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RESEARCH ARTICLE

Biomechanical behaviour of lizard osteoderms and skin under external loading

Loïc Kéver^{1,*}, Damien Olivier^{2,3}, Arsalan Marghoub⁴, Susan E. Evans⁵, Matthew K. Vickaryous⁶, Mehran Moazen⁴ and Anthony Herrel¹

ABSTRACT

Many species of lizards are partially enveloped by a dermal armour made of ossified units called osteoderms. Lizard osteoderms demonstrate considerable species-specific variation in morphology and histology. Although a physical/protective role (against predators, prey, conspecifics and impact loading during falls) is frequently advanced, empirical data on the biomechanics of lizard osteoderms are scarce, limiting our understanding of form–function relationships. Here, we report deformation recorded at the surface of temporal osteoderms during controlled external loading of preserved specimens of 11 lizard species (*Tiliqua rugosa*, *Tiliqua scincoides*, *Corucia zebrata*, *Pseudopus apodus*, *Timon lepidus*, *Matobosaurus validus*, *Broadleysaurus major*, *Tribolonotus gracilis*, *Tribolonotus novaeguineae*, *Heloderma horridum* and *Heloderma suspectum*). Based on the strain recorded *in situ* and from isolated osteoderms, the skin of the species investigated can be ranked along a marked stiffness gradient that mostly reflects the features of the osteoderms. Some species such as *T. rugosa* and the two *Heloderma* species had very stiff osteoderms and skin while others such as *T. lepidus* and *P. apodus* were at the other end of the spectrum. Histological sections of the osteoderms suggest that fused (versus compound) osteoderms with a thick layer of capping tissue are found in species with a stiff skin. In most cases, loading neighbouring osteoderms induced a large strain in the instrumented osteoderm, attesting that, in most species, lizard osteoderms are tightly interconnected. These data empirically confirm that the morphological diversity observed in lizard osteoderms is matched by variability in biomechanical properties.

KEY WORDS: Bone, Functional morphology, Squamata, Osteoderms, Stiffness, Strain gauges

INTRODUCTION

Animals can be subjected to traumatic events that damage their tissues, sometimes with lethal consequences (e.g. Jennings, 2009; Tanke and Currie, 1998; Thomas and Cole, 1996). Physical stresses are generated in biological tissues when an individual impacts the ground or objects in its environment, but damage may also occur


under the action of an opponent's or predator's claws, teeth, beak, tail, hoofs or horns, for example (e.g. Mukherjee and Heithaus, 2013; Song et al., 2011). As such strong loading regimes likely induce different patterns of loading, one could hypothesize that they probably create a wide range of species-specific selective pressures and, consequently, may have promoted the diversity of protective structures that have evolved in animals (see also Connors et al., 2019). For example, loading during agonistic interactions and falls likely constrained the cranial design of primates (e.g. Carrier and Morgan, 2014; Hylander and Johnson, 1997; Hylander et al., 1991).

A potential shielding strategy against traumatic events involves the incorporation of dermal bony plates also known as osteoderms (Vickaryous and Sire, 2009). Such plates are present in several distantly related vertebrate taxa (Yang et al., 2013), yet are particularly common and diverse among extant lizards (Vickaryous and Sire, 2009; Williams et al., 2021). Osteoderms with various shapes and types of organization have been reported for many lizard families (see Williams et al., 2021, for an exhaustive list) including scincids (e.g. Canei and Nonclercq, 2020; Oliver, 1951), lacertids (Arnold, 1973, 1989), gekkonids (e.g. Laver et al., 2020; Paluh et al., 2017), anguids (e.g. Strahm and Schwartz, 1977; Zylberberg and Castanet, 1985), gerrhosaurids and cordylids (e.g. Broeckhoven et al., 2018a; Marques et al., 2019).

Lizard osteoderms have classically been considered to have a protective function against predators (Williams et al., 2021), but this simplistic view has been challenged, notably by results from phylogenetically informed comparative studies on cordylids (Broeckhoven, 2022; Broeckhoven et al., 2018a,b; Stanley, 2013). Indeed, the covariation between osteoderm expression, distribution or morphology and ecological factors such as climate suggests that osteoderms likely have a multi-functional nature including thermoregulation (e.g. Clarac et al., 2019; but see Inacio Veenstra and Broeckhoven, 2022) and mineral storage (Broeckhoven and du Plessis, 2022). To date, our understanding of osteoderm function(s) suffers from a lack of data relating osteoderm structural features and their biomechanical and thermal properties. The sole empirical data available for lizard osteoderms were provided by Broeckhoven et al. (2015, 2017) and suggest that the resistance of cordylid skin to predator bites increases with osteoderm thickness. Simulations using single osteoderms reconstructed from high resolution micro-computed tomography (μ CT) scans have also proven insightful (Broeckhoven et al., 2017; Iacoviello et al., 2020), highlighting, for example, how vascularization or material density can impact stress magnitude and distribution. Surprisingly, two simpler yet fundamental questions have never been addressed: (1) are there interspecific differences in the *in toto* deformation of lizard osteoderms under external loading?; and (2) can these differences be quantified? Aside from improving our understanding of osteoderm function and evolution, answers to these questions would also be useful for applied research. Lizard osteoderms are

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regarded as valuable models for the development of biomimetic materials such as protective clothing (Broeckhoven et al., 2017; Iacoviello et al., 2020; Liang et al., 2021) but are rarely the topic of biomimetic studies. Insights into the biomechanics of the lizard body ‘armour’ and the forces that have driven the evolution of their diversity could provide valuable insights for the bioinspiration of protective materials.

In this study, we measured deformation in morphologically different osteoderms. To do so, we instrumented temporal osteoderms of formalin-fixed specimens from 11 species of lizards with different osteoderm shapes and distribution patterns (Fig. 1). Next, we applied two loading regimes roughly mimicking either a bite or an impact on different locations of the head. Our goals were to test: (1) whether loading applied at different locations of the head

generated strain in the instrumented osteoderm, (2) whether species differed in the relative stiffness of their osteoderm and skin, and (3) whether the two loading regimes (‘static’ versus ‘dynamic’) differed, thus providing insights into the morphological features and the patterns of organization driving variation in osteoderm stiffness.

MATERIALS AND METHODS

Sampling

Our experimental setup required specimens with at least one osteoderm that was larger than the smallest strain gauge available. We worked on temporal osteoderms because they are generally larger than the body and tail osteoderms and are not fused with skull bones. Data were collected for 11 species with 2–6 individuals per species (Table 1). All the specimens were adults that had been

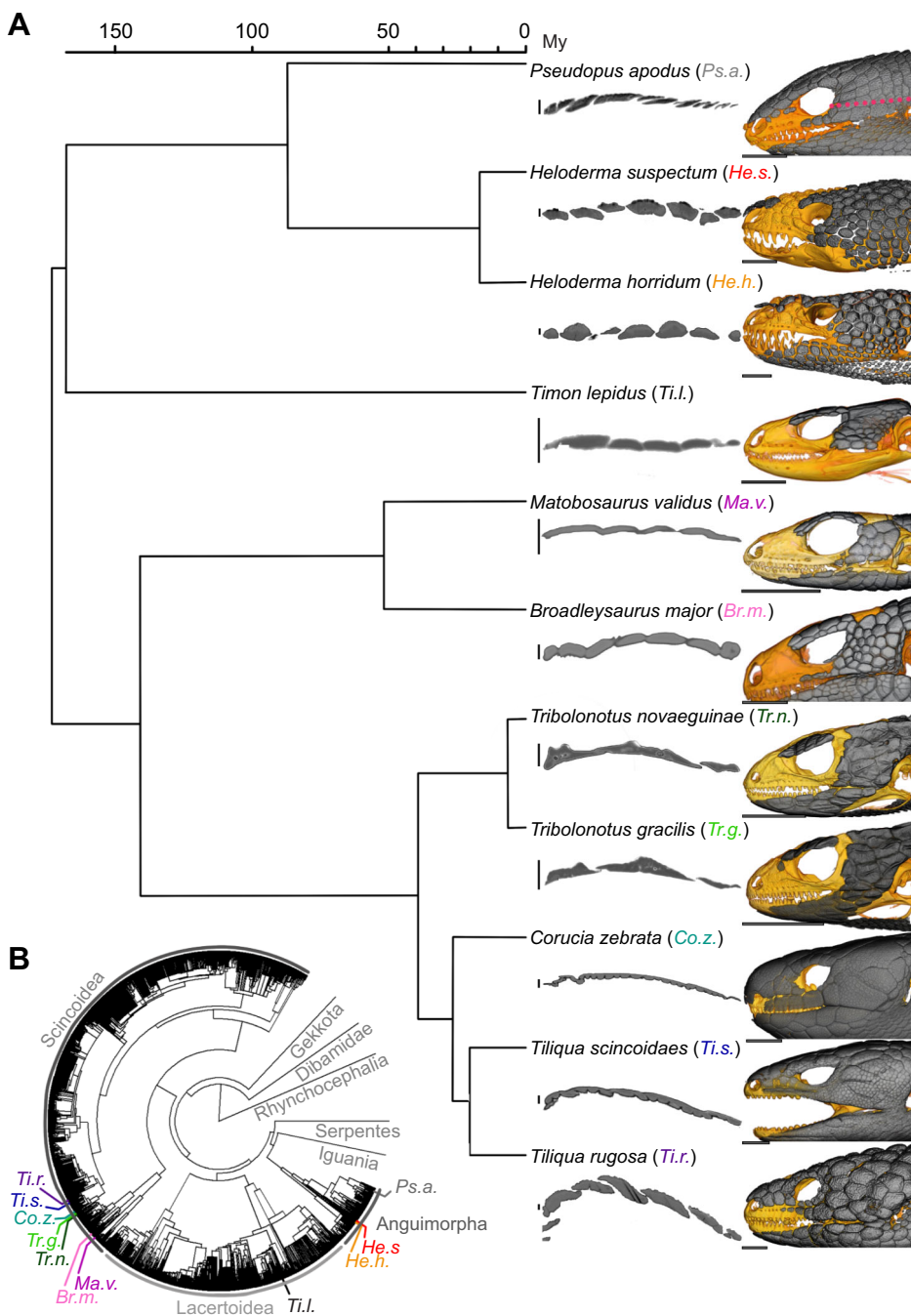


Fig. 1. Distribution, phylogenetic relationships and osteoderm morphology of the studied species. (A) Phylogenetic relationships (based on Tonini et al., 2016) and morphology of the temporal osteoderms in the 11 sampled species. A 3D reconstruction of the skull (orange–yellow) and head osteoderms (grey) of these species is shown on the right. Virtual longitudinal section (under the species names) through temporal osteoderms from a ventral view and along the rostro-caudal axis (illustrated by the dotted magenta line on the 3D reconstruction of the *Pseudopus apodus* head). Scale bars: 1 mm (vertical black bars) and 1 cm (horizontal black and grey bars). (B) Distribution of the sampled species within Lepidosauria. Except for *Pseudopus apodus* (MNHN, FUNEVOL virtual collection) and *Heloderma suspectum* (UCL, Evans Lab), micro-computed tomography (μ CT) raw images from the studied species were downloaded from MorphoSource [*Heloderma horridum*: ark:/87602/m4/M98538, oUTCT, funded by National Science Foundation (NSF) EF-0334961; *Timon lepidus*: ark:/87602/m4/M72278, Yale Peabody Museum, funded by NSF DBI-1701714, NSF DBI-1702263, oVert TCN, NSF DBI-1701769; *Broadleysaurus major*: ark:/87602/m4/M74672, California Academy of Sciences, funded by oVert TCN; *Matobosaurus validus*: ark:/87602/m4/M71913, California Academy of Sciences, funded by oVert TCN; *Tribolonotus novaeguinae*: ark:/87602/m4/M61910, University of Michigan Museum of Zoology, funded by NSF DBI-1701714, NSF DBI-1701735, oVert TCN; *Tribolonotus cf. gracilis* (listed as *Tribolonotus* sp. on MorphoSource): ark:/87602/m4/M40298, The oVert Thematic Collections Network, funded by oVert Thematic Collections Network (TCN), NSF DBI-1701714; *Corucia zebrata*: ark:/87602/m4/M43624, California Academy of Sciences Herpetology collection, funded by oVert Thematic Collections Network (TCN), NSF DBI-1701714, 1701870; *Tiliqua scincoides*: ark:/87602/m4/M74717, funded by NSF DBI-1701714, NSF DBI-1701870, NSF DBI-1701713, oVert TCN; *Tiliqua rugosa*: ark:/87602/m4/M48823, Florida Museum of Natural History (University of Florida), funded by oVert TCN: NSF DBI1701714].

Table 1. Family, species, specimen and collection ID, snout–vent length (SVL), head length (HL) and sex of the sampled specimens

Family	Species	Specimen	Collection ID	SVL (mm)	HL (mm)	Sex
Anguillidae	<i>Pseudopus apodus</i> (Pallas 1775)	Psa_1	Herrel psa1	375	39	U
		Psa_2	MNHN 2619	418	45	M
		Psa_3	Herrel psa2 (head)	NA	36	U
		Psa_4	MNHN 1939 245	387	37	U
		Psa_5	MNHN 1912 355	342	35	U
		Psa_6	MNHN 1884 519	371	32	U
Helodermatidae	<i>Heloderma horridum</i> (Wiegmann 1829)	Heh_1	Herrel heh2	317	53	F
		Heh_2	MNHN 1201	271	47	F
	<i>Heloderma suspectum</i> Cope 1869	Hes_1	MNHN 133138	259	44	U
		Hes_2	MNHN 1950 174	279	49	U
Lacertidae	<i>Timon lepidus</i> (Daudin 1802)	Til_1	MNHN 1978 963	161	44	M
		Til_2	MNHN 1918-91	196	52	M
		Til_3	Herrel til1	149	41	M
		Til_4	MNHN 1922 325	119	33	M
Gerrhosauridae	<i>Broadleysaurus major</i> (Duméril 1851)	Brm_1	MNHN 1050	194	38	M
		Brm_2	MNHN 2788	199	35	M
		Brm_3	MNHN 2786	211	34	M
	<i>Matobosaurus validus</i> (Smith 1849)	Mav_1	Herrel mav1	216	40	M
		Mav_2	MNHN: 1987 1813	209	48	M
Scincidae	<i>Corucia zebrata</i> Gray 1855	Coz_1	Herrel coz1	275	46	F
		Coz_2	Herrel coz2	211	45	F
		Coz_3	MNHN 1990-4733	270	58	F
		Coz_4	MNHN 4413	246	41	F
	<i>Tiliqua rugosa</i> (Gray 1825)	Tir_1	Herrel tir1	288	55	M
		Tir_2	Herrel tir2	275	44	F
		Tir_3	MNHN 9047	227	52	F
		Tir_4	MNHN 56-103	265	46	M
	<i>Tiliqua scincoides</i> (White 1790)	Tis_1	Herrel tis1	310	49	M
		Tis_2	Herrel tis2	230	47	M
		Tis_3	MNHN 94534	307	58	F
		Tis_4	MNHN 1982-511	284	54	F
		Tis_5	MNHN 1911-194	269	47	F
		Tis_6	MNHN 1904-140	250	45	F
	<i>Tribolonotus gracilis</i> de Rooij 1909	Trg_1	Herrel trg1	106	26	U
		Trg_2	Herrel trg2	109	24	U
		Trg_3	Herrel trg3	106	22	U
		Trn_1	Herrel trn1	103	22	U
	<i>Tribolonotus novaeguineae</i> (Schlegel 1834)	Trn_2	Herrel trn2	90	22	U

F, female; M, male; U, undetermined.

formalin fixed and preserved in 70% ethanol, and are housed at the Muséum National d'Histoire Naturelle (MNHN) in Paris. Small differences in the fixation protocol (e.g. time spent in formalin, formalin concentration) cannot be excluded for the specimens from the MNHN collections, causing the intraspecific variability to increase.

'In toto' experiment

Instrumentation of the target osteoderm

An osteoderm located in the temporal area was instrumented for every specimen (Fig. 2). The epidermis and the periosteum of the target osteoderm were removed with a scalpel and the surface of the osteoderm was cleaned with hydrogen peroxide, allowing the strain gauge to be glued to the external surface of the osteoderm using cyanoacrylate glue.

The smallest model of rectangular rosette strain gauge (item code: MMF402103; gauge total surface: 1.33 mm×2.9 mm; grid resistance: 350 Ω; gauge factors: 1.7–1.9) manufactured by Micro-Measurements (Vishay Measurements Group France, Chartres, France) was selected for these experiments. The rosette configuration was favoured because it bears three recording units and therefore can provide tensile, compressive and shear strain in every direction of the gauge plane which here corresponds to the external surface of the instrumented osteoderm.

Loading of the osteoderms

The target osteoderm was loaded by applying an external force orthogonally to the surface at 14 different locations of the animal head (Fig. 3A). Loading was applied directly on the instrumented osteoderm (when the osteoderm surface was not entirely covered by the gauge), on several neighbouring osteoderms, and to the rostral area. First, force was applied for a very short period of time (<0.1 s; Fig. 2B) using a piezoelectric hammer (Model 086E80, PCB Piezotronics S.A., Saint Aubin, France). As the force–time slope was very steep, we refer to this loading regime as 'dynamic' loading. Second, force was applied slowly and for a relatively long period of time (>0.5 s; Fig. 2B) using a metal pin mounted on a Kistler piezo-sensor. The signal generated by the piezo-electric sensor and amplified by a 5011B charge amplifier (Kistler Instrumente AG, Winterthur, Switzerland) provided the magnitude of the load. As the force was gently increased by hand, we will refer to this experimental set-up as the 'static' loading condition hereafter. For each loading location, the procedure was repeated on average 33 times for the static loading and 104 times for the dynamic loading. More data points were collected during the dynamic loading because of the apparent lower consistency in the recording of the force magnitude. In both cases, care was taken to administer forces across the widest possible range of magnitudes (0.01–36.5 N for the static loading and 0.01–5.3 N for the dynamic loading).

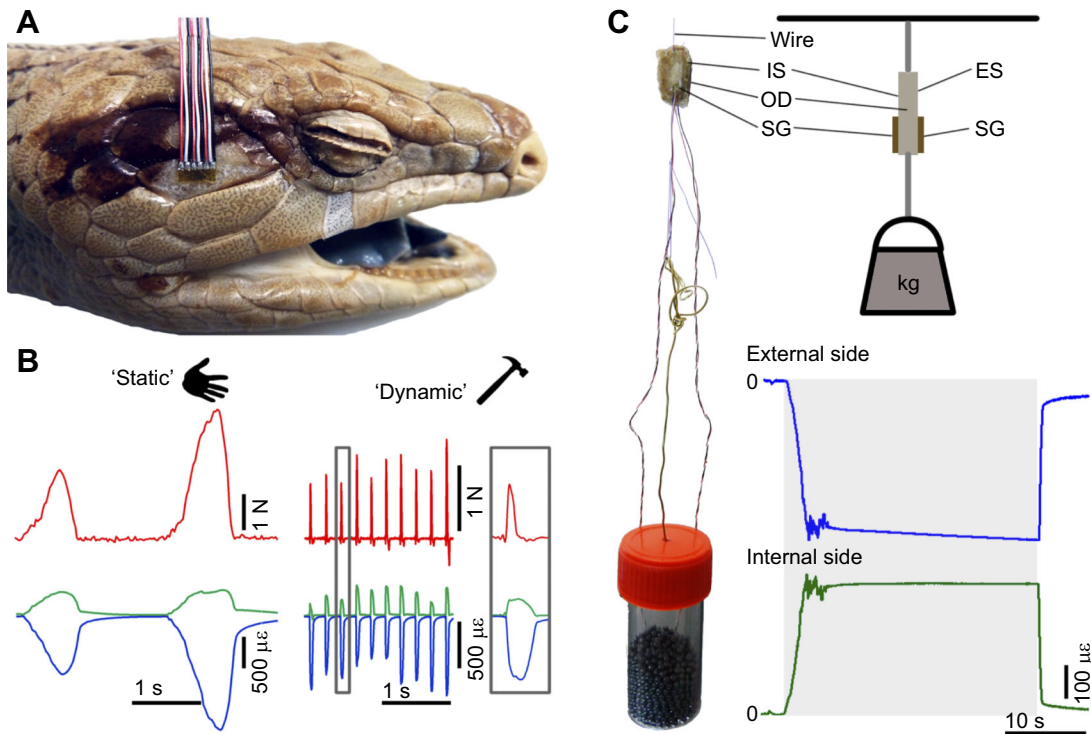


Fig. 2. Osteoderm loading and strain recordings. (A) Temporal osteoderm of a *T. scincoides* instrumented with a rectangular rosette strain gauge. Loading was obtained either by manually pressing a pin mounted on a piezoelectric sensor or by using a small piezoelectric hammer. (B) Illustration of the force (red), tensile strain (green) and compressive strain (blue) waveforms recorded during the experiment illustrated in A. The hand symbol illustrates the relatively 'static' loading obtained when the pin was pressed manually, whereas the hammer symbol means that the loading was more dynamic and was obtained by using the piezoelectric hammer. (C) 'Tensile' test experiment on a dissected osteoderm of *C. zebata*. An image (left) and schematic representation (top right) of the experimental setup are shown, with waveforms recorded from the external and internal sides of the osteoderms using single-element gauges. The grey area highlights the time period during which the osteoderm was loaded using a 200 g weight. ES, external side of the dissected osteoderm; IS, internal side of the dissected osteoderm; OD, osteoderm; SG, single-element gauge.

'Tensile test' experiment

Strain recorded *in toto* may not only reflect interspecific differences in the features of the osteoderms themselves. Indeed, the distribution of stresses between osteoderms may be dependent on the thickness and nature of the tissues underlying the osteoderms. Therefore, we tested whether similar trends could be observed in an experimental setup with fewer confounding factors. Six specimens (two *Pseudopus apodus*, one *Heloderma horridum*, one *Heloderma suspectum*, two *Tiliqua scincoides*, two *Tiliqua rugosa* and two *Corucia zebata*) with at least one temporal osteoderm that was large enough to drill small holes on its rostral and caudal ends (see Fig. 2C) were selected for this experiment. In each case, temporal osteoderms of one side of the head were dissected out and the largest osteoderm was cleaned of any underlying soft tissue and instrumented. Note that, because of the destructive nature of dissections, several specimens from the MNHN collections that are listed in Table 1 were not included in the sampling for this experiment.

Next, two single-element gauges (item code: MMF402183, Micro-Measurements; gauge total surface: 2.41 mm×2.6 mm; 350 Ω; gauge factor: 1.9) were glued to the dissected osteoderm, one on its external side and the other on its internal side. Single gauges can only provide compressive, tensile and shear strain along their long axis; here, the anteroposterior axis of the osteoderm. They were, however, appropriate considering the goal of this second set of experiments and the loading applied to the osteoderms.

Two small holes (0.8 mm) were drilled through the osteoderms. They were located along the anteroposterior axis of the

osteoderm near the edges. One of these holes was used to hang the osteoderm from a stand (Fig. 2C). The second hole was used to load the osteoderm with a series of weights (100 g, 200 g, 500 g and 700 g) using a thin wire. Consequently, the force vector resulting from the weight and the long axis of the gauge were aligned. The force applied to the osteoderm was obtained by multiplying the mass by the gravitational acceleration ($F=m \times g$). In a second experimental setup, the osteoderm was held horizontally using forceps and the weight vector was oriented perpendicular to the anteroposterior axis of the osteoderm (Fig. S1). This experiment was conducted on an even smaller sample of individuals because some osteoderms were too small to be held using forceps.

The instrumented osteoderms were prepared following the polishing protocol reported by Marghoub et al. (2022) and sectioned along the axis of the force vector in a way that ensured the drilled holes were in view. Sections performed at the level of the gauges were imaged using a KEYENCE Vhx 7000 Digital Microscope to illustrate the thickness and shape of the osteoderms, and the multi-partite morphology of compound osteoderms (characteristic of some skink species), and to document the presence and proportion of capping tissue (i.e. osteodermine) described by Marghoub et al. (2022) and Kirby et al. (2020).

Strain recording and analysis

The gauges were connected with a 2310 signal conditioning amplifier (Vishay, Malvern, PA, USA) set up to feed the gauges with a 1.4 V current. Changes in the resistance of the gauges caused

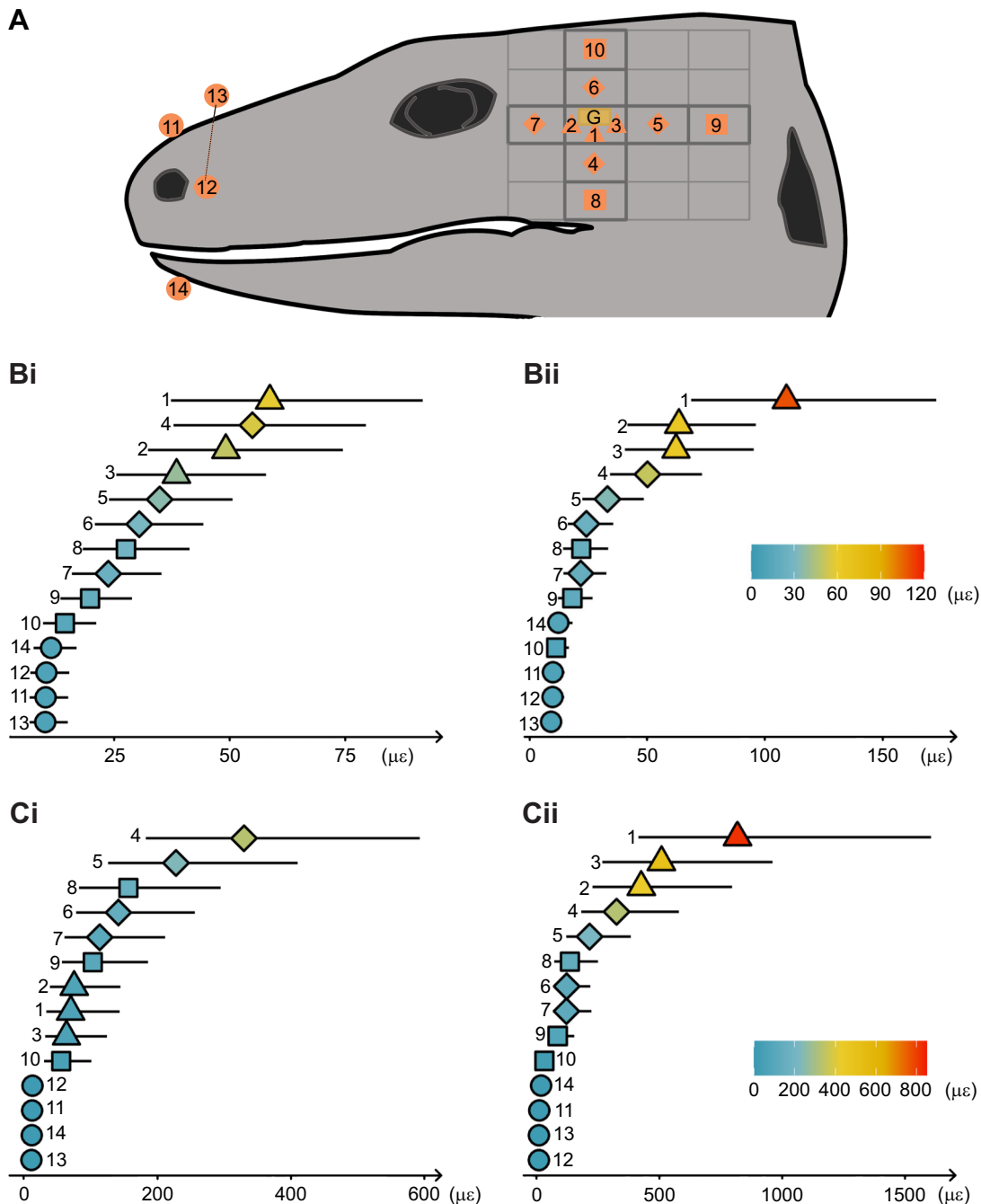


Fig. 3. Effect of the location of loading on the strain predicted for the instrumented osteoderm. (A) Location of the 14 different places where loading was applied (G, gauge location). Locations are indicated by a triangle when force was applied directly on the instrumented osteoderm, a diamond when the force was applied on an osteoderm adjacent to the instrumented osteoderm, a square when it was applied to an osteoderm separated from the instrumented osteoderm by one osteoderm, and a circle when it was applied near the rostral end of the head. (B) Tensile (i) and compressive strain (ii) predicted for each location in A by the linear mixed model using the 'dynamic' data, with a median force of 0.44 N (all species). (C) Tensile (i) and compressive strain (ii) predicted for each location in A by the linear mixed model using the 'static' data, with a median force of 8 N (all species). Strain magnitude is expressed using a colour gradient.

by changes in the gauge length resulted in voltage variations that were captured and amplified by the differential amplifier. The output signals from the amplifier were then converted from analog to digital and synchronized in a MP150 data acquisition and analysis system (BIOPAC Systems Inc., Goleta, CA, USA). For the *in toto* experiment, the force signal recorded from the piezoelectric sensors was synchronized with the recordings of strain using the MP150 data acquisition system. The output of the BIOPAC system was

recorded on a laptop using AcqKnowledge 4 software (BIOPAC Systems Inc.).

Using calibration data and a series of custom-written Igor (Wavemetrics, Inc., Portland, OR, USA) procedures, the force and strain signals were calibrated, transformed and analysed. For strain recorded from the rectangular rosette gauges (experiment 1), the data recorded from each element of the gauge were used to calculate the maximal principal strain (maximal tensile strain in the plane of

the gauge), the minimal principal strain (maximal compressive strain in the plane of the gauge), the shear strain (maximal–minimal principal strain), and the angle between the element A of the gauge and the maximal principal strain. The procedure used for the analysis first located the peaks of force then measured the maximal force and the largest strain associated with each peak. Strain values recorded from the tensile test experiment were measured manually using *AcqKnowledge* software. Here, the compressive and tensile strain caused by the attached weight were recorded as soon as the signal became stable.

Statistical analysis

We ran linear mixed models (LMMs) using the maximal tensile and compressive strain recorded at the level of the outer surface of the instrumented osteoderm and the following covariates: loading force, loading location, species, specimen and specimen head length. For each of the next four hypotheses, several models were run in R (<http://www.R-project.org/>) with the *lmer* function of the *lme4* package (Bates et al., 2018).

Hypothesis 1: for a given loading force, the largest strain amplitudes are always measured when the loading was applied directly on the instrumented osteoderm (versus other locations in the head). Here, we considered the covariates ‘location’, ‘loading force’ and their interaction as fixed factors and the covariates ‘species’, ‘specimen’ and ‘head length’ as random effects. The interaction between specimen and location was also considered as a random effect to deal with the pseudo-replications.

Hypothesis 2: there are interspecific differences in the stiffness of the ‘skin system’ (i.e. osteoderm and soft tissues of the skin). Here, the covariates species, loading force and their interaction were considered a fixed factor while the other covariates were modelled as random effects.

For the purpose of testing hypotheses 3 and 4, a variable ‘group’ was added to the dataset. This group variable had four different levels: (0) loading at the rostral end of the head (Fig. 3A; 11–14), (1) loading on osteoderms that are separated from the instrumented osteoderm by one osteoderm (Fig. 3A; 8–10), (2) loading on osteoderms that are adjacent to the instrumented osteoderm (Fig. 3A; 4–7), and (3) loading on the instrumented osteoderm (Fig. 3A; 1–3). *Matobosaurus validus* specimens were excluded from these models because we were not able to load the instrumented osteoderm directly (i.e. gauges occupied most of the osteoderm surface).

Hypothesis 3: there are interspecific differences in the stiffness of osteoderms. Here, we focused on strain measured for loading locations that are on the instrumented osteoderm (group 3) to exclude the effects of factors such as stress transmission between osteoderms and skull morphology, for example. The data obtained from the tensile test experiment were also used to test hypothesis 3.

Hypothesis 4: there are interspecific differences in the transmission of the force between neighbouring osteoderms. Here, we investigated relative differences in strain predicted for groups 0, 1, 2 and 3 within each species to determine which species transmit more force from the loading location to the instrumented osteoderm. For each model, the compressive and tensile strain predicted (using the median loading forces) for each combination of species and group were normalized between 0 and 1. This was achieved by dividing the strain values predicted for each combination of species and group by the highest values predicted for a group of the same species. These data provide a sense of the relative changes in the strain recorded in the instrumented osteoderm when the same loading force was moved from group 3 to 0. The underlying idea is

that in some species the strain could decrease by small increments (e.g. 100% of the strain when the instrumented osteoderm is loaded directly, 90% when the adjacent osteoderms are loaded, and 70% when the osteoderms separated from the instrumented osteoderm by one osteoderm are loaded), while in some other species the increments would be very large (e.g. 100%, 50% and 10%, respectively).

The following equations were used to generate the models:

$$\begin{aligned} \text{Hypothesis 1 : (STRAIN)} &\sim (\text{Loading_Force}) \times \text{Locations} \\ &+ (1|\text{Specimens}) + (1|\text{Specimens : Locations}) + (1|\text{Species}) \\ &+ (1|\text{HeadLength}), \end{aligned} \quad (1)$$

$$\begin{aligned} \text{Hypothesis 2 : (STRAIN)} &\sim (\text{Loading_Force}) \times \text{Species} \\ &+ (1|\text{Specimens}) + (1|\text{Specimens : Locations}) + (1|\text{Locations}) \\ &+ (1|\text{HeadLength}), \end{aligned} \quad (2)$$

$$\begin{aligned} \text{Hypothesis 3 and 4 : (STRAIN)} &\sim (\text{Loading_Force}) \times \text{Species} \\ &\times \text{Group} + (1|\text{Specimens}) + (1|\text{Specimens : Locations}) \\ &+ (1|\text{Locations}) + (1|\text{HeadLength}), \end{aligned} \quad (3)$$

where strain refers to tensile and compressive strain in static or dynamic manipulations.

The strain and loading force were log-transformed to fit a normal distribution of the residuals. Visual inspection of the model residuals did not reveal strong violations of parametric conditions (normality and homogeneity of variances). Small to moderate residuals correlations were still observed (from 0.2 to 0.4), which should not affect our interpretation of the models. We performed type III ANOVA to check whether the fixed effects and their interactions had significant impacts on the measured strain. Next, we used the models to predict the strain under a unique value of loading force. We chose the median applied force value observed in both the static and dynamic loading experiment.

RESULTS

Hypothesis 1: the largest strain amplitudes are always measured when loading is applied directly on the instrumented osteoderm

Loading force and loading location significantly impacted the maximal tensile and compressive strain in both the static and dynamic models (Table S1A; $P < 0.0001$). Compressive strain predicted for the static and dynamic data tended to confirm our hypothesis that loading of the instrumented osteoderm induced the largest strain amplitudes recorded at the outer surface of this structure (triangles in Fig. 3Bii,Cii). However, loading of the adjacent osteoderms (diamonds in Fig. 3Bii,Cii) can also generate large compressive strain in the instrumented osteoderm. This was mostly true when the osteoderm located at the ventral rim of the instrumented osteoderm (Fig. 3Bii and Ci, diamond 4) was loaded.

The highest tensile strain in the dynamic model was predicted for loading on the instrumented osteoderm (Fig. 3Bi). However, tensile strain with similar magnitude was predicted for loading on the osteoderm located at the ventral rim of the instrumented osteoderm (Fig. 3Bi, diamond 4). Considering the static data, the model even predicts greater tensile strain when the loading is applied on adjacent osteoderms (diamond in Fig. 3Ci) instead of the instrumented osteoderm (triangles in Fig. 3Ci).

For both the dynamic and static loading conditions, the models predict much smaller tensile and compressive strain in the instrumented osteoderm when the loaded osteoderm is separated by at least one osteoderm from the instrumented osteoderm (Fig. 3). In the case of loading at the rostral end of the head, the strain generally barely exceeded the background noise (Fig. 3B,C). It is also worth mentioning that loading osteoderms located ventrally or posteriorly to the instrumented osteoderm generally produced larger strain than loading osteoderms that were located anteriorly or dorsally (see location 4 versus 5, 6 and 7; and location 8 versus 9 and 10 in Fig. 3B,C).

Our data showed that the largest compressive strain was observed when the instrumented osteoderm was loaded directly even though loading of adjacent osteoderms also generated large compressive strain. Loading of the adjacent osteoderms produced tensile strain that was similar to (dynamic data) or even larger than (static data) that observed when the instrumented osteoderm was loaded.

Hypothesis 2: there are interspecific differences in the stiffness of the 'skin system'

The static and dynamic models confirmed interspecific variation in osteoderm strain (species effect: $P < 0.05$; Table S1B) considering all the loading locations. Species can be placed along a stiffness gradient based on the predicted tensile and compressive strain (Fig. 4). This gradient also becomes apparent when looking at the

strain predicted for groups 1–3 (Figs 5 and 6) and species order is highly similar using the different models (Figs 4, 5 and 6). The osteoderms of *T. rugosa* and the two *Heloderma* species were stiffer than those of *Timon lepidus*, *P. apodus*, *Tribolonotus gracilis*, *T. scincoides* and, in some instances, *C. zebra*, whereas the three-remaining species (*Tribolonotus novaeguineae* and the gerrhosaurids *B. major* and *M. validus*) showed intermediate strain values. Our second hypothesis was confirmed, with *T. rugosa* standing out as a species with a very stiff skin system whereas the skin of *T. lepidus*, *T. gracilis* and *P. apodus* appears the least stiff. However, the question remains as to whether these differences are due to the structure or mechanical behaviour of the osteoderm itself.

Hypothesis 3: there are interspecific differences in the stiffness of osteoderms

The group variable had a significant effect on strain in every model (Table S1C). When considered alone, species did not have a significant effect on strain. However, when considered in combination with loading force and/or group, its impact was always significant (Table S1C).

Very similar gradients of species were obtained with the four models (Figs 5, 6 and 7). With the exception of tensile strain predicted for *H. suspectum* during static loading (Fig. 7Bi), the tensile and compressive strain predicted for *T. rugosa* and the two

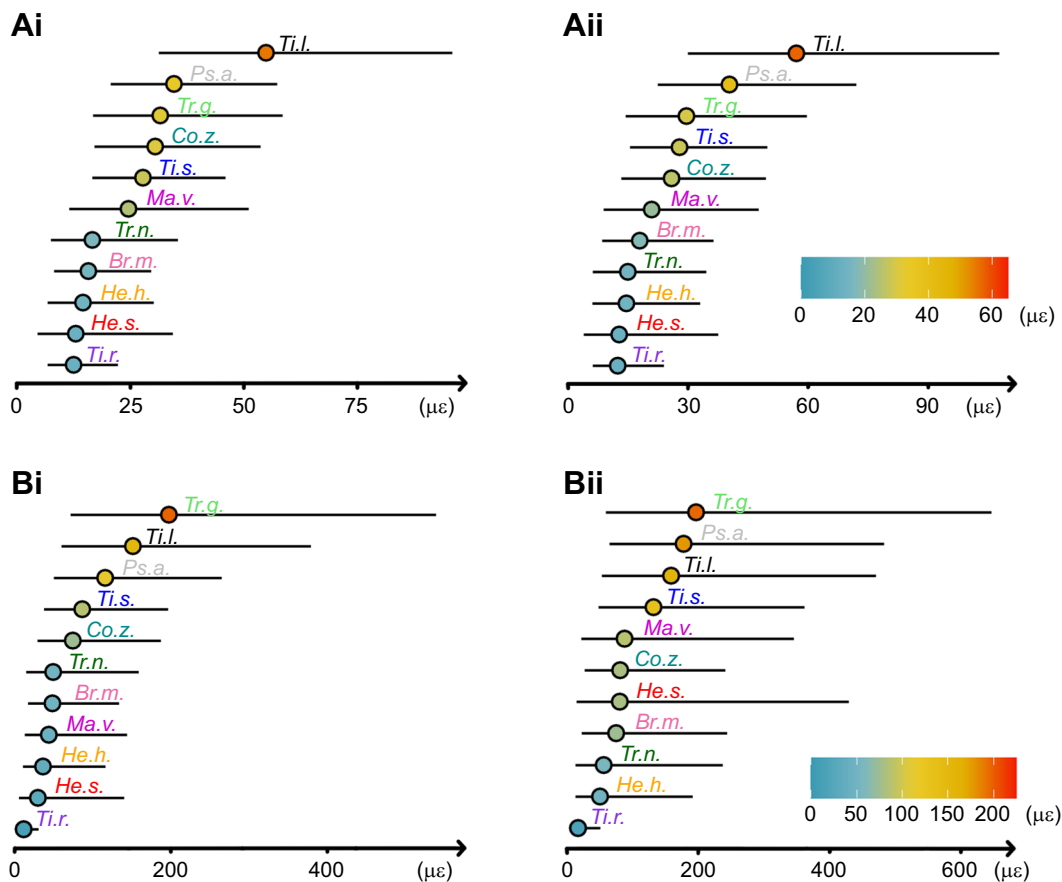


Fig. 4. *In toto* interspecific differences in the stiffness of the skin system. The graphs show the tensile (i) and compressive (ii) strain predicted by the linear mixed model for the instrumented osteoderm using (A) the dynamic data (median force of 0.44 N, all species) and (B) the static data (median force of 8 N, all species). Strain magnitude is expressed using a colour gradient. The locations were included in the model as a random covariate. *Br.m.*, *Broadleysaurus major*; *Co.z.*, *Corucia zebra*; *He.h.*, *Heloderma horridum*; *He.s.*, *Heloderma suspectum*; *Ma.v.*, *Matobosaurus validus*; *Ps.a.*, *Pseudopus apodus*; *Ti.r.*, *Tiliqua rugosa*; *Ti.s.*, *Tiliqua scincoides*; *Ti.l.*, *Timon lepidus*; *Tr.g.*, *Tribolonotus gracilis*; *Tr.n.*, *Tribolonotus novaeguineae*.

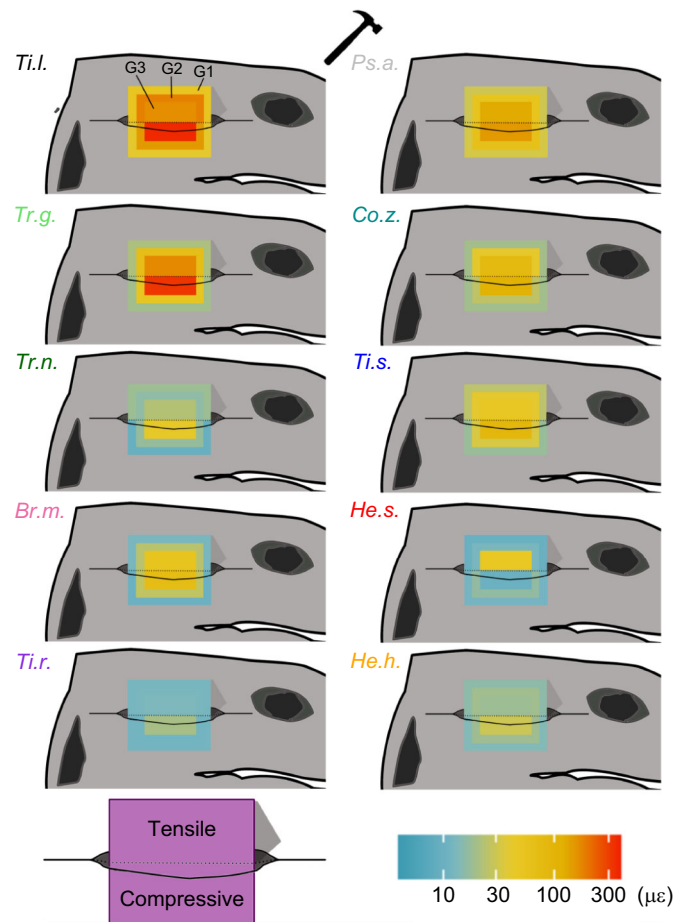


Fig. 5. Interspecific differences in stiffness depending on the location of dynamic loading. G1, loading on osteoderms that are separated from the instrumented osteoderm by one osteoderm; G2, loading on osteoderms that are adjacent to the instrumented osteoderm; G3, loading directly applied on the instrumented osteoderm. The upper half of each rectangle illustrates the predicted tensile strain while the lower half illustrates the predicted compressive strain. Strain magnitude is expressed using a colour gradient. *Br.m.*, *Broadleysaurus major*; *Co.z.*, *Corucia zebrata*; *He.h.*, *Heloderma horridum*; *He.s.*, *Heloderma suspectum*; *Ps.a.*, *Pseudopus apodus*; *Ti.r.*, *Tiliqua rugosa*; *Ti.s.*, *Tiliqua scincoides*; *Ti.l.*, *Timon lepidus*; *Tr.g.*, *Tribolonotus gracilis*; *Tr.n.*, *Tribolonotus novaeguineae*.

Heloderma species were always very small. In contrast, large compressive and tensile strain amplitudes were predicted for *T. lepidus*, *T. gracilis*, *P. apodus* and *C. zebrata* (except for tensile strain predicted under static loading). Intermediate values of strain were predicted for *B. major*, *T. scincoides* and *T. novaeguineae*. The species order along the stiffness gradients is very similar irrespective of whether all loading locations or direct loading is considered, suggesting that the features of the osteoderms strongly impact the biomechanical properties of the skin system.

The strain recorded on the external and internal sides of the dissected osteoderms during the tensile test experiment was several orders of magnitude (at least with the highest weights) larger in *C. zebrata*, *T. scincoides* and *P. apodus* than in the two *Heloderma* species or in *T. rugosa* (Fig. 8). The experimental setup, similar to a classical tensile test, confirmed the trends observed *in toto* even though, in the present case, osteoderms appear more elastic in *C. zebrata* and *T. scincoides* than in *P. apodus*. Notably, *P. apodus* is the only species for which we recorded tensile strain on the external surface of the osteoderm. For the other species, we recorded

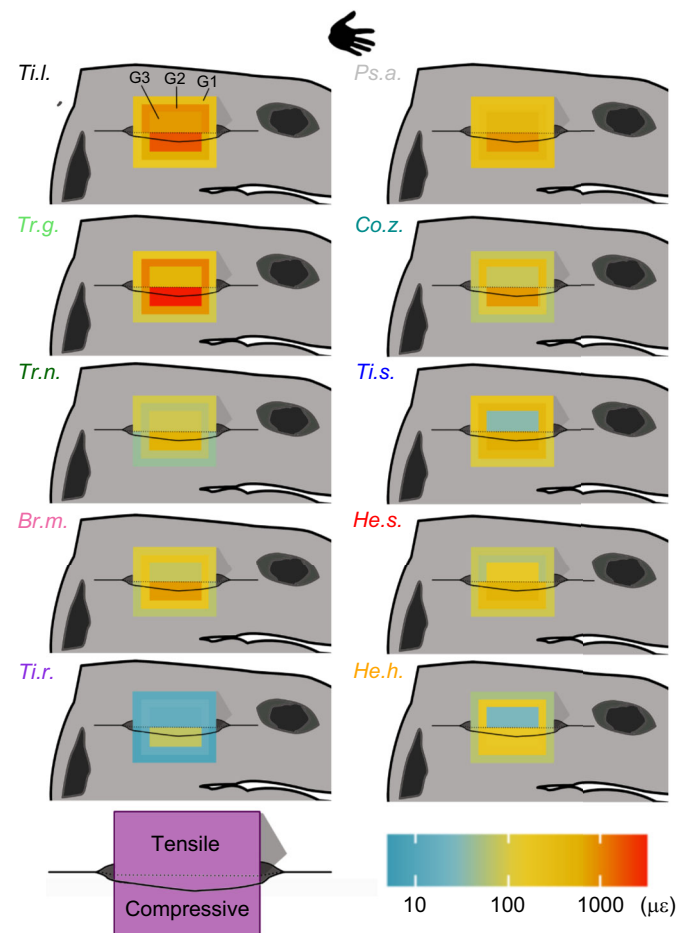


Fig. 6. Interspecific differences in stiffness considering the location of static loading. G1, loading on osteoderms that are separated from the instrumented osteoderm by one osteoderm; G2, loading on osteoderms that are adjacent to the instrumented osteoderm; G3, loading directly applied on the instrumented osteoderm. The upper half of each rectangle illustrates the predicted tensile strain while the lower half illustrates the predicted compressive strain. Strain magnitude is expressed using a colour gradient. *Br.m.*, *Broadleysaurus major*; *Co.z.*, *Corucia zebrata*; *He.h.*, *Heloderma horridum*; *He.s.*, *Heloderma suspectum*; *Ps.a.*, *Pseudopus apodus*; *Ti.r.*, *Tiliqua rugosa*; *Ti.s.*, *Tiliqua scincoides*; *Ti.l.*, *Timon lepidus*; *Tr.g.*, *Tribolonotus gracilis*; *Tr.n.*, *Tribolonotus novaeguineae*.

compressive strain on this side of the osteoderm. This was most likely explained by the more convex shape of their osteoderms (Fig. 8). On the internal side, tensile strain was recorded for every species.

Our third hypothesis was confirmed because there were clear interspecific differences in the strain recorded when the instrumented osteoderm was directly loaded. *Tiliqua rugosa* again stood out as having very stiff osteoderms whereas the temporal osteoderms of *T. lepidus*, *T. gracilis* and *P. apodus* were the least stiff. Based on the tensile test experiment, osteoderms of *T. scincoides* and *C. zebrata*, in particular, were also less stiff.

Hypothesis 4: there are interspecific differences in the transmission of force between neighbouring osteoderms

We also took advantage of the results provided by the models described for hypothesis 3 above ('Hypothesis 3: there are interspecific differences in the stiffness of osteoderms') to test our fourth hypothesis. During dynamic loading, large proportions of the forces causing tensile strain in the instrumented osteoderm were

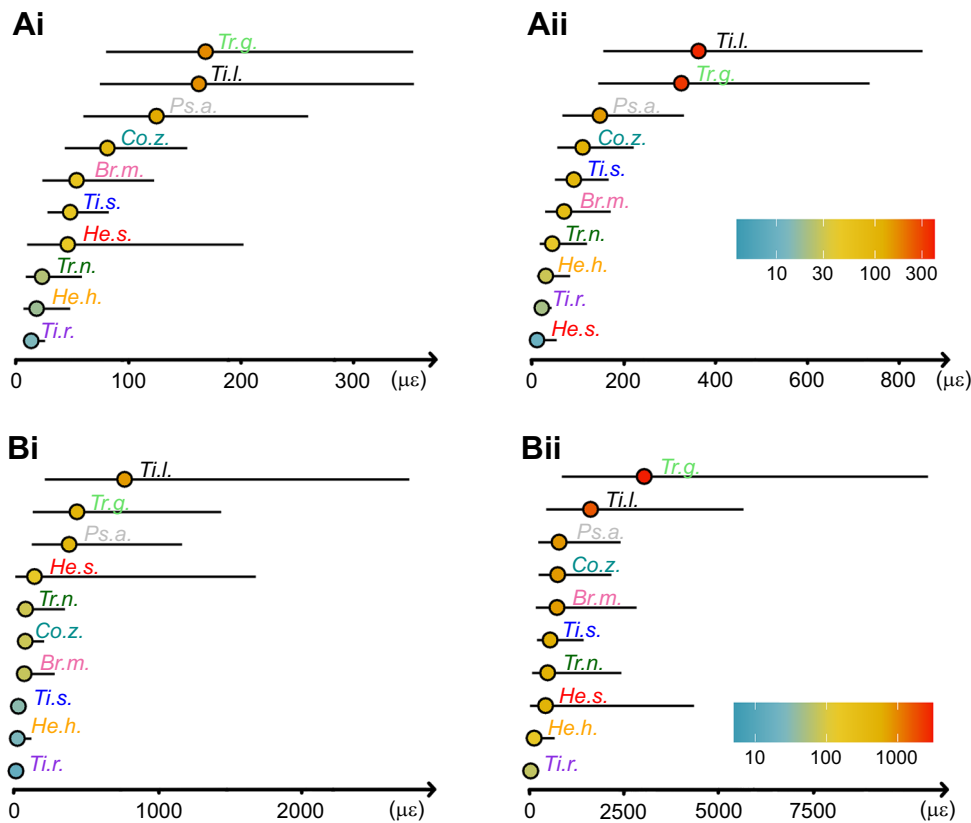


Fig. 7. In toto interspecific differences in the stiffness of the instrumented osteoderms. The graphs show the tensile (i) and compressive (ii) strain predicted by the linear mixed model considering only loading applied directly to the instrumented osteoderm *in toto* for (A) dynamic loading (median force of 0.44 N, all species) and (B) static loading (median force of 8 N, all species). Strain magnitude is expressed using a colour gradient. *Br.m.*, *Broadleysaurus major*; *Co.z.*, *Corucia zebtrata*; *He.h.*, *Heloderma horridum*; *He.s.*, *Heloderma suspectum*; *Ma.v.*, *Matobosaurus validus*; *Ps.a.*, *Pseudopus apodus*; *Ti.r.*, *Tiliqua rugosa*; *Ti.s.*, *Tiliqua scincoides*; *Ti.l.*, *Timon lepidus*; *Tr.g.*, *Tribolonotus gracilis*; *Tr.n.*, *Tribolonotus novaeguineae*.

transferred between the different groups of osteoderms in *T. rugosa* and *H. horridum* and to some extent in *T. lepidus*, *T. novaeguineae* and *T. scincoides* (upper half of schemes under the hammer shapes in Fig. 9). Forces causing compressive strain (lower half of schemes under the hammer shapes in Fig. 9) were transmitted in the two *Heloderma* species and in *T. rugosa*. In contrast, forces causing tensile and compressive strain were rapidly dissipated between the groups of osteoderms in *T. gracilis*, *C. zebtrata*, *P. apodus* and *B. major*.

During static loading, the forces causing tensile strain in the instrumented osteoderm seemed high in every species, at least between groups 1–3 (upper half of schemes under the hand shapes in Fig. 9). There were, however, differences in the transmission of forces causing compressive strain, with *H. suspectum*, *H. horridum*, *T. scincoides*, *P. apodus* and *T. lepidus* showing high levels of transmissions and the two *Tribolonotus* species, *B. major*, *C. zebtrata* and *T. rugosa* showing much lower levels of force transmission (lower half of schemes under the hand shapes in Fig. 9).

In addition to the interspecific differences, some species seemed to show differences in the transmission of the force depending on the type (static versus dynamic) of loading. In *T. rugosa*, more force was transmitted under dynamic loading (Fig. 9). This was especially striking when investigating the compressive strain. In *B. major*, *P. apodus* and *C. zebtrata*, force transmission seemed larger during static loading, especially when tensile strain was considered (Fig. 9).

DISCUSSION

During the last seven decades, strain gauges have been used extensively to quantify minute length changes in loaded human bones (Grassi and Isaksson, 2015). They have also been used to quantify the level of mechanical strain, subject to different loading regimes, across various bones of mammals (e.g. Hylander, 1984;

Thomason, 1991; Weijs and De Jongh, 1977) and, more recently, of fish (Markey et al., 2006) and reptiles (Dutel et al., 2021; Metzger et al., 2005; Porro et al., 2013, 2014; Ross et al., 2018; Smith and Hylander, 1985). Here, we used strain gauges to record maximal compressive, tensile and shear strain (shear strain is illustrated in Tables S1 and S2 and Figs S2 and S3) in temporal osteoderms of 11 lizard species. We observed that large strain amplitudes can be recorded when the instrumented osteoderm is directly loaded but also when force is applied to neighbouring osteoderms. We also recorded large strain amplitudes on the external and internal side of the dissected osteoderms during a tensile test. Although it may seem counterintuitive that we recorded compressive strain on the external surface for every species but *P. apodus*, this could be explained by the convex shape of the osteoderms (Fig. 8). A clear species gradient emerged from our analyses. The two helodermatids (*H. suspectum*, *H. horridum*) and *T. rugosa* had stiff skin and osteoderms. *Timon lepidus*, *P. apodus* and *T. gracilis* were at the other end of the spectrum, with less stiff skin and osteoderms. The osteoderms of *T. scincoides* and *C. zebtrata* had mechanical features more similar to those of *T. lepidus*, *P. apodus* and *T. gracilis*, and very large strain amplitudes were recorded for these two species when dissected osteoderms were subjected to a tensile test. The two gerrhosaurids (*B. major* and *M. validus*) and the scincid *T. novaeguineae* had temporal osteoderms with an intermediate stiffness.

Morpho-functional inferences

In most cases, the stiffness of the osteoderms appeared to be related to their overall shape. The relatively elastic osteoderms of *T. scincoides*, *T. lepidus*, *P. apodus* and *C. zebtrata* were thin and slightly to markedly elongated along the anteroposterior axis, whereas the stiff osteoderms of *T. rugosa* and the two helodermatids were proportionally thicker and more rounded (Fig. 1A).

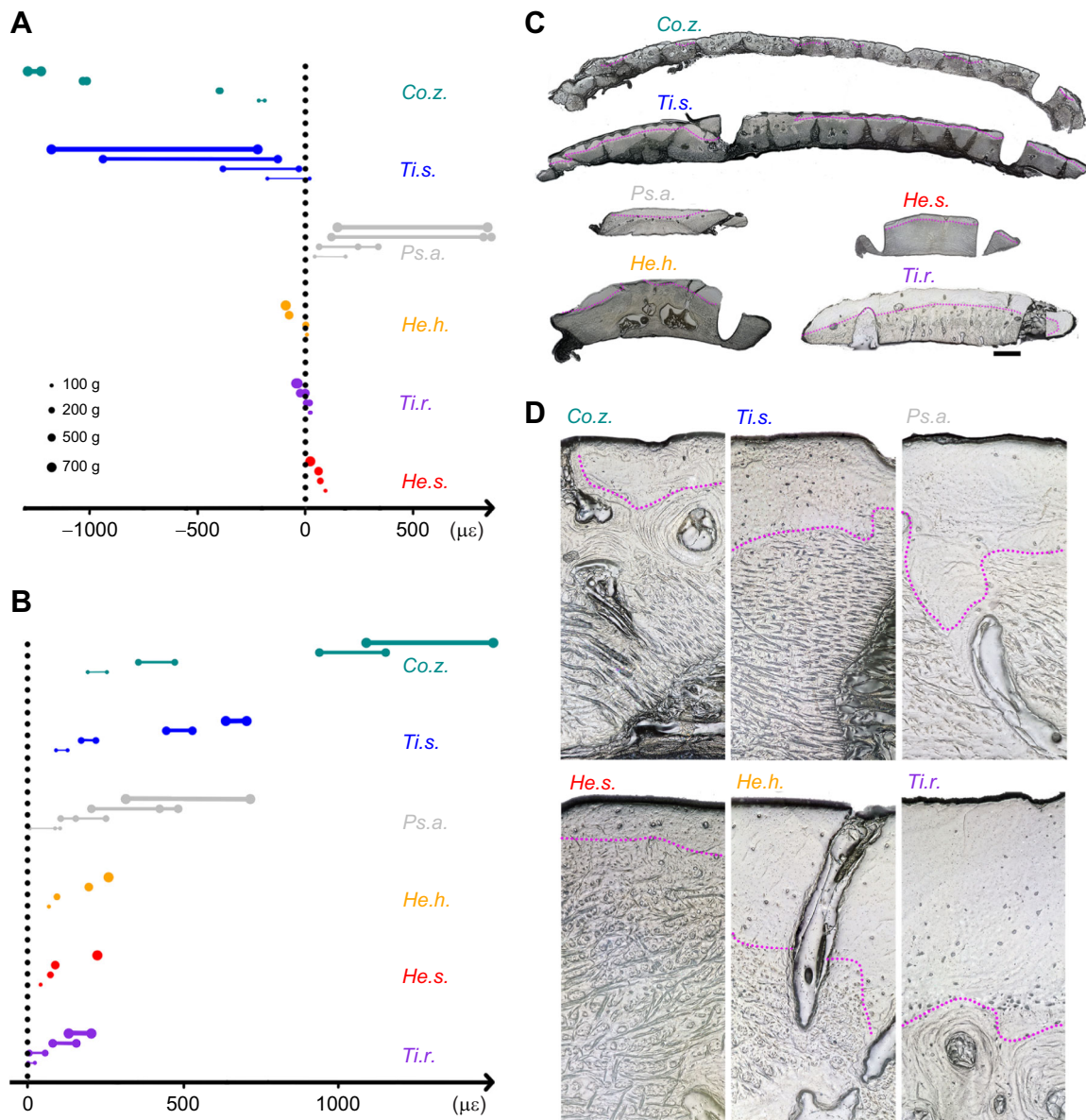


Fig. 8. Interspecific differences in the stiffness of dissected osteoderms. Tensile and compressive strain recorded from a subsample of dissected osteoderms. Here, the osteoderm was loaded along its long axis using different weights (100, 200, 500 and 700 g) and strain was recorded along that axis using single strain gauges placed on the external (A) and internal sides (B) of the osteoderm. (C) Histological cross-section of one dissected osteoderm per species. Scale bar: 1 mm. (D) Sections (500 μm width) of the osteoderms shown in C. Pink dotted lines highlight the edge of the capping tissue. Note that the layer of capping tissue is generally thicker in *H. suspectum* but the section presented here comes from the area that was trimmed to attach the strain gauge. Co.z., *Corucia zebrata*; He.h., *Heloderma horridum*; He.s., *Heloderma suspectum*; Ps.a., *Pseudopus apodus*; Ti.r., *Tiliqua rugosa*; Ti.s., *Tiliqua scincoides*.

The osteoderms of the two gerrhosaurids had an intermediate shape and biomechanical properties (Fig. 1A). Among cordylids, thicker osteoderms have been shown to provide better resistance against puncture during predator bites (Broeckhoven et al., 2015, 2017). We therefore propose that thickness, inherent mechanical properties and protection against puncture are correlated, and that the dermal armour of *T. rugosa* and the *Heloderma* species should better resist puncture.

Focusing on the five scincid species, at least three additional features of the gross morphology of the osteoderms likely explain why *T. rugosa* and *T. novaeguineae* had stiff osteoderms compared with *C. zebrata* and *T. scincoides*. First, the length of the overlapping section of adjacent osteoderms was greater in

T. rugosa than in all the other species, which may have helped increase the overall stiffness of the skin system (Fig. 1A). Second, the temporal osteoderms of *C. zebrata* and *T. scincoides* have a compound morphology (Fig. 1A), with each element being composed of several smaller bony pieces (termed osteodermite) sutured together with fibrous connective tissue (Williams et al., 2021). This compound organization likely contributed to a reduction in their overall stiffness. In contrast, the temporal osteoderms of *T. rugosa* and *T. novaeguineae* were consolidated elements without intervening fibrous sutures. Third, the midline keel observed in *T. novaeguineae* has previously been suggested to reduce vertical stress, at least in crocodylomorph osteoderms (Clarac et al., 2019).

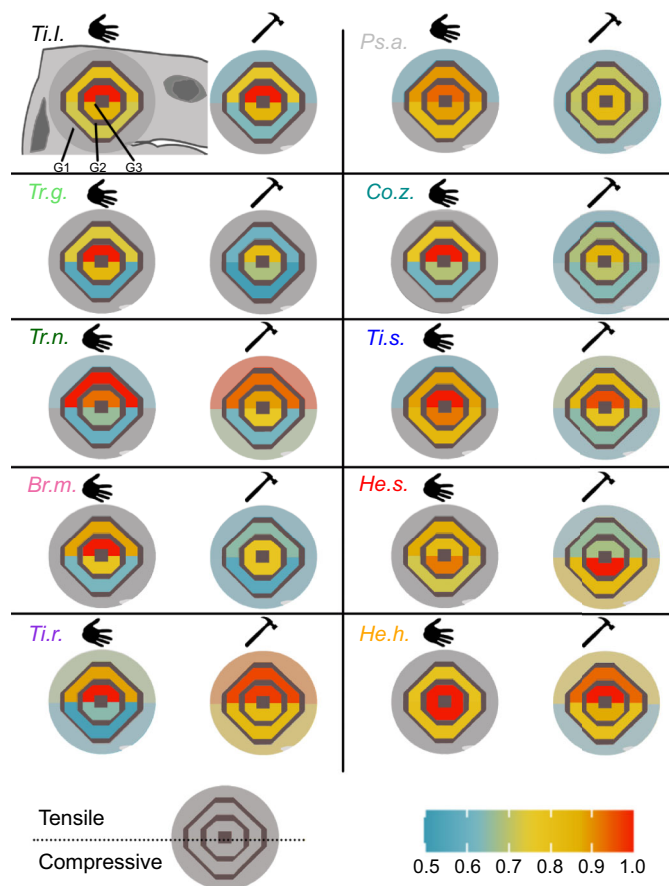


Fig. 9. Interspecific differences in the force transfer between adjacent osteoderms. Relative changes between groups 1, 2 and 3 were calculated within each species and colour coded to determine whether there were interspecific differences in the relative amount of stress transmitted to the neighbouring osteoderms. Colour bar: ratio between the strain values predicted for each combination of species and loading group and the highest values predicted for a group of the same species. *Br.m.*, *Broadleysaurus major*; *Co.z.*, *Corucia zebrata*; *He.h.*, *Heloderma horridum*; *He.s.*, *Heloderma suspectum*; *Ps.a.*, *Pseudopus apodus*; *Ti.r.*, *Tiliqua rugosa*; *Ti.s.*, *Tiliqua scincoides*; *Ti.l.*, *Timon lepidus*; *Tr.g.*, *Tribolonotus gracilis*; *Tr.n.*, *Tribolonotus novaeguineae*.

Unexpectedly, osteoderms from the two *Tribolonotus* species (Fig. 1A) demonstrated different loading properties (we recorded larger strain for *T. gracilis* as compared with *T. novaeguineae*), despite sharing a similar large size (relative to snout–vent length) and shape (with a strong, midline keel). Investigating structural features of osteoderms may help explain this discrepancy, as histological composition and structural heterogeneity likely alter resistance to external loads (see Iacoviello et al., 2020). For example, an enamel-like capping tissue with a high Young’s modulus has been reported in the dorsal osteoderms of some lizards (Kirby et al., 2020; Marghoub et al., 2022). The thickness of this tissue varies across species (including *H. suspectum* and *P. apodus*) but was absent from the osteoderms of *Varanus komodoensis*. For the temporal osteoderms, we observed a tissue similar to the capping tissue in every species investigated in the tensile test experiment although it was much thinner and sparsely distributed in *C. zebrata* (Fig. 8). The variation in osteoderm stiffness observed among these species underscores the need for systematic reporting of the presence, thickness, distribution and functional properties of this capping tissue. In particular, the morphological and functional diversity of scincid osteoderms stands

out as an important target for future evolutionary and biomimetic studies of body armour. The biomechanical and morphological data gathered here offer a first glimpse of the knowledge that will be gained from such studies and the features that could be mimicked to improve the stiffness or elasticity of materials used in the production of armour, vehicles or helmets, for example.

Using a drop weight impact test on biomimetically manufactured skin sheets of the dorsum of *H. suspectum*, *P. apodus* and *C. zebrata*, Liang et al. (2021) found that the skin of the last of these species absorbed the most energy. They suggested that it was related to the compound nature of the osteoderm. Our histology and tensile test showed that temporal osteoderms of *C. zebrata* are relatively elastic, compared with those of *P. apodus* and *H. suspectum*, and have only a thin layer of capping tissue (Fig. 8). These observations are particularly interesting given that *C. zebrata* is the only arboreal species in our sample and is potentially at risk of falling. However, the stiffness of *C. zebrata* osteoderms *in toto* was intermediate between that of *H. suspectum* and *P. apodus*. Moreover, the osteoderms of *C. zebrata* did not stand out when it came to their response to dynamic loading. As the shape, size and imbrication pattern differ between the temporal, ventral and dorsal skin areas of every species studied, investigating the ventral and dorsal osteoderms *in toto* and using tensile tests may help improve our understanding of the interspecific differences in the biomechanical properties and energy absorption of the lizard skin.

When we consider biomechanical properties of lizard osteoderms it is important to keep in mind that they are only one part of a complex skin system. Our study showed that external loading of neighbouring osteoderms generally causes larger tensile than compressive strain in the instrumented osteoderm, with the exception of *H. suspectum* (Fig. 9). Overall, however, our force transmission data did not reveal any obvious phylogenetic or morphological trends. For example, *H. horridum* showed a pattern of compressive and tensile strain distinct from that of the morphologically similar sister taxon *H. suspectum*. As differences during development have been reported for helodermatid osteoderms (Moss, 1969), we cannot exclude the possibility that intraspecific rather than interspecific variation caused the observed differences. Further, given the large amount of overlap between osteoderms, we were expecting a proportionally greater transmission of force between the temporal osteoderms of *T. rugosa* and *P. apodus* compared with the other studied species (Fig. 1A). However, neither species demonstrated relatively high levels of force transmission during static loading, and only *T. rugosa* showed high force transmission levels during the dynamic loading. Force transmission during dynamic loading was also relatively high in the two *Heloderma* species, but relatively low in *P. apodus*, *C. zebrata* and *T. gracilis*, which had stiff and elastic osteoderms, respectively. We therefore hypothesize that force transmission during dynamic loading is only partly impacted by the stiffness of the osteoderms. Force transmission between osteoderms is probably impacted by several other factors, including the presence and organization of the large collagen bundles (Sharpey’s fibres) that anchor the osteoderm within the skin (and interconnected osteodermites of compound osteoderms; see Williams et al., 2021). In order to better understand what role (if any) Sharpey’s fibres and other microstructural components play in the context of force transmission, future investigations integrating detailed histological descriptions and functional analyses are needed.

Other anatomical features of the head (e.g. muscle and bones of the skull) likely impacted our results as well. For example, the osteoderms located ventrally or posteriorly to the instrumented

osteoderm were more often positioned adjacent to jaw muscles, while the anterior and dorsal osteoderms generally covered parts of the skull. We observed that larger strain amplitudes were generally recorded from osteoderms that covered muscles (Fig. 3), suggesting that the nature of the underlying tissue could greatly impact the transmission of force between neighbouring osteoderms. As we worked on fixed specimens, the underlying muscles were probably stiffer than they would have been in live animals, thus possibly leading to an overestimation of the stiffness of the skin system *in toto*. However, live individuals may be able to increase the stiffness of their skin by contracting the jaw adductor muscles. Additional experiments should be conducted on anaesthetized and live specimens to investigate this topic further.

Evolutionary and ecological considerations

Based on our sample, it is impossible to formally assess the degree to which biomechanical properties of osteoderms are conserved among and between the different clades of lizards. It is, however, worth mentioning that we observed large differences in osteoderm stiffness between species of *Tiliqua* and *Tribolonotus* but not between the two helodermatids or the two gerrhosaurids.

Stiffer osteoderms could be advantageous during conspecific fighting and/or interspecific interactions with predators or prey by preventing lethal or sublethal injuries. However, they most likely represent higher maintenance and energetic costs, and may impair locomotor performance (e.g. higher mass resulting in a lower endurance, lower flexibility possibly reducing mobility in fossorial species). The trade-off observed between locomotor performance and the degree of armature in cordylid lizards (Losos et al., 2002) provides some evidence for this.

In our sampling, osteoderms were confined to the head only in *T. lepidus* but covered large/most parts of the body and head in the other species. *Timon lepidus* is likely the fastest lizard among the studied species (see Beck et al., 1995; Gans and Gasc, 1990; John-Alder et al., 1986; Main and Bull, 2000; Van Damme and Vanhooydonck, 2001; Vanhooydonck et al., 2014, for maximum sprint speeds), suggesting that evolutionary constraints related to speed are probably stronger than those favouring skin stiffness of the body in this species. Such differences in osteoderm distribution as well as those reported in this study for osteoderm stiffness could be related to interspecific differences in the balance between selective pressures associated with locomotor performance and protection against external loads.

Conspecific bites near or on the head have been reported for most of the studied species (e.g. Beck, 1990; Beck and Ramirez-Bautista, 1991; Jablonski, 2018; McCoy, 2006; Turner, 2010) or some of their close relatives (Pietruszka, 1988). In *T. rugosa*, conspecific bites to the head are common during male fights, often causing scale damage and sometimes breaking bones such as the mandible (Kerr and Bull, 2002; Murray and Bull, 2004). Considering the stiff temporal osteoderms of that species, we hypothesized that stiff temporal osteoderms evolved in species capable of strong bites during interactions with conspecifics. A rapid evaluation of the bite forces recorded for eight out of the 11 species investigated here (Fig. S4) suggests that the data do not unequivocally support this hypothesis because there is no obvious correlation between temporal osteoderm stiffness and bite force; *H. suspectum* had low bite forces despite having stiff temporal osteoderms while *T. rugosa*, *T. scincoides* and *C. zebrata* had strong bites of similar magnitude despite the differences in the stiffness of their osteoderms. In addition, both *Heloderma* species have stiff osteoderms although their fights are highly ritualized compared

with those observed in most other lizards (Beck and Ramirez-Bautista, 1991). We would expect ritualized fights to reduce the occurrence of biting during agonistic interactions. Data suggest that this could be true for *H. suspectum* where bites are occasional, but not for *H. horridum*, where the males can bite each other tenaciously and the dominant individual typically bites the jaw of the subordinate male at the end of each fighting bout, sometimes causing bleeding (Beck and Ramirez-Bautista, 1991). In this respect, it may be relevant that the lower jaw is covered with osteoderms in *H. horridum* but not in *H. suspectum* (Fig. 1). Again, more morphological, behavioural and biomechanical data from a much larger sample of species would be needed to formally test our hypothesis but the data available suggest that the functional roles of osteoderms are likely diverse.

Conclusions and perspectives

Temporal osteoderms show interspecific differences in their stiffness and those differences can be quantified using strain gauges. Specimens investigated here were all formalin-fixed which mostly likely increased the stiffness of the osteoderms and the underlying tissues. Therefore, differences in strain recorded for this study should be considered as relative differences. Future studies should explore strain in osteoderms of fresh specimens but also record strain during *in vivo* experiments to improve our understanding of the biomechanical properties and the ecological role of osteoderms. Our discussion also highlights the need for more ecological data in order to be able to interpret the functional role(s) of the lizard osteoderms.

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Competing interests

The authors declare no competing or financial interests.

Author contributions

Conceptualization: L.K., S.E.E., M.K.V., M.M., A.H.; Methodology: L.K., D.O., M.M., A.H.; Software: L.K., D.O.; Validation: D.O., A.H.; Formal analysis: L.K., D.O.; Investigation: L.K., A.M., A.H.; Resources: M.M., A.H.; Writing - original draft: L.K.; Writing - review & editing: D.O., A.M., S.E.E., M.K.V., M.M., A.H.; Visualization: L.K.; Supervision: M.M., A.H.; Project administration: M.M., A.H.; Funding acquisition: S.E.E., M.K.V., M.M., A.H.

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References

- Arnold, E. N. (1973). Relationships of the palaeartic lizards assigned to the genera *Lacerta*, *Algyroides* and *Psammotromus* (Reptilia: Lacertidae). *Bull. Br. Museum (Natural Hist. Zool)* **25**, 289-366.
- Arnold, E. N. (1989). Towards a phylogeny and biogeography of the Lacertidae: relationships within an old-world family of lizards derived from morphology. *Bull. Br. Museum (Natural Hist. Zool)* **55**, 209-257.
- Bates, D., Maechler, M., Bolker, B. and Walker, S. (2018). Fitting linear mixed-effects models using lme4. *J. Stat. Softw.* **67**, 1-48. doi:10.18637/jss.v067.i01
- Beck, D. D. (1990). Ecology and behavior of the Gila monster in southwestern Utah. *J. Herpetol.* **24**, 54-68. doi:10.2307/1564290
- Beck, D. and Ramirez-Bautista, A. (1991). Combat behavior of the beaded lizard, *Heloderma h. horridum*, in Jalisco, México. *J. Herpetol.* **25**, 481-484. doi:10.2307/1564773
- Beck, D. D., Dohm, M. R., Garland, T., Ramirez-Bautista, A. and Lowe, C. H. (1995). Locomotor performance and activity energetics of helodermatid lizards. *Copeia* **3**, 577-585. doi:10.2307/1446755

- Broeckhoven, C.** (2022). Intraspecific competition : a missing link in dermal armour evolution? *J. Anim. Ecol.* **91**, 1562–1566. doi:10.1111/1365-2656.13749
- Broeckhoven, C. and du Plessis, A.** (2022). Osteoderms as calcium reservoirs: Insights from the lizard *Ouroborus cataphractus*. *J. Anat.* **241**, 635–640. doi:10.1111/joa.13683
- Broeckhoven, C., Diedericks, G. and Mouton, P. I. F. N.** (2015). What doesn't kill you might make you stronger: functional basis for variation in body armour. *J. Anim. Ecol.* **84**, 1213–1221. doi:10.1111/1365-2656.12414
- Broeckhoven, C., du Plessis, A. and Hui, C.** (2017). Functional trade-off between strength and thermal capacity of dermal armor: insights from girdled lizards. *J. Mech. Behav. Biomed. Mater.* **74**, 189–194. doi:10.1016/j.jmbbm.2017.06.007
- Broeckhoven, C., El Adak, Y., Hui, C., Van Damme, R. and Stankowich, T.** (2018a). On dangerous ground: the evolution of body armour in cordylid lizards. *Proc. R. Soc. B Biol. Sci.* **285**, 20180513. doi:10.1098/rspb.2018.0513
- Broeckhoven, C., Mouton, P. I. F. N. and Hui, C.** (2018b). Proximate causes of variation in dermal armour: insights from armadillo lizards. *Oikos* **127**, 1449–1458. doi:10.1111/oik.05401
- Canei, J. and Nonclercq, D.** (2020). Morphological study of the integument and corporal skeletal muscles of two psammophilous members of Scincidae (*Scincus scincus* and *Eumeces schneideri*). *J. Morphol.* **282**, 230–246. doi:10.1002/jmor.21298
- Carrier, D. R. and Morgan, M. H.** (2014). Protective buttressing of the hominin face. *Biol. Rev.* **90**, 330–346. doi:10.1111/brv.12112
- Clarac, F., Goussard, F., De Buffrénil, V. and Sansalone, V.** (2019). The function(s) of bone ornamentation in the crocodylomorph osteoderms: a biomechanical model based on a finite element analysis. *Paleobiology* **45**, 182–200. doi:10.1017/pab.2018.48
- Connors, M., Yang, T., Hosny, A., Deng, Z., Yazdandoost, F., Massaadi, H., Eernisse, D., Mirzaeifar, R., Dean, M. N., Weaver, J. C. et al.** (2019). Bioinspired design of flexible armor based on chiton scales. *Nat. Commun.* **10**, 1–13. doi:10.1038/s41467-019-13215-0
- Dutel, H., Gröning, F., Sharp, A. C., Watson, P. J., Herrel, A., Ross, C. F., Jones, M. E. H., Evans, S. E. Fagan, M. J.** (2021). Comparative cranial biomechanics in two lizard species: impact of variation in cranial design. *J. Exp. Biol.* **224**, jeb234831. doi:10.1242/jeb.234831
- Gans, C. and Gasc, J.-P.** (1990). Tests on the locomotion of the elongate and limbless reptile *Ophisaurus apodus* (Sauria: Anguillidae). *J. Zool.* **220**, 517–536. doi:10.1111/j.1469-7998.1990.tb04731.x
- Grassi, L. and Isaksson, H.** (2015). Extracting accurate strain measurements in bone mechanics: a critical review of current methods. *J. Mech. Behav. Biomed. Mater.* **50**, 43–54. doi:10.1016/j.jmbbm.2015.06.006
- Hylander, W. L.** (1984). Stress and strain in the mandibular symphysis of primates: a test of competing hypotheses. *Am. J. Phys. Anthropol.* **64**, 1–46. doi:10.1002/ajpa.1330640102
- Hylander, W. L. and Johnson, K. R.** (1997). *In vivo* bone strain patterns in the zygomatic arch of macaques and the significance of these patterns for functional interpretations of craniofacial form. *Am. J. Phys. Anthropol.* **102**, 203–232. doi:10.1002/(SICI)1096-8644(199702)102:2<203::AID-AJPA5>3.0.CO;2-Z
- Hylander, W. L., Johnson, K. R. and Picq, P. G.** (1991). Masticatory–stress hypotheses and the supraorbital region of primates. *Am. J. Phys. Anthropol.* **86**, 1–36. doi:10.1002/ajpa.1330860102
- Iacoviello, F., Kirby, A. C., Javanmardi, Y., Moeendarbary, E., Shabanli, M., Tsolaki, E., Sharp, A. C., Hayes, M. J., Keevend, K., Li, J. H. et al.** (2020). The multiscale hierarchical structure of *Heloderma suspectum* osteoderms and their mechanical properties. *Acta Biomater.* **107**, 194–203. doi:10.1016/j.actbio.2020.02.029
- Inacio Veenstra, L. L. and Broeckhoven, C.** (2022). Revisiting the thermoregulation hypothesis of osteoderms : a study of the crocodylian *Paleosuchus palpebrosus* (Crocodylia: Alligatoridae). *Biol. J. Linn. Soc.* **135**, 679–691. doi:10.1093/biolinnean/blac001
- Jablonski, D.** (2018). Male-male combat in *Pseudopus apodus* (Reptilia: Anguillidae). *Russ. J. Herpetol.* **25**, 293–298. doi:10.30906/1026-2296-2018-25-4-293-298
- Jennings, A. R.** (2009). An analysis of 1,000 deaths in wild birds. *Bird Study* **8**, 26–31. doi:10.1080/00063656109475985
- John-Alder, H. B., Garland, T. J. and Bennett, A. F.** (1986). Locomotory capacities, oxygen consumption, and the cost of locomotion of the shingle-back lizard (*Trachydosaurus rugosus*). *Physiol. Zool.* **59**, 523–531. doi:10.1086/physzool.59.5.30156116
- Kerr, G. D. and Bull, C. M.** (2002). Field observations of aggressive encounters between male sleepy lizards (*Tiliqua rugosa*). *Herpetol. Rev.* **33**, 24–26.
- Kirby, A., Vickaryous, M., Boyde, A., Olivo, A., Moazen, M., Bertazzo, S. and Evans, S.** (2020). A comparative histological study of the osteoderms in the lizards *Heloderma suspectum* (Squamata: Helodermatidae) and *Varanus komodoensis* (Squamata: Varanidae). *J. Anat.* **236**, 1035–1043. doi:10.1111/joa.13156
- Laver, R. J., Morales, C. H., Heinicke, M. P., Gamble, T., Longoria, K., Bauer, A. M. and Daza, J. D.** (2020). The development of cephalic armor in the tokay gecko (Squamata: Gekkonidae: *Gekko gecko*). *J. Morphol.* **281**, 213–228. doi:10.1002/jmor.21092
- Liang, C., Marghoub, A., Kever, L., Bertazzo, S., Abzhanov, A., Vickaryous, M., Herrel, A., Evans, S. and Moazen, M.** (2021). Lizard osteoderms – morphological characterisation, biomimetic design and manufacturing based on three species. *Bioinspir. Biomim.* **16**, 066011. doi:10.1088/1748-3190/ac26d0
- Losos, J. B., Mouton, P. L. F. N., Bickel, R., Cornelius, I. and Ruddle, L.** (2002). The effect of body armature on escape behaviour in cordylid lizards. *Anim. Behav.* **64**, 313–321. doi:10.1006/anbe.2002.3051
- Main, A. R. and Bull, M.** (2000). The impact of tick parasites on the behaviour of the lizard *Tiliqua rugosa*. *Oecologia* **122**, 574–581. doi:10.1007/s004420050981
- Marghoub, A., Williams, C. J. A., Leite, J. V., Kirby, A. C., Kéver, L., Porro, L. B., Barrett, P. M., Bertazzo, S., Abzhanov, A., Vickaryous, M. et al.** (2022). Unravelling the structural variation of lizard osteoderms. *Acta Biomater.* **146**, 306–316. doi:10.1016/j.actbio.2022.05.004
- Markey, M. J., Main, R. P. and Marshall, C. R.** (2006). *In vivo* cranial suture function and suture morphology in the extant fish *Polypterus*: implications for inferring skull function in living and fossil fish. *J. Exp. Biol.* **209**, 2085–2102. doi:10.1242/jeb.02266
- Marques, M. P., Ceriaco, L. M. P., Stanley, E. L., Bandeira, S. A., Agarwal, I. and Bauer, A. M.** (2019). A new species of girdled lizard (Squamata: Cordylidae) from the Serra da Neve Inselberg, Namibe Province, southwestern Angola. *Zootaxa* **4668**, 503–524. doi:10.11646/zootaxa.4668.4.4
- McCoy, M.** (2006). *Reptiles of the Solomon Islands*. Sofia, Bulgaria: Pensoft Publisher.
- Metzger, K. A., Daniel, W. J. T. and Ross, C. F.** (2005). Comparison of beam theory and finite-element analysis with *in vivo* bone strain data from the alligator cranium. *Anat. Rec. Part A* **283A**, 331–348. doi:10.1002/ar.a.20167
- Moss, M.** (1969). Comparative histology of dermal sclerifications. *Acta Anat. (Basel)* **73**, 510–533. doi:10.1159/000143315
- Mukherjee, S. and Heithaus, M. R.** (2013). Dangerous prey and daring predators: a review. *Biol. Rev.* **88**, 550–563. doi:10.1111/brv.12014
- Murray, K. and Bull, C. M.** (2004). Aggressiveness during monogamous pairing in the sleepy lizard, *Tiliqua rugosa* : a test of the mate guarding hypothesis. *Acta Ethol.* **7**, 19–27. doi:10.1007/s10211-004-0092-2
- Oliver, J. A.** (1951). Ontogenetic changes in osteodermal ornamentation in skinks. *Copeia* **1951**, 127. doi:10.2307/1437541
- Paluh, D. J., Griffing, A. H. and Bauer, A. M.** (2017). Sheddable armour: identification of osteoderms in the integument of *Geckolepis maculata* (Gekkota). *African J. Herpetol.* **66**, 12–24. doi:10.1080/21564574.2017.1281172
- Pietruszka, R. D.** (1988). Observations on sexual dimorphism and social structure in the lizard *Angolosaurus skoogi* (Cordylidae) of the northern Namib Desert dunes. *South African J. Zool.* **23**, 47–51. doi:10.1080/02541858.1988.11448076
- Porro, L. B., Metzger, K. A., Iriarte-Diaz, J. and Ross, C. F.** (2013). *In vivo* bone strain and finite element modeling of the mandible of *Alligator mississippiensis*. *J. Anat.* **223**, 195–227. doi:10.1111/joa.12080
- Porro, L. B., Ross, C. F., Iriarte-Diaz, J., O'Reilly, J. C., Evans, S. E. and Fagan, M. J.** (2014). *In vivo* cranial bone strain and bite force in the agamid lizard *Uromastyx geyri*. *J. Exp. Biol.* **217**, 1983–1992. doi:10.1242/jeb.096362
- Ross, C. F., Porro, L. B., Herrel, A., Evans, S. E. and Fagan, M. J.** (2018). Bite force and cranial bone strain in four species of lizards. *J. Exp. Biol.* **221**, jeb180240. doi:10.1242/jeb.180240
- Smith, K. K. and Hylander, W. L.** (1985). Strain gauge measurement of mesokinetic movement in the lizard *Varanus exanthematicus*. *J. Exp. Biol.* **114**, 53–70. doi:10.1242/jeb.114.1.53
- Song, J., Ortiz, C. and Boyce, M. C.** (2011). Threat-protection mechanics of an armored fish. *J. Mech. Behav. Biomed. Mater.* **4**, 699–712. doi:10.1016/j.jmbbm.2010.11.011
- Stanley, E. L.** (2013). Systematics and morphological diversification of the Cordylidae (Squamata).
- Strahm, M. H. and Schwartz, A.** (1977). Osteoderms in the anguid lizard subfamily Diploglossinae and their taxonomic importance. *Biotropica* **9**, 58–72. doi:10.2307/2387862
- Tanke, D. H. and Currie, P. J.** (1998). Head-biting behavior in tetrapod dinosaurs: paleopathological evidence. *Gaia* **184**, 167–184.
- Thomas, N. J. and Cole, R. A.** (1996). The risk of disease and threats to the wild population. *Endanger. Species Updat* **13**, 23–27.
- Thomason, J. J.** (1991). Cranial strength in relation to estimated biting forces in some mammals. *Can. J. Zool.* **69**, 2326–2333. doi:10.1139/z91-327
- Tonini, J. F. R., Beard, K. H., Ferreira, R. B., Jetz, W. and Pyron, R. A.** (2016). Fully-sampled phylogenies of squamates reveal evolutionary patterns in threat status. *Biol. Conserv.* **204**, 23–31. doi:10.1016/j.biocon.2016.03.039

- Turner, G. S.** (2010). Natural history notes on the eastern blue-tongued skink *Tiliqua scincoides scincoides* from the basalt plains around Melbourne. *Vic. Nat.* **127**, 71-78.
- Van Damme, R. and Vanhooydonck, B.** (2001). Origins of interspecific variation in lizard sprint capacity. *Funct. Ecol.* **15**, 186-202. doi:10.1046/j.1365-2435.2001.00513.x
- Vanhooydonck, B., James, R. S., Tallis, J., Aerts, P., Tadic, Z., Tolley, K. A., Measey, G. J. and Herrel, A.** (2014). Is the whole more than the sum of its parts? Evolutionary trade-offs between burst and sustained locomotion in lacertid lizards. *Proc. R. Soc. B* **281**, 2132677. doi:10.1098/rspb.2013.2677
- Vickaryous, M. K. and Sire, J. Y.** (2009). The integumentary skeleton of tetrapods: origin, evolution, and development. *J. Anat.* **214**, 441-464. doi:10.1111/j.1469-7580.2008.01043.x
- Weijs, W. A. and De Jongh, H. J.** (1977). Strain in mandibular alveolar bone during mastication in the rabbit. *Arch. Oral Biol.* **22**, 667-675. doi:10.1016/0003-9969(77)90096-6
- Williams, C., Kirby, A., Marghoub, A., Kéver, L., Ostashevskaya-gohstand, S., Bertazzo, S., Moazen, M., Abzhanov, A., Herrel, A., Evans, S. E. et al.** (2021). A review of the osteoderms of lizards (Reptilia : Squamata). *Biol. Rev.* **97**, 1-19. doi:10.1111/brv.12788
- Yang, W., Chen, I. H., Gludovatz, B., Zimmermann, E. A., Ritchie, R. O. and Meyers, M. A.** (2013). Natural flexible dermal armor. *Adv. Mater.* **25**, 31-48. doi:10.1002/adma.201202713
- Zylberberg, L. and Castanet, J.** (1985). New data on the structure and the growth of the osteoderms in the reptile *Anguis fragilis* L. (Anguillidae, Squamata). *J. Morphol.* **186**, 327-342. doi:10.1002/jmor.1051860309