applied cumulative link mixed models (CLMM; 'clmm'
273 function of 'ordinal' package; Christensen 2015), where we also entered age as a covariate.

274 To analyse phenotypic sex ratio, first, we excluded those few individuals the gonads of
275 which were not unambiguously categorizable either as male or female (Table S2). Then we
276 analysed the proportion of phenotypic males using phenotypic sex as a binary response variable
277 in generalised linear mixed modelling procedures (GLMM) with binomial error distribution and
278 logit link ('glmmTMB' function of the 'glmmTMB' package; Brooks et al. 2017). To analyse
279 sex reversal in agile frogs, we could not apply the same modelling framework as for sex ratios
280 because of separation, i.e. sex-reversed individuals were absent in certain treatment groups
281 whereas in some others there was 100% sex reversal. Therefore, we applied six separate analyses
282 comparing the two elevated temperature treatments to their associated controls in each of the
283 three ontogenetic periods using Fisher's exact tests. The dependent variable was phenotype, i.e.
284 whether or not the individual was sex-reversed. We restricted these analyses to genetic females

since heat induces female-to-male sex reversal, and we detected no male-to-female sex reversal. Because of multiple testing, we corrected *P* values using the FDR method.

Results

Survival of agile frogs that were exposed to 30 °C during either the 1st or the 2nd larval period was significantly reduced (by 56 and 17 %, respectively; Fig. 2, Table 1, Table S3). Survival of common toads also significantly decreased upon exposure to 30 °C (by ca. 33 %), but only if this temperature was applied during the 2nd larval period (Fig.2, Table 2). Thermal treatments that exposed tadpoles to 30 °C in other larval periods (3rd in both species and 1st in common toads) and those involving 28 °C at any period did not affect survival in either species (Table 1-2).

Length of larval development of agile frogs was significantly prolonged by all thermal treatments applied in all larval periods (Fig. 3, Table 1 and S3). By contrast, in common toads, the length of larval development was not affected when tadpoles were exposed to 28 °C during the 1st larval period, but larvae that were exposed to this temperature during the 2nd and 3rd larval period developed faster compared to their control groups (Fig. 4, Table 2 and S3). When common toad tadpoles were exposed to 30 °C, their larval development was only shortened upon exposure during the 3rd larval period but remained unaffected if treated in the 1st or 2nd larval period (Fig 4, Table 2 and S3).

Body mass at metamorphosis was significantly reduced in agile frogs by the 28 °C thermal treatment if applied during the 1st larval period but was not affected if 28 °C was applied later on (Fig 3, Table 1). Exposure to 30 °C tended to decrease body mass at metamorphosis when applied in the 1st larval period and exerted a significant negative effect during the 2nd and 3rd larval period (Fig 3, Table 1 and S3). In common toads, both thermal treatments applied in all

larval periods resulted in significantly reduced body mass at metamorphosis (Fig 4, Table 2 and S3).

At dissection, net body mass of juvenile agile frogs was only increased in animals treated with 30 °C during the 2nd larval period, but remained unaffected in all other treatment groups (Fig 3, Table 1 and S3). Thermal treatments applied in any larval period did not affect the net body mass of common toads (Fig 4, Table 2 and S3). The number of days between metamorphosis and dissection positively affected net body mass at dissection in both species (Table S3).

The size of fat bodies was significantly smaller in juvenile agile frogs as a result of both thermal treatments, but only upon exposure during the 1st larval period and not during later periods (Fig 3, Table1). In juveniles of the common toad the size of fat bodies was unaffected by thermal treatments applied in any larval period (Fig 4, Table 2), and positively correlated with the age of juveniles (Table S3).

Phenotypic sex ratio in agile frogs was affected by exposure to elevated temperature: exposure to 28 °C during the 3rd larval period (but not in the earlier periods) caused a significant shift towards a male-biased sex ratio, and treatment with 30 °C in all larval periods resulted in highly male-biased sex ratios (Fig 3, Table 1, S2 and S3). Accordingly, the proportion of agile frog individuals that underwent heat-induced sex reversal was significantly higher (between 30 and 100 % of genetic females) at both temperatures and in all treatment periods compared to the respective control groups ($\leq$ 4.5 %, all $P \leq 0.012$; Fig 3, Table S2). In contrast, none of the thermal treatments applied in either larval period had any effect on the phenotypic sex ratio of juvenile common toads (Fig 4, Table 2, S2 and S3).

Discussion

Our results demonstrate that high temperatures experienced for six days during larval development can negatively affect the survival, growth, somatic and sexual development of amphibians, but the severity of these effects depends on the intensity and timing of thermal stress and can largely differ between species. Agile frogs proved to be more sensitive: in this species, all studied variables were affected by one or more heat treatments, and almost all of the resulting changes are likely disadvantageous for individual fitness and population viability (Fig. 5). In contrast, for common toads, the only consistent effect of thermal stress was reduced mass at metamorphosis and, in a few treatments, faster larval development, while we observed barely any effect on survival and no lasting developmental effects in juveniles (Fig. 5). These results highlight that even sympatric species that are relatively similar in their ecology may be affected very differently by heat waves.

Survival rate in both species was decreased by exposure to 30 °C, but only if tadpoles experienced it relatively early on during their development (during 1st and 2nd larval periods). Temperatures of around 30 °C throughout the entire larval development often resulted in decreased survival in earlier studies (Bellakhal et al. 2014, Goldstein et al. 2017, Phuge 2017, Lambert et al. 2018). Our results suggest that the adverse effect of elevated temperature on larval survival depends on the species and on the timing of exposure, indicating a peak in thermosenitivity during the early stages of larval development (in addition to the increased thermosensitivity of the final larval stages, directly before the onset of metamorphosis (Floyd 1983), which we did not study). This is in line with many previous studies suggesting that the earliest life stages of amphibians are the most susceptible to several stress factors such as chemicals, parasites, poor environmental conditions and pesticides (Ortiz-Santaliestra et al. 2006,

354 Holland et al. 2007, Crespi and Warne 2013, Mikó et al. 2017). The energetically costly cellular
355 repairing mechanisms and the maintenance and restoration of homeostasis during and after
356 thermal stress compromise higher-level functions that are necessary for survival (Williams et al.
357 2016). Furthermore, dissolved oxygen level in the water decreases with rising temperature
358 (Stefan et al. 2001, Fang and Stefan 2009), which in turn can cause hypoxia and oxidative stress
359 in tadpoles (Lushchak 2011, Freitas and Almeida 2016). High temperature may also accelerate
360 bacterial bloom in the water (Ferreira and Chauvet 2011), potentiating the accumulation of
361 opportunistic pathogens. All of these processes might contribute to mortality observed in
362 experiments involving thermal treatments and, under natural conditions, during or after heat
363 waves.

364 Timing of metamorphosis and body mass at metamorphosis are crucial components of
365 fitness in amphibians. Earlier metamorphosis allows for leaving the more hazardous aquatic
366 environment faster (Denver 1997), and allows for a longer post-metamorphic growth period
367 compared to late-metamorphosing individuals, which in turn leads to increased survival during
368 the first hibernation (Altwegg and Reyer 2003, Üveges et al. 2016). In the present study, the
369 simulated heat waves prolonged larval development in agile frogs but shortened it (when heat
370 was experienced in the late larval period) in common toads, whereas mass at metamorphosis
371 decreased after heat exposure in both species (although in agile frogs the latter effect was only
372 significant in a few treatment groups). According to the temperature-size rule (Kozłowski et al.
373 2004), high temperatures are associated with increased metabolic rates and accelerated
374 development in larval anurans (Álvarez and Nicieza 2002, McLeod et al. 2013, Courtney Jones
375 et al. 2015), which results in earlier metamorphosis at a smaller body size (Laugen et al. 2003,
376 Niehaus et al. 2006). Our results likely documented this relationship between development and

377 growth in common toads. However, in agile frogs, this relationship was disrupted by heat
378 treatments, most probably because the applied temperatures acted as severe stressors. This result
379 aligns with the observation that larvae of the common toad are more thermophilic than those of
380 agile frogs, as suggested by a higher critical thermal maximum and higher preferred temperatures
381 in the former than in the latter (Hettyey, personal communication).

382 Stress experienced early in life can have long-lasting consequences, such as small adult
383 size and limited energy reserves (Crespi and Warne 2013, Jonsson and Jonsson 2014). However,
384 in our study, the reduced mass at metamorphosis in heat-treated groups did not persist into
385 juvenility: after a few months of post-metamorphic growth, we found no differences in body
386 mass or fat reserves in either species. There were only two exceptions to this: in juvenile agile
387 frogs, fat bodies were smaller if they received either heat treatment in the 1st larval period, and
388 unexpectedly, their body mass was larger after exposure to 30 °C applied during the 2nd larval
389 period. The death of lighter individuals likely contributed to the equalization of juvenile body
390 mass among treatment groups, given that most individuals that died between the onset of
391 metamorphosis and dissection had a lower body mass at metamorphosis than conspecifics that
392 survived until the end of the experiment in both species (Welch's tests; agile frogs: t = -3.54, df =
393 32.0, P = 0.001, common toads: t = -9.30, df = 53.9, P < 0.001). A further contributing factor
394 may be compensatory growth (Squires et al. 2010, Hector et al. 2012). Nonetheless,
395 compensatory growth can have hidden costs (Stoks et al. 2006, De Block and Stoks 2008,
396 Murillo-Rincón et al. 2017), so that the lack of among-treatment differences in juvenile mass
397 does not necessarily indicate the absence of long-term malign consequences of high temperatures
398 experienced during larval life. Indeed, the majority of juvenile agile frogs completely lacked fat
399 bodies if they were exposed to heat during the 1st larval period. Fat bodies in amphibians are

major energy stores that are vital to survival (Scott et al. 2007) and regulate processes related to reproduction (Pierantoni et al. 1983, Girish and Saidapur 2000). Consequently, high temperatures experienced during early ontogeny may have long-lasting negative effects on the survival and reproductive potential of agile frogs, which may compromise population persistence. The observation that the size of fat bodies was not affected by thermal treatments in common toads confirms that these are more tolerant to high temperatures than agile frogs, and, more generally, reinforces the hypothesis that there is large among-species variation also in the long-term consequences of thermal stress.

Sex reversal can occur naturally in wild populations of agile frogs (Nemesházi et al. 2020) and other species (Alho et al. 2010, Lambert et al. 2019, Xu et al. 2021), but high temperature can increase its frequency in a wide range of ectothermic vertebrates (Baroiller and D'Cotta 2016, Ruiz-Garciá et al. 2021, Whiteley et al. 2021). In our study, six-day 30 °C heat waves caused male-biased sex ratios via sex reversal in agile frogs, and the same effect was induced by exposure to 28 °C in the 3rd larval period. These results align with previous studies documenting altered sex ratios in several anuran species where larvae were raised at high temperatures throughout their development (Ruiz-Garciá et al. 2021), and additionally suggest that the sensitivity of sex determination to elevated temperature increases close to the end of larval development. Our findings caution that heat waves lasting for only a few days during tadpole development can trigger sex reversal, which may have wide-ranging consequences including skewed sex ratios and lowered population viability (Bókony et al. 2017, Wedekind 2017, Nemesházi et al. 2021). However, our observation that the same thermal treatments did not affect phenotypic sex ratios in common toads suggests that there is considerable interspecific variation in the thermosensitivity of sexual development.

Heat treatment is a promising mitigation method against amphibian chytridiomycosis (Chatfield and Richards-Zawacki 2011, Geiger et al. 2011, Hettyey et al. 2019). Our results, however, underline the importance of pre-assessing the thermal sensitivity of each species, including that of their sexual development. Based on our results, thermal treatment at 30 °C could be applied for six days to common toads, which would likely lead to *Bd* clearance, or at least to a drastic suppression of *Bd* growth (Retallick and Miera 2007, Chatfield and Richards-Zawacki 2011, Geiger et al. 2011). This treatment could be recommended in specific situations, such as epizootic outbreaks, when the benefits clearly outweigh the costs arising from decreased body mass at metamorphosis, or when the latter can be compensated for (e.g. by supplemental feeding). In agile frogs, treatment with 28 °C during the 2nd larval period (days 12-18 after hatching) was the only treatment combination without adverse effects on most life-history traits and sexual development. Although this treatment also caused somewhat lengthened larval development, this cost may be negligible (especially so in captivity) considering the benefit of Bd clearance. Whether treatment with temperatures lower than 28 °C would be applicable without costs and still suppresses *Bd* growth sufficiently needs further investigation (Hettyey et al. 2019). A further possibility to explore is that under controlled conditions, capitalising on the feminizing effect of estrogens or other estrogenic chemicals might make thermal treatment of *Bd*-infected animals potentially suitable also for species with thermally sensitive sex determination (Kitano et al. 2012).

In conclusion, our study demonstrates that species can differ in a multitude of ways in how they are affected by short periods of elevated temperatures which are similar in magnitude to those occurring in natural water bodies during heat waves. Most importantly, we demonstrate that already 28 °C can have surprisingly severe consequences for larvae of a thermosensitive

446 anuran, where the strength of effects depends largely on the developmental stage of individuals
447 that become exposed to the heat. At the same time, even 30 °C experienced any time during
448 larval development does little harm to individuals of another sympatric species. Such species-
449 specific differences should be examined in a wide range of taxa and considered when evaluating
450 the impact of climate change on amphibians, and also in the development of mitigating methods
451 against chytridiomycosis.
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Data availability statement - The data that support the findings of this study are openly available in figshare at http://doi.org/ 10.6084/m9.figshare.17197847.

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Table 1: Agile frog responses to heat by the timing of exposure (1st, 2nd and 3rd larval period) and the applied temperature. Results represent pre-planned comparisons from the models shown in Table S2, comparing each period and temperature combination to the 19°C treatment in the corresponding period. Linear contrasts (*c*), associated standard errors (SE), t-values (z-values in case of Cox's proportional hazards model in the analyses of survival) and *P*-values adjusted using the FDR method are reported. Treatment groups that differed significantly (P < 0.05) from their corresponding controls are highlighted in bold.

Dependent variable	Period	Temperature (°C)	*c*	SE	*t* (or *z*)	*P*
Survival*	1	28	0.31	0.51	0.62	0.610
	2	28	-0.47	0.91	-0.51	0.610
	3	28	-0.43	0.65	-0.67	0.610
	1	**30**	**2.11**	**0.43**	**4.85**	**< 0.001**
	2	**30**	**1.56**	**0.65**	**2.41**	**0.048**
	3	30	0.60	0.52	1.16	0.490
Length of larval development (log(days))	**1**	**28**	**0.14**	**0.03**	**4.38**	**< 0.001**
	2	**28**	**0.05**	**0.01**	**4.58**	**< 0.001**
	3	**28**	**0.05**	**0.02**	**2.57**	**0.011**
	1	**30**	**0.24**	**0.04**	**6.24**	**< 0.001**
	2	**30**	**0.19**	**0.02**	**8.29**	**< 0.001**
	3	**30**	**0.14**	**0.02**	**9.12**	**< 0.001**
Body mass at metamorphosis (mg)	**1**	**28**	**-52.53**	**19.20**	**-2.74**	**0.020**
	2	28	7.75	11.20	0.69	0.490
	3	28	-23.12	13.50	-1.71	0.106
	1	30	-51.06	27.00	-1.89	0.089
	2	**30**	**-32.23**	**13.60**	**-2.37**	**0.036**
	3	**30**	**-34.01**	**12.30**	**-2.77**	**0.020**
Net body mass at dissection (g)	1	28	0.03	0.03	1.01	0.374
	2	28	0.07	0.03	2.05	0.123
	3	28	-0.07	0.03	-1.76	0.127
	1	30	0.09	0.05	1.73	0.127
	2	**30**	**0.13**	**0.03**	**3.97**	**< 0.001**
	3	30	-0.02	0.03	-0.63	0.528
Size of fat bodies**	**1**	**28**	**-1.76**	**0.47**	**-3.78**	**< 0.001**
	2	28	0.21	0.38	0.54	0.705
	3	28	0.47	0.41	1.15	0.375
	1	**30**	**-1.86**	**0.56**	**-3.31**	**0.003**

	2	30	-0.73	0.42	-1.71	0.175
	3	30	0.06	0.43	0.14	0.889
Phenotypic sex ratio (proportion of males)***	1	28	-0.22	0.47	-0.47	0.640
	2	28	0.72	0.45	1.62	0.127
	3	**28**	**3.83**	**1.06**	**3.60**	**0.002**
	1	**30**	**1.63**	**0.71**	**2.30**	**0.034**
	2	**30**	**1.54**	**0.55**	**2.80**	**0.011**
	3	**30**	**3.70**	**1.06**	**3.47**	**0.002**

*The linear contrast is the log(hazard ratio)

**The linear contrast is the log(cumulative odds ratio)

***The linear contrast is the log(odds ratio)

730

Table 2: Common toad responses to heat by the timing of exposure (1st, 2nd and 3rd larval period) and the applied temperature. Results represent pre-planned comparisons from the models shown in Table S2, comparing each period and temperature combination to the 19°C treatment in the corresponding period. Linear contrasts (*c*), associated standard errors (SE), t-values (z-values in case of Cox's proportional hazards model in the analyses of survival) and *P*-values adjusted using the FDR method are reported. Treatment groups that differed significantly ($P < 0.05$) from their corresponding controls are highlighted in bold.

Dependent variable	Period	Temperature (°C)	*c*	SE	*t* (or *z*)	*P*
Survival*	1	28	-0.06	0.40	-0.15	0.882
	2	28	0.88	0.60	1.46	0.433
	3	28	-0.21	0.56	-0.38	0.882
	1	30	-0.09	0.42	-0.23	0.882
	2	**30**	**1.51**	**0.57**	**2.66**	**0.047**
	3	30	0.12	0.53	0.23	0.882
Length of larval development (days)	1	28	0.03	1.55	0.02	0.986
	2	**28**	**-7.24**	**0.59**	**-12.21**	**< 0.001**
	3	**28**	**-7.11**	**0.81**	**-8.73**	**< 0.001**
	1	30	-0.48	0.96	-0.50	0.742
	2	30	-1.28	0.95	-1.36	0.263
	3	**30**	**-3.24**	**0.87**	**-3.76**	**< 0.001**
Body mass at metamorphosis (mg)	**1**	**28**	**-63.80**	**8.04**	**-7.93**	**< 0.001**
	2	**28**	**-63.20**	**7.89**	**-8.01**	**< 0.001**
	3	**28**	**-73.00**	**7.36**	**-9.92**	**< 0.001**
	1	**30**	**-54.00**	**7.29**	**-7.40**	**< 0.001**
	2	**30**	**-76.30**	**9.30**	**-8.02**	**< 0.001**
	3	**30**	**-73.50**	**9.32**	**-7.88**	**< 0.001**
Net body mass at dissection (g)	1	28	-0.39	0.17	-2.27	0.144
	2	28	-0.01	0.15	-0.06	0.950
	3	28	-0.02	0.12	-0.14	0.950
	1	30	-0.28	0.15	-1.95	0.156
	2	30	-0.29	0.20	-1.47	0.288
	3	30	0.11	0.13	0.83	0.615
Size of fat bodies**	1	28	-0.11	0.47	-0.23	0.941
	2	28	0.03	0.47	0.07	0.941
	3	28	-0.57	0.47	-1.21	0.675

	1	30	-0.09	0.47	-0.20	0.941
	2	30	-0.19	0.48	-0.39	0.941
	3	30	-0.97	0.46	-2.11	0.211
Phenotypic sex ratio (proportion of males)***	1	28	-0.67	0.48	-1.38	0.377
	2	28	-0.46	0.45	-1.01	0.377
	3	28	0.76	0.46	1.65	0.377
	1	30	-0.58	0.49	-1.19	0.377
	2	30	-0.23	0.48	-0.50	0.621
	3	30	0.55	0.49	1.13	0.377

*The linear contrast is the log(hazard ratio)

**The linear contrast is the log(cumulative odds ratio)

***The linear contrast is the log(odds ratio)

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741

742 Figure Legends

743 Figure 1: A schematic illustration of experimental treatments. Each horizontal bar represents one
744 treatment group. Striped bars represent periods when tadpoles were exposed to thermal
745 treatments. Orange (28 °C) and red (30 °C) bars symbolize heat treatments, while blue filling
746 represents maintenance at 19 °C. Treatments were identical in both species.

747

748 Figure 2: Survival of agile frogs (a) and common toads (b) during the experiment over time in
749 each treatment group. Blue lines represent controls maintained at 19 °C throughout, orange lines
750 represent treatment groups exposed to 28 °C, red lines represent treatment groups exposed to 30
751 °C; dotted lines represent individuals exposed to thermal treatments during the 1st larval period,
752 dashed lines those exposed during the 2nd larval period, and solid lines those exposed during the
753 3rd larval period. Numbered vertical lanes depict the respective larval periods when thermal
754 treatments were performed. Note that the experiment lasted longer for common toads than for
755 agile frogs.

756

757 Figure 3: Agile frog responses to thermal treatments in terms of the length of larval development,
758 body mass at metamorphosis, net body mass at dissection, the size of fat bodies and phenotypic
759 sex ratios in juveniles. In boxplots, horizontal lines and boxes represent medians and
760 interquartile ranges (IQR), respectively, while whiskers extend to IQR ± 1.5×IQR and dots
761 indicate more extreme data points.

762

763 Figure 4: Common toad responses to thermal treatments in terms of the length of larval
764 development, body mass at metamorphosis, net body mass at dissection, the size of fat bodies

765 and phenotypic sex ratios in juveniles. In boxplots, horizontal lines and boxes represent medians
766 and interquartile ranges (IQR), respectively, while whiskers extend to IQR ± 1.5×IQR and dots
767 indicate more extreme data points.
768
769 Figure 5: Summary of responses by the two species to the simulated heat waves in different
770 larval periods. Arrows show the direction of the observed change in the given variable relative to
771 the respective control group, not the advantageousness or harmfulness of the effect. Separation is
772 aided by different colours.

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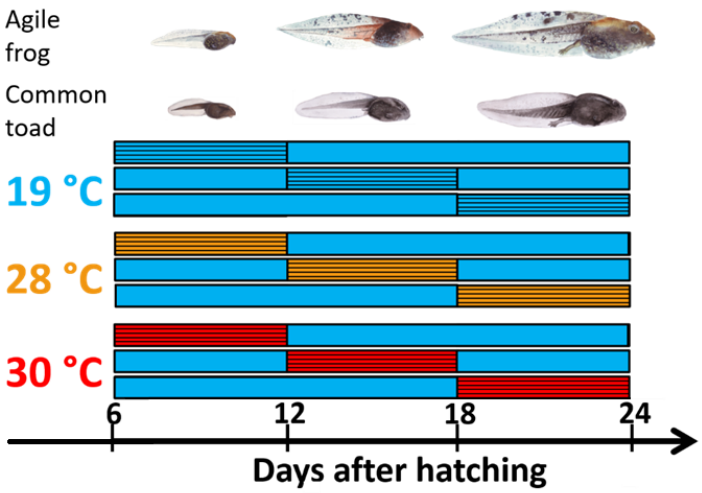
774 Ujszegi et al., Fig. 1



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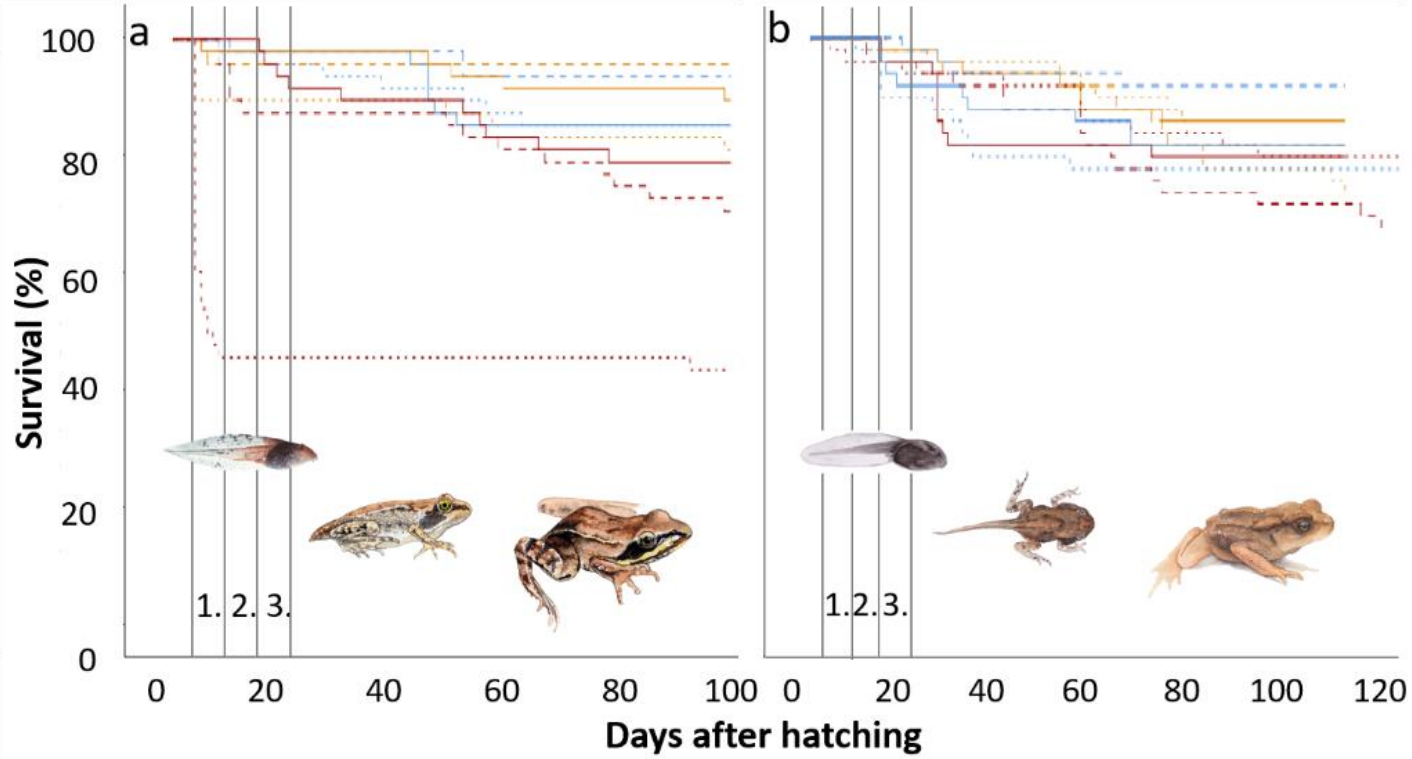
778 Ujszegi et al., Fig. 2



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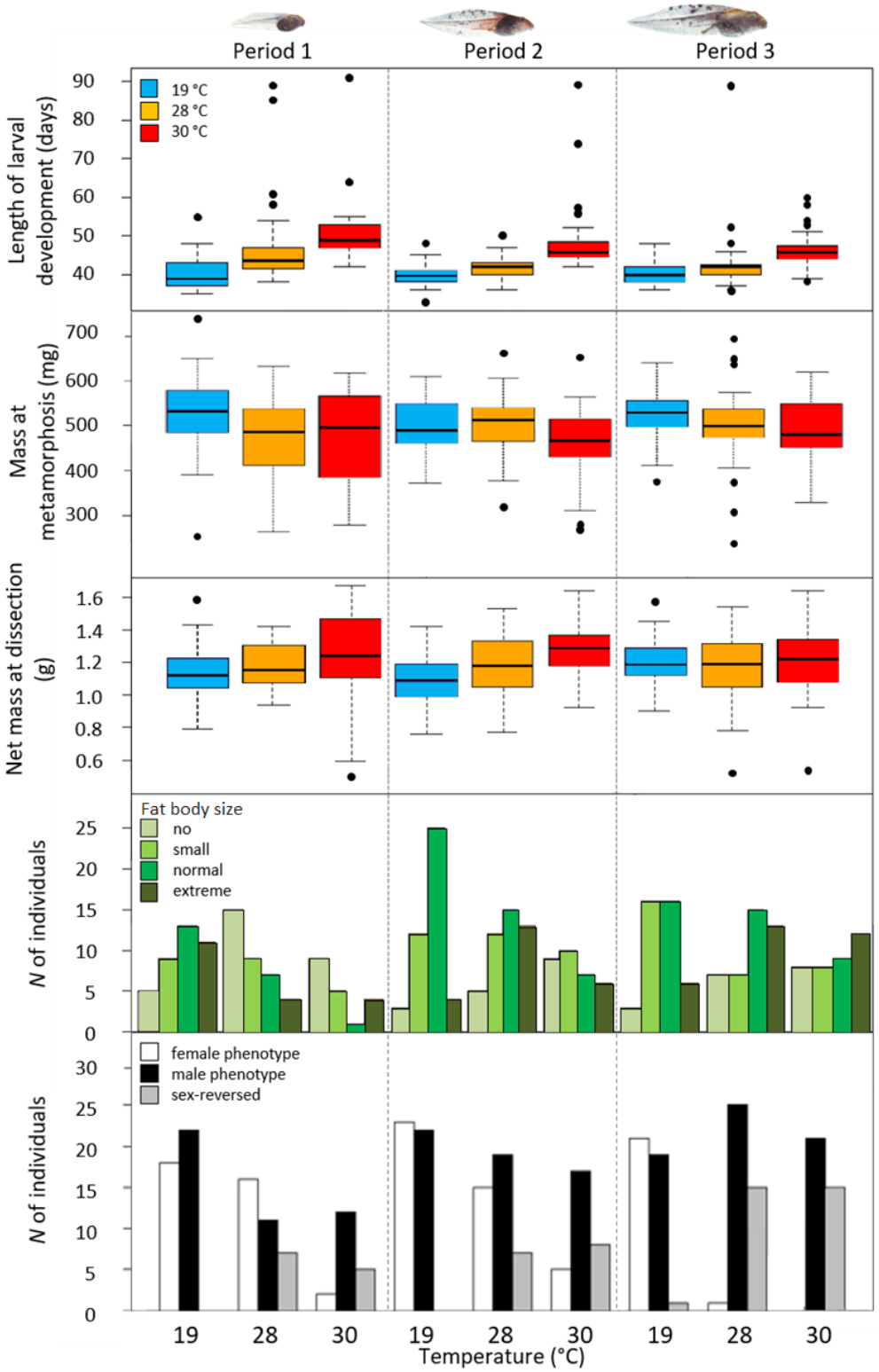
782 Ujszegi et al., Fig. 3



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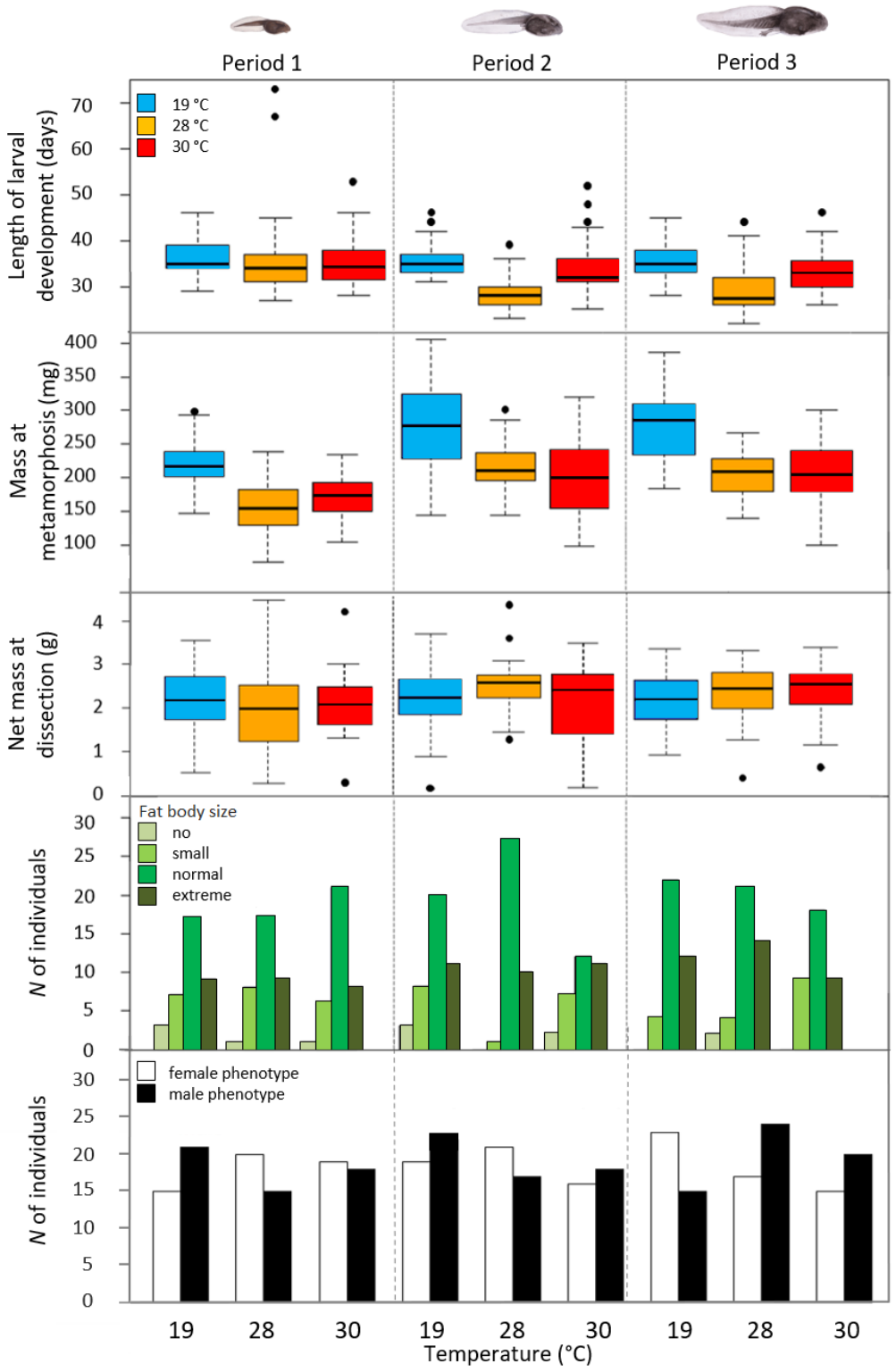
785 Ujszegi et al., Fig. 4



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788 Ujszegi et al., Fig. 5

Variables \ Treatment combinations per species	 Agile frog (*Rana dalmatina*)						 Common toad (*Bufo bufo*)					
	Period 1		Period 2		Period 3		Period 1		Period 2		Period 3	
	28 °C	30 °C	28 °C	30 °C	28 °C	30 °C	28 °C	30 °C	28 °C	30 °C	28 °C	30 °C
Survival		↓		↓						↓		
Length of larval development	↑	↑	↑	↑	↑	↑			↓		↓	↓
Body mass at metamorphosis	↓			↓		↓	↓	↓	↓	↓	↓	↓
Net body mass at dissection				↑								
Fat body size	↓	↓										
Ratio of phenotypic males		↑		↑	↑	↑						

789

Supporting Information to:

"Heat waves" experienced during larval life have species-specific consequences on life-history traits and sexual development in anuran amphibians

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Measurements validating temperature in heat treatments

We validated the heating setup by repeatedly measuring water temperature (± 0.1 °C) in tadpole containers of each position in each tray, as well as water temperature in the trays in which treatments took place. Before the experiment, we measured these temperatures ten times on two consecutive days with a Greisinger digital thermometer (GTH175/PT). After termination of the experiment, we repeated these measurements five times. To detect eventual temperature fluctuations during each treatment, twice per day we checked water temperature in all trays using the digital thermometer. Furthermore, data loggers (Onset HOBO Pendant Temperature/Light 8K Data Logger; one per each tray) recorded temperature in the trays every 30 minutes during the treatments. We did not measure temperature in the tadpole containers during the treatment periods in order to avoid stress and injury as a result of stirring the water.

We did not detect considerable temperature fluctuations during the treatments (Fig S1, Table S1), and temperature readings were very similar before and after the experiment in each container position. Temperature did vary somewhat among containers in different positions within trays (maximal temperature difference within a tray at 19 °C: 1.3 °C; at 28 °C: 1.5 °C; at 30 °C: 1.5 °C), but this variation was highly consistent over time. We calculated the difference between the actual (experienced by the tadpoles) and nominal temperature of the given treatment for each tadpole container by subtracting the nominal temperature from the mean water temperature (measured before and after the treatments in each container with the digital thermometer). This method minimised the disturbance caused to animals during the experiment while delivering accurate data on the temperatures experienced by the tadpoles.

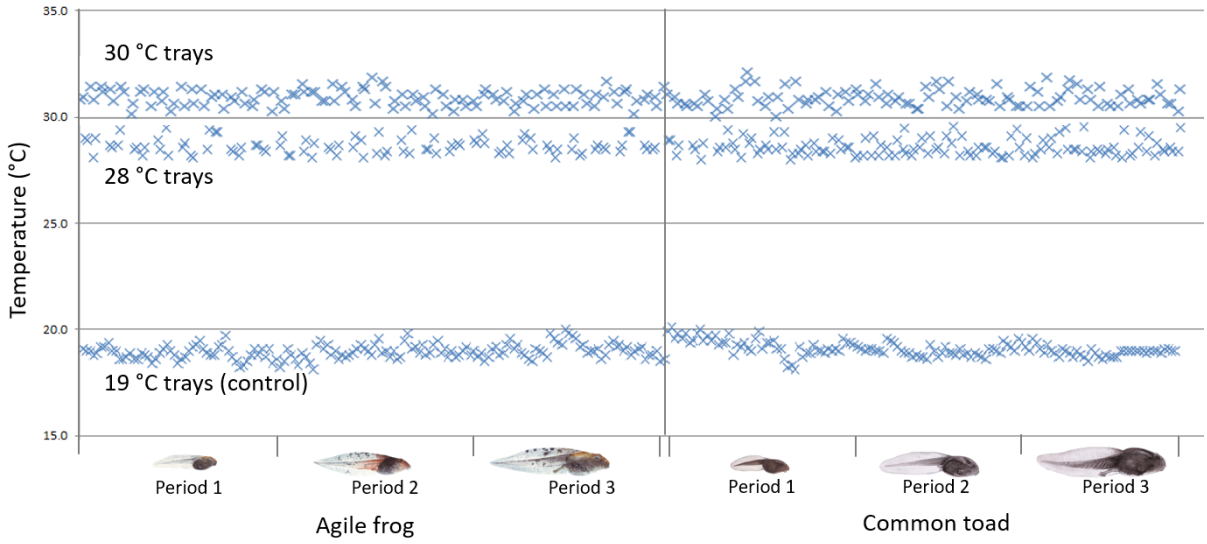

Figure S1 Water temperatures in the trays during treatment periods show minimal temperature fluctuations. Note that the water temperature in the heated trays was always warmer than the temperature in the tadpoles' containers (set to be as close to the nominal temperature as possible; Table S1).

Table S1 Minimal (T_{min}), maximal (T_{max}) and mean (T_{mean}) temperatures in the heated trays during temperature treatments. Mean diff. represents the average difference in water temperature between the tadpoles' containers and the trays, since water temperature in the heated trays was always warmer than the temperature in the tadpoles' containers.

Tray	T_{min} (°C)	T_{max} (°C)	T_{mean} (°C)	Mean diff. (± SE)
30°C /1	30.6	31.6	31.0	1.1 (± 0.13)
30°C /2	30.2	31.6	30.8	1.1 (± 0.10)
30°C /3	30.4	31.6	30.9	0.5 (± 0.09)
30°C /4	30.5	31.3	30.8	0.7 (± 0.09)
28°C /1	28.2	28.9	28.5	0.4 (± 0.06)
28°C /2	28.4	29.4	28.8	0.1 (± 0.07)
28°C /3	28.0	29.5	28.5	0.5 (± 0.06)
28°C /4	28.1	29.5	28.6	1.1 (± 0.08)

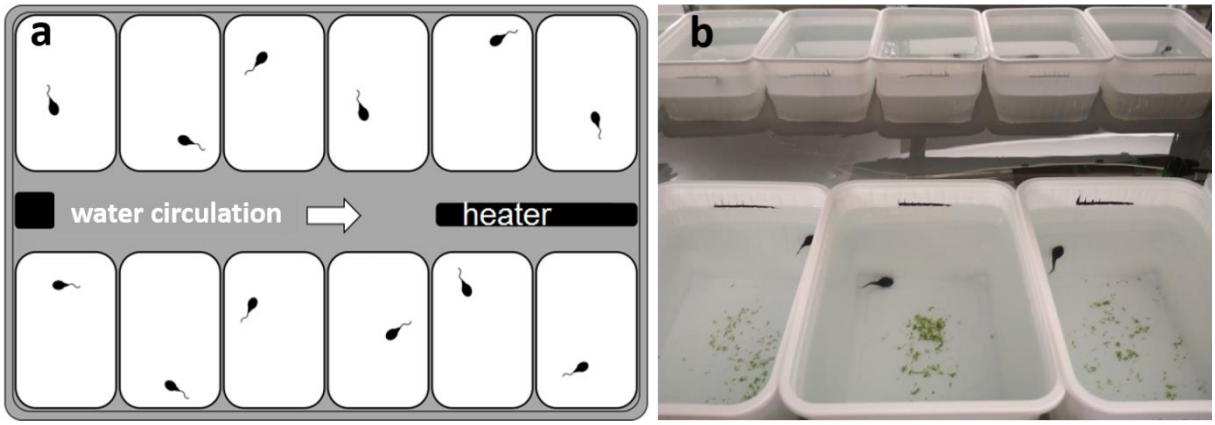

Figure S2 Schematic representation (a) and *in situ* photograph (b) of the heating system used in the thermal treatments.

Table S2 Phenotypic sex ratio (% males) in each treatment group. In case of agile frogs, genetic sex ratio and female-to-male sex-reversal rate (% of phenotypic males in genetic females) are also shown.

Species	Treatment group	Dissected (*N*)	Phenotypic sex ratio (% male)†	Genetic sex ratio (% male)‡	Female-to-male sex reversal (%)
Agile frog	Period 1				
	Control	40	55.0	55.0	0.0
	28 °C	39	52.9	32.4	30.4
	30 °C	19	89.5*	63.2	71.4
	Period 2				
	Control	45	48.9	48.9	0.0
	28 °C	46	63.4	46.3	31.8
	30 °C	32	83.3*	56.6	61.5
	Period 3				
	Control	41	48.8	46.3	4.5
	28 °C	43	97.5*	60.9	93.8
	30 °C	37	100.0*	58.3	100.0
Common toad	Period 1				
	Control	36	58.3		
	28 °C	35	42.9		
	30 °C	37	48.6		
	Period 2				
	Control	42	54.8		
	28 °C	39	43.6		
	30 °C	34	52.9		
	Period 3				
	Control	38	39.5		
	28 °C	41	58.5		
	30 °C	36	55.6		

*Sex ratios that differ significantly from 1:1 according to Fisher's exact tests

†Excluding those individuals whose gonads were not unambiguously categorizable either as male or female (3 agile frogs: 2 individuals at 28 °C in the 2nd larval period and 1 individual at 30 °C in the 2nd larval period; 2 common toads: 1 individual at 28 °C in the 2nd larval period and 1 individual at 30 °C in the 3rd larval period)

‡Excluding those individuals whose genetic sex was unknown due to contradiction between the Rds1 and Rds3 markers' results (at 28 °C 1 individual in the 1st, and 2 individuals in the 3rd larval period; at 30 °C 1 individual each in the 2nd and 3rd larval periods)

Table S3 Type-2 analysis-of-deviance tables of the statistical models. Significant effects ($P < 0.05$) are highlighted in bold. The covariate "T.diff" is the difference between mean temperatures in each tadpole container and the nominal temperature of the given treatment.

Dependent variable	Predictors	Agile frog			Common toad		
		χ^2	df	*P*	χ^2	df	*P*
Survival							
	Heat	40.90	2	< **0.001**	2.15	2	0.340
	Period	28.71	2	< **0.001**	4.71	2	0.095
	T.diff	0.36	1	0.551	0.08	1	0.783
	Heat×Period	5.48	4	0.241	5.93	4	0.204
Length of larval development							
	Heat	195.34	2	< **0.001**	208.73	2	< **0.001**
	Period	9.04	2	**0.011**	7.19	2	**0.028**
	T.diff	0.05	1	0.817	0.36	1	0.548
	Heat×Period	12.03	4	**0.017**	23.78	4	< **0.001**
Body mass at metamorphosis							
	Heat	17.44	2	< **0.001**	250.50	2	< **0.001**
	Period	6.88	2	< **0.001**	145.80	2	< **0.001**
	T.diff	0.02	1	0.988	2.66	1	0.103
	Heat×Period	8.67	4	0.070	7.18	4	0.127
Net body mass at dissection							
	Heat	7.84	2	**0.020**	1.83	2	0.399
	Period	1.43	2	0.488	4.08	2	**0.130**
	T.diff	0.23	1	0.631	0.75	1	0.386
	Age at dissection	80.86	1	< **0.001**	11.38	1	< **0.001**
	Heat×Period	14.71	4	< **0.001**	8.29	4	0.081
Size of fat bodies							
	Heat	7.20	2	**0.027**	2.18	2	0.337
	Period	7.03	2	**0.030**	1.13	2	0.567
	T.diff	0.07	1	0.790	2.46	1	0.117
	Age at dissection	2.56	1	0.109	6.87	1	**0.009**
	Heat×Period	17.87	4	**0.001**	2.60	4	0.627
Phenotypic sex ratio							
	Heat	22.53	2	< **0.001**	0.18	2	0.916
	Period	2.17	2	0.338	0.10	2	0.953
	T.diff	0.02	1	0.964	3.32	1	0.068
	Heat×Period	15.18	4	**0.004**	5.92	4	0.205