



Challenges of a novel range: Water balance, stress, and immunity in an invasive toad

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ARTICLE INFO

Keywords:

Invasive species
Dehydration
Corticosterone
Immunocompetence

ABSTRACT

Species introduced by human activities can alter the normal functioning of ecosystems promoting negative impacts on native biodiversity, as they can rapidly expand their population size, demonstrating phenotypic plasticity and possible adaptive capacity to novel environments. Twenty years ago, the guttural toad, *Sclerophrys gutturalis*, was introduced to a peri-urban area of Cape Town, with cooler and drier climatic characteristics than its native source population, Durban, South Africa. Our goal was to understand the phenotypic changes, in terms of physiology and immunity, of populations in native and novel environments. We evaluated body index (BI), field hydration level, plasma corticosterone levels (CORT), proportion of neutrophils: lymphocytes (N:L), plasma bacterial killing ability (BKA), and hematocrit (HTC) in the field, and after standardized stressors (dehydration and movement restriction) in males from the native and invasive populations. Toads from the invasive population presented lower BI and tended to show a lower field hydration state, which is consistent with living in the drier environmental conditions of Cape Town. Additionally, invasive toads also showed higher BKA and N:L ratio under field conditions. After exposure to stressors, invasive animals presented higher BKA than the natives. Individuals from both populations showed increased CORT after dehydration, an intense stressor for these animals. The highest BKA and N:L ratio in the field and after submission to stressors in the laboratory shows that the invasive population has a phenotype that might increase their fitness, leading to adaptive responses in the novel environment and, thus, favoring successful dispersion and population increase.

1. Introduction

Species introduced through human activities are present in many ecosystems and some of these species may become invasive and rapidly expand their populations (Torchin et al., 2003; Seebacher and Franklin, 2011). A high invasive potential is mainly due to their ability to produce integrated phenotypes reconciling pressures from many selective agents allowing for greater expansion efficiency (Reznick and Ghalambor, 2001; Lee, 2002; Sax et al., 2007; Jessop et al., 2013). Due to this rapid evolution and population increase, invasive species are currently recognized as one of the main threats to vertebrate populations (Mooney and Hobbs, 2000; Strauss et al., 2006) as they exploit the same resources used by local populations (Altieri et al., 2010), thus causing biodiversity losses (Pimentel, 2014). For this reason, invasive species provide an

excellent model for investigating how organisms mitigate the various challenges associated with rapid environmental change.

To understand how these species adapt to environmental changes in novel habitats, it is necessary to integrate information on physiological mechanisms and processes that regulate and integrate the various phenotypic characteristics optimizing the fitness of individuals (Wingfield, 2008; Chown et al., 2010; Martin et al., 2011; Cohen et al., 2012; Jessop et al., 2013). Some environmental challenges, as potential stressors, can activate the hypothalamic-pituitary-adrenal/interrenal (HPA/I) axis in vertebrates, resulting increased levels of plasma glucocorticoids (GCs), such as corticosterone (CORT). CORT is a metabolic hormone, and its secretion is increased in response to higher energy demands associated with environmental challenges, such as limited resources and/or divergent climatic conditions (Bonier et al., 2009).

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<https://doi.org/10.1016/j.cbpa.2020.110870>

Received 2 August 2020; Received in revised form 8 December 2020; Accepted 8 December 2020

Available online 13 December 2020

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Additionally, it can modulate several functions, such as reproduction, behavior, immunocompetence, and growth (Sapolsky et al., 2000; Dhabhar, 2014). The acute activation of the HPA/I axis results in an integrative and adaptive response to short-term stressors (Wingfield et al., 1997; Sapolsky et al., 2000; Romero and Wingfield, 2001; Sapolsky, 2002). For example, GCs are known to be immunomodulatory hormones and have bimodal effects on the immune system. Short-term increased GC secretion in response to acute stressors is associated with immunoenhancing effects, including increased presentation of antigens, effector cell function and production of antibodies and cytokines (Dhabhar and McEwen, 1996; Dhabhar, 1998; Dhabhar and McEwen, 1997; Madelaire et al., 2017). These immunoenhancing effects increase immune surveillance in the face of stressors, preparing the immune system for a rapid response to an immune challenge, such as injury or infection (Dhabhar et al., 1994, 1995, 1996; Dhabhar and McEwen, 1996; Dhabhar, 1998). On the other hand, long-term increase of GC secretion in response to chronic stressors is usually associated with immunosuppressive effects (Dhabhar and McEwen, 1997; Dhabhar, 2014; Titon et al., 2018), such as suppression of maturation, differentiation and proliferation of immune cells (Sterberg, 2006; Martin, 2009) in addition to triggering apoptosis in T and B cells (Sapolsky et al., 2000) and reduction of anti-inflammatory effects (Dhabhar, 2014). Additionally, exposure to chronic stressors can alter the ability of the HPI axis to respond to acute secondary stressors (Rich and Romero, 2005). In the wild, animals may be subjected to chronic and/or intense acute stressors, and additional stressors may lead to allostatic overload, affecting their ability to mount a rapid response and compromising their survival (Romero and Wingfield, 2015).

When compared with other tetrapods, amphibians have comparatively higher rates of transcutaneous evaporation (Toledo and Jared, 1993), and one of the challenges in a new environment may be the maintenance of adequate hydration levels. Dehydration affects locomotor performance and its underlying physiological determinants (e.g. Vimercati et al., 2018), which may impair ecologically relevant behaviors such as breeding, foraging, and escape from predators (Moore and Gatten Jr, 1989; Titon Jr and Gomes, 2017). Accordingly, dehydration represents a potent stressor for toads, activating the HPI axis and increasing CORT secretion (Barsotti et al., 2019), which may affect immune function, as observed in other ectothermic vertebrates (Moeller et al., 2013, 2017). Despite the general physiological characteristics assumed in the literature as potentially restrictive to the spread of originally mesic amphibians to more xeric novel environments, recent studies have described fast and integrated behavioral and physiological adaptations of invasive amphibians to unfavorable drier conditions (Tingley et al., 2012; Kosmala et al., 2017, 2020; Roznik et al., 2018). Cane toads (*Rhinella marina*) from the invasion front in the Australian desert, for example, show greater tolerance to dehydration and better locomotor performance when dehydrated than their native conspecifics (Kosmala et al., 2017). In addition, cane toads from drier areas in Australia show higher rates of water uptake than those from more mesic areas (Tingley et al., 2012). Moreover, the behavioral flexibility associated with shelter selection and adjustments of daily movement rates according to ambient moisture has been key for invasive cane toads to successfully colonize arid zones in Australia (Tingley and Shine, 2011).

Likewise, an invasive population of guttural toads (*Sclerophrys gutturalis*) has recently established (2000) in Cape Town, in a climate significantly drier and colder than that of the native source area of Durban, South Africa (Telford et al., 2019; Measey et al., 2020). Previous studies have shown that these invasive toads outperformed those from the native range in locomotor trials when dehydrated and were more efficient in minimizing rates of water loss through postural adjustments (Vimercati et al., 2018, 2019). However, the interaction between response to stress and functions modulated by this response, such as immunocompetence, has not yet been explored in this invasive population of the guttural toad. Despite an eradication program, the species is still spreading across the invaded area, in a small peri-urban area of Cape

Town (Measey et al., 2017; Davies et al., 2020). Therefore, it is important to study the adjustments in the physiological characteristics and life history of invasive guttural toads, in order to predict its potential for propagation within Cape Town. In addition, by comparing the invasive population with the native population, we can clarify information about the possible phenomena that may increase the fitness of this species in the invaded area.

In the present study, we aim to compare the stress response and its effects in immunocompetence of native and invasive populations of *S. gutturalis*. We hypothesize that invasive individuals of *S. gutturalis* are under chronic stress in Cape Town's drier environment, and thus show lower immunocompetence and a reduced ability to respond to an acute stressor when compared to individuals from native populations in the more humid region of Durban. To test our hypothesis, we compared blood samples collected in the field and after experimental manipulation in the laboratory of male individuals from invasive and native populations. The experiment consisted of dehydrating animals by 10% or 20%, followed by movement restriction for 24 h. Blood samples were collected before and after dehydration, and after restriction. We measured body index and hydration state in the field, hematocrit (HTC) as an estimate of hydration levels, corticosterone plasma levels (CORT) and neutrophil: lymphocyte (N:L) ratio as estimates of stress, and bacterial killing ability (BKA) as an estimate of immunocompetence. According to our hypotheses and experimental design employed we predicted that: 1) individuals of *S. gutturalis* from the invasive population will display higher CORT and N:L ratio, lower hydration state and lower BKA in the field when compared to the native population; 2) after submission to dehydration and movement restriction stress, toads from both populations will present increased CORT and N:L ratio, and lower BKA; 3) individuals from the native population will have higher CORT, N:L ratio and BKA after both dehydration and movement restriction.

2. Data collection

2.1. Study species, collection localities, and maintenance in captivity

Male *Sclerophrys gutturalis* were collected in Cape Town (87 m a.s.l., 34°01'S, 18°25'E) and Durban (75 m a.s.l., 29°47'S, 31°01'E), corresponding to invasive and native populations, respectively (see Telford et al., 2019).

In Cape Town, 25 invasive individuals were collected in the peri-urban Constantia region with prior permission from CapeNature (Permit No.: CN44-31-7259) and with the assistance of the Nature Conservation Corporation. The collections were carried out in private residences, where toads were found close to or immersed in bodies of water (e.g. fountains, ponds, and swimming pools: see Vimercati et al., 2017), between 17 and 27 January 2019 [Average temperature, dew point, and relative humidity for the collection days: 20.2 °C, 13.5 °C and 65.8%, respectively (Source: Weather Underground website, available at <https://www.wunderground.com/dashboard/pws/IWCAPECO2>)].

A total of 20 male *Sclerophrys gutturalis* individuals from the native population were collected in a similar peri-urban area of Durban North with prior permission from KZN Wildlife (Permit No.: 0P 4353/2018), between 16 and 23 February 2019 [Average temperature, dew point, and relative humidity for the collection days: 25.5 °C, 21.8 °C and 81.1%, respectively (Source: Weather Underground website, available at <https://www.wunderground.com/dashboard/pws/IKWAZULU12>)]. The collections were carried out on residential streets, some private residences, and a golf course.

The animals were collected at both sites during the reproductive period of the species (Vimercati et al., 2019). They were located by visual inspection, captured by hand and a blood sample was collected in the field (details below). Subsequently, animals were weighed using a portable balance (± 0.01 g, WTB 2000, Radwag, Radom, Poland) and placed individually in pots with water overnight to rehydrate. After rehydration, individuals had their bladders emptied by pressure in the

ventrum (Titon Jr and Gomes, 2017), and were then weighed again. Body mass of 100% hydrated individuals with empty bladders were considered as the standard mass (Ruibal, 1962; Titon Jr and Gomes, 2017). Field hydration state was calculated as the field mass divided by the standard mass after rehydration (Preest and Pough, 1989; Preest et al., 1992; Tracy et al., 2014). Animals of both populations were brought to Stellenbosch University and kept individually in plastic containers (43.0 (W) × 28.5 (L) × 26.5 (H) cm) with foliage, and water and crickets (*Acheta domesticus*) were provided ad libitum. The containers and water were checked daily, and the boxes cleaned when needed. Animals were kept in a climatic chamber with constant temperature (24 °C ± 1 °C), relative humidity (40% ± 1%) and 13 L:11D light regime. In order to allow for the recovery of blood volume, all individuals were kept in these conditions for at least one week before being submitted to stress by dehydration and movement restriction. After the experiments were finished, we measured animals' snout-vent length (SVL) in order to calculate the body index (BI), which was calculated as unstandardized residuals of a linear regression of standard mass as a function of SVL.

2.2. Collection of blood samples

Blood collection was performed by cardiac puncture with 1 mL syringes and previously heparinized 26Gx1/2" needles. Only blood samples collected within 3 min after the capture of the animal were considered in order to avoid possible influences of capture and manipulation (Romero and Reed, 2005; Tylan et al., 2020). The individuals were sampled once at the time of collection and twice in the experiment, for each sample, approximately 100 µL of blood was drawn. The minimal interval between the date of collection and the experiment in the laboratory was 1 week, enough time for the animals to recover the blood, since amphibians, especially terrestrial ones, have a high capacity to regulate blood volume in the face of dehydration and hemorrhage (Hillman, 2018). These samples were used to evaluate the corticosterone plasma levels (CORT), neutrophil:lymphocyte ratio (N: L ratio), bacterial killing ability (BKA) and hematocrit (HTC) collected in the field and after subjection to stressors in the laboratory.

All blood samples were identified and kept on ice until divided into three aliquots. One of these aliquots was used to mount two blood smear slides for leukocyte profile (≈4 µL), another for hematocrit (HTC) (≈5 µL) measurement and a third one was transferred to Eppendorf tubes and centrifuged (4 min at 604g) for plasma separation. Plasma samples were kept on ice until transferred to the -80 °C freezer the same night. 5–10 µL of plasma were used for hormonal and BKA analysis, respectively.

2.3. Dehydration and movement restriction

The animals were randomly divided into 3 groups with different levels of hydration: control (100% hydrated), 10% and 20% dehydration groups. An ANOVA test with Bonferroni adjustment was performed to assess whether there was a statistical difference between groups with respect to mass. The test showed that there was no difference in mass between groups for both populations (native population: $P = 0.441$; $F = 0.850$; invasive population: $P = 0.665$; $F = 0.281$). The animals in the control group remained with access to water throughout the period of data collection. The toads from the 10% and 20% dehydration groups were weighed in the laboratory again when fully hydrated and with empty bladders. They were then deprived of food two days before the experiment and of access to water the day before the experiment until the time of blood sampling, which allowed for natural evaporative water loss. To increase the rate of water loss for the animals from the 20% dehydration group, we replaced the plastic top of the box with a mesh. Both groups lost the desired mass in a period of 24 h. The next morning the toads were weighed to check if they reached the mass corresponding to the level of dehydration desired and then had a blood sample

collected. If the toads had not reached the desired mass, we would place them closer to the fan to dehydrate and reach the desired mass more quickly. Toads had their body mass reduced through cutaneous water loss by 10% or 20% of the initial body mass; for example, a toad with an initial mass of 100 g achieves a dehydration percentage of 10% or 20% with a mass of 90 g and 80 g, respectively. Once reaching the target mass, individuals had samples of blood collected. Toads from the control group were handled and weighed as those from the 10% and 20% dehydration groups in order to control for handling stress and, later, they were submitted to movement restriction stress.

To perform the movement restriction stress test, toads from all groups were placed in wet cloth bags inside their individual maintenance containers and remained in these conditions for a period of 24 h, following Assis et al. (2015). After this period, blood samples were collected to evaluate the variables: BKA, N:L ratio, CORT and HTC.

The collections and the laboratory experiment were authorized by the Research Ethics Committee: Animal Care and Use of Stellenbosch University (reference number #ACU-2019-8839).

2.4. Bacterial Killing Ability (BKA)

To perform this test, we followed the protocol of Assis et al. (2013). Briefly, plasma samples were diluted in amphibian Ringer's solution (10 µL plasma, 190 µL Ringer's) and then mixed with a 10 µL *Aeromonas hydrophila* working solution (1×10^5 microorganisms mL⁻¹, *A. hydrophila*, IOC/ FDA 110- 36). Positive controls consisted of 10 µL bacteria working solution in 200 µL Ringer's solution, and negative controls contained 210 µL Ringer's solution. Both controls and samples were incubated at 37 °C, optimal temperature for bacterial growth, for 60 min. We chose to use the optimal growth temperature of *A. hydrophila* instead of an ecologically relevant temperature for *S. gutturalis* because there is interspecific variation in temperatures that maximize BKA in bufonids (Moretti et al., 2019), and we did not access the thermal sensitivity curve of BKA for *S. gutturalis* previously to this study. In addition, most studies involving interaction between plasma and bacteria use the optimal growth temperature for the bacteria and this allows us to make comparisons with other studies. After incubation, 500 µL of TSB (Tryptic Soy Broth) was added to each sample, vortexed and pipetted (300 µL from each sample), in duplicate, into a 96-well plate. The microplate was incubated at 37 °C for 1 h, and thereafter the optical density of the samples was measured hourly in a plate spectrophotometer (Thermo, model Multiskan EX) at a wavelength of 595 nm, totaling four readings. The plasma BKA was evaluated at the beginning of the bacterial exponential growth phase and calculated according to the formula: $1 - [\text{optical density of sample} / \text{optical density of positive control}]$ (Assis et al., 2013). The optical density of the samples is the average of the two duplicates. The result of this division is the proportion of microorganisms that the toads' plasma was able to kill compared to the proportion of microorganisms in the positive control.

2.5. Leukocyte profile

One drop of blood (≈2 µL) was used to make two blood smear slides. One of them was stained with Giemsa's solution (10%) to perform differential counting leukocyte types. In an optical microscope with a magnification of 100×, one hundred leukocytes were counted on each slide and classified into neutrophils, lymphocytes, eosinophils, basophils and monocytes for each animal (Campbell, 2012).

2.6. Hormone dosage

Steroid hormones were initially extracted with ether according to Assis et al. (2015, 2019). Plasma CORT levels were determined using EIA kits (CORT number 501320; Cayman Chemical), according to the manufacturer's instructions and previous studies conducted with toads from the genus *Rhinella* (Assis et al., 2019; Titon et al., 2018; Madelaire

et al., 2019; Barsotti et al., 2019). The intra- and inter-assay variation were 11.22% and 3.35%, respectively. The sensitivity of the assays, calculated as 80% B/B0 curve value, was 22.84 pg.

Validation of the use of the corticosterone assay kit from Cayman Chemicals (cat number 501320; Cayman Chemical, Ann Arbor, Michigan) for *S. gutturalis* was conducted with a parallelism test, including baseline. Pooled plasma samples leftover (25 μ L of 10 subjects) were initially extracted with 5 mL of ethyl ether and followed the same procedures mentioned above (according to Assis et al., 2015, 2019). At the end, these pooled samples were resuspended and diluted in EIA buffer. The top standard of the corticosterone kit and the pooled plasma samples were used for a serial dilution (neat, 1:2, 1:4, 1:8, 1:16, 1:32, 1:64 and 1:128) and assayed on the same plate. The standard and sample corticosterone concentration were plotted as a function of Binding/Total Binding, and the 50% binding point was considered indicative of the best dilution factor to run the samples. The standard and sample curves were parallel, not crossing each other (Fig. 1- supp. file), corroborating the functionality of the assay for guttural toads. The best dilution factor for baseline pooled plasma samples from *S. gutturalis* corresponds to 1:2 (Fig. 1- supp. file).

2.7. Hematocrit (HTC)

The HTC was measured shortly after blood samples were collected and calculated as the ratio of blood cells to total blood volume after centrifugation of blood within a microhematocrit tube (4 min at 218g).

2.8. Statistical analysis

Data were initially submitted to descriptive statistics and the Shapiro-Wilk normality test. HTC (\log_{10+1}), BKA (ACOS), N:L ratio (square root) and CORT (\log_{10+1}) were transformed to adjust to the premise for parametric tests. Differences in field variables between invasive and native populations were verified by *t*-tests. A set of mixed ANCOVAs were performed to evaluate possible effects of body index as a covariate for the variables of the experiment. The body index was calculated as unstandardized residuals of a linear regression of standard mass as a function of snout-vent-length and were considered as a proxy of body condition to both populations.

In these ANCOVAs, physiological variables were considered as dependent variables, dehydration (control, 10% dehydration and 20% dehydration) and population (native and invasive) were considered as a between subject factor and restraint was considered as a within subject factor following a full factorial model. When the covariate effect was not

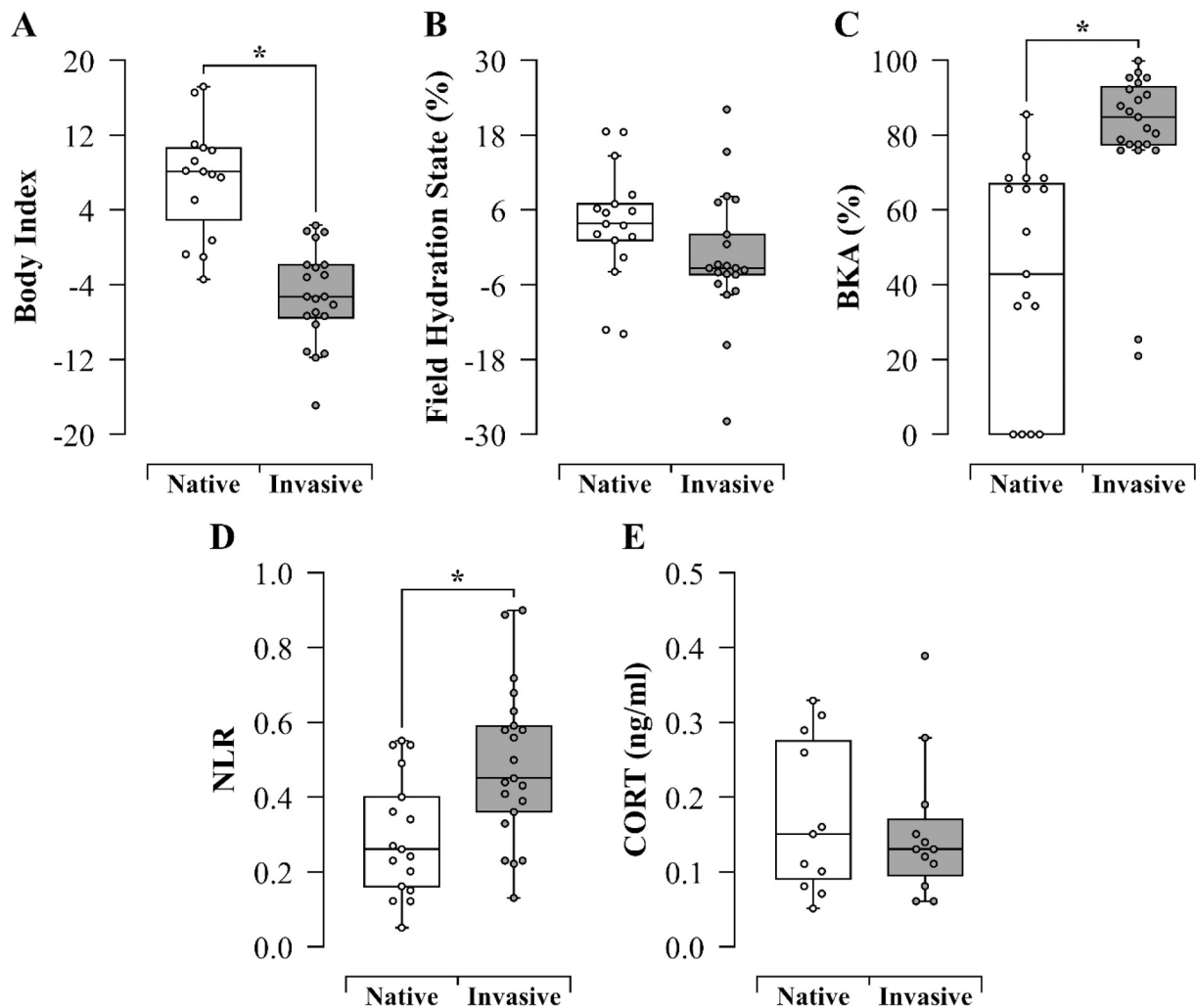


Fig. 1. Variables collected in the field from the invasive (Cape Town) and native (Durban) populations of *Sclerophrys gutturalis* males. Body index (A); Field Hydration State (B); Bacterial Killing Ability (C); Neutrophil: Lymphocyte Ratio (D); Corticosterone Plasma Levels (E). Boxplot inside lines indicate medians, lower and upper borders represent 1st and 3rd quartiles, respectively, whiskers represents upper and lower limits of 1.5 times inter-quartile range and circles represent individual data points. Asterisk (*) represents mean multiple comparisons with Bonferroni adjustment significant differences ($P \leq 0.05$).

present, the covariate was eliminated from the model. The analysis was followed by a means comparison test, using the Bonferroni adjustment. Statistical analyses were performed in SPSS 22 for Windows (IBM Corp., Armonk, NY).

3. Results

3.1. Comparison between invasive and native population of baseline samples

Individuals from Cape Town (invasive population) and Durban (native population) differed in baseline variables of field mass, standard body mass, body index, BKA and N:L ratio (Table 1). The field mass recorded at the time of collection ($t_{41} = -3.859$, $P < 0.001$) and the body condition ($t_{34} = -6.632$, $P < 0.001$, Fig. 1 A) were higher for the native population compared to the invasive population. In addition, the standard body mass was higher in individuals from the native population ($t_{39} = -4.767$, $P < 0.001$). Although not significant, there was a tendency of higher field hydration state for individuals from the native population ($t_{39} = -1.795$, $P = 0.080$; Fig. 1 B). Individuals from the native population showed lower BKA and N: L compared to those from the invasive population ($t_{40} = -5.525$, $P < 0.001$, $t_{39} = 2.316$, $P = 0.026$; Figs. 1 C and D, respectively). CORT did not differ between populations ($t_{20.182} = -0.811$, $P = 0.427$, Fig. 1 E).

Table 1

Descriptive statistics and statistical analysis (*t*-test) comparing field data (body mass, rehydrated mass, dehydration level in field, snout-vent length, body index, bacterial killing ability, neutrophil: lymphocyte ratio and corticosterone plasma levels) of invasive and native *Sclerophrys gutturalis* populations. Due to technical issues, not all variables were measured for all individuals.

Variables	Population	N	Mean \pm SD	t	DF	P-value
Field Mass (g)	Invasive	23	31.810 \pm 9.029	-3.859	41	<0.001
	Native	20	43.983 \pm 11.633			
Standard Body Mass (g)	Invasive	21	30.800 \pm 8.904	-4.767	39	<0.001
	Native	20	45.606 \pm 10.928			
Field Hydration State (%)	Invasive	21	-1.606 \pm 10.365	-1.795	39	0.080
	Native	20	3.815 \pm 8.868			
SVL (mm)	Invasive	25	73.336 \pm 7.296	-0.389	38	0.699
	Native	15	74.267 \pm 7.363			
Body Condition (g.mm-3)	Invasive	21	-5.162 \pm 5.075	-6.632	34	<0.001
	Native	15	7.227 \pm 6.110			
BKA Field (%)	Invasive	23	80.855 \pm 19.818	-5.525	40	<0.001
	Native	19	40.300 \pm 31.252			
N:L Ratio Field	Invasive	25	0.488 \pm 0.208	2.316	39	0.026
	Native	20	0.345 \pm 0.201			
CORT (ng/mL)	Invasive	12	0.154 \pm 0.095	-0.780	22	0.444
	Native	13	0.159 \pm 0.101			

SD: Standard deviation; DF: degrees of freedom; SVL: Snout-Vent Length; BKA: Bacterial Killing Ability; N:L Ratio: Neutrophil: Lymphocyte Ratio; CORT: Corticosterone plasma levels.

3.2. Experimental dehydration and movement restriction: comparison between invasive and native populations

Mixed ANOVA shows significant difference between populations for HTC and BKA ($F_{1,29} = 8.018$, $P = 0.008$, $F_{1,34} = 142.202$, $P < 0.001$, respectively, Tables 2 and 3). N:L ratio was affected by restraint, dehydration and an interaction between populations and restraint ($F_{1,33} = 87.737$, $P < 0.001$, $F_{2,33} = 4.739$, $P = 0.016$, $F_{1,33} = 10.117$, $P = 0.003$, respectively, Table 3). Individuals from the native population showed lower BKA than individuals from the invasive population both after dehydration and after movement restriction (control group: $P < 0.001$, 10% dehydration group: $P < 0.001$, 20% dehydration group after dehydration: $P = 0.010$, 20% dehydration group after restriction of movement: $P < 0.001$, Fig. 2). In addition, within the invasive population, toads dehydrated by 20% increased BKA compared to the control group ($P = 0.034$, Fig. 2). The populations had no changes in the N: L ratio after dehydration. Nonetheless, individuals from the invasive population showed increased N:L ratio after movement restriction in control, 10% and 20% dehydration groups ($P < 0.001$; Fig. 3). Individuals from the native population also showed higher N:L ratio after movement restriction in control and 10% dehydration groups ($P = 0.001$; $P = 0.012$, respectively), but not in the 20% dehydration group ($P = 0.417$, Fig. 3).

Toads from native and invasive populations did not differ in CORT in response to the treatments. CORT was higher in invasive toads dehydrated by 10% and 20% when compared with the control group ($P = 0.002$ and $P = 0.007$, respectively, Fig. 4). Invasive toads dehydrated by 10% presented higher CORT than those from the control group after movement restriction ($P = 0.019$, Fig. 4). In parallel, CORT was higher in native toads dehydrated by 20% when compared to the control group ($P = 0.017$, Fig. 4). Invasive and native toads showed lower CORT after movement restriction when compared to those measured after dehydration at 20% ($P = 0.005$ and $P = 0.022$, respectively, Fig. 4). The results also showed an interaction between restriction and group for CORT ($P = 0.003$, Table 3).

Native toads showed higher HTC than invasive toads after movement restriction in control and 20% dehydration groups ($P = 0.030$ and $P = 0.046$, respectively; Fig. 5).

4. Discussion

Guttural toads, *Sclerophrys gutturalis*, from the invasive population had a lower body index and standard body mass, and higher BKA and N: L ratio, compared to individuals from the native population under field conditions. These results are consistent with previous studies showing that the challenges of invasion can directly impact body condition of invasive animals (Brown et al., 2011; Vimercati et al., 2019). Cape Town is characterized by a Mediterranean climate with limited water availability in summer, compared to the native region of these invasive guttural toads, which could reduce the amount and diversity of arthropod prey for anurans (Vimercati et al., 2019). They are probably also restricted in hours of foraging activity, given the increased need to remain longer in microenvironments in contact with water. Additionally, the invasive population of *S. gutturalis* had lower field body mass and tended to show a lower field hydration state than those from the native range. Vimercati et al. (2018) compared the same two populations of *S. gutturalis* and found similar results, along with a positive relationship between hydration level and relative humidity. These results in conjunction suggest that these invasive toads live under stressful conditions of higher dehydration potential.

The invasive population of *S. gutturalis* had higher BKA and N: L ratio compare to the native population. Some studies have shown a lower immune response of toads on the invasion front compared to longer established populations (Brown et al., 2007; Brown and Shine, 2014; Llewellyn et al., 2012). However, the literature has shown diverse and contradictory data related to immunocompetence of invasive organisms.

Table 2

Descriptive statistics of hematocrit, bacterial killing ability, neutrophil: lymphocyte ratio and corticosterone plasma levels analyzed after the experiment of dehydration stress and movement restriction for *Sclerophrys gutturalis* invasive and native populations.

Variable	Restriction	Population	Group	N	Mean	SD		
HTC (%)	Pre	Invasive	Control	6	19.028	9.402		
			Dehydration 10%	7	27.701	7.443		
			Dehydration 20%	8	26.976	11.662		
			Native	Control	3	25.740	3.942	
				Dehydration 10%	6	42.666	10.665	
				Dehydration 20%	6	47.023	9.724	
		Post	Invasive	Control	6	18.631	7.251	
				Dehydration 10%	7	22.648	3.057	
				Dehydration 20%	8	19.795	13.680	
			Native	Control	3	30.553	4.809	
				Dehydration 10%	6	32.110	13.029	
				Dehydration 20%	6	25.945	9.977	
	BKA (%)	Pre	Invasive	Control	8	77.795	24.020	
				Dehydration 10%	7	69.611	19.498	
				Dehydration 20%	8	81.196	9.976	
				Native	Control	5	10.968	14.873
					Dehydration 10%	6	16.458	10.134
					Dehydration 20%	6	22.223	7.410
			Post	Invasive	Control	8	76.302	8.903
					Dehydration 10%	7	69.185	29.308
					Dehydration 20%	8	71.245	25.467
				Native	Control	5	24.516	13.573
					Dehydration 10%	6	18.085	10.120
					Dehydration 20%	6	20.833	11.053
N:L Ratio		Pre	Invasive	Control	7	0.391	0.191	
				Dehydration 10%	9	0.492	0.240	
				Dehydration 20%	8	0.301	0.307	
			Native	Control	5	0.480	0.204	
				Dehydration 10%	6	0.748	0.310	
				Dehydration 20%	6	0.546	0.202	
		Post	Invasive	Control	7	1.444	0.601	
				Dehydration 10%	9	1.202	0.371	
				Dehydration 20%	8	0.953	0.302	
			Native	Control	5	1.094	0.249	
				Dehydration 10%	6	1.253	0.632	
				Dehydration 20%	6	0.665	0.275	
CORT (ng/mL)	Pre	Invasive	Control	7	0.226	0.121		
			Dehydration 10%	9	0.900	0.505		
			Dehydration 20%	8	0.820	0.252		
		Native	Control	4	0.368	0.315		
			Dehydration 10%	6	0.841	0.329		
			Dehydration 20%	6	1.037	0.385		
	Post	Invasive	Control	7	0.379	0.267		

Table 2 (continued)

Variable	Restriction	Population	Group	N	Mean	SD	
			Dehydration 10%	9	0.778	0.333	
			Dehydration 20%	8	0.495	0.223	
			Native	Control	4	0.535	0.034
				Dehydration 10%	6	0.593	0.306
				Dehydration 20%	6	0.737	0.275

SD: Standard Deviation; HTC: Hematocrit; BKA: Bacterial Killing Ability; N:L Ratio: Neutrophil: Lymphocyte Ratio; CORT: Corticosterone plasma levels.

These organisms are exposed to new biotic and abiotic conditions, which exert novel selective pressures (Müller-Schärer et al., 2004; Carroll, 2008; White and Perkins, 2012; Brown et al., 2015), and favor the emergence of phenotypes that increase dispersal (Travis and Dytham, 2002; Shine et al., 2011; Brown et al., 2015; Courant et al., 2019) among other traits. Thus, it follows that diminished immune function could make invasive toads more susceptible to new pathogens, decreasing their survival and invasion success. Similar to our results, Brown et al. (2015) showed that offspring of invasive cane toads (*Rhinella marina*) in Australia have higher BKA, higher numbers of neutrophils and phagocytic activity on the invasion front when compared to offspring of toads from long-established populations bred under standard conditions. These results suggest genetically based shifts in immune response compromised by long-distance dispersal in free-ranging toads (Brown et al., 2015). Additionally, studies with shore crabs and birds show greater non-specific immune function of invasive individuals when compared to native ones (Keogh et al., 2017; Moller and Cassey, 2004). Still, according to Brown et al. (2015), the baseline levels of complement and neutrophils have been reinforced in cane toads on the invasion front, while the costs associated with its activation decreased. Thus, it is possible that invasive animals, such as *S. gutturalis*, tend to have increased constitutive immunity associated with increased surveillance. This would ensure a faster response to immunological challenges, reducing the scope of immune activation after infection and its costs associated with fever, anorexia, decreased locomotor activity, growth rates (Klasing and Korver, 1997; Spurlock, 1997; Lee and Klasing, 2004), and potentially dangerous systemic inflammatory response (Janeway Jr et al., 2001; Klein and Nelson, 1999; Lee and Klasing, 2004). It is unknown whether such responses are exaggerated in the invasive population as novel climates differ from the original, or if they are, at least in part, genetically determined in guttural toads. These hypotheses remain to be tested.

Blood leukocyte numbers and proportions have also been commonly used as indicators of physiological stress in vertebrates, including toads from the genus *Rhinella* (Davis et al., 2008; Campbell, 2015; Savage et al., 2016; Assis et al., 2015, 2017, 2019; Barsotti et al., 2017, 2019). In response to various stressors, including dehydration and malnutrition, the number of circulating neutrophils increases while the number of lymphocytes decreases, reflecting changes in leukocyte distribution and production (Davis et al., 2008; Savage et al., 2016; Barsotti et al., 2019). Moreover, increased N: L ratio has been shown to reflect more accurately chronic stress than increased glucocorticoid plasma levels (Davis et al., 2008; Swan and Hickman, 2014; Hickman, 2017). Thus, it is possible that anurans from the invading population in different environmental conditions and facing new challenges, may be more stressed, reflecting the neutrophilia and lymphopenia observed in our results.

Contrary to what we expected, toads from the invasive population show higher BKA both after dehydration and restriction protocols. Interestingly, the difference between populations regarding field BKA was maintained after captivity and all the experimental protocols. These results show that BKA's response was robust to exposure to acute stressors, at least within the period analyzed, suggesting an increase in

Table 3

Statistical analysis of mixed ANOVA, for repeated and independent measurements at two different times (for the control group: pre and post restriction, for the 10% and 20% dehydration groups: dehydration and post-restriction), as well as interaction between treatment and group.

Variable	Source	Type III SS	DF	MS	F	P
HTC	Intercept	128.843	1	128.843	4768.653	<0.001
	Restraint	0.007	1	0.007	0.387	0.539
	Population	0.217	1	0.217	8.018	0.008
	Group	0.116	2	0.058	2.152	0.134
	Restraint*Population	0.002	1	0.002	0.131	0.720
	Restraint*Group	0.027	2	0.013	0.786	0.465
	Restraint*Population*Group	0.069	2	0.034	1.999	0.154
	Population*Group	0.059	2	0.030	1.095	0.348
	Error (Restraint)	0.497	29	0.017		
	Error	0.784	29	0.027		
BKA	Intercept	274,526.693	1	274,526.693	1321.536	<0.001
	Restraint	719.941	1	719.941	3.898	0.057
	Population	29,540.097	1	29,540.097	142.202	<0.001
	Group	581.098	2	290.549	1.399	0.261
	Restraint*Population	221.591	1	221.591	1.200	0.281
	Restraint*Group	1.834	2	0.917	0.005	0.995
	Restraint*Population*Group	305.449	2	152.724	0.827	0.446
	Population*Group	1041.507	2	520.754	2.507	0.096
	Error (Restraint)	6279.394	34	184.688		
	Error	7062.926	34	207.733		
N:L Ratio	Intercept	56.426	1	56.426	1260.811	<0.001
	Restraint	2.196	1	2.196	87.737	<0.001
	Population	0.001	1	0.001	0.032	0.858
	Group	0.424	2	0.212	4.739	0.016
	Restraint*Population	0.253	1	0.253	10.117	0.003
	Restraint*Group	0.143	2	0.072	2.866	0.071
	Restraint*Population*Group	0.053	2	0.026	1.054	0.360
	Population*Group	0.023	2	0.011	0.255	0.777
	Error (Restraint)	0.826	33	0.025		
	Error	1.477	33	0.045		
CORT	Intercept	30.924	1	30.924	207.239	<0.001
	Restraint	0.237	1	0.237	5.013	0.032
	Population	0.137	1	0.137	0.919	0.344
	Group	2.369	2	1.185	7.939	0.001
	Restraint*Population	0.004	1	0.004	0.086	0.770
	Restraint*Group	0.679	2	0.340	7.182	0.003
	Restraint*Population*Group	0.024	2	0.012	0.254	0.777
	Population*Group	0.472	2	0.236	1.580	0.221
	Error (Restraint)	1.608	34	0.047		
	Error	5.073	34	0.149		

SS: Sum of Squares; DF: degrees of freedom; MS: Mean Square; HTC: Hematocrit; BKA: Bacterial Killing Ability; N:L Ratio: Neutrophil: Lymphocyte Ratio; CORT: Corticosterone plasma levels.

constitutive immunity in the invasive population.

Moreover, dehydrating toads by 20% increased CORT and BKA, which may indicate an immunostimulatory effect of CORT. Accordingly, increased CORT along with different immune variables after submission to an acute stressor has been repeatedly observed in anurans (Assis et al., 2015, 2017, 2019; Titon et al., 2018; Titon et al., 2019; Barsotti et al., 2017, 2019). Dhabhar et al. (1996) showed that acute stress results in a rapid reduction in circulating leukocyte numbers associated with a redistribution of blood leukocytes to other organs, tending to increase the immune response in some body compartments. An acute response to stressors is also associated with increased antigen presentation, effector cell function, antibody production and cytokines expression (Dhabhar and McEwen, 1997). Additionally, in response to short-term stressors and acute increase in glucocorticoids, there is an increase in activity of the complement system (Dhabhar and McEwen, 1996, 1997; Dhabhar, 1998), which is also consistent with higher BKA response in dehydrated toads.

According to our predictions, dehydration activated the HPI axis culminating in increased CORT, mainly when toads were dehydrated by 20%. This increase in CORT indicates that dehydration is a stressor for *S. gutturalis*, as has already been concluded for other vertebrate species (Arnhold et al., 2007; Cain and Lien, 1985; Moeller et al., 2017), including toads from genus *Rhinella* (Barsotti et al., 2019). Since CORT participates in the integrated physiological response of anurans to dehydration, due to its mineralotropic actions (Broillet et al., 1993;

Chen et al., 1998; Vera et al., 2017), it is possible that the increase of CORT observed in *S. gutturalis* may contribute to protection against possible deleterious effects caused by the loss of volume of extracellular fluid. Moreover, glucocorticoids potentiate the effects of angiotensin II (ANG II) mediating drinking behavior in mammals (Ganesan and Sumners, 1989; Sumners et al., 1991; Takei, 2000), and can similarly stimulate water seeking-behavior in anurans. Interestingly, hydrated individuals of *S. gutturalis* who had their CORT artificially increased, through transdermal application, showed an increase in water-seeking behavior (Madelaire et al., 2020). Considering that invasive toads tended to show lower hydration levels than native toads under field conditions, this higher efficiency in finding water due to increased CORT when dehydrated might favor the dispersal of the invasive Cape Town population.

Movement restriction represented a stressor for *S. gutturalis*, increasing the N: L ratio for both populations. Changes in leukocyte numbers and proportions in the blood are often used as indicators of infection and physiological stress in amphibians and other vertebrates (Assis et al., 2015, 2017, 2019; Barsotti et al., 2017, 2019; Davis et al., 2008; Campbell, 2015; Savage et al., 2016). Neutrophils and lymphocytes constitute the majority of blood leukocytes in amphibians (Savage et al., 2016) and the number of neutrophils increases and the lymphocyte decreases in the face of stressors, reflecting changes in leukocyte distribution and production (Davis et al., 2008; Savage et al., 2016). Assis et al. (2015) showed increased N: L ratio after restraint with

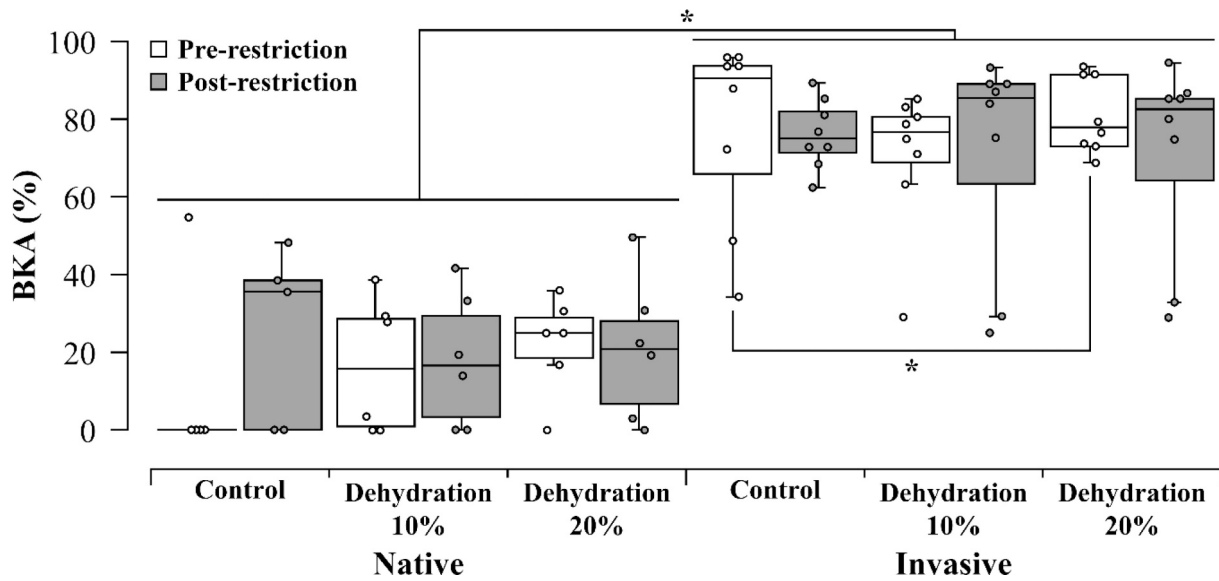


Fig. 2. Bacterial Killing Ability against *Aeromonas hydrophila* of *Sclerophrys gutturalis* males of the invasive and native population in two distinct moments for the control group: pre and post restriction, for the 10% and 20% dehydration groups: dehydration and post-restriction. Boxplot inside lines indicate medians, lower and upper borders represent 1st and 3rd quartiles, respectively, whiskers represents upper and lower limits of 1.5 times inter-quartile range and circles represent individual data points. Asterisk (*) represents mean multiple comparisons with Bonferroni adjustment significant differences ($P \leq 0.05$).

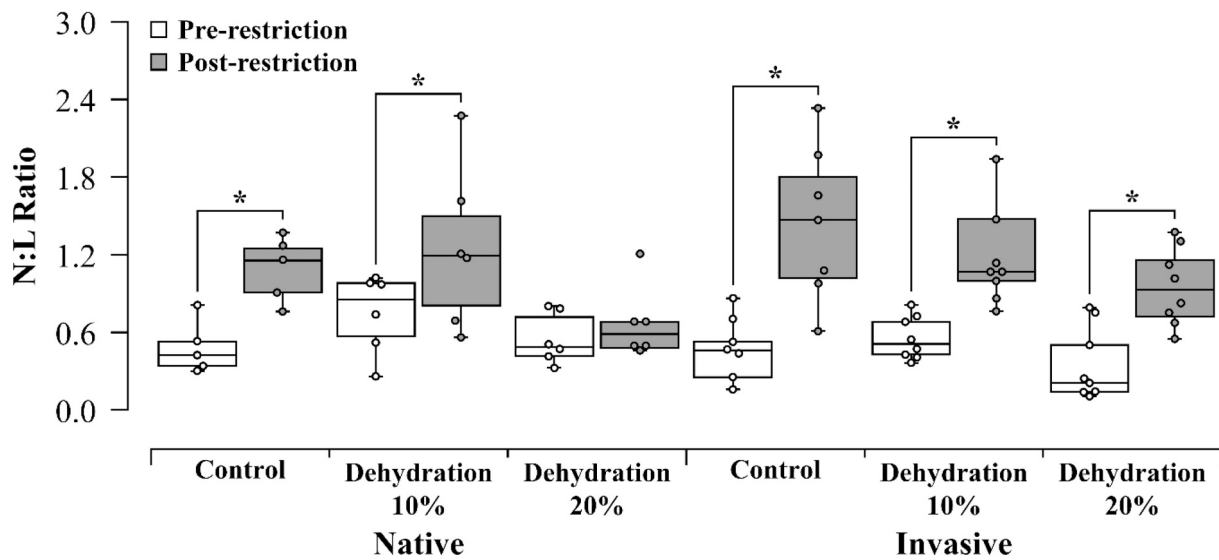


Fig. 3. Neutrophil: Lymphocyte ratio of *Sclerophrys gutturalis* males of the invasive and native population in two distinct moments (for the control group: pre and post restriction, for the 10% and 20% dehydration groups: dehydration and post-restriction). Boxplot inside lines indicate medians, lower and upper borders represent 1st and 3rd quartiles, respectively, whiskers represents upper and lower limits of 1.5 times inter-quartile range and circles represent individual data points. Asterisk (*) represents mean multiple comparisons with Bonferroni adjustment significant differences ($P \leq 0.05$).

movement restriction in *R. ictERICA*, but not in individuals subjected to less stressful restraint without movement restriction. Additionally, Barsotti et al. (2019) showed that *R. ornata* subjected to 20% dehydration showed a more pronounced increase in N: L than those subjected to 10% dehydration.

CORT was not increased after 24 h of movement restriction in hydrated native and invasive toads, in contrast with previous observations on other anuran species (Assis et al., 2015, 2019; Barsotti et al., 2019). Given that the N: L ratio increased after movement restriction in these toads, it is probable that a peak of increased CORT in response to this treatment occurred earlier than the time at which the blood sample was collected. Moreover, it is interesting to observe that, both invasive and native toads that were dehydrated by 20% and posteriorly submitted to movement restriction, CORT values after movement restriction were

lower than those measured after intense dehydration. Given that movement restriction was performed in damp cloth bags, toads may have rehydrated and reduced CORT during the period of movement restriction. Rehydration up to almost 100% has been previously observed in another toad (*Rhinella ornata*) under the same experimental conditions (Table 6- supp. file). Even so, these results suggest that movement restriction was not a stressor as intense as dehydration for this species. Likewise, intense dehydration is a more potent stressor than movement restriction in *R. ornata* (Barsotti et al., 2019), probably the only other anuran species from which there is data available on these two stressors.

In summary, invasive toads show lower standard field body mass, lower body index and tended to show lower hydration status in the field, which is consistent with life in the novel drier environmental conditions

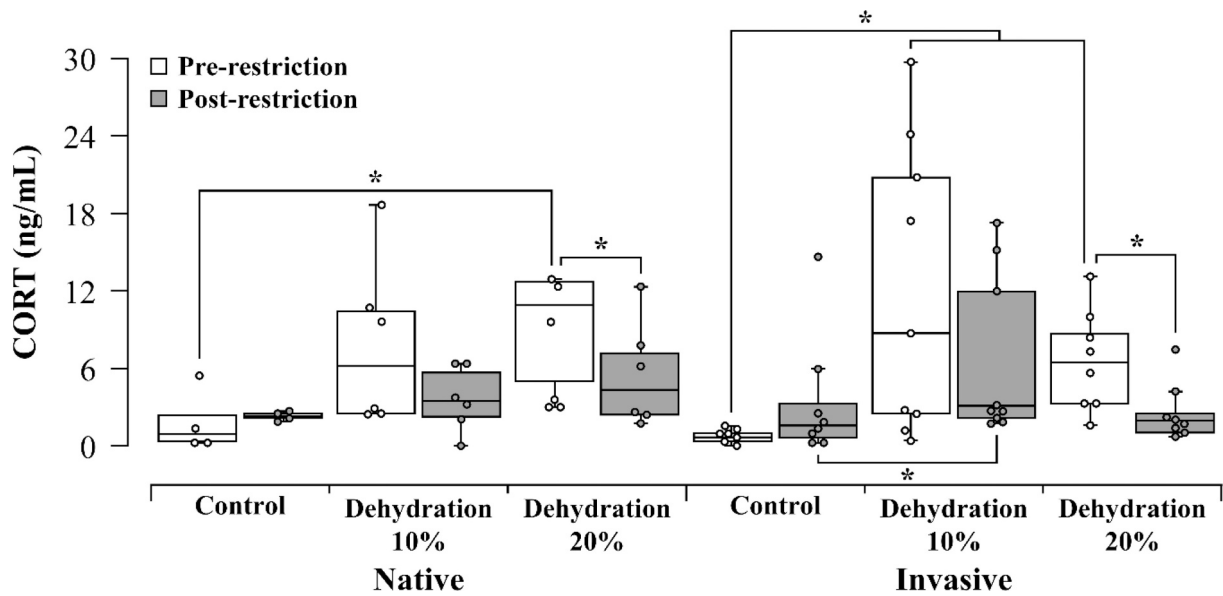


Fig. 4. Corticosterone plasma levels of *Sclerophrys gutturalis* males of the invasive and native population in two distinct moments (for the control group: pre and post restriction, for the 10% and 20% dehydration groups: dehydration and post-restriction). Boxplot inside lines indicate medians, lower and upper borders represent 1st and 3rd quartiles, respectively, whiskers represents upper and lower limits of 1.5 times inter-quartile range and circles represent individual data points. Asterisk (*) represents mean multiple comparisons with Bonferroni adjustment significant differences ($P \leq 0.05$).

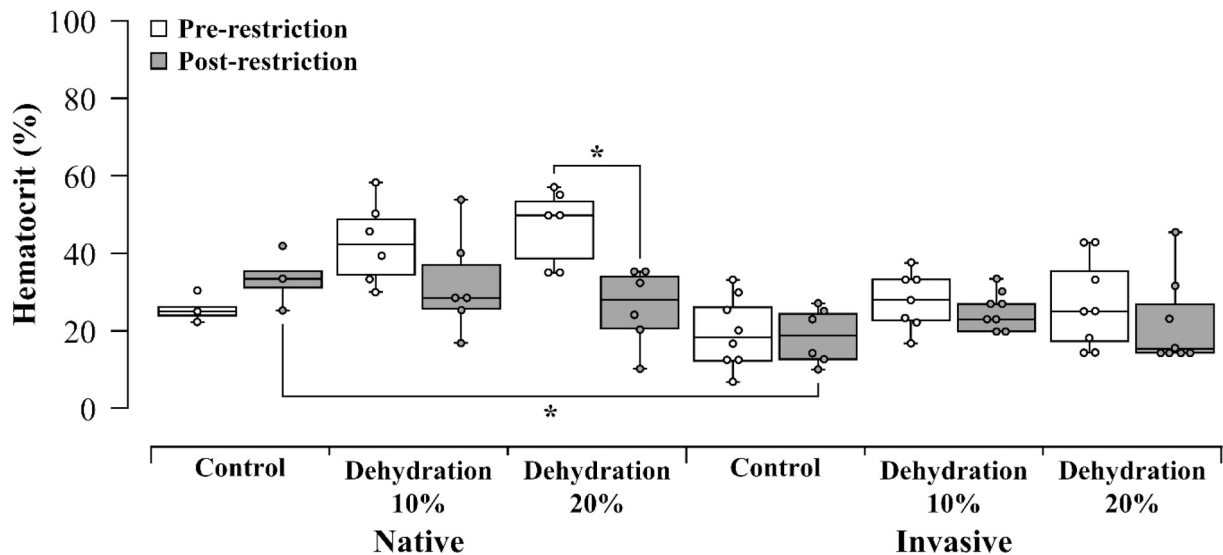


Fig. 5. Hematocrit of *Sclerophrys gutturalis* males of the invasive and native population in two distinct moments (for the control group: pre and post restriction, for the 10% and 20% dehydration groups: dehydration and post-restriction). Boxplot inside lines indicate medians, lower and upper borders represent 1st and 3rd quartiles, respectively, whiskers represents upper and lower limits of 1.5 times inter-quartile range and circles represent individual data points. Asterisk (*) represents mean multiple comparisons with Bonferroni adjustment significant differences ($P \leq 0.05$).

of Cape Town. Invading toads also showed higher N: L and BKA ratio in the field, compared to those from the native population, suggesting that invasive toads face stressful conditions and invest comparatively more in non-expensive components of their immune system. It is interesting to observe the persistence of inter-population differences in BKA, even after submission to acute stress (dehydration and movement restriction). Although these traits have favored the permanence and dispersion of *S. gutturalis* in the invasive range, it is important to emphasize that we have investigated short-term phenotypic differences between these populations. An analysis of genetic diversity suggests this invasive population is not derived from founder effects (Telford et al., 2019). These differences might be, at least in part, due to plastic responses to current environmental differences during the breeding season

(Vimercati et al., 2019), or fast adaptive responses after 20 years of their introduction in Cape Town (Vimercati et al., 2018). Additional studies, including rearing individuals from both populations under common garden conditions, would be necessary to understand the causes of this phenotypic divergence.

Funding

This work was supported by Fundação de Amparo à Pesquisa do Estado de São Paulo (FAPESP) through a grant for F.R.G (2014/16320-7), PhD scholarship to A.M.G.B (2018/23661-6) and postdoctoral fellowship awarded to C.B.M. (2018/00406-0). F.R. Gomes is a research fellow from the Brazilian Conselho Nacional de Desenvolvimento

Científico e Tecnológico (CNPq) #302308/2016-4. J. Measey is chief researcher at the Centre for Invasion Biology (C-I-B) based in the Department of Botany and Zoology at Stellenbosch University.

Declaration of Competing Interest

The authors declare no competing or financial interests.

Acknowledgements

We would like to thank Professor Hannes van Wyk for allowing the use of his laboratory and equipment. This study was supported by the Fundação de Amparo à Pesquisa do Estado de São Paulo (FAPESP) FRG (2013/00900-1 and 2014/16320-7), AMGB (2018/23661-6) and CBM (2018/00406-0). JM and CW would like to thank the DSI-NRF Centre of Excellence for Invasion Biology and Stellenbosch University.

Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.cbpa.2020.110870>.

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