

Two new species of the *Liolaemus elongatus-kriegi* complex (Iguania, Liolaemidae) from Andean highlands of southern Chile

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Abstract

The *elongatus-kriegi* complex is one of the most diverse clades of the *Liolaemus (sensu stricto)* subgenus of lizards. There are currently 29 species recognized in this group distributed between Chile and Argentina. Based on molecular evidence, there seem to be five main clades nested within this complex: the *elongatus*, *leopardinus*, *kriegi*, *petrophilus* and *punmahuida* clades. *Liolaemus buergeri* and *L. kriegi*, both of the *kriegi* clade, were believed to inhabit the surroundings of the Laja Lagoon, in the Biobío Region of Chile. Moreover, this Chilean population of *L. kriegi* was recently recognized as an undescribed taxon called “*Liolaemus* sp. A” based on molecular phylogenetics. In this work, we studied these two populations of the Laja Lagoon and provided the morphological diagnosis to describe them as two new species: *L. scorialis* sp. n. and *L. zabalai* sp. n., previously considered *L. buergeri* and “*L. kriegi* *Liolaemus* sp. A” respectively. Additionally, we identified another population of *L. scorialis* in the vicinity of La Mula Lagoon in the Araucanía Region of Chile. *Liolaemus scorialis* differs from almost all of the species of the *elongatus-kriegi* complex by its considerably smaller size. Nevertheless, without molecular data we cannot assign it to any particular subclade. *Liolaemus zabalai* belongs to the *kriegi* clade based on published molecular phylogenies. Finally, we provide some natural history data on both species and we document for the first time the presence of *L. neuquensis* in Chile from a museum specimen from La Mula Lagoon.

Keywords

Liolaemus buergeri, *Liolaemus kriegi*, new species, lizard, Laja Lagoon, Biobío

Introduction

Liolaemus is a diverse genus of South American lizards, with currently 245 species (Uetz and Hošek 2014) grouped into two subgenera: *Liolaemus (sensu stricto)* and *Eulaemus* (e.g. Laurent 1985, Schulte et al. 2000). Each of these subgenera has been divided into several groups based on phylogenetic relationships (Abdala 2007, Avila et al. 2006, Fontanella et al. 2012, Lobo 2005).

The *elongatus-kriegi* complex Cei (1979), is one of the most diverse groups of the *Liolaemus (sensu stricto)* subgenus with currently 29 species distributed in Chile and Argentina. In a phylogenetic study based on three mitochondrial genes, Morando et al. (2003) found that this complex is subdivided into three clades: *elongatus*, *kriegi* and *petrophilus*. Later, Avila et al. (2010a) based on one mitochondrial locus, added a fourth clade: the *punmahuida* clade. Finally, Esquerré et al. (2014) added a fifth clade comprising only Chilean endemic species: the *leopardinus* clade. An alternative classification has been proposed by Lobo (2005) and updated by Lobo et al. (2010b), based mainly on morphological and lifestyle traits, which classifies these species in three groups: *elongatus* (which includes the *capillitas* subgroup), *kriegi* and *leopardinus*, with a different arrangement compared with the molecular hypothesis (Table 1 and Table 2).

Currently, the *elongatus-kriegi* complex (Avila et al. 2012, Esquerré et al. 2014, Morando et al. 2003) or *elongatus*, *kriegi* and *leopardinus* groups (Lobo 2005, Lobo et al. 2010b) includes the following species: *Liolaemus antumalguen* Avila et al., 2010, *L. austromendocinus* Cei, 1974, *L. buergeri* Werner 1907, *L. burmeisteri* Avila et al. 2012, *L. carlosgarini* Esquerré et al. 2013, *L. capillitas* Hulse, 1979, *L. choique* Abdala et al. 2010, *L. cristiani* Núñez et al. 1991, *L. dicktracyi* Espinoza & Lobo, 2003, *L. elongatus* Koslowsky, 1896, *L. flavipiceus* Cei & Videla, 2003, *L. frassinettii* Núñez, 2007, *L. gununakuna* Avila et al. 2004, *L. heliodermis* Espinoza et al. 2000, *L. kriegi* Müller & Hellmich, 1939, *L. leopardinus* Müller & Hellmich, 1932, *L. parvus* Quinteros et al. 2008, *L. petrophilus* Donoso-Barros & Cei, 1971, *L. punmahuida* Avila et al. 2003, *L. ramonensis* Müller & Hellmich, 1932, *L. shitan* Abdala et al. 2010, *L. smaug* Abdala et al. 2010, *L. talampaya* Avila et al. 2004, *L. thermarum* Videla & Cei, 1996, *L. tregenzai* Pincheira-Donoso & Scolaro, 2007, *L. tulkas* Quinteros et al. 2008, *L. ubaghsi* Esquerré et al. 2014, *L. umbrifer* Espinoza & Lobo, 2003 and *L. valdesianus* Hellmich, 1950.

Liolaemus buergeri, of the *kriegi* clade, was described from El Planchón Volcano, Maule Region, Chile (Werner 1907). This species has been traditionally believed to be widely distributed in Chile and Argentina (Cei 1986, Pincheira-Donoso 2001). However, its current wide distribution is in part due to cases of misidentification and a lumping of cryptic species (Medina et al. 2013). Donoso-Barros (1970) extended the southern distribution of *L. buergeri* to the Andes of Talca, Maule Region, Chile (50 km S from El Planchón Volcano).

Table 1. Species of the *elongatus-kriegi* complex grouped by clades, based on mitochondrial molecular phylogenies. (1) Species included by Morando et al. (2003). (2) Species added by Avila et al. (2004). (3) Species added by Avila et al. (2010a). (4) Species added by Avila et al. (2012). (5) Species added fide Esquerré et al. (2014). *Liolaemus thermanum* is included in the *elongatus* clade by Avila et al. (2010a) but omitted by Avila et al. (2012).

<i>elongatus</i> clade	<i>kriegi</i> clade	<i>leopardinus</i> clade	<i>petrophilus</i> clade	<i>punmahuida</i> clade
<i>L. antumalguen</i> (3)	<i>L. buergeri</i> (1)	<i>L. frassinettii</i> (5)	<i>L. austromendocinus</i> (1)	<i>L. flavipiceus</i> (3)
<i>L. burmeisteri</i> (4)	<i>L. kriegi</i> (1)	<i>L. leopardinus</i> (5)	<i>L. capillitas</i> (1)	<i>L. punmahuida</i> (3)
<i>L. elongatus</i> (1)		<i>L. ramonensis</i> (5)	<i>L. dicktracyi</i> (2)	
<i>L. smaug</i> (4)		<i>L. ubaghsi</i> (5)	<i>L. gununakuna</i> (2)	
<i>L. thermanum</i> (3)		<i>L. valdesianus</i> (5)	<i>L. parvus</i> (3)	
			<i>L. petrophilus</i> (1)	
			<i>L. talampaya</i> (2)	
			<i>L. tulkas</i> (3)	
			<i>L. umbriifer</i> (2)	

Table 2. Species of the *elongatus-kriegi* complex by groups, based on morphological, skeletal and lifestyle traits phylogeny according to (1) Lobo (2005), (2) updated by Lobo et al. (2010b) and (3) fide Esquerré et al. (2013). The *capillitas* subgroup is nested into *elongatus* group (Lobo et al. 2010b).

<i>capillitas</i> subgroup	<i>elongatus</i> group	<i>kriegi</i> group	<i>leopardinus</i> group
<i>L. capillitas</i> (1)	<i>L. austromendocinus</i> (2)	<i>L. buergeri</i> (1)	<i>L. frassinettii</i> (2)
<i>L. dicktracyi</i> (1)	<i>L. carlosgarini</i> (3)	<i>L. cristiani</i> (1)	<i>L. leopardinus</i> (1)
<i>L. heliodermis</i> (1)	<i>L. elongatus</i> (1)	<i>L. kriegi</i> (1)	<i>L. ramonensis</i> (1)
<i>L. talampaya</i> (2)	<i>L. flavipiceus</i> (2)		<i>L. valdesianus</i> (1)
<i>L. tulkas</i> (2)	<i>L. gununakuna</i> (2)		
<i>L. umbriifer</i> (1)	<i>L. parvus</i> (2)		
	<i>L. petrophilus</i> (2)		
	<i>L. punmahuida</i> (2)		
	<i>L. thermanum</i> (2)		
	<i>L. tregenzai</i> (2)		

Later, Pincheira-Donoso (2001) extended the Chilean southern distribution of *L. buergeri* to the Batea-Mahuida Volcano (Araucanía Region, 240 km S from El Planchón Volcano) and pointed out that he also examined three specimens from the Laja Lagoon (Biobío Region, Chile, 150 km S from El Planchón Volcano); but Pincheira-Donoso and Núñez (2005) indicated that the specimens from Batea-Mahuida Volcano indeed correspond to *L. elongatus*, whereas the status of “*L. buergeri*” from the Laja Lagoon in Chile remains uncertain. In regards to Argentina, Cei (1986) stated that this species occurs in Mendoza and Neuquén Provinces, but Morando et al. (2003) and Medina et al. (2013), based on genetic and morphological evidence, respectively, indicated that several Argentinean populations attributed to *L. buergeri* correspond to at least three undescribed species.

Liolaemus kriegi, also of the *kriegi* clade, was described from Estancia El Cóndor, Río Negro Province, Argentina (Müller and Hellmich 1939a). Later, Donoso-

Barros (1966) extended its northern distribution to the Cordillera de Curicó, Maule Region, Chile, 650 km N of Estancia El Cóndor; and to the Laja Lagoon, Biobío Region, Chile, 400 km N of Estancia El Cóndor (Donoso-Barros 1974). Morando et al. (2003), based on mitochondrial genes, found three candidate species related to *L. kriegi*, all from Argentina and previously assigned to *L. buergeri*: *Liolaemus* sp. A (from Caviahue, Neuquén Province), *Liolaemus* sp. B (from Ranquil Norte, Neuquén Province) and *Liolaemus* sp. C (from Laguna Los Barros, Neuquén Province). Medina et al. (2013), in a morphological analysis of these populations, corroborated the status of candidate species of these *Liolaemus* sp., adding new localities for *Liolaemus* sp. A, including samples from the Laja Lagoon (Chile) which corresponds to the species previously identified as *L. kriegi* by Donoso-Barros (1974). Also, Medina et al. (2013) found another candidate species from Argentina (*Liolaemus* sp. D), previously identified as *L. buergeri* by Morando et al. (2003). Recently, Medina et al. (2014) in a new phylogenetic study based on mitochondrial and nuclear genes, corroborate the previous studies and provide strong evidence for *Liolaemus* sp. A as a candidate species, also based on samples from Chile (Laja Lagoon) and Argentina (several localities of Neuquén Province).

Here, we studied the taxonomic status of the southernmost currently-recognized Chilean population of “*Liolaemus buergeri*”, from the vicinity of the Laja Lagoon, Biobío Region; and of “*L. kriegi*/*Liolaemus* sp. A” from the same locality. This population of “*L. buergeri*” is described as a new species which differs greatly from *L. buergeri* and almost all species of the *elongatus-kriegi* complex by its small snout-vent length (less than 70.0 mm). Additionally, specimens of this new species are recorded from La Mula Lagoon, Araucanía Region, Chile. For “*L. kriegi*/*Liolaemus* sp. A”, we provide a full description and diagnosis of this new species belonging to the *kriegi* clade.

Materials and methods

We examined specimens of almost all Chilean species currently considered as belonging to the *Liolaemus elongatus-kriegi* complex. The morphological characters were examined according to Etheridge (1995), Lobo (2005), Abdala et al. (2010) and Avila et al. (2010a, 2012). Body measurements were taken with a digital vernier caliper (0.02 mm precision). Measurements are provided as mean \pm standard deviation ($\bar{x} \pm SD$). The Mann–Whitney U test was used to compare the new species and some related species. Scales were observed with different magnifying lenses and scalation and measurements were recorded on the right side of the specimen, unless otherwise indicated. Dorsal scales were counted between the occiput and the level of the anterior border of the hind limbs. Ventral scales were counted from mental scale to the anterior margin of cloacal opening. Stomach and intestinal contents were observed under a binocular microscope for one specimen of each new species. The specimens examined are listed in Appendix 1. Data for Argentinean species were taken from the literature. *Liolaemus ceii* is not accepted as valid species in this work (see discussion). Museum codes are as

follow: **MRC** (Museo Regional de Historia Natural, Concepción), **MZUC** (Museo de Zoología, Universidad de Concepción) and **SSUC** (Colección de Flora y Fauna Patricio Sánchez Reyes, Pontificia Universidad Católica de Chile).

Results

Liolaemus scorialis sp. n.

<http://zoobank.org/35B1E4BC-4EA1-4FEF-B025-B93D5C5A9CB9>

Fig. 1

Liolaemus buergeri (in part?), Pincheira-Donoso, 2001. Not. Mens. Mus. Nac. Hist. Nat., Chile, 346: 8.

Liolaemus buergeri (in part?), Pincheira-Donoso & Núñez, 2005. Pub. Oc. Mus. Nac. Hist. Nat., Chile, 59: 285.

Holotype. SSUC Re 617 (Fig. 1). Male collected 7 km NW of the summit of the Antuco Volcano, near the Laja Lagoon, Biobío Region, Chile (37°21'S – 71°23'W, 1450 m). Collected by J. Troncoso-Palacios, F. Urra and H. Díaz. 08/01/2014.

Paratypes. SSUC Re 615–16 two males and 612–614 three females (Figs 1 and 3). The same data as the holotype. MRC 675, 677, 680, 682. Four males. La Mula Lagoon (37°53'S – 71°22'W), Ralco National Reserve. Unknown coll. 01/12/2001.

Etymology. The species name refers to the habitat, which is composed of accumulations of igneous rocks from the Antuco Volcano, called “scoria” from the Greek “skoria”. We propose the common name “Slag Lizard” in English and “Lagarto del escorial” in Spanish.

Diagnosis. *Liolaemus scorialis* belongs to the *elongatus-kriegi* complex, but its specific assignation to a particular subclade is currently unknown since we have no molecular data for this new species, and molecular and morphological phylogenies for the *elongatus-kriegi* complex disagree in the arrangement of this complex subgroups (see discussion).

Below a wide diagnosis is provided on aspect of all species of the complex. *Liolaemus scorialis* differs from almost all species of the *elongatus-kriegi* complex by its size (maximum SVL = 69.9 mm), smaller than *L. antumalguen* (Table 3), *L. austromendocinus* (max. SVL = 103.0 mm, Espinoza et al. 2000), *L. buergeri* (Table 3, Fig. 2), *L. burmeisteri* (Table 3), *L. capillitas* (max. SVL = 93.0 mm, Espinoza et al. 2000), *L. choique* (Table 3), *L. dicktracyi* (max. SVL = 91.0 mm, Espinoza and Lobo 2003), *L. elongatus* (max. SVL = 94.7 mm, Avila et al. 2012), *L. flavipiceus* (Table 3, Fig. 2), *L. frassinettii* (max. SVL = 91.1 mm), *L. gununakuna* (max. SVL = 97.5 mm, Avila et al. 2004), *L. kriegi* (max. SVL = 101.0 mm; Avila et al. 2003), *L. leopardinus* (max. SVL = 98.2 mm), *L. petrophilus* (max. SVL = 100.0 mm; Espinoza et al. 2000), *L. punmahuida* (Table 3), *L. ramonensis* (max. SVL = 94.9 mm), *L. shitan* (max. SVL = 98.3 mm, Abdala et al. 2010), *L. talampaya* (max. SVL = 85.5 mm, Avila et al. 2004), *L. thermarum* (max. SVL = 85.0 mm, Videla and Cei 1996), *L. tregenzai* (Table 3),



Figure 1. *Liolaemus scorialis* sp. n. **A, B** Holotype, male **C, D** Paratype, female **E** Paratype, male **F** Paratype, female. All from the type locality, 7 km NW of the summit of the Antuco Volcano, near the Laja Lagoon, Biobío Region, Chile.

L. ubaghsi (max. SVL = 89.6 mm), *L. umbrifer* (max. SVL = 89.0 mm, Espinoza and Lobo 2003), *L. valdesianus* (max. SVL = 93.4 mm) and “*L. kriegi*/*Liolaemus* sp. A” (max. SVL = 92.0 mm, described below).

Liolaemus scorialis has probably been previously confused with *L. buergeri* (see discussion), but in addition to the size difference, *L. scorialis* differs from *L. buergeri* because the latter has a vertebral stripe on the tail, whereas the tail is ringed in *L. scorialis*. Moreover, *L. buergeri* has more midbody scales ($x = 89.4 \pm 5.5$, $n = 14$) than *L. scorialis* ($x = 82.0 \pm 4.7$, $n = 10$) (Mann–Whitney $U = 20.5$, $P < 0.01$, $DF = 21$) and more dorsal scales ($x = 84.1 \pm 4.4$) than *L. scorialis* ($x = 76.5 \pm 4.3$) (Mann–Whitney $U = 15.0$, $P < 0.01$, $DF = 21$); but *L. buergeri* has fewer ventral scales ($x = 118.7 \pm 4.7$) than *L. scorialis* ($x = 124.0 \pm 6.0$) (Mann–Whitney $U = 36.0$, $P = 0.05$, $DF = 21$).

Liolaemus scorialis is syntopic with “*L. kriegi*/*Liolaemus* sp. A”, but in addition to the size difference, the latter has more midbody scales ($x = 94.3 \pm 4.8$, $n = 8$) than it (Mann–Whitney $U = 1.5$, $P < 0.01$, $DF = 16$). Moreover, the dorsal scale count range of *L. scorialis* does not overlap with the range of “*L. kriegi*/*Liolaemus* sp. A” (Table 3).

Table 3. Scelation and morphological characteristics for the species of the *Liolaemus elongatus-kriegi* complex occurring near *L. scoriialis* sp. n. and *L. zabalalai* sp. n. distribution. Juvenile specimens examined are excluded. Source of data for not examined species are: *L. antumalguen* (Avila et al. 2010a), *L. burmeisteri* (Avila et al. 2012), *L. choique* (Abdala et al. 2010), *L. punnabuidai* (Avila et al. 2004) and *L. tregenzai* (Pincheira-Donoso and Sclaro 2007). (*) Medina et al. (2013). M = males; F = females.

	<i>L. antumalguen</i>	<i>L. buergeri</i> (M = 5, F = 9)	<i>L. burmeisteri</i>	<i>L. carlosgarini</i> (M = 6, F = 11)	<i>L. choique</i>	<i>L. flavipiceus</i> (M = 5, F = 10)	<i>L. punnabuidai</i>	<i>L. scoriialis</i> sp. n. (M = 7, F = 3)	<i>L. tregenzai</i>	<i>L. zabalalai</i> sp. n. (M = 3, F = 5)
Maximum SVL (mm)	107.8	96.2	85.2	68.8	90.7	95.8	96.0	69.9	90.2	92.0
Midbody scales	72–82	80–100	70–81	80–95	74–88	68–77	67–81	76–90	71–85	90–104
Dorsal scales	70–78	78–91	76–85	68–82	65–81	60–71	70–78	74–81	-	86–96
Ventral scales	105–118	111–125	99–110	112–124	118–135	93–105	-	115–131	-	116–122
Sexual dichromatism	Absent	Absent	Absent	Absent	Absent	Absent	Absent	Slight	Present	Slight
Cloacal region color (males)	Yellowish in some specimens but usually black	Yellowish	Yellowish	Yellowish	Yellowish	Reddish or yellowish in some specimens but usually black	Reddish or yellowish	Yellowish	-	Yellowish
Tail pattern	Absent	Vertebral line with diffuse rings in the tail base	Weak rings	Rings (marked or weak)	Absent	Absent or weak rings	Absent	Rings	Rings	Rings
Preloacal pores on males	3–4	3–4	0–5	0–3	3–4	0	0	3–4	0	3–4 (3–5*)

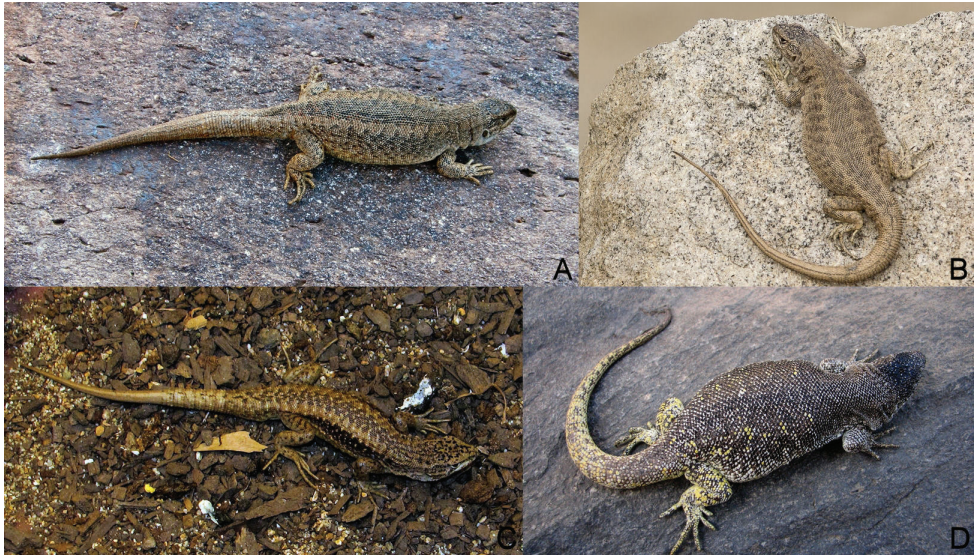


Figure 2. Chilean species of the *elongatus-kriegi* complex that live near the distribution of *Liolaemus scorialis* sp. n. and *L. zabalai* sp. n. **A** *L. buergeri* from El Planchón (type locality, photo by J. Troncoso-Palacios) **B** *L. buergeri* from Altos de Lircay (photo by R. Díaz) **C** *L. carlosgarini* from the road to the Maule Lagoon (type locality, photo by J. Troncoso-Palacios) **D** *L. flavipiceus* from the Maule Lagoon (photo by C. Garín).

There is a black lateral band running from the tip of snout to the groin in “*L. kriegel* *Liolaemus* sp. A”, whereas in *L. scorialis* there is a dark brown lateral band running from the shoulder to the groin.

Liolaemus scorialis differs from similar size species of the *elongatus-kriegi* complex as follows. *Liolaemus scorialis* differs from *L. cristiani* because the males of the latter lack preloacal pores and have reddish ventral coloration, whereas males of *L. scorialis* have 3–4 preloacal pores and no reddish ventral coloration.

Liolaemus scorialis differs from *L. heliodermis*, because the males of the latter have a black head and sulfur-yellow dorsum (Espinoza et al. 2000), an unique feature in the *Liolaemus* subgenus. Moreover, *L. heliodermis* has 62–69 midbody scales (Espinoza et al. 2000), whereas *L. scorialis* has 76–90.

Liolaemus scorialis differs from *L. parvus*, because the latter has 60–77 midbody scales and 96–113 ventral scales (Quinteros et al. 2008), whereas *L. scorialis* has 76–90 midbody scales and 115–131 ventral scales. *Liolaemus scorialis* has a ringed tail, whereas *L. parvus* has weak or absent rings on the tail (Quinteros et al. 2008).

Liolaemus scorialis differs from *L. smaug*, because the latter has marked sexual dichromatism with white spots dispersed on the dorsum of males and absent in females (Abdala et al. 2010), whereas both males and females of *L. scorialis* have white spots on the dorsum. *Liolaemus scorialis* has ringed tail, whereas *L. smaug* has weak or no rings on the tail (Abdala et al. 2010). Males of *L. smaug* have bright golden yellow dorsal color, a trait absent in *L. scorialis*.



Figure 3. Comparison of the ventral color pattern. **A** *Liolaemus scorialis* sp. n. from type locality, with immaculate gray ventral color **B** *L. carlosgarini* with light gray ventral color and dark inconspicuous spots dispersed.

Liolaemus scorialis differs from *L. tulkas*, because the males of the latter have 0–1 preloacal pores (Quinteros et al. 2008), whereas males of *L. scorialis* have 3–4 preloacal pores. Moreover, *L. tulkas* has 63–68 midbody scales (Quinteros et al. 2008), whereas *L. scorialis* has 76–90.

Liolaemus scorialis differs from *L. carlosgarini* (Fig. 2), because the males of the latter have 0–3 preloacal pores (present in 50% of the males, these are small and underdeveloped), whereas males of *L. scorialis* have 3–4 well developed preloacal pores. *Liolaemus scorialis* has more ventral scales ($x = 124 \pm 6.0$, $n = 10$) than *L. carlosgarini* ($x = 115 \pm 4.0$, $n = 17$) (Mann–Whitney $U = 11.0$, $P = 0.01$, $DF = 25$). Moreover, *L. scorialis* has brown dorsal color and immaculate gray ventral color, whereas *L. carlosgarini* has light brown-yellowish dorsal color and whitish ventral color with dark inconspicuous spots on the gular region and belly (Figs 2 and 3).

Description of the holotype. Adult male. SVL 62.3 mm. Tail length 101.5 mm (not autotomized). Axilla-groin length 26.3 mm. Head length (from the posterior border of the auditory meatus to the tip of the snout) 16.4 mm. Head width (distance between the two ear openings) 11.4 mm. Head height (at the level of ear openings) 6.9 mm. Forelimb length 21.1 mm. Hindlimb length 39.7 mm. Foot length 18.9 mm. Rostral scale wider

(2.5 mm) than high (1.0 mm). Two postrostrals. Four internasals. Hexagonal interparietal scale, with a central, small, and whitish spot marking the position of the parietal eye. Interparietal smaller than parietals, surrounded by six scales; nine scales between the interparietal and rostral (both excluded); 15 scales between occiput and rostral; orbital semicircle complete on the right side, formed by 13 scales, incomplete on the left side; 6–5 supraoculars (left-right); six superciliary scales. Frontal area is divided into six scales (two posterior, one in the center and three anterior); 2 scales between nasal and canthal; preocular separated from the lorilabials by one loreal scale; nasal in contact with the rostral, surrounded by seven scales. There is one row of lorilabials between the supralabials and the subocular. Seven supralabials, the fifth is curved upward without contacting the subocular. Four infralabial scales. Mental scale pentagonal, in contact with four scales; four pairs of postmental shields, the second is separated by two scales. Temporal scales are subimbricated and slightly keeled. There are ten temporal scales between the level of superciliary scales and the rictal level. Three projected scales on the anterior edge of the ear, which are small and do not cover the auditory meatus; auricular scale is wide and is restricted to the upper third of the meatus. Forty gulars between the auditory meatuses. Well developed “Y” shaped lateral neck fold and dorsolateral fold slightly developed. Antehumeral fold present. Midbody scales 88. Dorsal scales of the vertebral zone lanceolate, imbricate, keeled and without mucrons. Dorsal scales of the paravertebral fields more rounded, subimbricate, with more poorly developed keel, without mucrons and with interstitial granules between them. Dorsal scales of the vertebral zone are larger than the ventral scales. Dorsal scales of the paravertebral fields are similar in size to the ventral scales. Dorsal scales 81. Ventral scales are rhomboidal to rounded, smooth, imbricate, and without interstitial granules. Ventral scales 131. There are four preloacal pores. The suprafemoral scales are rhomboidal to rounded, imbricate, and smooth or slightly keeled. Infracemoral scales are rounded, smooth, and imbricate. Supra-antebrachials scales are rhomboidal to rounded, imbricate, and slightly keeled or smooth. Infra-antebrachials are rounded to rhomboidal, subimbricate with few interstitial granules, and smooth. The dorsal scales of the tail are rhomboidal, imbricate, keeled and some with mucrons. The ventral scales of the tail vary from rhomboidal to triangular, and are imbricate and smooth. Lamellae of the fingers: I: 10, II: 17, III: 21, IV: 23 and V: 13. Lamellae of the toes: I: 13, II: 18, III: 22, VI: 29 and V: 20.

Color of the holotype in life. Light brown head, with dark brown lines: a “Ω” shaped line between nasal scales and supraocular area, two short stripes on the posterior supraocular areas, an incomplete “O” shaped dark brown line surrounding the interparietal scale, six dark brown short lines on the occipital area. The temporal area is brown with two dark brown horizontal stripes; the ocular area and the cheeks are light gray. Subocular area is gray with two dark brown vertical lines on the middle and posterior edge. Background color of the dorsum is brown. A wide occipital band on the dorsum, formed by twelve transverse dark brown bars; some white scales on the posterior border of these bars. Dark brown lateral band with few yellowish scales dispersed into it, running from the shoulder to the groin; some white scales between the occipital and lateral bands; below the lateral band the flanks are yellowish. Limbs

are brown with dark brown spots and some white scales dispersed. Tail is brown with some white scales dispersed and dark brown rings. Posterior third of the tail is immaculate brown. Ventrally, the throat, belly, limbs and tail are immaculate gray. Rear portion of belly and thighs are yellowish. Preloacal pores orange.

Variation. There is no sexual dimorphism in size. In seven males: SVL: 57.4–69.9 mm. Axilla-groin distance: 21.4–28.7 mm. Head length: 15.1–17.2 mm. Head width: 11.2–13.0 mm. Head height: 6.4–8.9 mm. Foot length: 19.7–21.1 mm. Leg length: 37.1–46.2 mm. Arm length: 20.3–26.0 mm. Tail length: 101.6–111.3 mm ($n = 2$; autotomized in the rest). In three females: SVL: 57.3–65.6 mm. Axilla-groin distance: 25.6–32.8 mm. Head length: 15.3–15.8 mm. Head width: 11.1–12.1 mm. Head height: 6.2–6.7 mm. Foot length: 18.7–20.0 mm. Leg length: 37.2–39.0 mm. Arm length: 21.8–22.3 mm. Tail length 88.8–103.1 mm ($n = 2$; autotomized in the rest).

The variation of the scalation in *Liolaemus scorialis* is as follows. Midbody scales: 76–90 ($x = 82.0 \pm 4.7$). Dorsal scales: 74–81 ($x = 76.5 \pm 4.3$). Ventral scales 115–131 ($x = 124.0 \pm 6.0$). Fourth finger lamellae: 21–24 ($x = 22.7 \pm 1.1$). Fourth toe lamellae: 28–31 ($x = 29.2 \pm 1.4$). Supralabial scales: 6–7 ($x = 6.2 \pm 0.4$). Infralabial scales: 4–5 ($x = 4.7 \pm 0.5$). Preloacal pores in males: 3–4. Interparietal scale pentagonal or hexagonal, bordered by 5–9 scales ($x = 6.7 \pm 1.2$).

There is a slight sexual dichromatism, females have no yellowish color on the rear portion of belly and thighs. Males have the same color and pattern described for the holotype with variations only in shade. Females have the same color and pattern described for the holotype, but the background color of the dorsum can be brown or gray. One female lacks a wide occipital band because the transverse dark brown bars are not fused and it has an inconspicuous vertebral stripe. Also, in this female there are no lateral bands, since it has unfused vertical bars on the flanks. The tail has dark brown rings in both sexes. Males have orange preloacal pores. The coloration and pattern of the juveniles are unknown.

Distribution and natural history. The northern known distribution limit of the new species is the type locality, near the Laja Lagoon, 1450 m, Biobío Region, Chile (37°21'S – 71°23'W; Fig. 4). At the type locality, this new species was found inhabiting areas composed of sandy ground and volcanic sediments, where large accumulations of different sized igneous rocks protrude from the soil (Fig. 5). These sites correspond to a slag heap of solidified lava. The vegetational cover is low, consisting mainly of high-Andean forbs with species such as *Echium vulgare* and *Verbascum thapsus*, as well as the bush *Ephedra chilensis*. It is an abundant lizard of saxicolous habits. It was observed to be active between 9h00 and 18h00, taking refuge under the volcanic rocks. Also, we observed specimens in several places near the slopes of Antuco Volcano (37°23'S – 71°23'W, 1320 m; 37°23'S – 71°23'W, 1270 m; 37°23'S – 71°25'W, 1074 m) in similar environments. Near the Laja Lagoon, at its upper altitudinal limit (1450 m), this species was found in syntopy with *Phymaturus vociferator* Pincheira-Donoso, 2004. At 1320 m, it was found in syntopy with "*L. kriegii* *Liolaemus* sp. A" and *Diplolaemus sexcinctus* Cei et al., 2003. At its lower altitudinal limit (1074 m), it was found in syntopy with *L. lemniscatus* Gravenhorst, 1838 and *L. tenuis* (Duméril & Bibron, 1837).

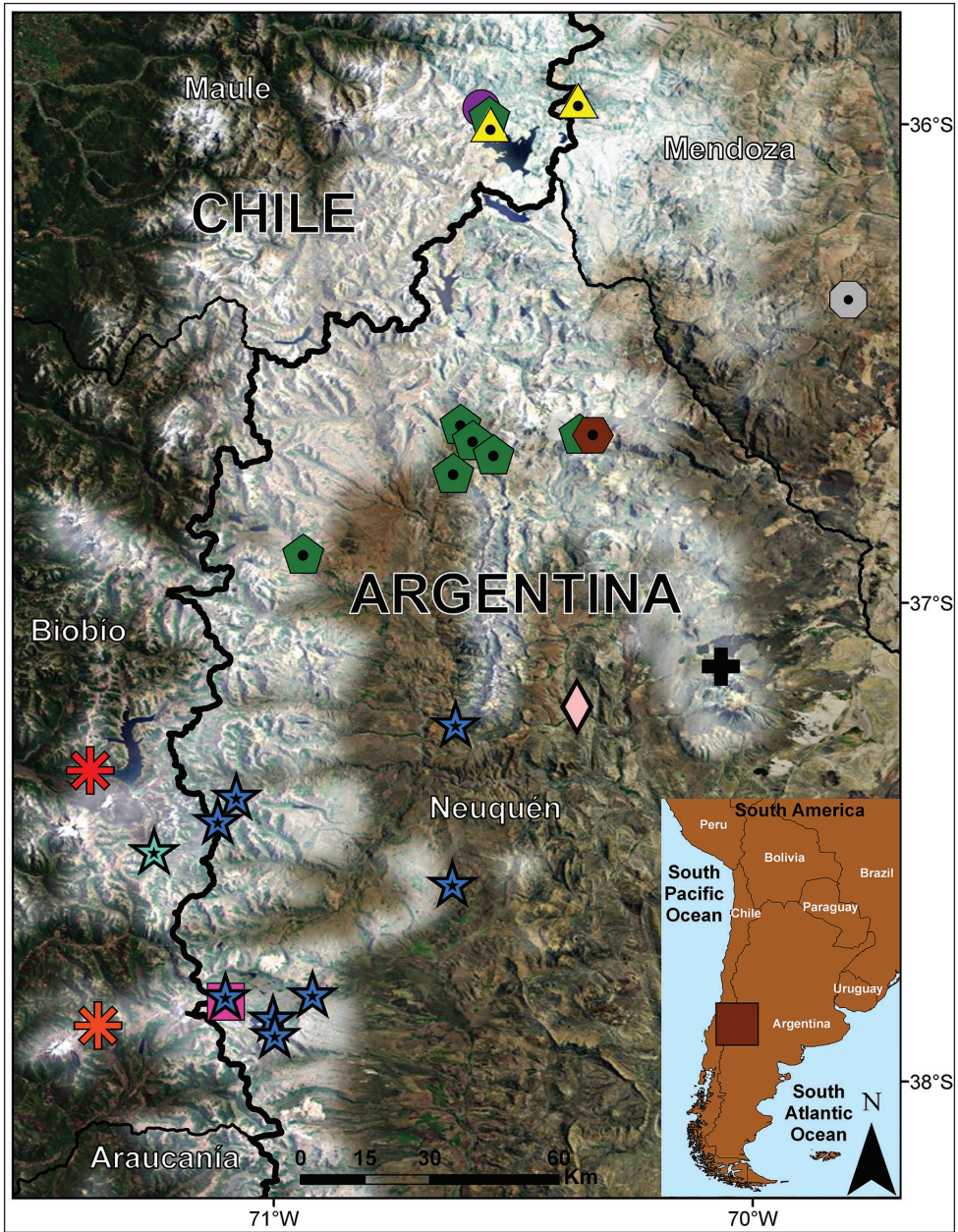


Figure 4. Distributional map for *Liolaemus scorialis* sp. n., *L. zabalai* sp. n. and the species of the *elongatus-kriegi* complex that inhabit in proximity of its. Asterisk: *L. scorialis* (red = near Laja Lagoon, type locality; orange = La Mula Lagoon). Star: *L. zabalai* sp. n. (light green = road to Los Barros, type locality; blue = distribution in Argentina). Purple circle: *L. carlosgarini*. Yellow triangle: *L. flavipiceus*. Green pentagon: *L. buergeri*. Gray octagon: *L. choique*. Brown hexagon: *L. antumalguen*. Black cross: *L. punmahuida*. Light pink diamond: *L. burmeisteri*. Pink square: *L. tregenzai*.



Figure 5. View of the type locality of *Liolaemus scorialis* sp. n., composed mainly of scoria volcanic rock.

Its southern limit of distribution is in La Mula Lagoon (La Araucanía Region, Chile), 48 km South from the Antuco Volcano (37°53'S – 71°22'W), 1600 m. We have no data for vegetation or environment in La Mula Lagoon. In this location, according to the Herpetological Catalog of the Museo de Historia Natural of Concepción (unpublished), *L. scorialis* occurs in syntopy with *L. pictus* (Duméril & Bibron, 1837). However, this report probably actually refers to *L. septentrionalis* Pincheira-Donoso & Núñez, 2005 (fide Vera-Escalona et al. 2012). The Museo de Historia Natural of Concepción also listed an unidentified species of *Liolaemus* (labeled as *Liolaemus monticola* sp., see discussion) and the snake *Tachymenis chilensis* Schlegel, 1837, from La Mula Lagoon.

The intestinal and stomach contents were examined; plant and insect remains were found in the intestine, along with a large number of nematodes of an unidentified species. No remains were found in the stomach. At the time of capture (January) two females had three embryos each and one female had several small oocytes.

***Liolaemus zabalai* sp. n.**

<http://zoobank.org/063D3CC3-0606-4CC4-8216-8F6B2B38CC3C>

Fig. 6

Liolaemus kriegi, Donoso-Barros, 1974. Bol. Soc. Biol. Concepción, 47: 287.

Liolaemus kriegi (in part), Cei, 1986. Mus. Reg. Scien. Nat. Torino, 4: 230.

Liolaemus sp?, Torres-Pérez, 1997. Not. Biol., 5(4): 146.

Liolaemus kriegi (in part), Pincheira-Donoso, 2001. Not. Mens. Mus. Nac. Hist. Nat., Chile, 346: 11.

Liolaemus sp. A, Morando et al., 2003. Syst. Biol., 52: 179.

Liolaemus kriegi (in part), Pincheira-Donoso & Núñez, 2005. Pub. Oc. Mus. Nac. Hist. Nat., Chile, 59: 289.

Liolaemus kriegi (in part), Mella, 2005. Guía Camp. Rep. Chil. Zon. Cent., p. 64.

Liolaemus sp. A, Medina et al. Cuad, 2013. Herp. 27(1): 27.

Liolaemus sp. A, Medina et al., 2014. Biol. J. Linnean Soc. 113: 256.

Holotype. SSUC Re 602 (Fig. 6). Near Los Barros, Laja Lagoon, Biobío Region, Chile. (37°31'S – 71°15'W, 1460 m). Collected by J. Troncoso-Palacios, F. Urra and H. Díaz. 07/01/2014.

Paratypes. SSUC Re 598. Adult male. SSUC Re 597, 599, 600–01. Four adult females. The same data as the holotype (Figs 6 and 8). MZUC 35607, 39567. One male and one female. Malleco, Antuco Volcano, Los Barros. Unknown coll.

Etymology. This species is named after Patricio Zabala, collection manager of the “Colección de Flora y Fauna Patricio Sánchez Reyes, Pontificia Universidad Católica de Chile” (SSUC). We dedicate this species to him because of his support of herpetological research in Chile, allowing us to review and deposit material in SSUC, and especially for his friendship.

Diagnosis. *Liolaemus zabalai* belongs to the *kriegi* clade of the *elongatus-kriegi* complex and is closely related to some undescribed species: *Liolaemus* sp. C and *Liolaemus* sp. D; being more distant from the currently described species *L. buergeri*, *L. kriegi* and *L. tregenzai* (Fig. 7). According to Medina et al. (2014), in regards to the species of the *kriegi* clade *L. zabalai* is sympatric only with *L. tregenzai* at the Copahue Volcano.

With respect to the species of the *kriegi* clade, *Liolaemus zabalai* differs from *L. tregenzai* because the latter has 71–85 midbody scales and the males have no preloacal pores (Pincheira-Donoso and Scolaro 2007), whereas *L. zabalai* has 90–104 midbody scales and the males have 3–5 preloacal pores. In addition, the green-bluish ventral color of *L. tregenzai* is completely absent in *L. zabalai*. The uncorrected pairwise difference (cyt-b) between the species is 3.09% (Medina et al. 2014).

Liolaemus zabalai differs from *L. kriegi* in that the latter reaches 101.1 mm SVL, has reddish cloacal coloration in both sexes and has an unringed tail (Avila et al. 2003), whereas *L. zabalai* is smaller (max. SVL = 92.0 mm), has yellowish cloacal coloration in both sexes and has a ringed tail (in specimens with original tails). The uncorrected pairwise difference between these species is 3.79% (Medina et al. 2014).

Liolaemus zabalai differs from *L. buergeri* in that the latter has fewer dorsal scales (78–91; $x = 84.1 \pm 4.4$, $n = 14$) than *L. zabalai* (86–96; $x = 89.4 \pm 3.2$, $n = 8$) (Mann–Whitney $U = 19.5$; $P = 0.01$, $DF = 20$). *Liolaemus zabalai* has more loreal scales between the nasal and the subocular (4–6; $x = 4.3 \pm 0.6$, $n = 8$) than *L. buergeri* (3–4; $x = 3.3 \pm 0.5$, $n = 14$) (Mann–Whitney $U = 11.0$; $P < 0.01$, $DF = 20$). Also, *L. buergeri* has a vertebral stripe on the tail, whereas *L. zabalai* has a ringed original tail. The limbs in *L. zabalai* are black with dispersed light brown spots, whereas *L. buergeri* has brown limbs with dispersed black spots (Fig. 8). *Liolaemus zabalai* and *L. buergeri* share basically the same dorsal coloration pattern, but this is noticeably more marked and darker in *L. zabalai* (Fig. 8, see discussion). Based on the cyt-b locus, the uncorrected average pairwise difference between *L. zabalai* and *L. buergeri* is 2.94% (Medina et al. 2014), greater than the values reported for other *Liolaemus* widely accepted as valid species



Figure 6. *Liolaemus zabalai* sp. n. **A, B** Holotype, male **C, D** Paratype, female **E** Paratype, male **F** Paratype, female. All from the type locality, near Los Barros, Laja Lagoon, Biobío Region, Chile.

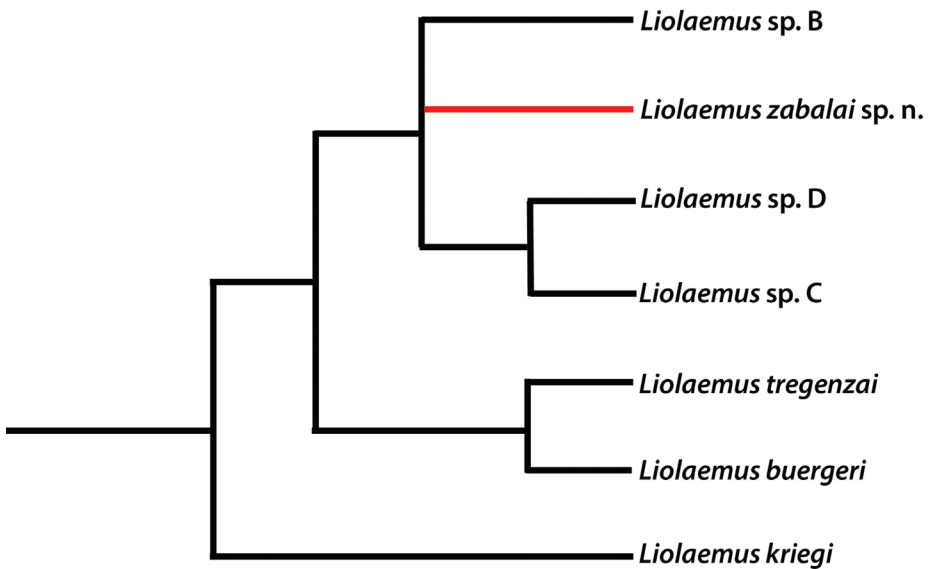


Figure 7. Phylogenetic position of *Liolaemus zabalai* sp. n. in the *kriegi* clade, based on cytochrome-b (cyt-b) locus according to Medina et al. (2014).

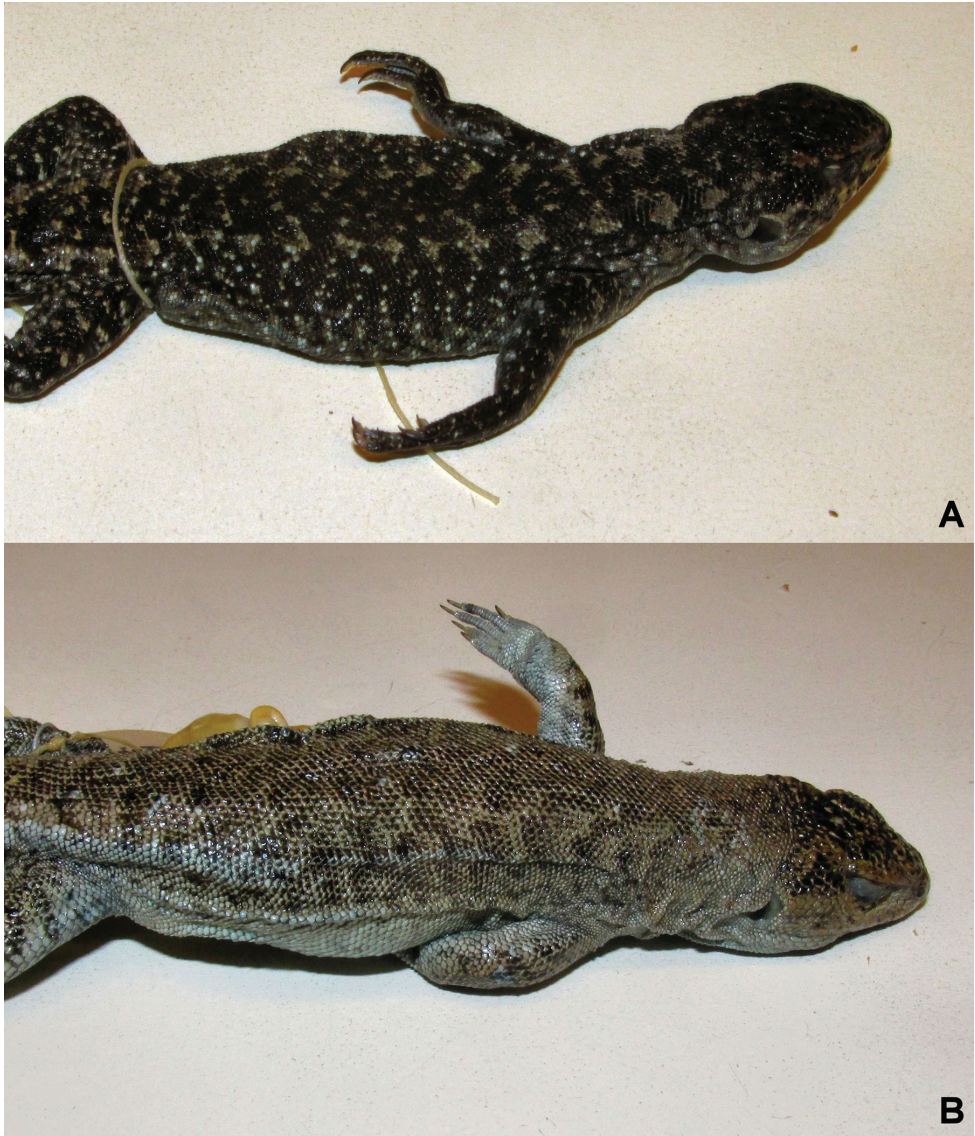


Figure 8. Comparison of the dorsal color pattern. **A** *Liolaemus zabalai* sp. n. with marked color pattern and **B** *L. buergeri*, diffuse color pattern.

(see discussion). Also, *L. zabalai* can vocalize, a feature only documented for *L. chilienensis* in the entire genus *Liolaemus* (Labra et al. 2013). Finally, although the ranges overlap, males of *L. buergeri* have 3–4 ($x = 3.3$) precloacal pores, whereas males of *L. zabalai* have 3–5 ($x = 3.9$) precloacal pores (Medina et al. 2014).

Compared to the other species of the *elongatus-kriegi* complex that occur near the known distribution of *Liolaemus zabalai*, the new species may be diagnosed as follows. Males of *L. zabalai* have precloacal pores, whereas males of *L. flavipiceus* and *L. pun-*

mahuída lack them (Table 3). *L. zabalai* is larger than *L. scorialis*; and *L. zabalai* has more midbody scales than *L. antumalguen*, *L. burmeisteri* and *L. choique* (Table 3).

Description of the holotype. Adult male. SVL: 90.3 mm. Tail length: 92.3 mm (autotomized). Axilla-groin length 39.7 mm. Head length (from the posterior border of the auditory meatus to the tip of the snout) 22.2 mm. Head width (distance between the two ear openings) 16.5 mm. Head height (at the level of ear openings) 11.7 mm. Forelimb length 28.5 mm. Hindlimb length 47.1 mm. Foot length 23.4 mm. Rostral scale wider (4.5 mm) than high (2.2 mm). Two postrostrals. Four internasals. Heptagonal interparietal scale, with a central, small, and whitish central spot marking the position of the parietal eye. Interparietal smaller than right parietal, but bigger than left parietal, surrounded by eight scales: nine scales between the interparietal and the rostral; 14 scales between occiput and rostral; orbital semicircle complete on both sides (formed by 13 scales); 5 supraoculars on both sides; seven superciliary scales. Frontal area is divided into six scales (three posterior, one anterior-left, two anterior-right); 2 scales between nasal and canthal; preocular separated from the lorilabials by one loreal scale; nasal in contact with the rostral, surrounded by six scales. There is one row of lorilabials between the supralabials and the subocular. Seven supralabials, the fourth is curved upward without contacting the subocular. Five infralabial scales. The mental scale is pentagonal and is in contact with four scales. Four pairs of postmental shields, the second is separated by two scales. Temporal scales are subimbricated and smooth or slightly keeled. Nine temporal scales between the level of superciliary scales and the rictal level. Two projected scales on the anterior edge of the ear, which are small and do not cover the auditory meatus. There is no differentiated auricular scale. Forty-two gulars between auditory meatus. Well developed “Y” shaped lateral neck fold with antehumeral and posthumeral folds developed. Dorsolateral fold slightly developed. Midbody scales 90. Dorsal scales on the vertebral zone are lanceolate to rounded, subimbricate, keeled and without mucrons. Dorsal scales on the paravertebral fields are more rounded, subimbricate, smooth or with less developed keels, without mucrons and there are interstitial granules between them. Dorsal scales are smaller than the ventral scales. Dorsal scales 86. Ventral scales are rhomboidal, smooth, subimbricate, and with few interstitial granules. Ventral scales 122. There are three preloocal pores. The suprafemoral scales are rhomboidal, imbricate, and smooth or keeled. Infracemoral scales are lanceolate to rhomboidal, smooth, and subimbricate and with few interstitial granules. Supra-antebrachials scales are rhomboidal to rounded, subimbricate, and keeled or smooth. Infra-antebrachials are rounded to rhomboidal, subimbricate, and smooth. The dorsal scales of the tail are lanceolate to rectangular, subimbricate, keeled or smooth and with few interstitial granules. The ventral scales of the tail vary from lanceolate to triangular, and are subimbricate and smooth. Lamellae of the fingers: I: 11, II: 16, III: 20, IV: 22 and V: 15. Lamellae of the toes: I: 12, II: 16, III: 21, VI: 27 and V: 18.

Color of the holotype in life. Black head, with some light brown spots on the supraocular and snout areas. The scales located behind the orbital semicircles are light brown; but the interparietal scale, parietal scales and the scales in contact with the

parietal scales are black. Superciliary scales are light brown with black spots. Temporal scales are light brown; cheeks light gray with some black spots. Subocular is gray with a black vertical line on the middle. Background color of the dorsum is light brown. Wide occipital band on the dorsum, formed by twelve transverse black bars. Very few whitish scales dispersed on the dorsum. Black lateral band bearing a few dispersed whitish scales, running from the tip of snout to the groin. Flanks below lateral band are light brown. Limbs black with dispersed light brown spots. Tail light brown with inconspicuous vertebral stripe in the regenerated zone; occipital black band ends in the first fifth of the tail, remainder with some dispersed black spots and a black vertebral stripe. Throat, belly and ventral surfaces of limbs whitish with dispersed inconspicuous dark dots. Rear portion of the belly and the thighs are yellowish. Ventrally, tail is whitish with a dark gray ventral stripe and diffuse dark gray rings from the cloaca to the midpoint of the tail. Precloacal pores orange.

Variation. In three males: SVL: 72.6–90.3 mm. Axilla-groin distance: 32.7–38.6 mm. Head length: 17.6–22.2 mm. Head width: 14.2–16.5 mm. Head height: 9.2–11.7 mm. Foot length: 21.5–23.0 mm. Leg length: 42.1–47.2 mm. Arm length: 24.6–28.5 mm. Tail length: 102.0 mm in one specimen (autotomized in the rest). In three females: SVL: 71.8–90.2 mm. Axilla-groin distance: 32.9–42.7 mm. Head length: 17.9–19.5 mm. Head width: 13.9–16.6 mm. Head height: 9.4–11.1 mm. Foot length: 20.6–24.2 mm. Leg length: 41.5–48.8 mm. Arm length: 24.8–29.4 mm. Tail length: 105–115 mm (in two specimens without autotomized tails).

The variation of the scalation in *Liolaemus zabalai* is as follows. Midbody scales: 90–104 ($x = 94.3 \pm 4.8$). Dorsal scales: 86–96 ($x = 89.4 \pm 3.2$). Ventral scales 116–122 ($x = 119.5 \pm 2.1$). Fourth finger lamellae: 19–22 ($x = 20.9 \pm 1.0$). Fourth toe lamellae: 26–27 ($x = 26.8 \pm 0.5$). Supralabial scales: 6–7 ($x = 6.6, \pm 0.5$). Infralabial scales: 4–5 ($x = 4.6 \pm 0.5$). Interparietal scale pentagonal, hexagonal or heptagonal, bordered by 5–8 scales ($x = 7.3 \pm 1.1$). Precloacal pores in males: 3–4.

There is slight sexual dichromatism; males are slightly darker than females. In general, all specimens have the pattern and color described for the holotype. One female has rusty-colored scales dispersed on the flanks, paravertebral fields and groin. In all specimens, the ventral surface of the throat, belly and limbs are whitish with dark marked or inconspicuous dots dispersed; there is a fragmented midventral stripe on the belly of two specimens. Males and females have a yellowish coloration in the posterior portion of the belly and the thighs (faint in some females). The tail has black rings, marked or diffuse, with a fragmented vertebral stripe in all specimens with complete original tails. Males have orange precloacal pores. The coloration and pattern of the juveniles are unknown.

Distribution and natural history. To our knowledge, in Chile this species is only found in the surroundings of the Laja Lagoon. The type locality is near Los Barros, Laja Lagoon, Biobío Region, Chile (37°31'S – 71°15'W, 1460 m, Fig. 9); but we also saw specimens (not collected) on the road to the Laja Lagoon at two localities (37°23'S – 71°23'W, 1320 m; 37°23'S – 71°22'W, 1390 m). The new species was found inhabiting areas of sandy soil with rocks of small and medium size. The vegetational cover is low, consisting mainly of *Ephedra chilensis*. It is an abundant lizard of saxicolous habits. This species was observed active between 11h00 and 18h00, taking refuge in



Figure 9. View of the type locality of *Liolaemus zabalai* sp. n.

cavities under the rocks. Near Los Barros, at its upper altitudinal limit (1460 m), this species was found in syntopy with *Diplolaemus sexcinctus*. At the lower altitudinal limit (1320 m), it was found in syntopy with *Liolaemus scorialis*, *Phymaturus vociferator* and *D. sexcinctus*. Two specimens of *L. zabalai* vocalized (squealed) in several occasions in response to the manipulation.

Liolaemus zabalai is also found in Argentina (where it has been called “*Liolaemus* sp. A”) at several localities in Neuquén Province (Morando et al. 2003, Medina et al. 2013, 2014).

An analysis of the intestinal contents performed on one specimen, showed that this species is omnivorous, but feeds mainly on plants. At the time of capture (January) the females had no embryos, but three had several small oocytes.

Discussion

In this work, the taxonomic status of two Chilean populations of the *Liolaemus elongatus-kriegi* complex from the Laja Lagoon have been clarified, here newly described as *L. zabalai* (previously confused with *L. kriegi* and also designed as *Liolaemus* sp. A) and *L. scorialis*. Pincheira-Donoso (2001) recorded two species of the *L. elongatus-kriegi* complex from the same location: *L. kriegi* and *L. buergeri*. Even though we did not examine the three specimens of “*L. buergeri*” listed by Pincheira-Donoso (2001), we believe that these correspond to *L. scorialis*, since the aspect of this new species resem-

bles *L. buergeri* (although it is notably smaller than it) and we did not find additional species of the *elongatus-kriegi* in the vicinity of Laja Lagoon. Also, Troncoso-Palacios et al. (2012) published several photographs of specimens from a population of “*L. buergeri*” from Los Humos, Libertador Bernardo O’Higgins Region, Chile, but unfortunately those specimens were not collected. This population is completely isolated from other populations of *L. buergeri* and some specimens exhibit a completely black ventral coloration, a feature absent in other populations of *L. buergeri* (Donoso-Barros 1966, Pincheira-Donoso and Núñez 2005). A more conclusive study in regard to this population should be conducted. Besides, there is diverse evidence supporting the existence of at least three more undescribed species currently assigned to *L. buergeri* in Argentina (Medina et al. 2013, 2014, Morando et al. 2003).

Assigning *Liolaemus scorialis* to any of the groups (Lobo 2005, Lobo et al. 2010b) or clades (Morando et al. 2003, Avila et al. 2012) proposed for such a diverse lineage of Patagonian lizards is a difficult task, especially taking into account that the phylogenetic studies based on morphological and molecular data disagree, and unfortunately we do not have molecular data for *L. scorialis*. However, it is unlikely that *L. scorialis* belongs to the *leopardinus* group-clade, because it completely lacks “leopard-like” dorsal spots, a distinctive feature of these lizards (Lobo 2005). Also, it is unlikely that *L. scorialis* belongs to the *capillitas* group, because species of this group share two synapomorphies absent in *L. scorialis*: spots in the shoulder region and a red coloration in the cloacal zone (Abdala et al. 2010, Lobo 2005). The *petrophilus* clade (Avila et al. 2012, Morando et al. 2003) includes all species of the *capillitas* group (with the exception of *L. heliodermis*, not sampled) plus *L. austromendocinus*, *L. gununakuna*, *L. parvus* and *L. petrophilus*. However, with the exception of *L. petrophilus* and *L. gununakuna*, all species of the *petrophilus* clade have fewer than 82 midbody scales (Abdala et al. 2010, Avila et al. 2004, Espinoza and Lobo 2003, Quinteros et al. 2008), whereas *L. scorialis* has 76–90 midbody scales. In regards to the *punmahuida* clade (Avila et al. 2010a), included into the *elongatus* group by Lobo et al. (2010b), both species of this clade (*L. flavipiceus* and *L. punmahuida*) have red coloration in the cloacal zone and males lack precloacal pores (Avila et al. 2003, Cei and Videla 2003), features absent in *L. scorialis*. *Liolaemus scorialis* is probably related to the *elongatus* or *kriegi* clades, as some species of these clades occur in the vicinity or in the type locality of *L. scorialis* and have similar counts of midbody, dorsal and ventral scales. Also, some of these species have white dorsal dots, rings on the tail and yellow in the cloacal zone (Abdala et al. 2010, Avila et al. 2010a, 2012, Cei 1986) like *L. scorialis*. A molecular phylogeny including *L. scorialis* is required to clarify this.

In the case of *Liolaemus zabalai* of the *kriegi* clade, the uncorrected pairwise differences between it and other species of the *kriegi* clade are 2.94–3.79%, almost at the limit of the value (3%) proposed for identify candidate species in *Liolaemus* (Breitman et al. 2012). In comparison, other *Liolaemus* lizards widely accepted as valid species show a lower level of differentiation for the mitochondrial gene *cyt-b*, for example: *L. martorii* Abdala, 2003 vs. *L. morenoi* Etheridge & Christie, 2003, 2.73% (Avila et al. 2010b); *L. riojanus* Cei, 1979 vs. *L. multimaculatus* (Duméril & Bibron, 1837), 1.23% (Avila et al. 2009); *L. chacabucoense* Núñez & Scolaro, 2009 vs. *L. kingii* (Bell, 1843), 2.22% (Breitman 2013). *Liolaemus zabalai* can vocalize, a trait only documented for *L. chiliensis*

(Labra et al. 2013) and also taken as diagnostic feature in *Liolaemus* (Pincheira-Donoso and Núñez 2005: 232) and the closely related genus *Phymaturus* (Lobo et al. 2010a: 118). Regarding the morphological diagnosis included in previous studies, Pincheira-Donoso and Núñez (2005) reviewed two specimens of *L. kriegi* from Laja Lagoon (here described as *L. zabalai*), which they described and provided the following diagnosis “the species is very similar to *L. buergeri*, differing in that the latter has a lighter color, brown or dark brown; in combination with a smaller number of keeled scales on the dorsum” (Pincheira-Donoso and Núñez 2005: 289, our translation). Here, we find the same color difference, and expand the differences in scalation; although we found no differences in the number of dorsal scales. Medina et al. (2013) recorded a similar maximum SVL (86.3 mm) compared to us (92.0 mm). Also, Medina et al. (2013) based on a discriminant analysis of several continuous and meristic characters, reported that *L. zabalai* (designated as “*Liolaemus* sp. A” in its study) has sexual dimorphism, with a sample of 21 females and 23 males. We were unable to replicate the statistical analysis to confirm this sexual dimorphism because our sample is small (5 females and 3 males). Also, Medina et al. (2013) recorded 3–5 precloacal pores in the males ($n = 23$), whereas we recorded only 3–4 ($n = 3$). Eventhough we found *Liolaemus scorialis* and *L. zabalai* in syntopy, *L. scorialis* was found mainly in a solid lava slag heap (where it was the only species recorded in this environment), whereas *L. zabalai* was found in bushy-rocky environments together with specimens of *L. scorialis* and other lizards. Regarding the population of “*L. kriegi*” from Cordillera de Curicó in Chile, 35°10'S (Donoso-Barros 1966), we have doubts about its real identity, especially considering that according to Medina et al. (2014) *L. kriegi* is distributed south of 38°40'S latitude (coordinates transformed by us).

Torres-Pérez (1997) recorded two *Liolaemus* sp. from Laja Lagoon. He pointed that one of them has 92 midbody scales, brown color and precloacal pores in males. It is difficult to try an identification, but the midbody scale count match with *L. zabalai*. Torres-Pérez (1997) indicated that the other *Liolaemus* sp. has no precloacal pores. It match only with *L. chillanensis* Müller & Hellmich, 1932, recorded in the Laja Laogon (Pincheira-Donoso and Núñez 2005).

In this study, *Liolaemus ceii* is considered a junior synonym of *L. kriegi*. This synonymy was recommended by Morando et al. (2003) because they did not find genetic evidence to differentiate both species. Recently, Medina et al. (2014) performed a wider genetic study and found that these two species form one lineage, called “*Liolaemus kriegi* + *ceii*”. Because individuals from both type localities show some morphological differences, they proposed two hypothesis: (1) *L. ceii* and *L. kriegi* constitute two species, for which different environments prompted relatively rapid and recent morphological divergence with insufficient time for molecular differentiation; and (2) they are conspecific and show clinal morphological variation owing to local adaptations (Medina et al. 2014). However, the published literature regarding *L. ceii* and *L. kriegi* (Cei 1986, Donoso-Barros 1971) does not include enough morphological comparison between them. We believe that for the moment *L. ceii* should be considered as a junior synonym of *L. kriegi*, because published morphological evidence to support *L. ceii* as full species is insufficient and the results of genetic studies (Medina et al. 2014, Morando et al. 2003) do not support to *L. ceii* as full species.

Liolaemus chillanensis was included in the *elongatus* clade by Avila et al. (2010a) and Avila et al. (2012) based on mitochondrial DNA data generated by Torres-Pérez et al. (2009), but at least part of the specimens used as vouchers were misidentified (Troncoso-Palacios, unpublished data). Therefore, in this study we do not consider *L. chillanensis* as a member of the *elongatus-kriegi* complex and we excluded it from our comparisons. Also, we examined one male of *Liolaemus monticola* ssp. (MRC 676) syntopic with *L. scorialis* in La Mula Lagoon, and identified it as *L. neuquensis* Müller & Hellmich, 1939, a species described from Copahue Volcano (Müller and Hellmich 1939b), 15 km E from La Mula Lagoon; being the first record of *L. neuquensis* in Chile.

In summary, this work describes two new species of the *elongatus-kriegi* complex lizards from the vicinity of the Laja Lagoon, in southern Chile, one probably confused with *L. buergeri*: *L. scorialis* and the other with a history of mis-identifications as *L. kriegi* or *Liolaemus* sp. A, for which we provide the formal name *L. zabalai*. Nonetheless, there is certainly still much to discover about the diversity of this group of Patagonian lizards.

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Appendix I

Specimens examined. Museum codes are as follow: LCUC (Laboratorio de Citogenética, Facultad de Ciencias, Universidad de Chile), MNHN-CL (Museo Nacional de Historia Natural, Chile), MRC (Museo de Historia Natural de Concepción), MZUC (Museo de Zoología de la Universidad de Concepción) and SSUC (Colección de Flora y Fauna Patricio Sánchez Reyes, Pontificia Universidad Católica de Chile).

Liolaemus buergeri. LCUC 2311. El Planchón, 2370 m. M. Lamborot & M.E. Manzur colls. 07/01/1996. SSUC Re 434–37. El Planchón, road to Teno Lagoon. J. Troncoso-Palacios, L. Negrete & R. Barros colls. January, 2012. SSUC Re 171–180. Maule Lagoon. F. Ferri coll. 20/02/2011.

Liolaemus carlosgarini. MNHN-CL 4531–67. Road to Maule Lagoon. C. Garín coll. 22/02/2008. SSUC Re 181–189, 349. Road to Maule Lagoon. F. Ferri coll. 20/02/2011.

Liolaemus cristiani. SSUC Re 537. El Peine. J. Troncoso-Palacios coll. 28/11/2011.

Liolaemus flavipiceus. MNHN-CL 2118, 2120. Maule Lagoon. C. Veloso & S. Silva colls. MNHN-CL 2167, 2170. Maule Lagoon. J.C. Torres-Mura & H. Núñez. MNHN-CL 4399–07. Laguna del Maule, aguas abajo, 2153 m. C. Garín & G. Lobos colls. 03/03/2008. SSUC Re 169–70. Maule Lagoon. F. Ferri coll. 20/02/2011.

Liolaemus frassinettii. LCUC 800–01. Cantillana. Unknown coll. 14/04/1983. SSUC Re 80. Altos de Cantillana. F. Torres coll.

Liolaemus leopardinus. MNHN-CL 3437–3439. El Colorado. H. Núñez, C. Garín, V. Meriggio, S. Fox & S. Perea colls. 06/01/2001. MNHN-CL 4025, 4027–28. Farellones. C. Veloso coll. 11/01/1988. MNHN-CL 4890–91. El Colorado. D. Esquerré, M. Palma, S. Fox & E. Santoyo colls. February, 2012. SSUC Re 364. Farellones. F. Ferri coll. 12/10/2010. SSUC Re 365. Farellones. F. Ferri coll. 13/02/2011. SSUC Re 366–67. Farellones. F. Ferri, M.L. Carrevedo & J. Troncoso-Palacios colls. 25/01/2012.

Liolaemus neuquensis. MRC 676. La Mula Lagoon, Araucanía Region, Chile. Unknown coll.

Liolaemus ramonensis. MNHN-CL 4007–08, 4012, 4015–17. Quebrada de Macul. C. Veloso & P. Espejo colls. 06/03/1987.

Liolaemus scorialis. SSUC Re SSUC Re 612–17. 7 km NW of the summit of the Antuco Volcano, near the Laja Lagoon, Biobío Region, Chile. J. Troncoso-Palacios, F. Urrea & H. Díaz colls. 08/01/2014. MRC 675, 677, 680, 682. La Mula Lagoon, Ralco National Reserve. Unknown coll. 01/12/2001.

Liolaemus ubaghsi. MNHN-CL 3808–16. Chapa Verde. H. Núñez, C. Garín & D. Pincheira-Donoso colls. 22–23/05/2003. MNHN-CL 1601. Chapa Verde. M. Elgueta coll. SSUC Re 491–92. Tranque Barahona, O'Higgins Region, Chile. R. Thomsom & G. Ugalde colls. 15/04/2008.

Liolaemus valdesianus. SSUC Re 129. Cajón del Maipo. Unknown coll. SSUC Re 363. Lo Valdés. F. Ferri coll. 10/01/2011. SSUC Re 559. El Yeso. C. Garín coll. 20/02/2013.

Liolaemus zabalai. SSUC Re 597–602. Near Los Barros, Laja Lagoon, Biobío Region, Chile. Collected by J. Troncoso-Palacios, F. Urra and H. Díaz. 07/01/2014. MZUC 35607, 39567. Malleco, Volcán Antuco, Los Barros. Unknown coll.