Differences in morphology and in composition and release of parotoid gland secretion in introduced cane toads (*Rhinella marina*) from established populations in Florida, USA

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Abstract
Cane toads are highly toxic bufonids invasive in several locations throughout the world. Although physiological changes and effects on native predators for Australian populations have been well documented, Florida populations have received little attention. Cane toads were collected from populations spanning the invaded range in Florida to assess relative toxicity, through measuring morphological changes to parotoid glands, likelihood of secretion, and the marinobufagenin (MBG) content of secretion. We found that residual body indices increased in individuals from higher latitude populations, and relative parotoid gland size increased with increasing toad size. There was no effect of latitude on the allometric relationship between gland size and toad size. We observed an increase in likelihood of secretion by cane toads in the field with increasing latitude. Individuals from southern and northern populations did not vary significantly in the quantity of MBG contained in their secretion. Laboratory-acclimated cane toads receiving injections of epinephrine were more likely to secrete poison with increasing dose, although there was no difference in likelihood of secretion between southern and northern populations. This suggests that differences between populations in the quantities of epinephrine released in the field, due to altered hypothalamic sensitivity upon disturbance, may be responsible for the latitudinal effects on poison secretion. Our results suggest that altered pressures from northward establishment in Florida have affected sympathetic sensitivity and defensive mechanisms of cane toads, potentially affecting risk to native predators.

Keywords
bufadienolide, epinephrine, invasion, phenotypic plasticity, *Rhinella marina*, sympathetic sensitivity
1 | INTRODUCTION

Cane toads (Rhinella marina) are large and highly toxic members of the family Bufonidae and are native to South America (Acevedo et al., 2016; Zug & Zug, 1979). Due to their voracious feeding responses, cane toads were introduced into Australia in the early to mid-1900s as a means of controlling sugar cane pests (Lever, 2001; Phillips et al., 2006) and were released in Florida (Miami) in the United States prior to 1955 (Krakauer, 1968). They have dispersed in these locations with invasion rates of up to 55 km/year in Australia (Phillips et al., 2007). Additionally, R. marina reproduce in large numbers, and females may lay up to 30,000 eggs per clutch in invaded areas (Hagman & Shine, 2006). However, the extent of cane toad dispersal and reproductive capacity in the United States, where they have established in higher latitude temperate areas (Mittan & Zamudio, 2019) is not well known.

Much of the devastation to native Australia fauna and predators caused by invasive cane toads has been due to their highly toxic secretions, common to bufonids and many other amphibians (Garg et al., 2007), that significantly impact the survival of native predators from the areas they invade (Shine, 2010). Toad poison is a cocktail of toxins (Hayes et al., 2009), the main components of (Mailho-Fontana et al., 2014) and in other cutaneous macroglands synthesized and stored in the large parotoid glands on the shoulders (Mailho-Fontana et al., 2014) and in other cutaneous macroglands in the skin (Mailho-Fontana et al., 2018). The secretion contains a cocktail of toxins (Hayes et al., 2009), the main components of which belong to a family of compounds known as bufadienolides (BDs) (Chen & Kovaříková, 1967; Lever, 2001). These steroidal compounds, derived from cholesterol (Porto & Gros, 1971), are more broadly classified as cardiotonic steroids (Sousa et al., 2017; Steyn & van Heerden, 1998). These compounds exert their effects by binding to the enzyme Na+/K+-ATPase to induce sustained contraction. Marinobufagenin (MBG) is a potent cardiotonic compound found in the secretions of R. marina (Sciani et al., 2013) and has been shown to induce cell death in cardiac myocytes (Liu et al., 2012).

Altered selective pressures may affect various physiological aspects in amphibians, including the potency of poisonous secretions. Members of the toad species Bufo bufo occupying regions of elevated anthropogenic disturbance differ in toxicity compared to those in less disturbed agricultural areas (Bókony et al., 2019), and individuals of Bufo boreas exposed to increased predation cues prior to metamorphosis had higher concentrations of more toxic compounds in their secretions (Benard & Fordyce, 2003). Altered morphological and physiological changes have been documented in R. marina at the invasion front when compared to longer established populations (Friesen & Shine, 2019; Gardner et al., 2020). Parotoid gland size has been shown to increase with body size (snout-vent length (SVL)) in R. marina (Phillips et al., 2007; Phillips & Shine, 2005).

In Australia, this allometric increase is greater in toads at the invasion front than in long-established populations (Phillips & Shine, 2005). This may indicate that cane toads produce more toxic secretions when invading new habitats, possibly to deter predators early in the invasion process, as many predators rapidly learn to avoid preying on toads (Greenlees et al., 2010).

Prior to the introduction of R. marina, Australia had no toad species (Beckmann & Shine, 2009; Hayes et al., 2009). In contrast, potential predators that may encounter cane toads invading Florida have evolved in sympatry with other toad species, such as the southern toad, Anaxyrus terrestris, which also possess BDs in their toxic secretions (Mohammadi et al., 2016). Invertebrate predators in Florida are capable of consuming cane toad eggs with low mortality risk (Punzo & Lindstrom, 2001) relative to larval anuran predators and fish. Other native predators, including several snake species, opossums, and birds (Meshaka, 2011) have been observed consuming adult toads with little or no ill-effect.

Individuals from edge populations of R. marina in Australia have shown altered behavior to novel stressors, such as decreased escape behavior and an increased likelihood of secreting poison during simulated predation events (Hudson et al., 2017). Although these behaviors have yet to be observed for toads in Florida, toads near the northern edge populations have shown elevated baseline corticosterone concentrations and attenuated responses to a restraint challenge when compared to individuals from southern core populations (Assis et al., 2020). This pattern of attenuated response to novel stressors has also been documented under laboratory conditions for Florida cane toads (Gardner et al., 2020). Although the risk of R. marina poisoning among native predators in Australia has been well documented (Crossland et al., 2008; Greenlees et al., 2006; Letnic et al., 2008; Phillips & Shine, 2006), that risk has not been reported quantitatively for the Florida cane toad invasion. Here, we assess morphological, behavioral, and physiological characteristics reflecting toxicity risk of R. marina from well-established sites in southern Florida and from sites stretching northward to the invasion front. We measured the size of the parotoid glands, the likelihood of secretion, and the concentration of MBG in samples of secretion obtained in the field. We then collected toads to assess the epinephrine-induced secretion response under controlled laboratory conditions. We chose to use MBG concentration as an index of overall toxicity of cane toads from field samples obtained from southern and northern populations due to the potency and abundance of this compound in the integumentary secretion of R. marina. We also document morphological and physiological differences between the source populations in southern Florida and established populations occurring closer to the invasion front in northern Florida.

2 | METHODS

2.1 | Field sampling, 2018

Approximately 20 toads from nine populations (established from 1955 to 1991 (U. S. Geological Survey, 2020)) across a south–north latitudinal gradient representing the current invasive range in Florida were collected from May 10 to 19, 2018. Toads were collected from approximately 2,000–2,400 hr and placed into plastic bags immediately upon capture (Figure 1) (for exact numbers collected from each
population, see Table 1). One hour following capture, toads were removed from the bags and their mass, sex, and snout-vent-length (SVL) were recorded. Toads were then photographed next to a ruler for scale, and the images were used to measure gland size (right parotoid gland), as well as to indicate whether a toad was secreting poison following the capture and handling period. Measurements of total length, width, and area of glands were performed using ImageJ software.

### 2.2 Field sampling, 2019

In May of 2019, toads from a southern source population in Miami (n = 10), as well as two northern “edge” populations, New Port Richey (NPR) and Deland (n = 10 NPR, n = 7 DeLand), were captured. Secretion samples were collected into cryovials by gentle compression of the parotoid glands (Toledo et al., 1992) and immediately frozen in liquid nitrogen. Sex, SVL, mass, and parotoid gland measurements were recorded for each toad sampled.

### 2.3 Marinobufagenin content of samples

Samples were kept frozen from the time of field collection through their transport to the laboratory. The lids of vials containing parotoid secretion were punctured with a 16G needle and the contents freeze-dried (LABCONCO Freezone 4.5). The dry contents were removed after 24 hr, and the dry mass of each whole sample was recorded. A portion of each sample was transferred to a glass vial and diluted with 0.5ml/mg of a 90:10 methanol/water UV grade solution. The samples were then sonicated for 30 min, rotated, and sonicated for another 15 min. Following sonication, the samples were centrifuged for 15 min at 3,000 rpm to separate undissolved solid material. The supernatant was then collected and stored for analysis in a glass instrument vial (1.5 ml) at −20°C.

High-performance liquid chromatography (HPLC) was carried out on a Shimadzu LC-2030, equipped with an Econosil C18 column (250 x 4.6 mm, i.d.; particle size: 10 µm) from Alltech. The column was maintained at room temperature and run at a pump speed of 1 ml/min. The mobile phases consisted of A = milli-Q water with 0.1% trifluoracetic acid and B = acetonitrile +0.1% TFA. Gradient methodology was employed using the following method: 20% B for 1 min then raised to 42% over 42 min. It was then raised to 95% B over 0.1 and held there for 4.9 min. B was lowered to 20% in 0.1 min and held for 3.9 min to return the column to equilibrium.

A commercial standard of marinobufagenin (MBG; Cayman Chemicals, catalog # 20798) was reconstituted in methanol. The peak was first identified at 40.45 min. For verification, test samples separate from those analyzed were spiked with commercially acquired marinobufagenin, and the additional peak height was used to confirm the identity of the marinobufagenin peak using the elution protocol described.

The MBG peak for each sample occurred at approximately 40.45 min per the elution gradient described above. The area of each peak was divided by the mass of the dry sample and the peak/mass ratio was used in further analysis. Peak tailing was minimal after the
### Table 1: Cane toad morphological data (2018)

| Variable              | Location     | Number of individuals | Range         | Median | Mean   | St. dev | St. err |
|-----------------------|--------------|-----------------------|---------------|--------|--------|---------|---------|
| **Mass (g)**          | Homestead    | 21                    | 70.00 to 292.00 | 137.00 | 141.43 | 57.18   | 12.48   |
|                       | Miami        | 20                    | 70.00 to 271.00 | 118.50 | 131.40 | 50.55   | 11.30   |
|                       | Ft. Lauderdale | 20                | 44.00 to 158.00 | 93.50  | 98.70  | 31.5    | 7.04    |
|                       | Naples       | 20                    | 53.00 to 211.00 | 112.00 | 109.60 | 39.06   | 8.73    |
|                       | Ft. Myers    | 25                    | 76.00 to 165.00 | 117.00 | 119.72 | 28.64   | 5.73    |
|                       | Port St. Lucie | 20                | 38.00 to 235.00 | 102.50 | 110.90 | 44.6    | 9.97    |
|                       | Lake Placid  | 20                    | 53.00 to 150.00 | 90.50  | 96.25  | 30.48   | 6.82    |
|                       | New Port Richey | 20            | 63.00 to 308.00 | 100.25 | 122.25 | 60.29   | 13.48   |
|                       | DeLand       | 13                    | 44.00 to 386.00 | 123.00 | 153.00 | 98.32   | 27.27   |
| **SVL (mm)**          | Homestead    | 21                    | 91.90 to 148.00 | 111.60 | 112.53 | 13.65   | 2.98    |
|                       | Miami        | 20                    | 90.00 to 189.70 | 108.40 | 113.00 | 21.06   | 4.71    |
|                       | Ft. Lauderdale | 20                | 79.20 to 120.40 | 98.90  | 99.68  | 9.71    | 2.17    |
|                       | Naples       | 20                    | 87.50 to 132.10 | 107.20 | 106.50 | 11.94   | 2.67    |
|                       | Ft. Myers    | 25                    | 92.50 to 122.70 | 108.50 | 108.21 | 8.75    | 1.75    |
|                       | Port St. Lucie | 20               | 70.30 to 121.60 | 103.10 | 101.99 | 11.66   | 2.61    |
|                       | Lake Placid  | 20                    | 81.50 to 115.90 | 98.90  | 97.79  | 10.85   | 2.43    |
|                       | New Port Richey | 20            | 83.50 to 145.20 | 97.80  | 102.63 | 16.42   | 3.67    |
|                       | DeLand       | 13                    | 75.10 to 148.40 | 105.90 | 109.34 | 18.26   | 5.06    |
| **Residual Body Index** | Homestead  | 21                    | −0.12 to 0.09  | 0.02   | 0.00   | 0.06    | 0.01    |
|                       | Miami        | 20                    | −0.54 to 0.10  | <0.01  | −0.02  | 0.14    | 0.03    |
|                       | Ft. Lauderdale | 20                | −0.12 to 0.07  | <0.01  | −0.01  | 0.05    | 0.01    |
|                       | Naples       | 20                    | −0.17 to 0.07  | <0.01  | −0.04  | 0.06    | 0.01    |
|                       | Ft. Myers    | 25                    | −0.11 to 0.05  | <0.01  | −0.01  | 0.04    | 0.01    |
|                       | Port St. Lucie | 20               | −0.09 to 0.29  | <0.01  | 0.01   | 0.08    | 0.02    |
|                       | Lake Placid  | 20                    | −0.06 to 0.08  | 0.01   | 0.01   | 0.04    | 0.01    |
|                       | New Port Richey | 20            | −0.10 to 0.13  | 0.05   | 0.04   | 0.05    | 0.01    |
|                       | DeLand       | 13                    | −0.08 to 0.16  | 0.03   | 0.04   | 0.06    | 0.02    |
| **Gland width (cm)**  | Homestead    | 21                    | 1.10 to 1.94   | 1.5    | 1.54   | 0.24    | 0.05    |
|                       | Miami        | 20                    | 1.06 to 1.78   | 1.38   | 1.37   | 0.18    | 0.04    |
|                       | Ft. Lauderdale | 20                | 0.81 to 1.64   | 1.31   | 1.28   | 0.19    | 0.04    |
|                       | Naples       | 20                    | 0.97 to 2.01   | 1.39   | 1.42   | 0.25    | 0.06    |
|                       | Ft. Myers    | 25                    | 1.13 to 3.30   | 1.41   | 1.51   | 0.42    | 0.08    |
|                       | Port St. Lucie | 20               | 1.04 to 1.86   | 1.5    | 1.49   | 0.23    | 0.05    |
|                       | Lake Placid  | 20                    | 0.94 to 1.71   | 1.36   | 1.37   | 0.23    | 0.05    |
|                       | New Port Richey | 20            | 1.06 to 2.00   | 1.28   | 1.39   | 0.28    | 0.06    |
|                       | DeLand       | 13                    | 0.84 to 2.05   | 1.25   | 1.31   | 0.33    | 0.09    |

(Continues)
optimal gradient elution protocol was established. However, area under the curve was chosen over peak height to accommodate any minor peak tailing or broadening. MBG sample concentrations were determined by the use of a four-parameter standard curve. Using the integrated peak value and a five-point calibration plot, the concentration of MBG (µg/mg) in solution was determined. This concentration was then back-calculated relative to the amount of solution used for sample preparation and adjusted for the dry weight of each sample prior to extraction.

### 2.4 | Sympathetic sensitivity

In May 2019, 58 additional toads \((n = 28 \text{ Miami}, \text{ and } n = 30 \text{ NPR})\) were collected and maintained under laboratory conditions, detailed by Gardner et al. (2020). Following acclimation, the toads were divided into five treatment groups: normal saline solution (NSS) and NSS with 0.125 µm, 0.25 µm, 0.5 µm, or 1 µm epinephrine gram of body per mass. Each treatment group was comprised of six toads collected from Miami and NPR (with the exception of the 0.125 µm treatment group, which had only four toads from Miami). Following injections, each toad was placed in a plastic bin (60.96 x 46.99 x 40 cm) and monitored for signs of poison secretion.

### 2.5 | Statistical analyses

To compare the morphological parameters in Florida populations of cane toads with those observed in a previous study by Phillips and Shine (2005), a model using the lm function in R (version 3.6.3), using latitude coordinates (decimal degrees) as a covariate and sex as a factor, was used to assess differences in log_{10}-transformed SVL measurements of toads. Log_{10}-transformed body masses of toads collected from the nine populations sampled during 2018 were then regressed against SVL measurements (also log_{10}-transformed) to obtain residual body condition index measures (Denoël et al., 2002). This measure follows a normal distribution that is independent of SVL (Plăiaşu et al., 2010) and has commonly been used to assess

| Variable          | Location    | Number of individuals | Range       | Median   | Mean   | St. dev | St. err |
|-------------------|-------------|-----------------------|-------------|----------|--------|---------|---------|
| Gland length (cm) | Homestead   | 21                    | 2.10 to 4.40| 2.89     | 2.91   | 0.53    | 0.12    |
|                   | Miami       | 20                    | 1.92 to 3.25| 2.64     | 2.70   | 0.33    | 0.07    |
|                   | Ft. Lauderdale | 20                | 1.78 to 2.72| 2.38     | 2.38   | 0.24    | 0.05    |
|                   | Naples      | 20                    | 2.16 to 3.09| 2.48     | 2.56   | 0.27    | 0.06    |
|                   | Ft. Myers   | 25                    | 2.05 to 3.51| 2.46     | 2.57   | 0.37    | 0.07    |
|                   | Port St. Lucie | 20              | 1.77 to 3.40| 2.74     | 2.70   | 0.43    | 0.10    |
|                   | Lake Placid | 20                    | 1.34 to 3.19| 2.47     | 2.46   | 0.40    | 0.09    |
|                   | New Port Richey | 20             | 2.08 to 3.83| 2.47     | 2.62   | 0.44    | 0.10    |
|                   | DeLand      | 13                    | 1.98 to 3.45| 2.81     | 2.76   | 0.41    | 0.11    |
| Gland area        | Homestead   | 21                    | 1.61 to 5.14| 2.67     | 2.98   | 0.96    | 0.21    |
|                   | Miami       | 20                    | 1.57 to 3.68| 2.23     | 2.39   | 0.60    | 0.13    |
|                   | Ft. Lauderdale | 20               | 1.08 to 2.82| 1.93     | 1.94   | 0.47    | 0.11    |
|                   | Naples      | 20                    | 1.43 to 4.69| 2.29     | 2.40   | 0.71    | 0.16    |
|                   | Ft. Myers   | 25                    | 0.93 to 3.74| 2.24     | 2.27   | 0.62    | 0.12    |
|                   | Port St. Lucie | 20             | 1.12 to 4.31| 2.64     | 2.67   | 0.82    | 0.18    |
|                   | Lake Placid | 20                    | 1.34 to 3.47| 2.22     | 2.23   | 0.60    | 0.13    |
|                   | New Port Richey | 20            | 1.29 to 5.21| 1.83     | 2.30   | 1.01    | 0.23    |
|                   | DeLand      | 13                    | 1.21 to 4.60| 2.45     | 2.52   | 0.96    | 0.27    |
| Gland size        | Homestead   | 21                    | −0.2 to 0.41| 0.06     | 0.09   | 0.16    | 0.03    |
|                   | Miami       | 20                    | −0.23 to 0.22| −0.03    | −0.01  | 0.12    | 0.03    |
|                   | Ft. Lauderdale | 20             | −0.42 to 0.08| −0.11    | −0.12  | 0.12    | 0.03    |
|                   | Naples      | 20                    | −0.24 to 0.32| −0.03    | −0.02  | 0.13    | 0.03    |
|                   | Ft. Myers   | 25                    | −0.24 to 2.34| −0.04    | 0.11   | 0.52    | 0.10    |
|                   | Port St. Lucie | 20            | −0.39 to 0.28| 0.05     | 0.03   | 0.17    | 0.04    |
|                   | Lake Placid | 20                    | −0.30 to 0.18| −0.03    | −0.06  | 0.14    | 0.03    |
|                   | New Port Richey | 20          | −0.30 to 0.40| −0.12    | −0.05  | 0.19    | 0.04    |
|                   | DeLand      | 13                    | −0.20 to 0.41| 0.02     | −0.01  | 0.19    | 0.05    |
RESULTS

3.1 Morphology and likelihood of secretion (2018)

There was no effect of latitude (−0.008 [± 0.004], t_{175} = −2.06, p = .08) or sex (0.002 [± 0.008], t_{176} = 0.24, p = 1.0 for 110 males compared to 69 females) on sizes of cane toads collected during 2018 (values represent mean [±] standard error). BI of male toads measured in 2018 did not differ from females (−0.006 [± 0.010], t_{175} = −0.638, p = 1.0), and BI was also not affected by toad size (0.055 [±0.09], t_{175} = 0.59, p = 1.0). BI did however increase in collected toads by 0.020 (± 0.005) (t_{175} = 3.87, p < .01) with each degree of increasing latitude among the cane toad populations. Parotoid gland size was not affected by BI (0.57 [± 0.24], t_{174} = 2.37, p = .08), or latitude (−0.010 [± 0.017], t_{174} = 0.58, p = 1.0). BI was also not different for male compared to female toads (0.011 [± 0.034], t_{174} = 0.32, p = 1.0), although gland size increased significantly (1.92 [± 0.30], t_{174} = 6.50, p < .01) with increasing toad size (for the full list of sample sizes, means, standard errors, data ranges, and medians for morphological data collected in 2018 see Table 1). There was no effect of BI (0.200 [95% C.L. = 0.0005–72.84], z_{173} = 0.55, p = 1.0), gland size (2.48 [95% C.L. = 0.54–11.28], z_{173} = 1.20, p = 1.0), or toad size (2.09 [95% C.L. = 0.0007–6.02276], z_{173} = 0.19, p = 1.0) on the likelihood of a toad to secrete poison upon capture and handling. Male toads were also no more likely to secrete poison compared to female cane toads (3.03 [95% C.L. = 1.11–8.24], z_{173} = 2.22, p = .13); however, cane toads were 1.94 (95% C.L. = 1.29–2.91; z_{173} = 3.24, p < .01) times more likely to secrete poison with each degree of increasing latitude (Figure 2).

3.2 MBG concentration and sympathetic sensitivity (2019)

MBG concentrations assessed using HPLC were not affected by BI (247.34 [± 185.45], t_{21} = 1.33, p = 1.0), toad size (469.32 [± 317.21], t_{21} = 1.48, p = .92), or gland size (−157.10 [± 100.70], t_{21} = −1.56, p = .80). MBG concentrations for Miami cane toads were not different compared to those from NPR (47.38 [± 20.87], t_{21} = −2.27, p = .21), or DeLand (−7.43 [± 29.57], t_{21} = 0.25, p = 1.0). MBG concentrations were also not different for NPR compared to DeLand cane toads (54.81 [± 30.91], t_{21} = 1.77, p = .54) (for the full list of...
sample sizes, means, standard errors, data ranges, and medians for morphological and MBG concentration data collected in 2019 see Table 2). Although laboratory-acclimated toads \( n = 28 \) Miami, 30 NPR) were 9.54 (95% C.L. = 1.80–50.58; \( z_{SS} = 2.71, p = .01 \)) times as likely to secrete poison with increasing epinephrine dose (Figure 3), the likelihood of cane toads secreting poison was not affected by locality (0.76 (95% C.L. = 0.23–2.55), \( z = −0.45, p = 1.0 \)).

4 | DISCUSSION

The cane toad range we examined in Florida extends approximately 480 km northward from the original introduction site in southern Florida. Previous studies have indicated that cane toad size (log10-transformed SVL) and relative parotoid gland size increases in populations near the expanding edge of the invaded range in Australia (Phillips & Shine, 2005), where cane toads have established approximately 1,600 km to the northwest (Brown et al., 2015) and 2,200 km to the southeast (Urban et al., 2008) since their original introduction in 1935. We found that neither cane toad size nor the relationship with gland size was affected by latitude for Florida populations. Although initially our results seem contradictory to the results of Phillips and Shine (2005), the time since colonization for these populations ranges from approximately 65 years for the Homestead and Miami populations to 29 years for the northern populations such as NPR (U. S. Geological Survey, 2020). Although it is unclear from our current data if the Florida populations initially showed the same pattern of increased size and gland morphology upon their initial establishment throughout the state (as time since colonization was shown to significantly affect body size and gland morphology in invasive Australian populations), it is possible that the Florida populations we sampled have now been present for long enough that increased body or parotoid gland sizes no longer afford a selective advantage (Phillips & Shine, 2005).

Unlike body and parotoid gland sizes, we did observe a significant effect of latitude on BI. Studies comparing growth rates of amphibians against latitudinal or altitudinal gradients have shown that individuals from higher latitude populations show higher growth efficiency compared to those from lower latitudes (Lindgren & Laurila, 2005), perhaps as a mechanism to mitigate the effects of cooler temperatures in higher latitudes (Berven, 1982). In addition, invasive species have been postulated to escape from the pathogens and parasites of their native range upon introduction to novel habitats (Lee and Klasing, 2004), with newly invading populations having

| Variable | Location | Number of individuals | Range        | Median   | Mean     | St. dev | St.err |
|----------|----------|-----------------------|--------------|----------|----------|---------|--------|
| Mass (g) | Miami    | 10                    | 97.00 to 183.00 | 144.00   | 144.20   | 30.96   | 9.79   |
|          | New Port Richey | 10               | 87.00 to 224.00 | 168.50   | 168.30   | 42.14   | 13.33  |
|          | DeLand   | 7                    | 17.00 to 169.00 | 86.00    | 76.43    | 59.35   | 22.43  |
| SVL (mm) | Miami    | 10                    | 96.70 to 126.70 | 113.15   | 111.48   | 12.57   | 3.97   |
|          | New Port Richey | 10               | 96.60 to 137.20 | 115.15   | 116.56   | 12.57   | 3.97   |
|          | DeLand   | 7                    | 57.20 to 117.20 | 97.70    | 87.01    | 24.93   | 9.42   |
| Residual Body Index | Miami    | 10                    | −0.12 to 0.06  | <0.01    | <0.01    | 0.05    | 0.02   |
|          | New Port Richey | 10               | −0.06 to 0.11  | 0.02     | 0.01     | 0.05    | 0.02   |
|          | DeLand   | 7                    | −0.06 to 0.01  | −0.03    | −0.02    | 0.02    | 0.01   |
| Gland Width (cm) | Miami    | 10                    | 1.11 to 2.06   | 1.63     | 1.57     | 0.28    | 0.09   |
|          | New Port Richey | 10               | 1.35 to 2.04   | 1.79     | 1.78     | 0.20    | 0.06   |
|          | DeLand   | 7                    | 0.88 to 1.77   | 1.35     | 1.31     | 0.43    | 0.16   |
| Gland length (cm) | Miami    | 10                    | 2.22 to 3.71   | 2.89     | 2.96     | 0.49    | 0.15   |
|          | New Port Richey | 10               | 2.33 to 3.81   | 3.27     | 3.24     | 0.42    | 0.13   |
|          | DeLand   | 7                    | 1.28 to 3.31   | 2.61     | 2.37     | 0.85    | 0.32   |
| Gland area | Miami    | 10                    | 1.78 to 4.39   | 2.70     | 3.00     | 0.84    | 0.27   |
|          | New Port Richey | 10               | 1.80 to 4.44   | 3.50     | 3.28     | 0.84    | 0.27   |
|          | DeLand   | 7                    | 0.61 to 3.37   | 1.92     | 1.94     | 1.22    | 0.46   |
| Gland Size | Miami    | 10                    | −0.29 to 0.19  | −0.07    | −0.03    | 0.16    | 0.05   |
|          | New Port Richey | 10               | −0.26 to 0.20  | 0.08     | 0.04     | 0.14    | 0.04   |
|          | DeLand   | 7                    | −0.75 to 0.15  | −0.14    | −0.25    | 0.39    | 0.15   |
| MBG (µg/g dry secretion) | Miami    | 10                    | 44.2 to 164.10 | 131.50   | 123.40   | 38.83   | 12.28  |
|          | New Port Richey | 10               | 5.3 to 153.50  | 80.75    | 74.30    | 49.74   | 15.73  |
|          | DeLand   | 7                    | 23.8 to 188.50 | 131.90   | 113.37   | 52.30   | 19.77  |
a lower parasitic load due to less "carry-over" (Torchin et al., 2003). This "lag" has been observed in Australian cane toad populations undergoing expansion (Phillips et al., 2010). Although the effects of parasitic burden on BI in amphibians are variable (Moretti et al., 2014, altered burdens and other pressures associated with northward expansion (Phillips et al., 2010). Although parasitic burdens have not been well documented in the cane toad populations of Florida, altered burdens and other pressures associated with northward establishment could further explain the elevated BI in the northern populations.

Although we found no differences in parotoid gland size relative to body size in toads from different populations we sampled from Florida, there was an increase in the likelihood that cane toads from higher latitudes would secrete poison upon handling or disturbance. Physiologically elevated glucocorticoid levels induce phenylethanolamine-N-methyltransferase (PNMT) (an enzyme in chromaffin cells of the adrenal) to synthesize epinephrine from norepinephrine (Wurtman, 2002). Baseline corticosterone levels of the cane toads collected in 2018 were shown to increase with increasing latitude (Assis et al., 2020). Elevated corticosterone levels may have contributed to the amount of epinephrine available stored and subsequently released upon capture in toads from this study and therefore to the likelihood of their secreting poison in the field. Mechanisms such as modified methylation patterns (Szym et al., 2005) or altered receptor expression (Martin et al., 2017) that modulate sympathetic sensitivity are affected by early life experiences (Yao et al., 2008). As development of cane toad tadpoles occurs more slowly in cooler temperatures (Wijethunga et al., 2016), and tadpoles have been observed to readily cannibalize each other during development (Crossland and Shine, 2011), prolonged development or other altered selection pressures on northern Florida populations may underlie these sympathetic differences. Additionally, Australian cane toads replenishing secretions following manual compression of the parotoid glands were shown to disperse more slowly than individuals that had not secreted poison (Blennerhassett et al., 2019). This combined with the increased number of potential predators (Meshaka, 2011; Punzo & Lindstrom, 2001) may contribute to range limits of cane toads in the United States.

We found no differences in MBG concentrations of cane toads from northern populations (NPR and Deland) compared to the southern Miami population. Adults have been shown to possess five primary compounds (arenobufagin, bufalin, marinobufagin, resibufagenin, and telocinobufagin) (Hayes et al., 2009). These compounds have differing toxicities, with bufalin and telocinobufagin having higher IC50 values relative to the other BDs (Kamano et al., 1998). Captive-raised Australian cane toads exposed to nonlethal predator cues during larval development were shown to shift poison content toward increased amounts of bufalin compared to other compounds following metamorphosis, suggesting investment into synthesis of more lethal compounds in the presence of stronger predation cues (Hagman et al., 2009). Further research evaluating whether synthesis of more lethal compounds is higher in northern populations, complementing the higher likelihood of secretion when disturbed, would be of interest.

While the likelihood of secretion in laboratory-acclimated toads collected during 2019 increased with increasing doses of epinephrine, there was no difference in likelihood of secretion between northern (NPR) and southern (Miami) toad populations. The similarities in sympathetic sensitivity may have resulted from the toads being removed from the field and acclimated to the same laboratory environment prior to epinephrine injection, although altered corticosterone responses to novel stressors were still observed between toads from these populations in the study of metabolism and immune response by Gardner et al. (2020). Further studies to assess whether differences in sympathetic sensitivity between northern and southern populations are related to epinephrine release could help to elucidate the responses to injected epinephrine.

The results of this study indicate that differences in sympathetic sensitivity have arisen in cane toad populations experiencing northward dispersal in Florida. It remains unclear whether BD concentrations have been affected by the northern latitude or whether there are trade-offs between MBG and other BDs in the parotoid gland secretion. As MBG was the only BD we examined, future studies quantifying a more diverse array of BDs in cane toad secretions are needed to more accurately assess overall toxicity, and the resulting risk to native predators. Quantifying methylation patterns from populations spanning the invaded range in Florida and assessing factors leading to the differences in sympathetic
sensitivity would also provide further insights, as would ecological studies assessing predation risk. Greater sampling of individuals across life-history stages is also necessary to determine how the volume and composition of secretion changes ontogenetically or by region.

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CONFLICT OF INTEREST
None declared.

AUTHOR CONTRIBUTION
Steven T Gardner: Conceptualization (lead); Formal analysis (lead); Investigation (equal); Methodology (equal); Visualization (lead); Writing-original draft (lead); Writing-review & editing (equal). Megen Kepas: Conceptualization (supporting); Formal analysis (supporting); Investigation (equal); Methodology (equal); Validation (equal); Writing-review & editing (equal). Casey Simons: Investigation (supporting); Methodology (supporting); Validation (equal); Writing-review & editing (supporting). L M Horne: Investigation (supporting); Methodology (supporting); Writing-review & editing (supporting). Alan H. Savitzky: Conceptualization (supporting); Formal analysis (supporting); Project administration (equal); Supervision (equal); Methodology (equal); Visualization (lead); Writing-original draft (lead); Writing-review & editing (equal). Mary T. Mendonca: Conceptualization (supporting); Supervision (equal); Writing-review & editing (equal).

DATA AVAILABILITY STATEMENT
The data and files supporting the results are archived in Mendeley Data: https://doi.org/10.17632/xr6rpmnc75.1.

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