

Research article

urn:lsid:zoobank.org:pub:9A36B75B-C0B4-4CB3-9BD2-B91704457A5B

A morphological and molecular review of the genus *Goniurosaurus*, including an identification key

Hai Ngoc NGO¹, Huy Quoc NGUYEN², Hieu Minh TRAN³, Hanh Thi NGO⁴,
 Minh Duc LE⁵, Laurenz Rafael GEWISS⁶, Mona van SCHINGEN-KHAN⁷,
 Truong Quang NGUYEN^{8,*} & Thomas ZIEGLER^{9,*}

^{1,2}Vietnam National Museum of Nature, Vietnam Academy of Science and Technology,
 18 Hoang Quoc Viet Road, Hanoi, Vietnam.

^{2,8}Graduate University of Science and Technology, Vietnam Academy of Science and Technology,
 18 Hoang Quoc Viet Road, Hanoi, Vietnam.

^{3,8}Institute of Ecology and Biological Resources, Vietnam Academy of Science and Technology,
 18 Hoang Quoc Viet Road, Hanoi, Vietnam.

⁴Department of Genetics, Faculty of Biology, University of Science, Vietnam National University
 Hanoi, 334 Nguyen Trai Road, Hanoi, Vietnam.

^{4,5}Central Institute for Natural Resources and Environmental Studies, Vietnam National University
 Hanoi, 19 Le Thanh Tong, Hanoi, Vietnam.

⁵Department of Environmental Ecology, Faculty of Environmental Sciences, University of Science,
 Vietnam National University Hanoi, 334 Nguyen Trai Road, Hanoi, Vietnam.

⁵Department of Herpetology, American Museum of Natural History,
 Central Park West at 79th Street, New York, USA.

^{1,6,9}Institute of Zoology, University of Cologne, Zùlpicher Strasse 47b, 50674 Cologne, Germany.

^{1,6,9}Cologne, Germany Cologne Zoo, Riehler Straße 173, 50735 Cologne, Germany.

⁷Federal Agency for Nature Conservation, CITES Scientific Authority, Konstantinstrasse 110,
 53179 Bonn, Germany.

*Corresponding authors: nqt2@yahoo.com; ziegler@koelnerzoo.de

¹Email: ngohai2709@gmail.com

²Email: huynguyen17295@gmail.com

³Email: tranminhhieu_t60@hus.edu.vn

⁴Email: ngohanhhus@gmail.com

⁵Email: minh.le.cres@gmail.com

⁶Email: lgewiss@smail.uni-koeln.de

⁷Email: Mona.van.Schingen-Khan@bfn.de

¹urn:lsid:zoobank.org:author:91BE5B7A-2ECC-4FEC-B087-C47BC074492A

²urn:lsid:zoobank.org:author:0B5CC3D1-D8D5-41CD-9DC3-9F9B7FA3CE82

³urn:lsid:zoobank.org:author:FA82E6B8-3B50-4E6C-90FB-3FF1DD98CBD7

⁴urn:lsid:zoobank.org:author:5F0DA41F-0566-4738-A2F1-0A7D7E6A37DB

⁵urn:lsid:zoobank.org:author:785D605E-6A6F-4A67-BD18-7F40F426FC2C

⁶urn:lsid:zoobank.org:author:9AF54639-2BA0-47DE-BDC3-7816F57971A8

⁷urn:lsid:zoobank.org:author:ECC52912-B0CA-4C7C-AB16-BCAF83866E19

⁸urn:lsid:zoobank.org:author:822872A6-1C40-461F-AA0B-6A20EE06ADBA

⁹urn:lsid:zoobank.org:author:5716DB92-5FF8-4776-ACC5-BF6FA8C2E1BB

Abstract. The genus *Goniurosaurus* (tiger geckos) currently consists of 23 species distributed in China, Japan and Vietnam. Several species complexes and recent discoveries of cryptic species pose challenges to the species identification, which is crucial to effectively implement the recent listing of the species from China and Vietnam in CITES Appendix II and the species from Japan in CITES Appendix III. Based on the results of our field work in northern Vietnam and data compiled from literature, we herein provide a taxonomic review of the genus *Goniurosaurus*. Our phylogenetic analyses showed that all recorded populations of tiger geckos from Vietnam, which were found to be monophyletic with low intra-specific genetic divergences, are assigned to one of the four species: *G. catbaensis*, *G. huuliensis*, *G. lichtenfelderi* or *G. luii*. Both genetic and morphological analyses confirm that the species from China and Vietnam can be split into three major groups. Based on the newly collected data, we provide an extended morphological description of the Vietnamese species. In addition, we provide an identification key for all *Goniurosaurus* species from China, Japan and Vietnam in order to assist authorities in the enforcement of the recent CITES listing.

Keywords. CITES enforcement, morphology, molecular phylogeny, taxonomy, tiger geckos.

Ngo H.N., Nguyen H.Q., Tran H.M., Ngo H.T., Le M.D., Gewiss L.R., Schingen-Khan M. van, Nguyen T.Q. & Ziegler T. 2021. A morphological and molecular review of the genus *Goniurosaurus*, including an identification key. *European Journal of Taxonomy* 751: 38–67. <https://doi.org/10.5852/ejt.2021.751.1379>

Introduction

The eublepharid genus *Goniurosaurus* Grismer, Viets & Boyle, 1999 (tiger geckos) currently consists of 23 species associated with granitic or karst formations in Southeast and East Asia. This genus shows a high level of local endemism and most tiger gecko species have been only recorded from a single locality, within unique ecosystems or an isolated archipelago (Orlov *et al.* 2008; Ziegler *et al.* 2008; Yang & Chan 2015; Honda & Ota 2017; Zhou *et al.* 2018; Ngo *et al.* 2019a; Qi *et al.* 2020a, 2020b; Zhu *et al.* 2020a, 2020b). The complex topography and variable climatic conditions throughout the distribution range of the genus may account for the disjunct distribution and occupation of different ecological niches by distinct populations. These factors may have driven natural selection, morphological diversification and phylogenetic evolution within this genus (Vitt *et al.* 1997; Clements *et al.* 2006; Sexton *et al.* 2009; Gomes *et al.* 2016; Liang *et al.* 2018; Qi *et al.* 2020a, 2020b). According to recent morphological and molecular analyses, the genus *Goniurosaurus* is split into four species groups, namely the *G. lichtenfelderi* group with five species from both mainland and islands in China and Vietnam; the *G. kuroiwaie* group containing six species from the Ryukyu Archipelago, Japan; the *G. luii* group comprising eight species distributed throughout islands and mainland in Vietnam and China; and the *G. yingdeensis* group consisting of four species from China (Nguyen *et al.* 2009; Nguyen 2011; Wang *et al.* 2013; Honda & Ota 2017; Liang *et al.* 2018; Qi *et al.* 2020a, 2020b; Zhu *et al.* 2020a, 2020b). However, the systematics of the genus *Goniurosaurus* remains challenging due to ongoing discoveries of further cryptic species and the fact that there is not a single genetic marker that covers all 23 species, precluding a complete generic phylogeny of the group. Five subspecies of the *G. kuroiwaie* group from Japan were reinstated at full species status and a new species, *G. sengokui* (Honda & Ota, 2017), was recently discovered from the Ryukyu Archipelago (Honda & Ota 2017). In China, two taxa, previously identified as *G. luii* Grismer, Viets & Boyle, 1999, were described as distinct species, namely *G. kadoorieorum* Yang & Chan, 2015 and *G. kwangsiensis* Yang & Chan, 2015 (Yang & Chan 2015), and a sister species of *G. araneus* Grismer, Viets & Boyle, 1999, namely *G. gezhi* Zhu, He & Li, 2020, was recently discovered (Zhu *et al.* 2020a). These three species occur in Southwest Guangxi Province (Zhu *et al.* 2020a). Four additional species, namely *G. kwanghua* Zhu & He, 2020 and *G. zhoui* Zhou, Wang, Chen & Liang, 2018, belonging to the *G. lichtenfelderi* group, as well as *G. gollum* Qi, Wang, Grismer, Chen, Lyu & Wang, 2020 and *G. varius* Qi, Grismer, Lyu,

Zhang, Li & Wang, 2020 of the *G. yingdeensis* group, were recently described from China (Zhou *et al.* 2018; Qi *et al.* 2020a, 2020b; Zhu *et al.* 2020b).

To date, five species of *Goniurosaurus* are reported from Vietnam, viz. *G. araneus*, *G. catbaensis* Ziegler, Nguyen, Schmitz, Stenke & Rösler, 2008, *G. huuliensis* Orlov, Ryabov, Nguyen, Nguyen & Ho, 2008, *G. lichtenfelderi* (Mocquard, 1897) and *G. luii* (Grismer *et al.* 1999; Vu *et al.* 2006; Orlov *et al.* 2008; Ziegler *et al.* 2008; Nguyen *et al.* 2009; Nguyen 2011). All species were described based only on a few specimens and a small set of diagnostic characters. As such, phenotypic variability among these species may lead to the misidentification of taxa. For example, Orlov & Darevsky (1999) described a new species (*G. murphyi*) based on a juvenile specimen, which was subsequently synonymized with *G. lichtenfelderi* by Grismer (2000). Ngo *et al.* (2016) documented another case with indistinct morphological differences between the newly described *G. kadoorieorum* and *G. luii*.

Due to habitat degradation and over-harvesting for the pet trade, wild populations of species of *Goniurosaurus* from China and Vietnam have been subject to severe declines (Stuart *et al.* 2006; Yang & Chan 2015; Ngo *et al.* 2019b). Therefore, all Chinese and Vietnamese tiger gecko species have recently been included in CITES (Convention on International Trade in Endangered Species of Wild Fauna and Flora) Appendix II (CITES 2019; Ngo *et al.* 2019b). To effectively enforce relevant national and international regulations and to enable sustainable trade in CITES listed species, a detailed guideline for species identification is an essential prerequisite, especially as trade in the morphologically similar species from Japan is currently regulated under a lower protection level, namely CITES Appendix III (CITES Notification No. 2020/068). A proper identification guidance will help to prevent certain taxa from being traded under a wrong name to circumvent trade restrictions.

In this study, phylogenetic analyses were performed across all recorded populations of *Goniurosaurus* from Vietnam and along with other species from China and Japan using a fragment of the mitochondrial Cytochrome b gene. Based on newly collected data from Vietnam, we expanded morphological descriptions of each species and evaluated intra- and inter-specific morphological variations among the species of *Goniurosaurus*. We further compared morphological data of Vietnamese tiger geckos with those from China and identified characters that can distinguish the Japanese species. We thereby provide an identification key for all *Goniurosaurus* species from China, Japan and Vietnam, in order to assist CITES enforcement authorities in species identification.

Material and methods

Field surveys

Field surveys were conducted in June and August 2014, May 2015, July 2016, June and July 2017, April 2018, from April to September 2019, as well as in June and July 2020 in Vietnam. Surveys were conducted for *G. catbaensis* in Hai Phong City and Quang Ninh Province, for *G. huuliensis* in Lang Son Province, for *G. lichtenfelderi* in Hai Duong, Bac Giang and Quang Ninh provinces, and for *G. luii* in Cao Bang and Lang Son provinces. Animals were captured by hand and subsequently released at the same sites after taking measurements and photos in life. Coordinates of each captured individual were recorded using a GPS device (Garmin GPSmap64, WGS84 datum) and can be shared upon request to the authors.

In addition, several vouchered specimens and small tissue samples of *Goniurosaurus* deposited in collections of the Institute of Ecology and Biological Resources (IEBR), Vietnam National Museum of Nature (VNMN) and Hanoi University of Science (HUS), Hanoi, Vietnam, were examined for morphological characters and molecular phylogeny.

Molecular data and phylogenetic analyses

All taxa of the genus *Goniurosaurus* from China, Japan, and Vietnam, for which Cytochrome b data were available in GenBank, were included in the study. Three species, *Gekko chinensis* (Gray, 1842) (NC027191), *Coleonyx mitratus* Peters, 1863 (AB853481) and *Holodactylus africanus* Boettger, 1893 (AB853482), were used as outgroups following Honda *et al.* (2014).

Small tissue samples were collected from the tail tip of 18 wild specimens from representative localities in Vietnam and preserved separately in 70% ethanol (Merck, Germany). The mitochondrial DNA Cytochrome b (cytb) gene was selected for DNA sequencing. The total genomic DNA was extracted using GeneJet Genomic DNA Purification (ThermoFisher Scientific, Lithuania), following protocols by the manufacturer's instructions. PCR reactions were performed using HotStar Taq Mastermix (Qiagen, Germany) to amplify a fragment of approximately 1000 bps of cytb. The primer pair used for this study was L14731 (5'-TGGTCTGAAAAACCATTGTTG-3'; Honda *et al.* 2014) and GoniR1 (5'-CTACGGGCTGTCCTCCGATTCAGGTT-3'; this study). The PCR volume consisted of 21 µl: 2 µl of each primer, 5 µl water, 10 µl of Taq mastermix and 1–2 µl DNA template depending on the quality of DNA in the final extraction solution. The PCR was performed at 95°C for 15 minutes, followed by 35 cycles, for 30 s at 95°C, 45 s at 48°C and 60 s at 72°C with a final elongation step for 6 minutes at 72°C. Negative and positive controls were used for all DNA extractions and PCR reactions.

PCR products were visualized using electrophoresis through a 1% agarose gel, marker 1kb, 1X TBE, stained with ethidium bromide and photographed under UV light. Successful amplifications were purified using GeneJet PCR Purification Kit (ThermoFisher Scientific, Lithuania). Cleaned PCR products were sent to 1st Base (Malaysia) for sequencing in both directions.

The obtained sequences were aligned in Sequencher ver. 5.4 (Gene Codes Corp, Ann Arbor, MI, USA) and afterwards aligned using ClustalX ver. 2.1 (Thompson *et al.* 1997) with default settings. Data were analyzed using maximum parsimony (MP) as implemented in PAUP*4.0b10 (Swofford 2001) and Bayesian inference (BI) as implemented in MrBayes ver. 3.2 (Ronquist *et al.* 2012). Settings for these analyses followed Le *et al.* (2006), except that the number of generations in the Bayesian analysis was increased to 1×10^7 . For the maximum likelihood (ML) analysis, we used IQ-TREE ver. 1.6.7.1 (Nguyen *et al.* 2015) with a single model and 10000 ultrafast bootstrap replications. The optimal model for nucleotide evolution was set to TPM1uf+I+G as selected by Jmodeltest ver. 2.1.4 (Posada *et al.* 1998). As the phylogenetic relationships between species and species groups were well resolved and virtually every important node received high statistical support from all analyses, we opted not to partition our data by codon positions.

For the Bayesian analysis, we used the optimal model determined by Jmodeltest with parameters estimated by MrBayes ver. 3.2.7. Four Markov chains, one cold and three heated utilizing default heating values, were sampled every 1000 generations. Log-likelihood scores of sample points were plotted against generation time to detect stationarity of the Markov chains. The burn-in value was set to 50 in the BI analysis, as -lnL scores reached stationarity after 50000 generations in both runs. Nodal support was evaluated using Bootstrap replication (BP) as estimated in PAUP*4.0b10 and ultrafast BP in IQ-TREE ver. 1.6.7.1 and posterior probability (PP) in MrBayes ver. 3.2. BP ≥ 70 (Hillis & Bull 1993) and ultrafast BP (UBP) and PP $\geq 95\%$ are regarded as strong support for a clade (Ronquist *et al.* 2012; Nguyen *et al.* 2015). The uncorrected pairwise distance (p) were calculated in PAUP*4.0b10.

Morphological analyses

A total of 486 live individuals and 54 museum specimens of four species from Vietnam were examined for morphological data, comprising 194 individuals of *G. catbaensis* (21 juveniles, 93 females and 80 males), 80 individuals of *G. huuliensis* (02 juveniles, 46 females and 32 males), 178 individuals of

G. lichtenfelderi (14 juveniles, 72 females and 92 males), and 88 individuals of *G. luii* (11 juveniles, 43 females and 34 males). Detailed descriptions of each species were based on newly collected data combined with previous literature for the morphological variation, such as for *G. catbaensis* from Ziegler *et al.* (2008) and Nguyen (2011), for *G. huuliensis* from Orlov *et al.* (2008) and Nguyen (2011), for *G. lichtenfelderi* from Grismer (2000), Grismer *et al.* (2002) and Nguyen (2011), and for *G. luii* from Grismer *et al.* (1999), Vu *et al.* (2006) and Nguyen (2011). No specimen of *G. araneus* was investigated in this study, however, we included morphological data of eight specimens from the type locality in Cao Bang Province, Vietnam and Guangxi Province, China from the literature (Grismer *et al.* 1999; Chen *et al.* 2014). The sex of each collected specimen was determined based on the presence (in males) or absence (in females) of large swollen hemipenial bulges. Lizards were categorized into two age classes based on the snout-vent length (*G. lichtenfelderi*: SVL < 80 mm = juveniles, SVL ≥ 80 mm = adults, while three other species of *Goniurosaurus* were sorted with SVL < 85 mm = juveniles, SVL ≥ 85 mm = adults).

Measurements were taken with dial calipers to the nearest 0.1 mm at the right side of each individual.

Abbreviations

AD	=	diameter of auditory meatus
AG	=	axilla to groin length, from posterior edge of forelimb insertion to anterior edge of hindlimb insertion
BH	=	maximum body height, from top of dorsal body to belly
BW	=	maximum body width, greatest width of torso, taken at level of midbody
CH	=	cheek height, from posterior edge of labial to top of head at parietal region
ED	=	diameter of eye, greatest diameter of orbit
EE	=	eye to ear distance, from posterior margin of eye to posterior margin of ear
FLL	=	forelimb length, from axilla to the tip of the fourth finger
HH	=	maximum head height
HL	=	head length, from the tip of snout to posterior edge of occiput
HLL	=	hindlimb length, from groin to the tip of the fourth toe
HW	=	maximum head width
IO1	=	interorbital distance, distance between anteriormost points of eyes
IO2	=	interorbital distance, distance between posteriormost points of eyes
ML	=	mouth length, from tip of snout to last posterior labial edge
MW	=	mouth width, distance between last posterior labial edges on each side
ND	=	supranasal distance, distance between nares
SE	=	snout to eye distance, measured from tip of snout to anteriormost point of eye
SVL	=	snout-vent length, from tip of snout to vent
TaL	=	tail length, from vent to tip of tail
WT	=	maximum tail width

Scale counts

CIL	=	eyelid fringe scales or ciliaria
DTR	=	dorsal tubercle rows at midbody
GP	=	gular scales bordering the postmentals
GST	=	granular scales surrounding dorsal tubercles
IFL	=	infralabials
IN	=	postrostrals or internasals
LD1	=	subdigital lamellae under the first finger
LD4	=	subdigital lamellae under the fourth finger
LT1	=	subdigital lamellae under the first toe

LT4	=	subdigital lamellae under the fourth toe
MB	=	scales around midbody
N	=	nasal scales surrounding nare
P-IN	=	post-internasals
PAT	=	postcloacal tubercles
PM	=	postmentals
PO	=	preorbital scales
PP	=	precloacal pores
SPL	=	supralabials
TL	=	paravertebral tubercles between limb insertions

Statistical analyses were performed by using the software environment R.3.1.2 (RStudio Team 2018). Shapiro-Wilk's test was used to test the assumption of normality. Kruskal-Wallis's test was performed to determine differences concerning the SVL of sex and age classes among species, as well as the extracted PC1 and PC2 values (mentioned below) of a principal component analysis among four tiger gecko species in Vietnam except of *G. araneus*. For both tests we applied $p=0.05$. The SVL variable was excluded from subsequent analyses due to high collinearity with other dimensions. We further performed a Principal Component Analysis (PCA) of \log_{10} -transformed raw data of 18 remaining morphometric characters (except TaL and WT variables, due to many regenerated tails), using the packages “factoextra” and “FactoMinerR” to detect variances among the four selected Vietnamese species via contribution percentage of the PC1 and other PCs scores (Kassambara 2017). Morphometric variation of each species was illustrated by representatively clustered ellipse spaces with different coded-color within a spatial coordinate of the first two most important dimensions (PC1 and PC2) in the PCA analysis, to visually evaluate the overlap among the four species. In addition, we identified the overall difference in meristic characters among the 17 tiger gecko species native to China and Vietnam assigned to three groups (*G. lichtenfelderi*, *G. luei* and *G. yingdeensis*) by using a Multiple Correspondence Analysis (MCA). The meristic variation among the three groups was also visualized by convex ellipses within a spatial coordinate of the first two most important dimensions (Dim1 and Dim2) in the MCA analysis.

Results

Molecular phylogeny

The final matrix consisted of 949 bp aligned characters with 48 sequences of 14 ingroup and three outgroup taxa. The alignment contained no gaps. In total, 404 characters were found to be parsimony informative. The MP analysis produced 624 most parsimonious trees with 1404 steps (CI=0.54, RI = 0.85). Our phylogenetic analyses recovered a generally similar topology to those reported by Liang *et al.* (2018) and Zhu *et al.* (2020a, 2020b). Specifically, the *Goniurosaurus lichtenfelderi* group and the *G. luei* group form a monophyletic group with high nodal values from all analyses and the *G. kuroiwae* group is sister to the other two. In addition, each species group was also strongly corroborated as monophyletic from all analyses (Fig. 1). Within each species group, the results supported by this study are more similar to those generated by Liang *et al.* (2018), because Zhu *et al.* (2020a) used a different mitochondrial marker, the 16S ribosomal RNA gene. Nonetheless, we found several noted discrepancies between the two studies. Specifically, our phylogenetic analyses showed that *G. kuroiwae* (Namiye, 1912) is polyphyletic with high nodal support for each of the clades. In addition, *G. luei* was strongly corroborated as monophyletic in our Bayesian and MP analyses (BP=93%; PP=98%), but only weakly recovered as monophyletic in Liang *et al.* (2018) (BP=67%; PP=64%). *Goniurosaurus lichtenfelderi* was recovered as monophyletic with perfect nodal support from all analyses in our study, but *G. hainanensis* Barbour, 1908 is polyphyletic. On the other hand, both species are not monophyletic in Liang *et al.* (2018). The positions of *G. zhoui* and *G. bawanglingensis* Grismer, Haitao, Orlov & Anajeva, 2002 are interchanged in the trees supported by the two studies with strong nodal support from Liang *et al.* (2018) and weak corroboration from

ours. All surveyed sub-populations of *Goniurosaurus* from Vietnam were placed in four separate clades corresponding to four distinct species, *G. catbaensis*, *G. huiliensis*, *G. luii* and *G. lichtenfelderi* with significantly statistical support values, except for the ML analysis of *G. luii* (UBP=88%). Intra-specific genetic divergences of these species were relatively low: *G. catbaensis* (0.7–1.6%), *G. huiliensis* (0.4–1.2%), *G. lichtenfelderi* (0–0.7%), and *G. luii* (0–1.9%) (Fig. 1; Supp. file 1: Table S1).

Taxonomic accounts

Class Squamata Oppel, 1811
Order Gekkota Cuvier, 1817
Family Eublepharidae Boulenger, 1883
Genus *Goniurosaurus* Grismer, Viets & Boyle, 1999

***Goniurosaurus araneus* Grismer, Viets & Boyle, 1999**

Diagnosis

Body splayed and gracile, SVL 108.9–124.0 mm; external nares bordered by 6–8 nasal scales; supraorbital region with a row of enlarged tubercles; outer surface of upper eyelid composed of small granular

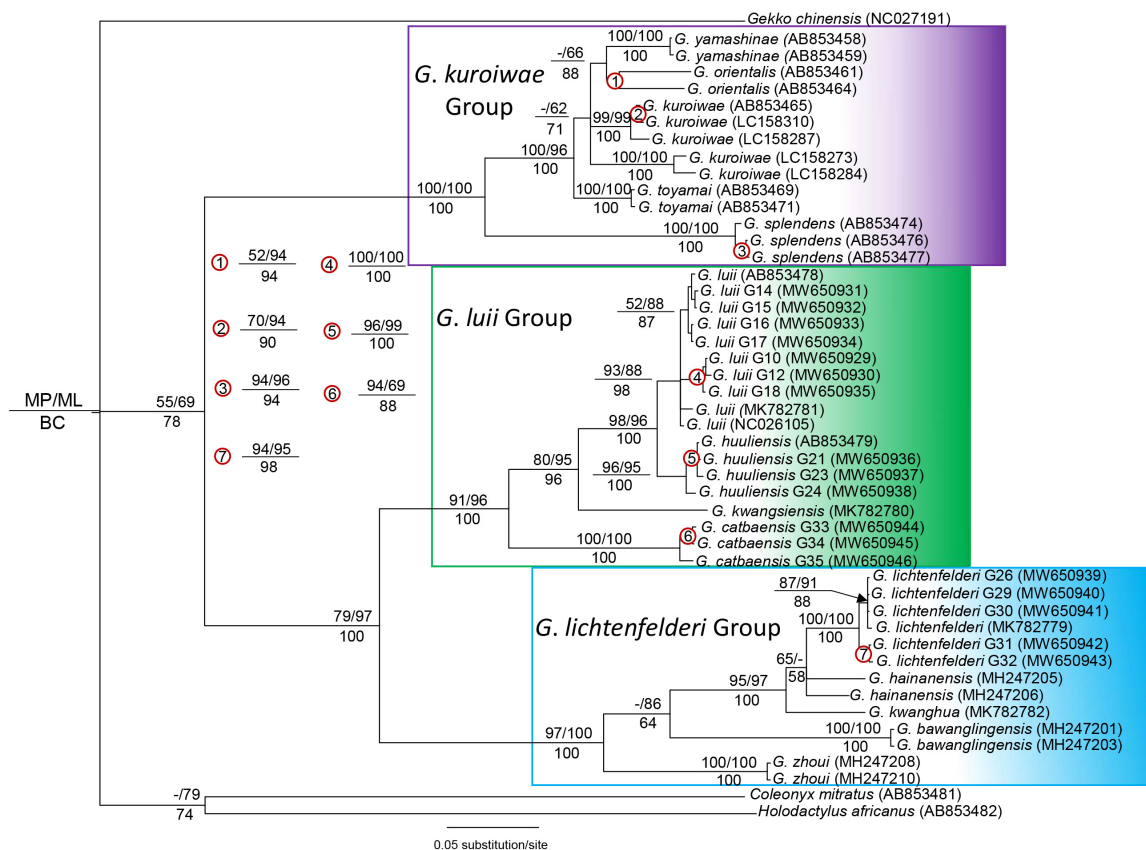


Fig. 1. Phylogram containing *Goniurosaurus* Grismer, Viets & Boyle, 1999 representatives from China, Japan and Vietnam based on the Bayesian analysis. Numbers above and below branches are MP/ML ultrafast bootstrap values and combined Bayesian posterior probabilities, respectively. Dash represents <50% value.

scales, one-half the size of those on top of head; internasal single; supralabials 8–10; infralabials 8 or 9; preorbital scales 13–18; eyelid fringe scales 52–67; postmentals 4–6; dorsal body scales elongate; paravertebral tubercles 29–38; scales around midbody 129–147; scales surrounding dorsal tubercles 10–14; axillary pockets deep; subdigital lamellae under fourth toe 23–25; precloacal pores in males 18–23; iris dark brown; dorsal ground color of head, body and limbs immaculate dull yellow-gray; nuchal loop posteriorly protracted (in V-shape); dorsal body bands between limb insertions 3, wide, edged anteriorly and posteriorly by wide dark brown bands; light band on tail base in width of 15 or 16 granular scales; ground color of tail black, caudal bands 5, white, completed ventrally; ventral surface of head, body and limbs dull white and immaculate (Grismer *et al.* 1999; Nguyen 2011; Chen *et al.* 2014).

Remarks

During our field work in northern Vietnam, including the type locality in Cao Bang Province, no specimen of *Goniurosaurus araneus* was seen. Therefore, the diagnosis was solely based on the descriptions of Grismer *et al.* (1999) and Chen *et al.* (2014).

Goniurosaurus catbaensis Ziegler, Nguyen, Schmitz, Stenke & Rösler, 2008

Fig. 2

Diagnosis

Body splayed and gracile, SVL 89.3–125.3 mm; external nares bordered by 6–8 nasal scales; supraorbital region with a row of enlarged tubercles; outer surface of upper eyelid composed of granular scales, about the same size of those on top of head and with a row of 6–10 enlarged tubercles; internasals absent; supralabials 8–11; infralabials 7–10; eyelid fringe scales 45–56; postmentals 2–5; gular region below lower jaws without enlarged tubercles; paravertebral tubercles 31–38; scale rows around midbody 112–127, granular scales surrounding tubercles 8–11; axillary pockets deep; subdigital lamellae under fourth toe 22–25; precloacal pores in males 16–23; iris orange-brown; dorsal ground color of head, body and limbs gray-brown to pale brown and mottled with dark brown blotches; nuchal loop thin, posteriorly protracted (in V-shape); dorsal body bands between limb insertions 3–4, thin, yellow, without dark spots; light band on tail base in width of 8–9 scales; ground color of tail black, caudal bands 5, white, completed ventrally; ventral surface of head, body and limbs dull white and immaculate, gular region with brown spots (modified after Ziegler *et al.* 2008; Nguyen 2011).

Description (Supp. file 1: Table S2)

Body splayed and gracile, adult males: SVL 92.9–125.3 mm (mean \pm SE: 112.3 \pm 0.8 mm, n=80), TaL 7.2–97.9 mm (75.2 \pm 1.9 mm); adult females: SVL 89.3–122.1 mm (111.8 \pm 0.8 mm, n=93), TaL 17.5–98.3 mm (70.3 \pm 1.7 mm); juveniles: SVL 53.4–78.8 mm (68.8 \pm 1.8 mm, n=21), TaL 11.8–61.2 mm (48.1 \pm 2.6 mm) (Supp. file 1: Table S2); head triangular, wider than neck, covered by uniform granular scales interspersed with tubercles in temporal and occipital regions; scales on rostrum slightly larger and flatter; enlarged supraorbital tubercles in a conspicuous row; middorsal portion of rostral partially sutured dorsomedially, bordered laterally by first supralabial on each side, dorsolaterally by prenasal on each side, and dorsally by two supranasals; internasal (postrostral) scales absent; external nares bordered by 6–8 nasals: anteriorly by prenasal and supranasal, dorsally by supranasal, posteriorly by two slightly enlarged postnasals and 1 or 2 smaller granular scales, and ventrally by prenasal; prenasals with long recurved ventral portion; supranasals triangular, meeting in midline behind rostral suture; preorbital scales 10–13; supralabials 8–11; infralabials 7–10; eyes relatively large, pupils vertical; eyelid fringe scales 45–56, those of upper eyelid slightly enlarged; outer surface of upper eyelid composed of granular scales of about the same size of those on top of head, including a row of 6–10 enlarged tubercles; fold of skin originating from suborbital region extends posteroventrally across angle of jaw; external auditory meatus elliptical with long axis directed dorsoventrally; tympanum deeply recessed; mental triangular,

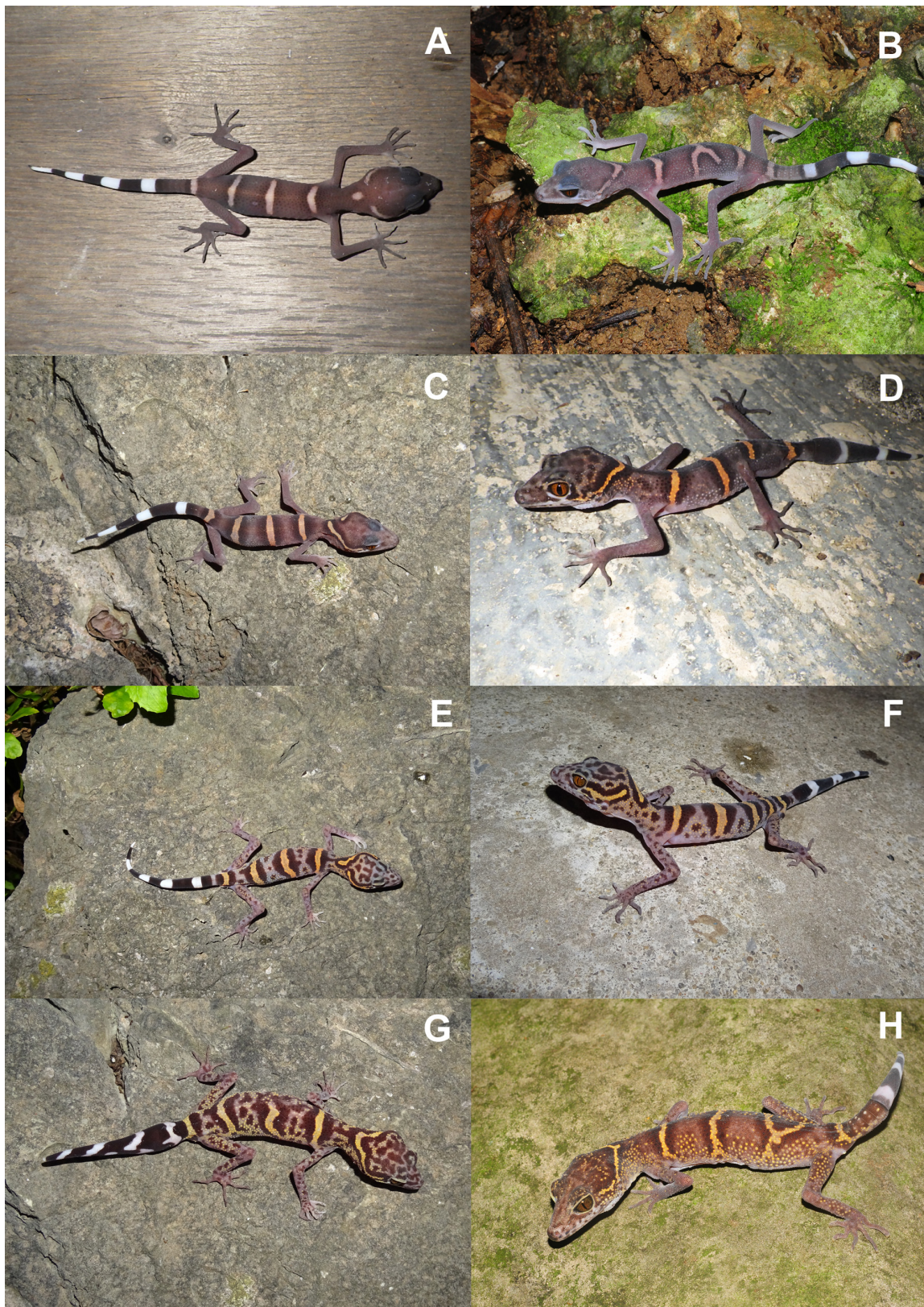


Fig. 2. *Goniurosaurus catbaensis* Ziegler, Nguyen, Schmitz, Stenke & Rösler, 2008. A. Juvenile (SVL=55 mm). B. Juvenile (SVL=69 mm). C. Juvenile (SVL=75 mm). D. Adult (SVL=91 mm). E. Adult (SVL=99 mm). F. Adult (SVL=105 mm). G. Adult (SVL=116 mm). H. Adult (SVL=125 mm).

bordered laterally by first infralabial on each side and posteriorly by 2–5 postmentals; postmentals bordered by 6–10 gular scales; gular region below lower jaws without enlarged tubercles; gular scales juxtaposed and granular, abruptly grading posteriorly into flat hexagonal pectoral scales and even larger ventral scales.

Neck narrower than body, covered with uniform granular scales interspersed with several sharply pointed conical tubercles on nape; tubercles on flanks conical, those of vertebral region somewhat lower in profile; dorsal body tubercles surrounded by 9–11 granular scales; dorsal tubercle rows at midbody 19–24; paravertebral tubercles between limb insertions 31–38, distinct vertebral row of tubercles absent; scales around midbody 112–127; larger ventral scales grade abruptly into smaller granular scales immediately anterior to vent at level of preanal pores; males with 18–23 precloacal pores in a transverse continuous series; region posterior to vent covered by flat juxtaposed scales and great hemipenial bulges, containing 2 or 3 enlarged postcloacal tubercles laterally on each side at level of vent; tail long and thin, thick at base, anteriorly with whorls; dorsal tail scales flat, smooth, up to 1.5 times the size of dorsal scales, arranged in more or less regular transverse rows; light band on tail base in width of 8–9 granular scales and with 7–8 tubercles in a transversal series; subcaudals larger than those on dorsal surface of tail.

Limbs relatively long and thin, covered dorsally with granular scales interspersed with several tubercles and ventrally with flat juxtaposed to subimbricate scales; dorsal granular scales grade into slightly flattened subimbricate scales on top of pes and manus; hind limbs slightly larger than forelimbs; larger granular scales on ventral surface of pes and manus; axillary pockets deep; subdigital lamellae wide, 9–11 under first finger, 18–21 under fourth finger, 9–12 under first toe, 22–25 under fourth toe; digits laterally compressed, increasing in length from first to fourth, fifth shorter than fourth (modified after Ziegler *et al.* 2008; Nguyen 2011).

Coloration in life (Fig. 2)

Dorsal ground color of head, body and limbs grey brown (in animals with SVL 53.4–105 mm) and blotches chestnut brown (in animals with SVL 110–125.3 mm), juveniles without small blotches, adults mottled with few circular blotches on body and limbs and long dark brown blotches on head; few yellow conical tubercles on franks of neck and body, and limbs in adults; iris light orange or red brown; five bands on the dorsal ground, thin, immaculate without dark spots, cream in juveniles and light orange or yellow in adults, all edged anteriorly and posteriorly by thin dark brown bands, including one thin nuchal loop extending from posterior corners of eyes and posteriorly protracted (in V-shape), three body bands between limb insertions, and another one on tail base; ground color of tail dark brown, and grey brown at mottled tail base; 3–5 immaculate white caudal bands, edged anteriorly and posteriorly in black; ventral surface of head, body and limbs dull white, juveniles absolutely immaculate, but adults with few dark spots on limbs, weak brown lateral spotting in gular region, venter and limbs.

Goniurosaurus huuliensis Orlov, Ryabov, Nguyen, Nguyen & Ho, 2008

Fig. 3

Diagnosis

Body splayed and gracile, SVL 97.2–134.6 mm; external nares bordered by 6–8 nasal scales; supraorbital region with a row of enlarged tubercles; outer surface of upper eyelid composed of granular scales, about one half the size of those on top of head and without enlarged tubercles; internasal 1 or 2 (rarely absent); supralabials 9–12; infralabials 9–12; preorbital scales 14–20; eyelid fringe scales 51–59; postmentals 2–4; gular region below lower jaws with enlarged tubercles; paravertebral tubercles 31–37; scale rows around midbody 118–130, granular scales surrounding tubercles 11–13; axillary pockets deep; subdigital lamellae under fourth toe 21–25; precloacal pores in males 25–30; iris reddish brown; dorsal ground color of head, body and limbs dark brown, without small dark brown blotches (dark blotches present

only on lower zone of flanks); nuchal loop thin, posteriorly protracted (in V-shape); dorsal body bands between limb insertions 3, thin, immaculate yellow; gular region with brown spots (modified after Orlov *et al.* 2008; Nguyen 2011).

Description (Supp. file 1: Table S2)

Body splayed and gracile, adult males SVL 97.2–132.2 mm (mean \pm SE: 118.9 \pm 1.4 mm, n=32), TaL 36.7–108.6 mm (72.7 \pm 3.7 mm), adult females SVL 97.4–134.6 mm (121.1 \pm 1.2 mm, n=46),

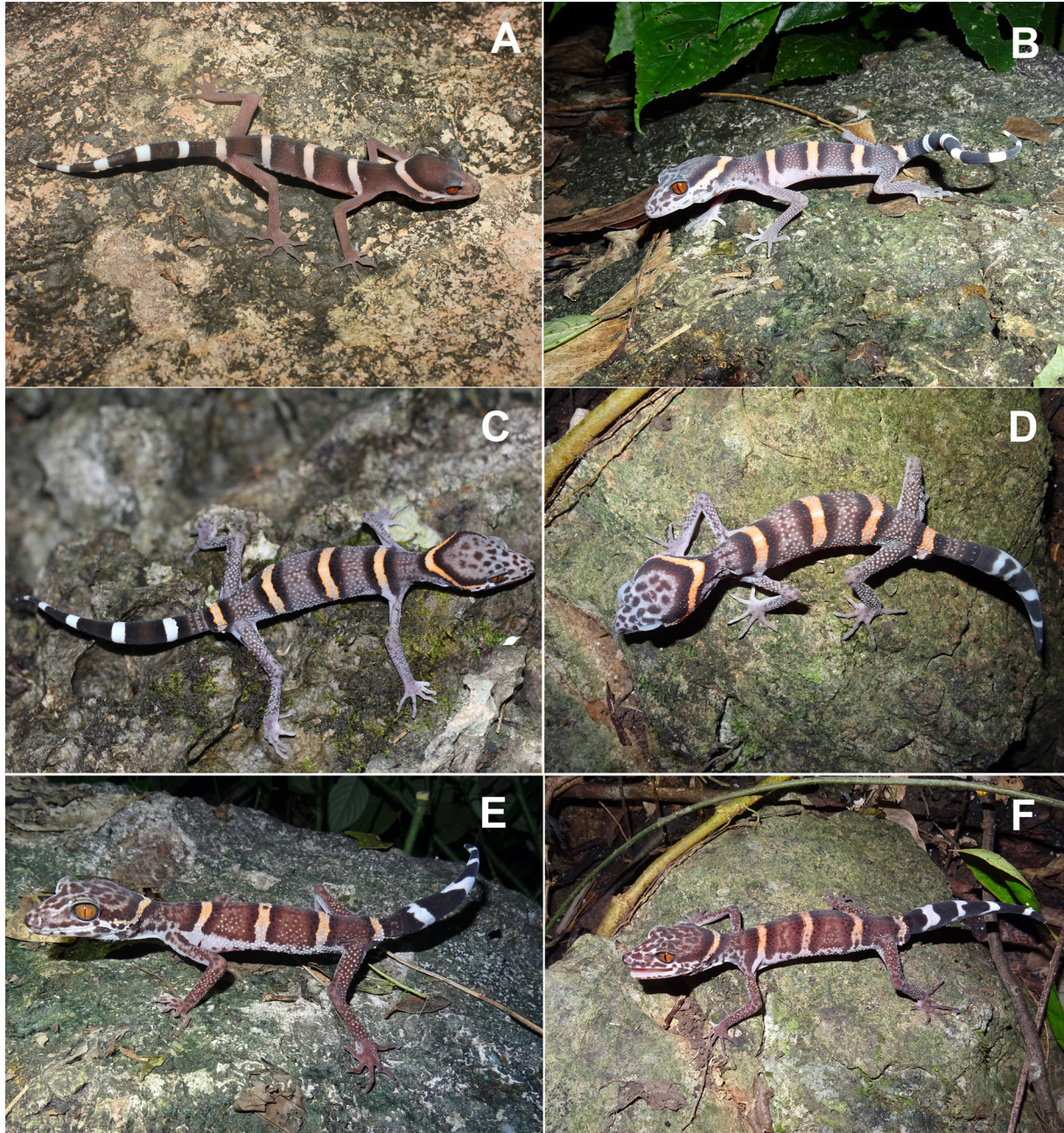


Fig. 3. *Goniurosaurus huuliensis* Orlov, Ryabov, Nguyen, Nguyen & Ho, 2008. **A.** Juvenile (SVL=74.4 mm). **B.** Adult (SVL=97 mm). **C.** Adult (SVL=108 mm). **D.** Adult (SVL=116 mm). **E.** Adult (SVL=125 mm). **F.** Adult (SVL=132 mm).

TaL 48.4–106.7 mm (78.2 ± 2.7 mm), juveniles SVL 73.5–74.4 mm ($n=2$), TaL 62.5–62.7 mm (Supp. file 1: Table S2); head triangular, wider than neck, covered by uniform granular scales interspersed with tubercles in temporal and occipital regions; scales on rostrum slightly larger and flatter; enlarged supraorbital tubercles in a conspicuous row; middorsal portion of rostral partially sutured dorsomedially, bordered laterally by first supralabial on each side, dorsolaterally by prenasal on each side, and dorsally by internasal and two supranasals; internasal 1–3 (rarely absent); external nares bordered by 5–7 nasals (prenasal, anterior and posterior supranasals, 2 slightly enlarged postnasal, and 1 or 2 granular scales); preorbital scales 14–20; supralabials 9–12, grading into granular scales posteriorly; infralabials 9–12; eyes large, pupils vertical; eyelid fringe scales 51–59, those of upper eyelid slightly enlarged; outer surface of upper eyelid composed of granular scales of about one half the size of those on top of head, without enlarged tubercles; fold of skin originating from suborbital region extends posteroventrally across angle of jaw; external auditory meatus elliptical with long axis directed dorsoventrally; tympanum deeply recessed; mental triangular, bordered laterally by first infralabial on each side and posteriorly by 2–4 postmentals; postmentals bordered by 7–10 gular scales; gular region below lower jaws with enlarged tubercles; gular scales juxtaposed and granular, grading posteriorly into flat hexagonal pectoral scales and even larger ventral scales.

Neck narrower than body, covered with uniform granular scales interspersed with several sharply pointed conical tubercles on nape; tubercles on flanks conical, those of vertebral region somewhat flatter; dorsal body tubercles surrounded by 11–13 granular scales; dorsal tubercle rows at midbody 19–24; paravertebral tubercles between limb insertions 31–37, distinct vertebral row of tubercles absent; scales around midbody 118–130; ventral scales large; males with 25–30 precloacal pores in continuous series, females without precloacal pores (but pitted scales present); region posterior to vent covered by flat juxtaposed scales and greatly swollen, containing 1 or 2 enlarged tubercles on each side at level of vent; tail thick at base; light band on tail base in width of 9–12 granular scales and with 8–10 tubercles in transversal series.

Limbs relatively long and thin, covered dorsally with granular scales interspersed with several tubercles and ventrally with flat juxtaposed to subimbricate scales; dorsal granular scales grade into slightly flattened subimbricate scales on top of pes and manus; hind limbs larger than forelimbs; axillary pockets deep; subdigital lamellae wide, 10 or 11 under first finger, 18–21 under fourth finger, 11 or 12 under first toe, 21–25 under fourth toe; digits laterally compressed, increasing in length from first to fourth, fifth shorter than fourth (modified after Orlov *et al.* 2008; Nguyen 2011).

Coloration in life (Fig. 3)

Dorsal ground color of head, body and limbs signal brown (juveniles), grey-brown (young adults, SVL: 97.2–116 mm) and signal brown (adults), without small dark brown blotches (dark blotches present only on lower zone of flanks); dull white tubercles on dorsal body, limbs in juveniles, few orange tubercles on limbs in adults; iris orange or red brown; five bands on the dorsal ground, thin, immaculate without dark spots, slight bisque in juveniles, orange brown or yellow in adults, all edged anteriorly and posteriorly by thin dark brown bands, including one thin nuchal loop extending from posterior corners of eyes and posteriorly protracted (in V-shape), three body bands between limb insertions, and another one on tail base; ground color of tail dark brown, and signal brown at mottled tail base; 3–6 immaculate white caudal bands; ventral surfaces of head, body and limbs dull white and immaculate except for few dark brown spots on margin regions of gular and limbs.

Goniurosaurus luii Grismer, Viets & Boyle, 1999
Fig. 4

Diagnosis

Body splayed and gracile, SVL 86.5–126.5 mm; external nares bordered by 5–8 nasal scales; supraorbital region with a row of enlarged tubercles; outer surface of upper eyelid composed of granular scales, about one half the size of those on top of head and without enlarged tubercles; internasals 1 or 2; supralabials 8–12; infralabials 8–11; preorbital scales 13–16; eyelid fringe scales 46–61; postmentals 2–6; gular region below lower jaws with enlarged tubercles; paravertebral tubercles 29–38; scale rows around midbody 119–144, granular scales surrounding tubercles 9–14; axillary pockets deep; subdigital lamellae under fourth toe 20–26; precloacal pores in males 23–32; iris brown or bright orange; dorsal ground color of head, body and limbs pale brown to grey brown, mottled with small dark brown blotched; nuchal loop thin, posteriorly protracted (in V-shape); dorsal body bands between limb insertions 3, thin, immaculate yellow; gular region, belly, and ventral surface of limbs with brown spots (Grismer *et al.* 1999; Vu *et al.* 2006; Nguyen 2011).

Description (Supp. file 1: Table S2)

Body splayed and gracile, males: SVL 88.8–123.0 mm (mean \pm SE: 109.3 \pm 1.4 mm, n=34), TaL 2.8–96.8 mm (72.5 \pm 3.8 mm); adult females: SVL 86.5–126.5 mm (112.4 \pm 1.3 mm, n=43), TaL 43.3–102.2 mm (72.2 \pm 2.1 mm); juveniles: SVL 55.0–84.8 mm (74.8 \pm 3.1 mm, n=11), TaL 44.2–68.9 (58.7 \pm 2.7 mm) (Supp. file 1: Table S2); head triangular, wider than neck, covered by uniform granular scales interspersed with tubercles in temporal and occipital regions; scales on rostrum slightly larger and flatter; enlarged supraorbital tubercles in a conspicuous row; middorsal portion of rostral partially sutured dorsomedially, bordered laterally by first supralabial on each side, dorsolaterally by prenasal on each side, and dorsally by 1 or 2 internasal and two supranasals; internasals 1 or 2; external nares bordered by 5–8 nasal scales (prenasal, anterior and posterior supranasals, 2 slightly enlarged postnasal, and 1–3 granular scales); preorbital scales 13–16; supralabials 8–12, grading into granular scales posteriorly; infralabials 8–11; eyes large, pupils vertical; eyelid fringe scales 46–56, those of upper eyelid slightly enlarged; outer surface of upper eyelid composed of granular scales, about one half the size of those on top of head and without enlarged tubercles; fold of skin originating in suborbital region extends posteroventrally across angle of jaw; external auditory meatus elliptical with long axis directed dorsoventrally; tympanum deeply recessed; mental triangular, bordered laterally by first infralabial on each side and posteriorly by 2–6 postmentals; postmentals bordered by 6–11 gular scales; gular region below lower jaws with enlarged tubercles; gular scales juxtaposed and granular, abruptly grading posteriorly into flat hexagonal pectoral scales and even larger ventral scales.

Neck narrower than body, covered with uniform granular scales interspersed with several sharply pointed conical tubercles on nape; tubercles on flanks conical, those of vertebral region somewhat more flat; dorsal body tubercles surrounded by 11–13 granular scales; dorsal tubercle rows at midbody 20–24; paravertebral tubercles between limb insertions 29–38, distinct vertebral row of tubercles absent; scale rows around midbody 119–144; ventral scales large; males with 24–32 precloacal pores in a transverse continuous series, females without precloacal pores (but pitted scales present); region posterior to vent covered by flat juxtaposed scales and greatly swollen, containing 1–3 enlarged tubercles on each side at level of vent; tail thick at base, light band on tail base in width of 9–13 granular scales with 9–10 tubercles in transversal series; ventral caudals of tail base larger than dorsal caudals.

Limbs relatively long and thin, covered dorsally with granular scales interspersed with several tubercles and ventrally with flat juxtaposed to subimbricate scales; dorsal granular scales grade into slightly flattened subimbricate scales on top of pes and manus; hind limbs larger than forelimbs; axillary pockets deep; subdigital lamellae wide, 9–12 under first finger, 17–22 under fourth finger, 10–12 under first toe,

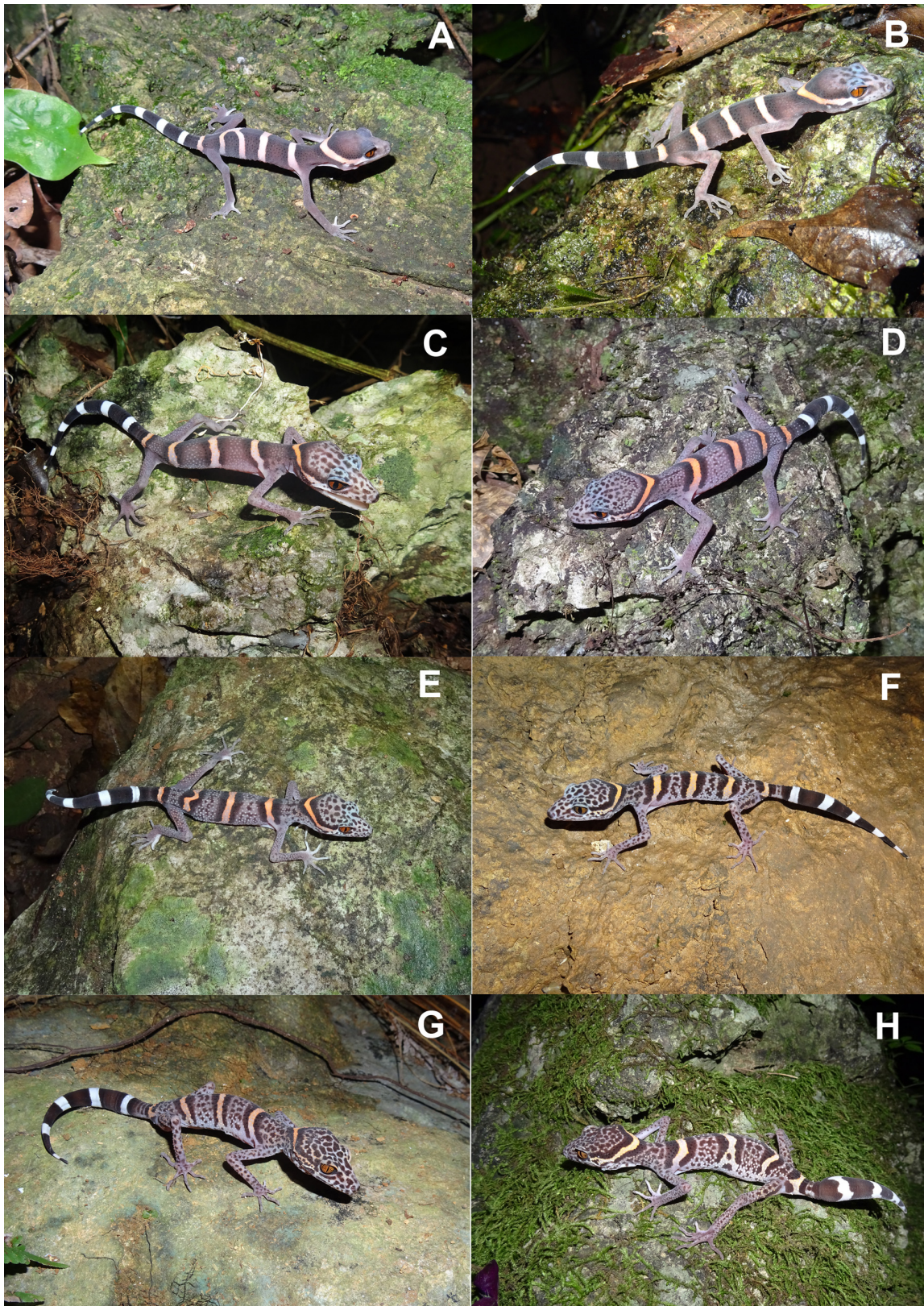


Fig. 4. *Goniurosaurus luii* Grismer, Viets & Boyle, 1999. A. Juvenile (SVL=61 mm). B. Juvenile (SVL=70 mm). C. Juvenile (SVL=80 mm). D. Adult (SVL=89 mm). E. Adult (SVL=96 mm). F. Adult (SVL=104 mm). G. Adult (SVL=116 mm). H. Adult (SVL=126 mm).

20–26 under fourth toe; digits laterally compressed, increasing in length from first to fourth, fifth shorter than fourth (Grismer *et al.* 1999; Vu *et al.* 2006; Nguyen 2011).

Coloration in life (Fig. 4)

Dorsal ground color of head, body and limbs pale brown to grey brown, mottled with small dark brown circle blotches, juveniles without small dark brown blotches (except juveniles $SVL \geq 72$ mm initially present on head); iris red brown or bright orange; five bands on dorsal ground, thin, immaculate without dark spots, slight bisque in juveniles, orange brown or yellow in adults, all edged anteriorly and posteriorly by thin dark brown bands, including one thin nuchal loop extending from posterior corners of eyes and posteriorly protracted (in V-shape), three body bands between limb insertions, and another one on tail base; ground color of tail nearly solid black, and grey brown at mottled tail base; 3–6 immaculate white caudal bands; ventral surfaces of head, body and limbs dull white with a few dark brown spots in gular region, on belly and limbs.

Goniurosaurus lichtenfelderi (Mocquard, 1897)

Fig. 5

Diagnosis

Body robust; SVL 80.6–113.5 mm; external nares bordered by 5–10 nasal scales; supraorbital region with a row of slightly enlarged tubercles; outer surface of upper eyelid composed of granular scales, about one-half the size of those on top of head and without enlarged tubercles; internasals 1–5 (rarely 1:2; 2:2 or 2:3); supralabials 7–10; infralabials 6–9; preorbital scales 12–19; eyelid fringe scales 43–58; postmentals 2–6; paravertebral tubercles 22–33; scale rows around midbody 117–131, granular scales surrounding tubercles 10–13; axillary pockets shallow; subdigital lamellae under fourth toe 17–21; precloacal pores in males 25–33, in females 17–21; dorsal ground color of head, body and limbs dark purple-brown, without small dark brown blotches; transverse body bands 4, nuchal loop thin, posteriorly rounded, in U-shape; dorsal body bands between limb insertions 2, thin, light yellow; gular region and without dark spots; ventral surfaces of head, body and limbs dull white with a few dark dots on margin regions (modified after Grismer 2000; Grismer *et al.* 2002; Nguyen 2011).

Description (Supp. file 1: Table S2)

Body robust, adult males: SVL 80.6–113.5 mm (mean \pm SE: 97.8 \pm 0.7 mm, n=92), TaL 6.2–84.8 mm (61.5 \pm 1.7 mm); adult females: SVL 81.0–105.5 mm (96.0 \pm 0.7 mm, n=72), TaL 22.7–81.2 mm (58.4 \pm 1.4 mm); juveniles: SVL 41.1–77.1 mm (65.4 \pm 2.9 mm, n=14), TaL 20.9–64.7 mm (48.3 \pm 3.4 mm) (Supp. file 1: Table S2); head triangular, wider than neck, covered by uniform granular scales interspersed with tubercles in temporal and occipital regions; scales on rostrum slightly larger and flatter; enlarged supraorbital tubercles in a conspicuous row; middorsal portion of rostral partially sutured dorsomedially, bordered laterally by first supralabial on each side, dorsolaterally by prenasal on each side, and dorsally by 1 or 2 internasals and two supranasals; internasals 1–5 (rarely 1:2; 2:2 or 2:3); external nares bordered by 5–10 nasals; preorbital scales 12–18; supralabials 7–10, grading into granular scales posteriorly; infralabials 6–9; eyes large, pupils vertical; eyelid fringe scales 47–58, those of upper eyelid slightly enlarged; outer surface of upper eyelid composed of granular scales of about one-half the size of those on top of head, without enlarged tubercles; fold of skin originating in the suborbital region extends posteroventrally across angle of jaw; external auditory meatus elliptical; tympanum deeply recessed; mental triangular, bordered laterally by first infralabial on each side and posteriorly by 2–5 postmentals; postmentals bordered by 7–10 gular scales; gular region below lower jaws without enlarged tubercles; gular scales juxtaposed and granular, abruptly grading posteriorly into flat hexagonal scales and even larger ventral scales.

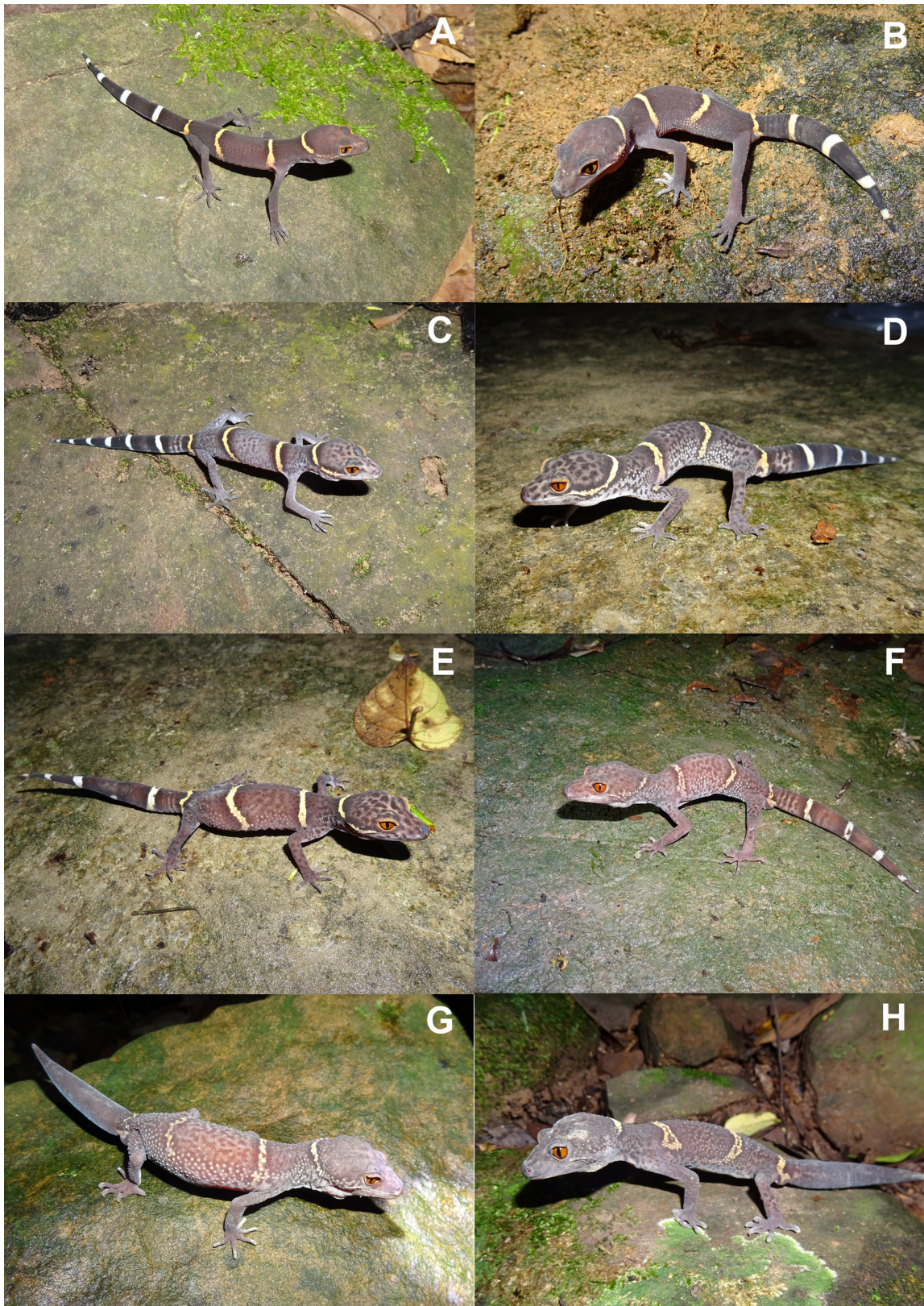


Fig. 5. *Goniurosaurus lichtenfelderi* (Mocquard, 1897). **A.** Juvenile (SVL=55 mm). **B.** Juvenile (SVL=63 mm). **C.** Juvenile (SVL=72 mm). **D.** Adult (SVL=80 mm). **E.** Adult (SVL=91 mm). **F.** Adult (SVL=97 mm). **G.** Adult (SVL=105 mm). **H.** Adult (SVL=113 mm).

Neck narrower than body, covered with uniform granular scales interspersed with several sharply conical tubercles on nape; tubercles on flanks conical, those of vertebral region somewhat more flat; dorsal body tubercles surrounded by 10–13 granular scales; dorsal tubercle rows at midbody 15–22; paravertebral tubercles between limb insertions 22–33, distinct vertebral row of tubercles absent; scales around midbody 117–130; ventral scales large; males with 24–33 precloacal pores in a transverse continuous series, females without distinct precloacal pores (but 25–33 pitted scales present); region posterior to vent covered by flat juxtaposed scales and greatly swollen, containing 1 (rarely 2) enlarged tubercles on each side at the level of vent; tail thick at base; light band on tail base in width of 4–7 scales and with 6–10 tubercles in a transversal series.

Limbs robust, covered dorsally with granular scales interspersed with several tubercles and ventrally with flat juxtaposed to subimbricate scales; dorsal granular scales grade into slightly flattened subimbricate scales on top of pes and manus; hind limbs larger than forelimbs; axillary pockets shallow; subdigital lamellae wide, 6–9 under first finger, 13–17 under fourth finger, 6–10 under first toe, 16–20 under fourth toe; digits laterally compressed, increasing in length from first to fourth, fifth shorter than fourth (modified after Grismer 2000; Grismer *et al.* 2002; Nguyen 2011).

Coloration in life (Fig. 5)

Dorsal ground color of head, body and limbs grey and grey brown in juveniles without blotches and chestnut brown in adults with dark brown blotches (in animals with SVL > 80 mm); iris orange or redish brown; four bands on the dorsal ground, thin, yellow in both juveniles and adults, immaculate without dark spots (few dark brown spots present in some animals), all edged anteriorly and posteriorly by thin dark brown bands, including one thin nuchal loop extending from posterior corners of eyes and posteriorly rounded (in U-shape), two body bands between limb insertions, and another one on tail base; ground color of tail dark brown, and grey brown at mottled tail base; 3–5 immaculate white caudal bands except first band slightly yellow, and some animals with regenerated tail present immaculate grey brown without white caudal bands; ventral surfaces of head, body and limbs dull white with a few dark dots on margin regions of belly, limbs, gular region, but immaculate dull white in juveniles (SVL ≤ 65 mm) without dark spots.

Morphological comparisons

Our morphological comparisons of the tiger gecko species (except for *G. araneus*) from Vietnam revealed an overall significant difference in the snout-vent length (SVL). As the result, *G. huuliensis* had the largest SVL, while *G. lichtenfelderi* had the shortest SVL in both adult males and females, compared to the other tiger geckos ($P < 0.05$). However, the SVL of *G. catbaensis* was not significantly different from that of *G. luii* ($P > 0.05$, Supp. file 2: Fig. S1).

The PCA of 18 selected morphometric characters estimated the first (PC1) and second principal (PC2) components to explain 82.5% (74.5% and 8.0%, respectively) of the variance among the four investigated species of *Goniurosaurus* in Vietnam (Supp. file 2: Fig. S2). The extracted PC1 and PC2 scores of the PCA differ significantly among the four species (Kruskal-Wallis's test, $P < 0.05$), of which the values and morphometric spaces of *G. lichtenfelderi* are not concordant with the three remainders (Fig. S2). Three morphological factors, namely head length (HL), head width (HW), and mouth width (MW) highly account for the overall difference in the PC1 score, while the PC2 score is mainly explained by the body height (BH) and body width (BW) (Supp. file 2: Fig. S2).

Regarding the comparisons of the 17 species of *Goniurosaurus* from China and Vietnam, the multiple correspondence analysis (MCA) clustered them separately into three groups (*G. lichtenfelderi* group, *G. luii* group and *G. yingdeensis* group) represented with different meristic spaces on the spatial coordinate of Dim 1 and Dim 2 (Fig. 6A, Table 1). The characters of fourth toe (LT4), body limb shape

Table 1 (continued on the next two pages). Meristic characters (minimum–maximum) of 17 species of *Goniurosaurus* Grismer, Viets & Boyle, 1999 in China and Vietnam. Data of *G. araneus* Grismer, Viets & Boyle, 1999 from Grismer *et al.* (1999) and Chen *et al.* (2014); *G. bawanglingensis* Grismer, Haitao, Orlov & Anajeva, 2002 from Grismer *et al.* (2002); *G. catbaensis* Ziegler, Nguyen, Schmitz, Stenke & Rösler, 2008, *G. huiliensis* Orlov, Ryabov, Nguyen, Nguyen & Ho, 2008, *G. luii* Grismer, Viets & Boyle, 1999 and *G. lichtenfelderi* (Mocquard, 1897) from this study; *G. gezhi* Zhu, He & Li, 2020 from Zhu *et al.* (2020a); *G. gollum* Qi, Wang, Grismer, Chen, Lyu & Wang, 2020 from Qi *et al.* (2020b); *G. hainanensis* Barbour, 1908 from Grismer *et al.* (1999, 2002); *G. kadoorieorum* Yang & Chan, 2015 and *G. kwangsiensis* Yang & Chan, 2015 from Yang & Chan (2015); *G. kwanghua* Zhu & He, 2020 from Zhu *et al.* (2020b); *G. liboensis* Wang, Yang & Grismer, 2013 from Wang *et al.* (2013); *G. varius* Qi, Grismer, Lyu, Zhang, Li & Wang, 2020 from Qi *et al.* (2020a); *G. yingdeensis* Wang, Yang & Cui, 2010 from Wang *et al.* (2010) and Qi *et al.* (2020b); *G. zhelong* Wang, Jin, Li & Grismer, 2014 from Wang *et al.* (2014) and Qi *et al.* (2020b); and *G. zhoui* Zhou, Wang, Chen & Liang, 2018 from Zhou *et al.* (2018).

<i>Goniurosaurus luii</i> group									
	<i>G. araneus</i>	<i>G. catbaensis</i>	<i>G. gezhi</i>	<i>G. huiliensis</i>	<i>G. kadoorieorum</i>	<i>G. kwangsiensis</i>	<i>G. liboensis</i>	<i>G. luii</i>	
Enlarged row of supraorbital tubercles (0) absent. (1) present	1	1	1	1	1	1	1	1	1
Scales of upper eyelid to top of head ½ size (1) of those on the top of the head or equal in size (2)	1	2	–	1	2	2	2	1	1
Deep axillary pockets (0) absent. (1) present	1	1	1	1	1	1	1	1	1
Body and limb (1) splayed–gracile (2) compact–robust	1	1	1	1	1	1	1	1	1
Posterior nuchal loop (1) protracted (2) rounded	1	1	1	1	1	1	1	1	1
Number of Body bands	4	4	4	4	4	4	4	4	4
Dorsal body bands (1) immaculate (2) maculate	1	1	1	1	1	1	1	1	1
Dark borders of body band (1) wide (2) narrow	1	1	1	1	1	1	1	1	1
Adult ground color (1) mottled (2) immaculate	2	1	1	2	1	1	1	1	1
Lateral spotting on belly (0) absent (1) present	0	1	–	1	1	1	0	1	1
IN	1–2	0–1	0–1	0–3	2	1–2	2–3	1–2	1–2
P–IN	–	0–2	3–5	0–7	3–9	1–2	3	2–6	2–6
SPL	8–10	8–11	9–10	9–12	10–11	8–10	9–11	8–12	8–12
IFL	8–9	7–10	8–10	9–12	9	7–9	9–12	8–11	8–11
N	6–8	6–8	6–7	5–7	6–7	6–7	8–9	5–8	5–8
PM	4–6	2–5	3–5	2–4	4–5	3–6	3–5	2–6	2–6

Table 1 (continued).

<i>Goniurosaurus luii</i> group									
	<i>G. araneus</i>	<i>G. catbaensis</i>	<i>G. gezhi</i>	<i>G. huiliensis</i>	<i>G. kadoorieorum</i>	<i>G. kwangsiensis</i>	<i>G. liboensis</i>	<i>G. luii</i>	
GP	7–9	6–10	7	7–10	8–11	7–9	9–14	6–11	
PO	13–18	10–1	15–19	14–20	15–19	15–19	16–18	13–16	
CIL	52–67	45–56	44–52	51–59	47–55	52–58	52–59	46–56	
MB	129–147	112–127	123–151	118–130	124–132	122–128	127–129	119–144	
GST	10–14	9–11	10–12	11–13	11–13	10–13	10–13	11–13	
TL	32–38	31–38	32–39	31–37	30–34	27–32	27–28	29–38	
DTR	21–22	19–24	20–21	19–24	22–24	20–22	23–24	20–24	
LD1	9–12	9–11	9–11	10–11	10–11	10–12	9–10	9–12	
LD4	19–21	18–21	20–21	18–21	17–19	18–21	17–19	17–22	
LT1	9–14	9–12	9–12	11–12	10–11	11–13	11–12	10–12	
LT4	23–24	22–25	21–25	21–25	21–24	22–27	23–26	20–26	
PP	18–23	18–23	18–20	25–30	26–28	31–33	23	24–32	
PAT	3–6	2–3	2	1–2	1–2	1–2	2–4	1–3	

<i>Goniurosaurus lichtenfelderi</i> group									
	<i>G. bavanglingensis</i>	<i>G. hainanensis</i>	<i>G. kwanghua</i>	<i>G. lichtenfelderi</i>	<i>G. zhoui</i>	<i>G. gollum</i>	<i>G. varius</i>	<i>G. yingdeensis</i>	<i>G. zhelongi</i>
Enlarged row of supraorbital tubercles (0) absent.	0	0	1	1	1	0	0	1	1
(1) present									
Scales of upper eyelid to top of head ½ size (1) of those on the top of the head or equal in size (2)	1	2	1	1	2	2	2	1	1
Deep axillary pockets (0) absent. (1) present	1	0	0	0	1	1	1	1	1
Body and limb (1) splayed–gracile (2) compact–robust	2	2	2	2	2	1	1	1	1
Posterior nuchal loop (1) protracted (2) rounded	2	2	2	2	1	2	2	2	2
Number of body bands	4	3	3	3	4	4	4	4	4

Table 1 (continued).

	<i>Goniurosaurus lichenfelderi</i> group										<i>Goniurosaurus yingdeensis</i> group				
	<i>G.</i> <i>bawanglingensis</i>	<i>G.</i> <i>hainanensis</i>	<i>G.</i> <i>kwanghua</i>	<i>G.</i> <i>lichenfelderi</i>	<i>G.</i> <i>zhoui</i>	<i>G.</i> <i>gollum</i>	<i>G.</i> <i>varius</i>	<i>G.</i> <i>yingdeensis</i>	<i>G.</i> <i>zhelongi</i>						
Dorsal body bands (1) immaculate (2) maculate	2	2	2	1	2	1	1	1	1						
Dark borders of body band (1) wide (2) narrow	2	2	2	2	1	1	1	1	1						
Adult ground color (1) mottled (2) immaculate	1	1	1	1	1	1	1	1	1						
Lateral spotting on belly (0) absent (1) present	0	0	0	1	0	1	1	1	1						
IN	–	–	0–1	1–5	1	1	1–2	2–3	1–2						
P–IN	–	–	0–2	2	2–4	2	3–4	2–6	3						
SPL	8–10	7–10	7–9	7–10	8–9	10	7–10	8–10	7–10						
IFL	7–11	6–9	6–8	6	7–9	10	8–9	8–10	6–9						
N	–	–	9–10	5–10	8	8–9	7–9	7–11	6–8						
PM	2–3	2–5	3–5	2–5	3–4	2–3	3–4	2–4	4–6						
GP	–	–	4–8	7–10	5–7	7–8	6–8	5–7	7–9						
PO	12–18	14–19	18–21	12–18	15–20	15–17	11–16	16–20	13–17						
CIL	56–67	55–70	47–49	47–58	49–62	59–63	50–56	46–64	42–53						
MB	104–133	95–125	109–118	117–130	130–140	121–128	101–110	102–115	99–109						
GST	9–13	11–15	11–12	10–13	11	9–11	8–12	8–12	9–12						
TL	32–36	23–32	25–26	22–33	24–32	25–26	27–29	25–33	28–33						
DTR	–	–	20–21	15–22	19–22	16–17	21–24	20–25	23–28						
LD1	–	–	9–10	6–9	9–10	10	7	8	7–8						
LD4	–	–	18–19	13–17	14–16	13/14	15–17	18–19	15–17						
LT1	–	–	9–10	6–10	9–11	15/16	8	11–12	7–9						
LT4	18–22	18–23	18–19	17–20	19–22	22/23	18/21	19–24	17–22						
PP	37–46	24–31	28	24–33	36–38	10–11	10	10–13	9–12						
PAT	–	–	1–2	1–2	2–3	2	2	2	2						

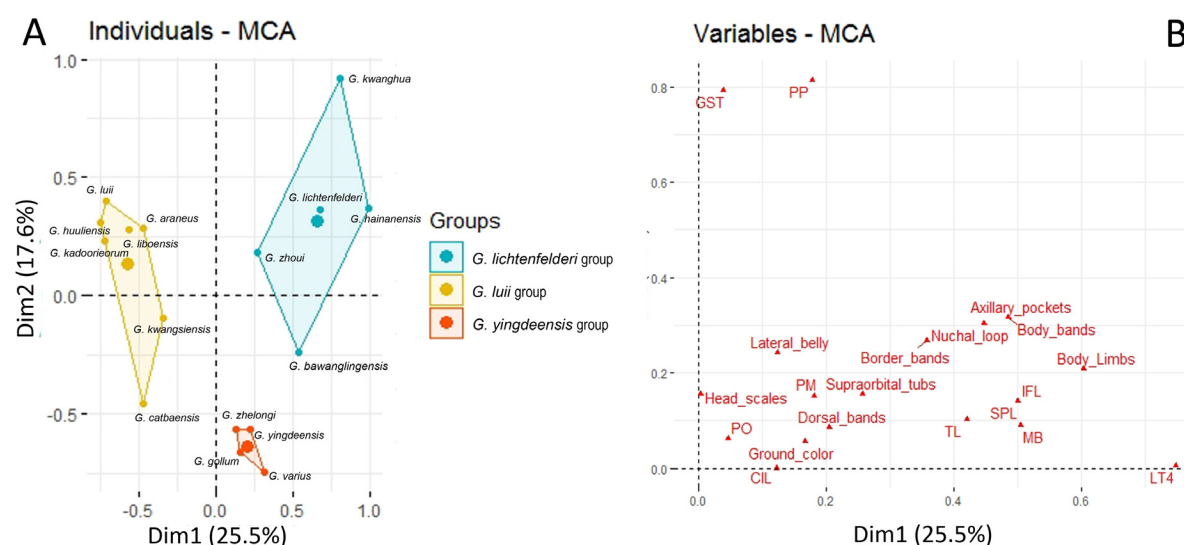


Fig. 6. **A.** Multiple correspondence analysis (MCA) on meristic variation among 17 recorded species of *Goniurosaurus* Grismer, Viets & Boyle, 1999 in China and Vietnam. **B.** Correlation between meristic variables and principal dimensions (Dim1 and Dim2).

and body bands highly account for the Dim1 score, whereas the characters of precloacal pores (PP) and granular scales surrounding dorsal tubercles (GST) are the most correlated with the Dim 2 score (Fig. 6B).

Key to the species of *Goniurosaurus* (Fig. 7, Table 1)

Modified from Grismer *et al.* (1999, 2002), Vu *et al.* (2006), Orlov *et al.* (2008), Ziegler *et al.* (2008), Wang *et al.* (2010, 2013, 2014), Nguyen (2011), Chen *et al.* (2014), Yang & Chan (2015), Honda & Ota (2017), Zhou *et al.* (2018, 2020a, 2020b), Qi *et al.* (2020a, 2020b) and the present study.

1. Precloacal pores in males present, claws are sheathed by scales 7
 - Precloacal pores absent, unsheathed claws (*G. kuroiwa* group) 2
2. Yellow brown to gold iris; a single scale at the base of each digit which is occasionally slightly enlarged *G. yamashinae* (Okada, 1936)
 - Blood-red iris; one to three enlarged scales at the base of each digit on the limbs 3
3. Adults without interspace mottling 4
 - Adults with interspace mottling 5
4. Robust body stature; dorsal body tubercles between the limb insertions in cross-section are triangular to elliptical and sharply keeled anteriorly; 34–42 paravertebral tubercles; ventral scales are juxtaposed and sharply raised *G. toyamai* Grismer, Ota & Tanaka, 1994
 - Slender body stature; dorsal body tubercles between the limb insertions in cross-section are smooth or very weakly keeled; 22–29 paravertebral tubercles; ventrals are flat, wide, and subimbricate to imbricate *G. splendens* (Nakamura & Uéno, 1959)
5. Dorsal banding absent or incomplete *G. kuroiwa* (Namiye, 1912)
 - Prominent dorsal pattern consisting of transverse bands between the nape of the neck and the caudal constriction 6

6. No such reddish or pinkish tint in dorsal pattern and iris *G. orientalis* (Maki, 1931)
 - Reddish or pinkish tint to some extent in dorsal pattern and iris *G. sengokui* (Honda & Ota, 2017)
7. Precloacal pores in males less than 16 (*G. yingdeensis* group) 8
 - Precloacal pores in males more than 16 11
8. Scales around midbody 121–128; longitudinal dorsal tubercle rows at midbody 16–17
 - *G. gollum* Qi, Wang, Grismer, Chen, Lyu & Wang, 2020
 - Scales around midbody 99–115; longitudinal dorsal tubercle rows at midbody 20–28 9
9. Nuchal loop and body bands with small dark blotches; enlarged row of supraorbital tubercles absent; trunk of body usually with a longitudinal light vertebral stripe
 - *G. varius* Qi, Grismer, Lyu, Zhang, Li & Wang, 2020
 - Nuchal loop and body bands without dark blotches; enlarged row of supraorbital tubercles present; trunk of body without a longitudinal light vertebral stripe 10
10. Tubercles between orbits present; gular scales bordering the postmentals 2–4; preorbital scales 5–7
 - *G. yingdeensis* Wang, Yang & Cui, 2010
 - Tubercles between orbits absent; gular scales bordering the postmentals 4–6; preorbital scales 7–9 *G. zhelongi* Wang, Jin, Li & Grismer, 2014
11. Body and limbs robust (*G. lichtenfelderi* group) 12
 - Body and limbs splayed gracile (*G. luii* group) 16
12. Precloacal pores in males 37–46 *G. bawanglingensis* Grismer, Haitao, Orlov & Anajeva, 2002
 - Precloacal pores in males fewer than 37 13
13. Nuchal loop protracted posteriorly (in V-shape); number of body bands 4; axillary pockets deep; precloacal pores in males more than 33 *G. zhoui* Zhou, Wang, Chen & Liang, 2018
 - Nuchal loop rounded posteriorly (in U-shape); number of body bands 3; axillary pockets shallow; precloacal pores in males fewer than 33 14
14. Eyelid fringe scales 55–70; enlarged row of supraorbital tubercles absent; scales of upper eyelid to top of head equal of those on the top of the head in size *G. hainanensis* Barbour, 1908
 - Eyelid fringe scales 47–58; enlarged row of supraorbital tubercles present; scales of upper eyelid to top of head ½ of those on the top of the head in size 15
15. Lateral spotting on belly absent; preorbital scales 18–21; scales around midbody 109–118; subdigital lamellae under the fourth finger 18–19 *G. kwanghua* Zhu & He, 2020
 - Lateral spotting on belly present; preorbital scales 12–18; scales around midbody 117–130; subdigital lamellae under the fourth finger 13–17 *G. lichtenfelderi* (Mocquard, 1897)
16. Internasal absent *G. catbaensis* Ziegler, Nguyen, Schmitz, Stenke & Rösler, 2008
 - Internasal present 17
17. Postcloacal tubercles 3–6 *G. araneus* Grismer, Viets & Boyle, 1999
 - Postcloacal tubercles 1–3 18
18. Adult body length (SVL) > 126 mm; dorsum without dark blotches
 - *G. huuliensis* Orlov, Ryabov, Nguyen, Nguyen & Ho, 2008
 - Maximum body length (SVL) ≤ 126 mm; dorsum with dark blotches 19

19. Dorsum with many small dark blotches *G. luii* Grismer, Viets & Boyle, 1999
 – Dorsum with rarely scattered dark blotches 20
20. Outer surface of upper eyelid composed of granular scales, about the same size of those on top of head and with enlarged tubercles *G. gezhi* Zhu, He & Li, 2020
 – Enlarged row of supraorbital tubercles present; scales of upper eyelid to top of head ½ of those on the top of the head in size 21
21. Precloacal pores in males 31–33 *G. kwangsiensis* Yang & Chan, 2015
 – Precloacal pores in males fewer than 31 22
22. Lateral spotting on belly present; nasal scales 6–7; paravertebral tubercles between limb insertions 30–34; precloacal pores 26–28 *G. kadoorieorum* Yang & Chan, 2015
 – Lateral spotting on belly absent, nasal scales 8–9; paravertebral tubercles between limb insertions 27–28; precloacal pores 23 *G. liboensis* Wang, Yang & Grismer, 2013

Discussion

Taxonomic review

Our phylogenetic analyses strongly suggest that populations recorded from Vietnam indeed belong to four known species, namely *G. catbaensis*, *G. huuliensis*, *G. lichtenfelderi* and *G. luii* (Grismer *et al.* 1999; Vu *et al.* 2006; Orlov *et al.* 2008; Ziegler *et al.* 2008; Nguyen *et al.* 2009; Nguyen 2011). Despite extensive surveys over the last two decades, *G. araneus* has not been recorded in Vietnam, leaving its accurate type locality in Cao Bang Province, northern Vietnam ambiguous (Grismer *et al.* 1999; Ngo *et al.* 2016). It is possible that the species has been extirpated from Vietnam as a result of over-exploitation to supply the international pet trade or that the species has never occurred at the documented type locality (Grismer *et al.* 1999; Ngo *et al.* 2016). Similarly, *G. luii* has not been recorded again at its type locality in China (Grismer *et al.* 1999; Stuart *et al.* 2006; Yang & Chan 2015).

Molecular results supported by our study confirm that *G. murphyi*, described by Orlov & Darevsky (1999), is a junior synonym of *G. lichtenfelderi* (Grismer 2000). All mainland populations of *G. lichtenfelderi* found in granitic forests are conspecific with those from the granitic offshore islands in Bai Tu Long National Park, Quang Ninh Province, northern Vietnam (type locality), with an intraspecific genetic distance of less than 0.7% (Supp. file 1: Table S1). However, a few issues need to be resolved in the future. In particular, *G. hainanensis* was recovered as polyphyletic by Liang *et al.* (2018) and our phylogenetic analyses. Moreover, Zhu *et al.* (2020a) suggested that *G. luii* and *G. kadoorieorum* are polyphyletic. We highly recommend that they may be synonymized. Samples assigned to *G. kuroiwaie* were not recovered as monophyletic in our phylogenetic analysis. It is possible that more cryptic species from the Japanese group will be discovered. Further studies with additional samples, especially from type localities, should be undertaken to clarify these taxonomic problems.

Regarding morphological analyses, six morphometric characters, namely snout-vent length, head length, head width, mouth width, body height and body width strongly supported the overall difference among the four species in Vietnam and five characteristics, namely fourth toe, body and limb shapes, body bands, precloacal pores and granular scales surrounding dorsal tubercles, mainly accounted for the variation among the 17 tiger geckos from China and Vietnam. Six species of the *G. kuroiwaie* group from Japan were not included in this study. However, the Japanese *G. kuroiwaie* group can be distinguished from Chinese and Vietnamese species by the absence of precloacal pores and unsheathed claws (Wang *et al.* 2014; Yang & Chan 2015; Honda & Ota 2017). We highly recommend that all these traits be considered as diagnostic characters for species of *Goniurosaurus*.

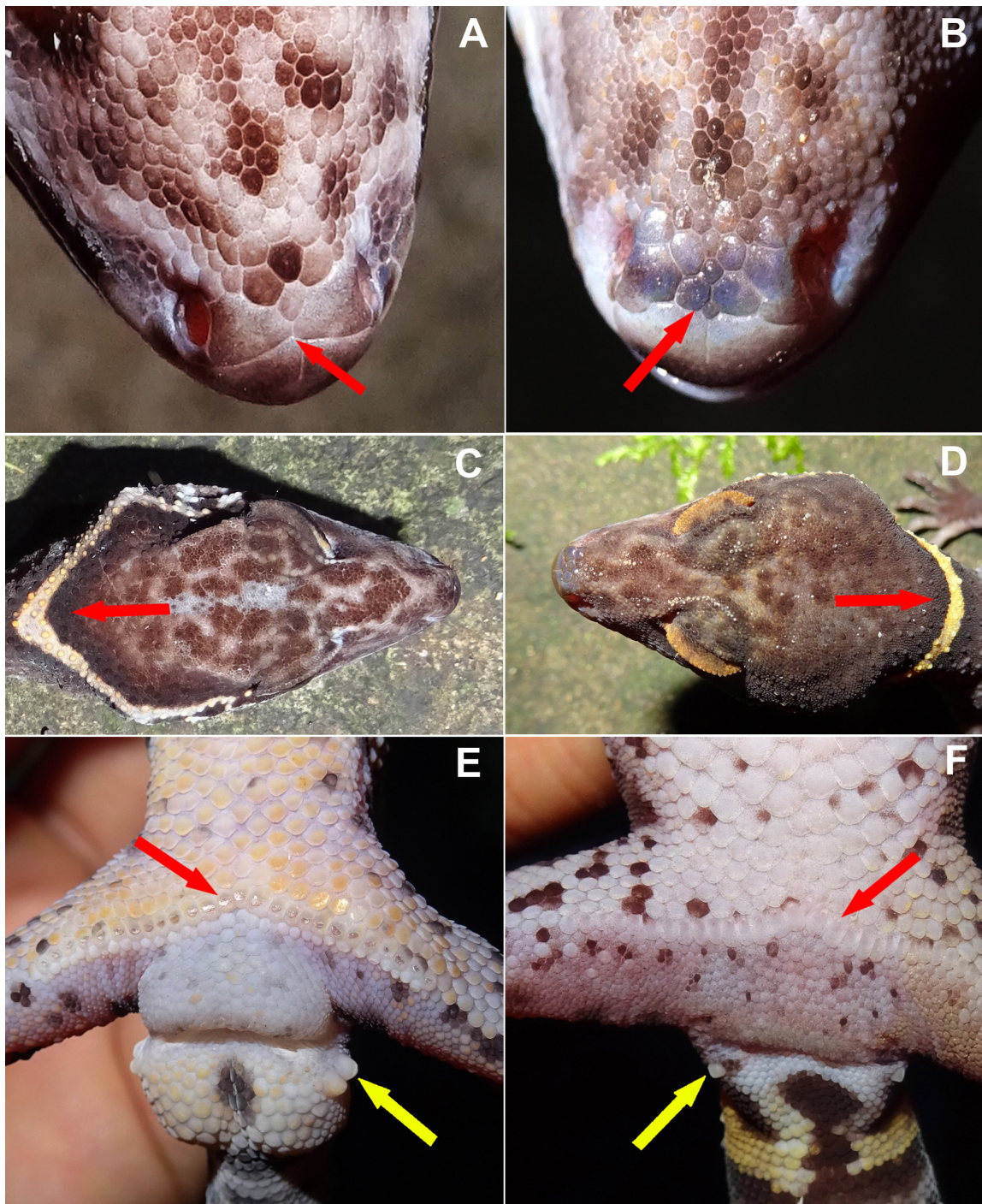


Fig. 7. A few key characters for species identification in the genus *Goniurosaurus* Grismer, Viets & Boyle, 1999. **A.** Snout tip of *G. catbaensis* Ziegler, Nguyen, Schmitz, Stenke & Rösler, 2008 with lacking postrostral (internasal) and two supranasals are in contact with each other. **B.** Snout tip of *G. lichtenfelderi* (Mocquard, 1897) with few internasals. **C.** Nuchal loop of *G. huuliensis* Orlov, Ryabov, Nguyen, Nguyen & Ho, 2008 protracted posteriorly (in V-shape). **D.** Nuchal loop of *G. lichtenfelderi* rounded posteriorly (in U-shape). **E.** Precloacal pores and large swollen hemipenial bulges of *G. luii* Grismer, Viets & Boyle, 1999 in males, and two postcloacal tubercles. **F.** Indistinct precloacal pores of *G. lichtenfelderi* in females and only one postcloacal tubercle on each site. Yellow arrows point to postcloacal tubercles, and red arrows to other scales.

Based on our identification key in accordance with the morphological and phylogenetic analyses, all tiger gecko species were grouped into four separate species groups, in agreement with the previous findings (Liang *et al.* 2018; Qi *et al.* 2020a, 2020b; Zhu *et al.* 2020a, 2020b). Liang *et al.* (2018), Qi *et al.* (2020a, 2020b) and our genetic analyses revealed that *G. bawanglingensis* and *G. zhoui* are embedded within the *G. lichtenfelderi* group, while they were previously placed as sister taxa in the *G. luii* group due to superficial morphological similarities as a result of adaptation to karst habitat (Grismer *et al.* 2002; Zhou *et al.* 2018). In accordance with these genetic findings, we found that the two karst-adapted species are morphologically most similar to three remaining granite-stream-adapted species of the *G. lichtenfelderi* group. Further comparative studies in southern China and Japan may help to resolve the phylogenetic relations of the genus *Goniurosaurus*.

Due to the high degrees of adaptation to specific microhabitats and local endemism, the genus *Goniurosaurus* could serve as a model system to study the evolution in lizards. In Vietnam, all species of *Goniurosaurus* have been found in isolated geographic ranges and none of them occurs in sympatry. For example, the distribution of *G. luii* stretches to the north of the Lang Son and Cao Bang provinces (Ngo *et al.* 2016), while *G. huuliensis* has been recorded in karst forests in the south of Lang Son Province. *Goniurosaurus catbaensis* is currently known only from offshore islands in Cat Ba National Park and the Ha Long Bay. *Goniurosaurus lichtenfelderi* can only be found in granitic habitats at several mainland localities and on some offshore islands in the Bai Tu Long Archipelago, which is contiguous with Ha Long Bay in the Gulf of Tonkin (Orlov *et al.* 2008; Gawor *et al.* 2016; Ngo *et al.* 2019a). *Goniurosaurus araneus* has not yet been re-discovered at its type locality in Cao Bang Province, and we only recorded *G. luii* inhabiting limestone karst forests of Cao Bang Province (Grismer *et al.* 1999; Vu *et al.* 2006; Ngo *et al.* 2016). The allopatric distribution has also been documented among species of *Goniurosaurus* from China. To date, neither the four species from the Hainan Archipelago, nor the five species of the *G. luii* group from Guangxi Province, China, nor the four known species of the *G. yingdeensis* group from Guangdong Province were recorded to occur in sympatry (Grismer *et al.* 2002; Zhou *et al.* 2018; Zhu *et al.* 2020a, 2020b). A large river (Zuojiang River) is regarded as the potential geographic barrier between the two species *G. luii* and *G. araneus* within Nonggang Nature Reserve, China (Chen *et al.* 2014). Likewise, geographic barriers (e.g., rivers and canyons) are considered to prevent a genetic exchange between members of the *G. yingdeensis* group (Qi *et al.* 2020a). The global rising of the sea level during the last melting period of glacial ice has shaped the myriad of archipelagoes (Clements *et al.* 2006; Ziegler *et al.* 2008; Liang *et al.* 2018; Ngo *et al.* 2019a). Thus, the oceanic barrier known as an important geographic feature has constrained current distributions and limits the genetic exchange of insular tiger gecko species, such as *G. catbaensis*, species from the Hainan Archipelago and *G. kuroiwaie* group (Ziegler *et al.* 2008; Honda & Ota 2017; Liang *et al.* 2018; Zhu *et al.* 2020b). However, the last glacial maximum period occurring around 10 000–50 000 years ago was not sufficiently long for speciation events within *Goniurosaurus* (Clements *et al.* 2006; Sterling *et al.* 2006; Ziegler *et al.* 2008). Indeed, by employing a molecular dating method, Liang *et al.* (2018) estimated that the latest speciation events among tiger geckos took place approximately 2.8–2.9 Ma, in the late Pliocene, between *G. yamashinae*–*G. orientalis*, and *G. huuliensis*–*G. luii*. It is likely that the global warming during the Pliocene promoted the diversification in the genus, as also reported in the crocodile newts (genus *Tylostotriton* Anderson, 1871) in Asia (Bernardes *et al.* 2020). Thus, the genus *Goniurosaurus* might offer a unique opportunity to investigate mechanisms of speciation and evolutionary adaptation, that are expected to highly correlate with a pre-dominantly allopatric mode of diversification.

Implications for conservation

The taxonomy of several existing species complexes requires further investigation and new species are still being discovered. Nevertheless, wild populations of the genus *Goniurosaurus* are under high risks of extinction, due to habitat destruction and exploitation for national and international trade. Of

the 23 described tiger gecko species, 12 have been considered threatened (listed as VU, EN and CR) and one data deficient (DD) in the IUCN Red List (Ngo *et al.* 2019b). Assessments for the remaining species are still lacking. In order to better regulate the international trade in the species, all species of *Goniurosaurus* from China and Vietnam were recently listed in CITES Appendix II and the Vietnamese species were also protected under the Governmental Decree 06/2019/ND-CP (Group IIB). Trade data in the US showed that about half of the number of traded tiger geckos were only assigned to the genus level (Ngo *et al.* 2019b). This fact could be a consequence of the highly similar morphology among tiger gecko species and lacking guidance to distinguish the species (Ngo *et al.* 2019b). Besides, the conservation status and national protection status are not assessed equally among species. Therefore, the present identification guide will aid to prevent tiger geckos from being traded under a wrong name to circumvent legislations. The detailed descriptions of morphology provided in this study can assist local, national, and international authorities, such as local rangers, CITES authorities, custom officers, in enforcing international regulations. The identification key can also be useful for scientists and breeders to accurately identify tiger gecko species. However, morphological identification of captive lineages should be treated with reservation, as some breeders have produced hybrids (L. Grismer, pers. comm.), and thus molecular identification tools should be consulted as well in these cases.

After the CITES listing, four new species, namely *G. gezhi*, *G. gollum*, *G. kwanghua* and *G. varius* were recently described from China, which are automatically listed in Appendix II of the Convention (Qi *et al.* 2020a, 2020b; Zhu *et al.* 2020a, 2020b). As the Japanese species appear to be similarly threatened and to prevent a shift towards international trade, Janssen & Shepherd (2019) and Ngo *et al.* (2019b) highly recommended that the Japanese species be also included in the CITES Appendices. During the time of writing, the Japanese Ministry for Environment announced to list all six endemic species of *Goniurosaurus* from Japan in CITES Appendix III to prevent over-harvesting of wild animals for trafficking activities. As a result, the listing of all Japanese species in Appendix III has come into force since 14 February 2021 (CITES Notification No. 2020/068).

Acknowledgments

For supporting fieldwork and issuing relevant permits, we thank the authorities of the Cat Ba National Park (CBNP), Hai Phong City, Huu Lien Nature Reserve, Lang Son Province, Bai Tu Long National Park, Quang Ninh Province, Tay Yen Tu Nature Reserve, Bac Giang Province, Yen Tu Nature Reserve and the Management Department of Ha Long Bay, Quang Ninh Province. We are thankful to K.X. Nguyen (CBNP), M.L. Pham (MDHLB), N.H. Nguyen, C.T. Pham, T.Q. Phan (IEBR, Ha Noi), L.T. Pham (Thai Nguyen University), D.T. Le, H. Tran (Ha Noi National University of Education), for assistance in the field. We are grateful to T. Pagel and C. Landsberg (Cologne Zoo), S.V. Nguyen, (IEBR, Hanoi), L.V. Vu and T.T. Nguyen (VNMN, Hanoi) for their support of conservation-based biodiversity research in Vietnam. This research was supported by the Vietnam Academy of Science and Technology (Project Code KHCBS.02/20-22) and the Ministry of Science and Technology's Program 562 (grant no. ĐTDL. CN-64/19). Field surveys were partially funded by Cologne Zoo, the Wildlife Conservation Society ('WCS') John Thorbjarnarson Fellowship for Reptile Research Grant, the Mohamed bin Zayed Species Conservation fund (Project: 170515492) to H.N. Ngo. Field equipment was supported by the Idea Wild. Cologne Zoo is partner of the World Association of Zoos and Aquariums (WAZA): Conservation Project 07011, 07012 (Herpetodiversity Research, Amphibian and Reptilian Breeding and Rescue Stations). Research of Hai Ngoc Ngo in Germany is funded by the German Academic Exchange Service (DAAD).

References

Bernardes M., Le M.D., Nguyen T.Q., Pham C.T., Pham A.V., Nguyen T.T., Rödder D., Bonkowski M. & Ziegler T. 2020. Integrative taxonomy reveals three new taxa within the *Tylototriton asperrimus* complex (Caudata, Salamandridae) from Vietnam. *ZooKeys* 935: 121–164.
<https://doi.org/10.3897/zookeys.935.37138>

- Chen T.B., Meng Y.J., Jiang K., Li P.P., Wen B.H., Lu W., Lazell J. & Hou M. 2014. New record of the leopard gecko *Goniurosaurus araneus* (Squamata: Eublepharidae) for China and habitat portioning between geographically and phylogenetically close leopard geckos. *IRCF Reptiles & Amphibians* 21 (1): 16–27.
- Clements R., Sodhi N.S., Schilthuizen M. & Ng P.K.L. 2006. Limestone karsts of Southeast Asia: imperiled arks of biodiversity. *BioScience* 56: 733–742.
[https://doi.org/10.1641/0006-3568\(2006\)56\[733:LKOSAI\]2.0.CO;2](https://doi.org/10.1641/0006-3568(2006)56[733:LKOSAI]2.0.CO;2)
- CITES. 2019. E CoP18 Proposal Number 27. Available from <https://cites.org/sites/default/files/eng/cop/18/prop/19032019/E-CoP18-Prop-27.pdf> [accessed 25 Apr. 2021].
- Gawor A., Pham T.C., Nguyen Q.T., Nguyen T.T., Schmitz A. & Ziegler T. 2016. The herpetofauna of the Bai Tu Long National Park, northeastern Vietnam. *Salamandra* 52 (1): 23–41.
- Gomes V., Carretero M.A. & Kaliontzopoulou A. 2016. The relevance of morphology for habitat use and locomotion in two species of wall lizards. *Acta Oecologica* 70: 87–95.
<https://doi.org/10.1016/j.actao.2015.12.005>
- Grismer L.L. 2000. *Goniurosaurus murphyi* Orlov and Darevsky: a junior synonym of *Goniurosaurus lichtenfelderi* Mocquard. *Journal of Herpetology* 34 (3): 486–488. <https://doi.org/10.2307/1565379>
- Grismer L.L., Viets B.E. & Boyle L.J. 1999. Two new continental species of *Goniurosaurus* (Squamata: Eublepharidae) with a phylogeny and evolutionary classification of the genus. *Journal of Herpetology* 33 (3): 382–393. <https://doi.org/10.2307/1565635>
- Grismer L.L., Shi H.T., Orlov N.L. & Ananjeva N.B. 2002. A new species of *Goniurosaurus* (Squamata: Eublepharidae) from Hainan Island, China. *Journal of Herpetology* 36: 217–224.
<https://doi.org/10.2307/1565994>
- Hillis D.M. & Bull J.J. 1993. An empirical test of bootstrapping as a method for assessing confidence in phylogenetic analysis. *Systematic Biology* 42: 182–192. <https://doi.org/10.1093/sysbio/42.2.182>
- Honda M. & Ota H. 2017. On the live coloration and partial mitochondrial DNA sequences in the topotypic population of *Goniurosaurus kuroiwaie orientalis* (Squamata: Eublepharidae), with description of a new subspecies from Tokashikijima Island, Ryukyu Archipelago, Japan. *Asian Herpetological Research* 8: 96–107. <https://doi.org/10.16373/j.cnki.ahr.170003>
- Honda M., Kurita T., Toda M. & Ota H. 2014. Phylogenetic relationships, genetic divergence, historical biogeography and conservation of an endangered gecko, *Goniurosaurus kuroiwaie* (Squamata: Eublepharidae), from the Central Ryukyus, Japan. *Zoological Science* 31: 309–320.
<https://doi.org/10.2108/zs130201>
- Janssen J. & Shepherd C.R. 2019. Trade in endangered and critically endangered Japanese herpetofauna endemic to the Nansei Islands warrants increased protection. *Current Herpetology* 38 (1): 99–109.
<https://doi.org/10.5358/hsj.38.99>
- Kassambara A. 2017. *Practical Guide to Principal Component Methods in R: PCA, M(CA), FAMD, MFA, HCPC, factoextra*. Multivariate Analysis II, STHDA.com.
- Le M., Raxworthy C.J., McCord W.P. & Mertz L. 2006. A molecular phylogeny of tortoises (Testudines: Testudinidae) based on mitochondrial and nuclear genes. *Molecular Phylogenetics and Evolution* 40: 517–531. <https://doi.org/10.1016/j.ympev.2006.03.003>
- Liang B., Zhou R.B., Liu Y.L., Chen B., Grismer L.L. & Wang N. 2018. Renewed classification within *Goniurosaurus* (Squamata: Eublepharidae) uncovers the dual roles of a continental island (Hainan) in species evolution. *Molecular Phylogenetics and Evolution* 127: 646–654.
<https://doi.org/10.1016/j.ympev.2018.06.011>

- Ngo H.N., Ziegler T., Nguyen T.Q., Pham C.T., Nguyen T.T., Le M.D. & van Schingen M. 2016. First population assessment of two cryptic tiger geckos (*Goniurosaurus*) from Northern Vietnam: Implications for conservation. *Amphibian & Reptile Conservation* 10 (1): 34–45.
- Ngo N.H., Le Q.T., Pham L.M., Le D.M., van Schingen M. & Ziegler T. 2019a. First record of the Cat Ba tiger gecko, *Goniurosaurus catbaensis*, from Ha Long Bay, Quang Ninh Province, Viet Nam: Microhabitat selection, potential distribution, and evidence of threats. *Amphibian & Reptile Conservation* 13 (2): 1–13.
- Ngo N.H., Nguyen Q.T., Phan Q.T., van Schingen M. & Ziegler T. 2019b. A case study on trade in threatened tiger geckos (*Goniurosaurus*) in Vietnam including updated information on the abundance of the endangered *G. catbaensis*. *Nature Conservation* 33: 1–19.
<https://doi.org/10.3897/natureconservation.32.33590>
- Nguyen L.T., Schmidt H.A., von Haeseler A. & Bui MQ. 2015. IQ-TREE: a fast and effective stochastic algorithm for estimating maximum-likelihood phylogenies. *Molecular Biology and Evolution* 32: 268–274. <https://doi.org/10.1093/molbev/msu300>
- Nguyen T.Q. 2011. *Systematics, Ecology, and Conservation of the Lizard Fauna in Northeastern Vietnam, with Special Focus on Pseudocalotes (Agamidae), Goniurosaurus (Eublepharidae), Sphenomorphus and Tropidophorus (Scincidae) from this Country*. PhD thesis, University of Bonn, Germany.
- Nguyen V.S., Ho T.C. & Nguyen Q.T. 2009. *Herpetofauna of Vietnam*. Edition Chimaira, Frankfurt am Main.
- Orlov N.L. & Darevsky I.S. 1999. Description of a new mainland species of *Goniurosaurus* genus, from the north-eastern Vietnam. *Russian Journal of Herpetology* 6: 72–78.
- Orlov N.L., Ryabow S.A., Nguyen T.T., Nguyen Q.T. & Ho T.C. 2008. A new species of *Goniurosaurus* (Sauria: Gekkota: Eublepharidae) from north Vietnam. *Russian Journal of Herpetology* 15: 229–244.
- Posada D. & Crandall K.A. 1998. MODELTEST: testing the model of DNA substitution. *Bioinformatics* 14: 817–818. <https://doi.org/10.1093/bioinformatics/14.9.817>
- Qi S., Grismer L.L., Lyu Z.T., Zhang L., Li P.P. & Wang Y.Y. 2020a. A definition of the *Goniurosaurus yingdeensis* group (Squamata, Eublepharidae) with the description of a new species. *ZooKeys* 986: 127–155. <https://doi.org/10.3897/zookeys.986.47989>
- Qi S., Wang J., Grismer L.L., Chen H.-H., Lyu Z.-T. & Wang Y.-Y. 2020b. The stoer hobbit of Guangdong: *Goniurosaurus gollum* sp. nov., a cave-dwelling leopard gecko (Squamata, Eublepharidae) from South China. *ZooKeys* 991: 137–153. <https://doi.org/10.3897/zookeys.991.54935>
- Ronquist F., Teslenko M., van der Mark P., Ayres D.L., Darling A., Höhna S., Larget B., Liu L., Suchard M.A. & Huelsenbeck J.P. 2012. MrBayes 3.2: efficient Bayesian phylogenetic inference and model choice across a large model space. *Systematic Biology* 61: 539–542.
<https://doi.org/10.1093/sysbio/sys029>
- RStudio Team. 2018. RStudio: integrated development for R. RStudio, Inc., Boston, MA, USA. Available from <http://www.rstudio.com/> [accessed 24 Apr. 2021].
- Sexton J.P., McIntyre P.J., Angert A.L. & Rice K.J. 2009. Evolution and ecology of species range limits. *Annual Review of Ecology, Evolution and Systematics* 40: 415–436.
<https://doi.org/10.1146/annurev.ecolsys.110308.120317>
- Stuart B.L., Rhodin A.G.J., Grismer L.L. & Hansell T. 2006. Scientific description can imperil species. *Science* 312 (5777): 1137. <https://doi.org/10.1126/science.312.5777.1137b>

- Sterling E.J., Hurley M.M. & Le D.M. 2006. *Vietnam: A Natural History*. Yale University Press, New Haven and London.
- Swofford D.L. 2001. PAUP* Phylogenetic Analysis Using Parsimony (* and Other Methods), version 4. Sinauer Associates, Sunderland, Massachusetts, USA.
- Thompson J.D., Gibson T.J., Plewniak F., Jeanmougin F. & Higgins D.G. 1997. The CLUSTAL_X windows interface: flexible strategies for multiple sequence alignment aided by quality analysis tools. *Nucleic Acids Research* 25: 4876–4882. <https://doi.org/10.1093/nar/25.24.4876>
- Vitt L.J., Caldwell J.P., Zani P.A. & Titus T.A. 1997. The role of habitat shifts in the evolution of lizard morphology: evidence from tropical *Tropidurus*. *Proceedings of the National Academy of Sciences of the United States of America* 94 (8): 3828–3832. <https://doi.org/10.1073/pnas.94.8.3828>
- Vu N.T., Nguyen Q.T., Grismer L.L. & Ziegler T. 2006. First record of the Chinese leopard gecko, *Goniurosaurus luii* (Reptilia: Eublepharidae) from Vietnam. *Current Herpetology* 25: 93–95. [https://doi.org/10.3105/1345-5834\(2006\)25\[93:FROTCL\]2.0.CO;2](https://doi.org/10.3105/1345-5834(2006)25[93:FROTCL]2.0.CO;2)
- Wang Y.Y., Yang J.H. & Cui R.F. 2010. A new species of *Goniurosaurus* (Squamata: Eublepharidae) from Yingde, Guangdong Province, China. *Herpetologica* 66: 229–240. <https://doi.org/10.1655/09-046r2.1>
- Wang Y.Y., Yang J.H. & Grismer L.L. 2013. A new species of *Goniurosaurus* (Squamata: Eublepharidae) from Libo, Guizhou Province, China. *Herpetologica* 69: 214–226. <https://doi.org/10.1655/herpetologica-d-12-00084>
- Wang Y.Y., Jin M.J., Li Y.L. & Grismer L.L. 2014. Description of a new species of *Goniurosaurus* (Squamata: Eublepharidae) from the Guangdong Province, China, based on molecular and morphological data. *Herpetologica* 70: 309–322. <https://doi.org/10.1655/herpetologica-d-13-00080>
- Yang J.H. & Chan B.P. 2015. Two new species of the genus *Goniurosaurus* (Squamata: Sauria: Eublepharidae) from southern China. *Zootaxa* 3980 (1): 67–80. <https://doi.org/10.11646/zootaxa.3980.1.4>
- Zhou R.B., Wang N., Chen B. & Liang B. 2018. Morphological evidence uncovers a new species of *Goniurosaurus* (Squamata: Eublepharidae) from the Hainan Island, China. *Zootaxa* 4369 (2): 281–291. <https://doi.org/10.11646/zootaxa.4369.2.8>
- Zhu X.Y., Chen G.Y., Román-Palacios C., Li Z. & He Z.Q. 2020a. *Goniurosaurus gezhi* sp. nov., a new gecko species from Guangxi, China (Squamata: Eublepharidae). *Zootaxa* 4852 (2): 211–222. <https://doi.org/10.11646/zootaxa.4852.2.6>
- Zhu X.Y., Shen C.Z., Liu Y.F., Chen L., Li Z. & He Z.Q. 2020b. A new species of *Goniurosaurus* from Hainan Island, China based on molecular and morphological data (Squamata: Sauria: Eublepharidae). *Zootaxa* 4772 (2): 349–360. <https://doi.org/10.11646/zootaxa.4772.2.6>
- Ziegler T., Nguyen T.Q., Schmitz A., Stenke R. & Rösler H. 2008. A new species of *Goniurosaurus* from Cat Ba Island, Hai Phong, northern Vietnam (Squamata: Eublepharidae). *Zootaxa* 1771: 16–30. <https://doi.org/10.11646/zootaxa.1771.1.2>

Manuscript received: 16 November 2020

Manuscript accepted: 3 March 2021

Published on: 31 May 2021

Topic editor: Rudy Jocqué

Desk editor: Pepe Fernández

Printed versions of all papers are also deposited in the libraries of the institutes that are members of the *EJT* consortium: Muséum national d'histoire naturelle, Paris, France; Meise Botanic Garden, Belgium; Royal Museum for Central Africa, Tervuren, Belgium; Royal Belgian Institute of Natural Sciences, Brussels, Belgium; Natural History Museum of Denmark, Copenhagen, Denmark; Naturalis Biodiversity Center, Leiden, the Netherlands; Museo Nacional de Ciencias Naturales-CSIC, Madrid, Spain; Real Jardín Botánico de Madrid CSIC, Spain; Zoological Research Museum Alexander Koenig, Bonn, Germany; National Museum, Prague, Czech Republic.

Supplementary files

Supp. file 1. Supplementary tables. <https://doi.org/10.5852/ejt.2021.751.1379.4325>

Table S1. Pair-wise genetic divergence between species included in this study.

Table S2. Morphological (minimum–maximum (mean \pm standard deviation)) and meristic characters (minimum–maximum (number of specimens)) of four tiger geckos in Vietnam (except *Goniurosaurus araneus*). Length given in mm.

Supp. file 2. Supplementary figures. <https://doi.org/10.5852/ejt.2021.751.1379.4327>

Fig. S1. Snout-vent length (SVL) of four tiger geckos in Vietnam including *Goniurosaurus catbaensis*, *Goniurosaurus huuliensis*, *Goniurosaurus lichtenfelderi*, *Goniurosaurus luii*. A. Juveniles. B. Adult males. C. Adult females.

Fig. S2. A. Principal component analysis (PCA) on morphological variation amongst four recorded tiger geckos in Vietnam. B. Scatterplots of principal component scores for the first and second principal axes.

ZOBODAT - www.zobodat.at

Zoologisch-Botanische Datenbank/Zoological-Botanical Database

Digitale Literatur/Digital Literature

Zeitschrift/Journal: [European Journal of Taxonomy](#)

Jahr/Year: 2021

Band/Volume: [0751](#)

Autor(en)/Author(s): Ngo Hai Ngoc, Nguyen Huy Quoc, Tran Hieu Minh, Le Minh Duc, Gewiss Laurenz Rafel, Schingen-Khan Mona von, Nguyen Truong Quang, Ziegler Thomas

Artikel/Article: [A morphological and molecular review of the genus Goniurosaurus, including an identification key 38-67](#)