INTRODUCTION

Regionalization in biogeography allows division of the globe into areas according to different criteria, such as combination of biotic elements, physiognomy and climate. This regionalization constitutes a hierarchical system which comprises many categories of areas, from realms to biogeographic districts (Ebach, Morrone, Parenti, & Viloria, 2008), and is the most widely used system to address distribution and conservation aspects (Morrone, 2018).

America is divided into three regions: Nearctic, Neotropical and Andean (Morrone, 2004, 2006, 2015), with two transition zones being recognized between them: The Mexican TZ (Halffter, 1976; Morrone & Márquez, 2001; Morrone, 2015) and the South American TZ (Van...
der Hamer & Cleef, 1983; Morrone, 2004). The South American TZ (hereafter STZ) stretches across Argentina, Bolivia, Chile, Peru, Ecuador and Colombia and is bounded in the south by the Andean region and in the north and east by the Neotropical region (Morrone, 2014). The STZ comprises seven provinces: Páramo, Desert, Puna, Atacama, Cuyan High Andean, Monte and Comechingones (Martínez, Arana, Oggero, & Natale, 2017; Morrone, 2014; Morrone & Ezcurra, 2016). According to Morrone (2018), there is not a sole procedure to assign areas to any particular category; therefore, recognition of a particular area as belonging to a region or a TZ can be assessed using different approaches. For example, Morrone (2006) determined the belonging or placing of a biogeographic province (BP) into a TZ by applying panbiogeographic analyses, where a TZ can be identified by the presence of nodes or areas where different generalized tracks overlap. Other methods to define the belonging or placing of a BP into a region or a TZ could be providing a quantitative measure of the proportion of its biogeographical elements. Roig-Juñent, Tognelli, and Morrone (2008) designated a group of taxa belonging to a biogeographic area and sharing a common origin as biogeographic element (BE). The term biogeographic element refers to the origin of a specific taxon based on its phylogeny. The concept of BE allows delimitation of areas showing predominance of a particular biogeographic element (e.g., the biota of the Neotropical region presents an elevated proportion of taxa of neotropical origin). In contrast, transition zones (TZ), which are areas located on borders between geographic regions and represent areas of biotic overlap, are promoted by historical and ecological changes that allow the mixture of taxa of different biogeographic origins (Ferro & Morrone, 2014; Morrone, 2006). Hence, those biogeographic provinces that belong to a TZ, in accord with the definition given above, show no predominance of any particular BE.

The proportions of the different BEs represented in an area have been used in several studies. For example, Van der Hamer and Cleef (1983) found that the genera of woody plants associated with Nothofagus forests of the Austral-Antarctic region (Andean) comprised the following percentages of BEs: 70% were Austral-Antarctic, 25% Neotropical and 5% Holarctic elements, whereas in the Neotropical region, there were 85% of Neotropical, 10% of Holarctic and 5% of Austral-Antarctic elements. With respect to transition zones, these authors found that 50% of the biota of the Puna Biogeographic province, which is considered part of Morrone’s STZ (2004, 2006), has a neotropical origin. This percentage varies in the southern area of Puna where, according to Simpson (1983), there are more species of austral origin (Andean). Another province that Morrone (2006) considered as belonging to the STZ is the Monte where, according to Solbrig et al. (1977), 60% of plant species have neotropical origin. Morrone (2015) defines the Andean region, which borders with the southernmost part of the STZ, and divides it into three subregions: Central Chilean, Sub-Antarctic and Patagonian Steppe. Katinas, Morrone, and Crisci (1999) hypothesized that the biota occurring in the Andean region has a composite origin; particularly that of the Patagonian Steppe appears to be linked to that of the Neotropics. The Patagonian Steppe presents a mixture of neotropical and Andean BEs, its flora is biogeographically related to Andean BEs, along with elements of the Monte province (Hauman, 1947). Among Arthropoda taxa, both the Neotropical (Flores, 1997, 2000; Flores & Triplehorn, 2002; Lescano, Elizalde, Werenkraut, Pirk, & Flores, 2016) and Andean BEs (Morrone, 2013; Roig-Juñent, 2000) can be found. The Patagonian Steppe therefore shows a mixture of unknown proportions of different BEs, which is typical of TZs located between biogeographic regions (Morrone, 2006). The definition of the Patagonian Steppe has been based mainly on its floristic composition (Cabrera, 1971; Hauman, 1947; Roig, 1998; Soriano, 1956). Several authors, mainly focused on vegetation studies, recognized natural areas within this province, such as Hauman (1947) (Eastern and Western), Soriano (1956) (Sub-Andean, Western, Central, San Jorge Gulf and Fuegian) and Roig (1998) (Payunia, Septentrional, Central, Meridional and San Jorge Gulf). Taking into account insect distribution, Roig-Juñent (1994) recognized three areas (Western, Austral and Central Patagonia), Morrone, Roig-Juñent, and Flores (2002) recognized three areas (Payunia, Central and Fuegian) and Domínguez, Roig-Juñent, Tassin, Ocampo, and Flores (2006) recognized five areas (Western Patagonia, Payunia, Sub-Andean, Austral and Central Patagonia). Furthermore, in a study based on plant, mammal and mostly insect distribution data, Patagonia is divided into two provinces: Sub-Andean and Central Patagonia (Morrone, 2001). Recently, Morrone (2015) thoroughly revised the nomenclature for this area as well as its divisions.

The main aims of this contribution were to assess whether the Patagonian Steppe BP belongs to the Andean region or whether it forms the southernmost part of the STZ, and to provide an up-to-date vector map (i.e., as a polygon shape-file) of the Patagonian Steppe and its biogeographic region-alization mainly based on Roig (1998) and Domínguez et al. (2006).

2 | MATERIALS AND METHODS

2.1 | Taxa

We compiled information from 177 genera of reptili ans, amphibians, insects, arachnids and plants that inhabit the Patagonian Steppe (Supporting Information Table S1). We worked at the generic taxonomic level in order to make data comparable, because we found that other taxonomic levels,
such as family or species, are not sufficiently informative: Families may have elements of different origins and species alone are not informative if they are not accompanied by other species of the same genus.

### 2.2 Allocation to a biogeographic element

In hypothesizing the origin of the different taxa (genera) included in this study and to assign them to a BE (when this information was not available from previous studies), the following data were considered: (a) current distribution of each genus, (b) current distribution of the sister group or related genera (in cases of unsolved phylogenies) and (c) the phylogenetic relationship with the remaining genera within the monophyletic unit considered (e.g., whether it is a tribe, subfamily, etc.).

On the basis of this information, each genus was allocated to a biogeographic element according to the definitions proposed by Roig‐Juñent et al. (2008): **Pangeic**: A South American genus belonging to a family or taxon of higher rank that originated before the breakup of the Pangea; the genus is endemic to South America but the taxon of higher rank presents other genera found in disjoint relictual areas (in continents other than South America), the Pangeic genera occurring in South America generally constitute basal groups within the family or taxon of higher rank. **Holarctic**: A South American genus most commonly distributed in the Andes Mountains or other parts of South America, whose sister group is distributed in the Holarctic realm, specifically the Nearctic. These genera could have originated recently with the South American and North American connection through the Panama Isthmus. When difficulty appears to differentiate this BE from those of Pangea, it becomes necessary to consider the phylogenetic relationships within the taxon of higher rank to which the genus belongs. A Pangeic genus will belong to higher taxa (e.g., a family), whose South American representatives will constitute the sister group of the Holarctic representatives; on the other hand, a Holarctic genus will belong to higher taxa whose South American representatives are part of a large group of Holarctic representatives, in which the South American genus is a crown taxon. **Gondwanic**: A South American genus belonging to a family or taxon of higher rank that originated in Gondwana, after the separation of Laurasia; the members of the group of higher rank are found in Gondwana fragments and are absent from the Holarctic realm. Within this category, Roig‐Juñent et al. (2008) recognized five different elements depending on the age of origin of the taxon with respect to Gondwana's fragmentation; three of these elements are recognized in the Patagonian Steppe. **Endemic Gondwanic**: A South American genus endemic to Southern South America and whose sister group inhabits tropical areas of South America and other areas such as the Paleotropics. The age of these taxa can be hypothesized as prior to Gondwana's breakup, when tropical areas were connected. **Neotropical Gondwanic**: A genus distributed mainly in the Neotropics, whose sister group could also be found in the Neotropics, or in the Neotropics and Tropical Africa, but not in other tropical areas of the world; these taxa could have originated when tropical South America and Africa were still united, but formed a separate unit from the remaining portion of tropical Gondwana, such as India. **Patagonian Gondwanic**: A South American genus distributed mainly in the Andean region, whose sister group is most frequently found in the south Antarctic forests of South America, or other austral areas of the world, such as Australia, Tasmania and New Zealand; these taxa originated in temperate areas of Gondwana, when a southern connection existed via Antarctica.

### 2.3 Allocation criteria

On the basis of previous studies (Solbrig et al., 1977; Van der Hammer & Cleef, 1983; Simpson, 1983), it should be noted that a BP is considered a part of a particular region when one of its constituent BEs surpasses 70% (e.g., 70% in the Andean region and 85% in the Neotropical region), whereas a BP having no predominance of any BE and, on the contrary, showing similar percentages (lower than 50%) of its main BEs, should be considered as belonging to a TZ.

### 2.4 A shapefile for the Patagonian Steppe

The boundaries of the Patagonian Steppe BP and its five subprovinces are shown in Figure 1 and are also presented as a polygon shapefile as Supporting Information. These boundaries are based mainly on the map by Roig (1998) because it is the most detailed map of the Patagonian Steppe to date (scale 1:5,000,000). In addition, the map includes the Chilean part of the Patagonian Steppe, which was absent in Soriano (1956), Roig (1998) and Leon, Bran, Collantes, Paruello, and Soriano (1998). The Chilean sector was redrawn from maps by Artigas (1975), Boelcke, Moore, and Roig (1985), Morrone et al. (2002) and distribution data belonging to the CEI (Colección Entomológica IADIZA).

### 3 RESULTS

One hundred and seventy-seven genera were assigned to a BE (Table 1, Supporting Information Table S1). We obtained a similar proportion of both Patagonian Gondwanic and Neotropical Gondwanic elements which represent almost 86% of the total genera analysed, whereas 10% belonged to the Pangeic element and 2% was assigned to the Endemic Gondwanic element (Table 1). For the areas within the Patagonian Steppe, percentages of genera belonging to...
each BE are shown in Figure 2. The most abundant BE in Western, Payunia and Central subprovinces was Neotropical Gondwanic, whereas Patagonian Gondwanic BEs predominated in Sub-Andean and Austral subprovinces.

4 | DISCUSSION

As previously mentioned, inclusion of a BP in any given region is defined by the percentage of its constituent BEs. Based on previous studies, we propose that an area can be considered a part of a particular region when one of its constituent BEs surpasses 70%. If the percentage value of the most common BE is lower than 70%, it does not indicate a clear belonging to any given region. This is the case for the Monte BP, where 60% of its BEs have neotropical origin (Solbrig et al., 1977) or the Puna BP, where Van der Hammer and Cleef (1983) considered that 50% of its biota have also neotropical origin. With regard to the Patagonian Steppe, located south of the Monte and considered by Morrone (2006, 2015) to be the southern border of the STZ, we found it shows a similar proportion of Patagonian Gondwanic and Neotropical Gondwanic BEs, which could lead us to propose the Patagonian Steppe as the southernmost province belonging to the STZ (Figure 3) and not to the Andean region as is currently proposed (Morrone, 2015). Previous studies show that in the Patagonian Steppe, an important proportion of BEs does not belong to the Andean region. A panbiogeographic approach based on plant, fungus and animal taxa showed

TABLE 1  Number of genera per order that were assigned to a biogeographic element. Numbers in parenthesis indicate percentages of the total

| Class/Order   | EG  | Ne  | NG  | Pa  | PG  | Total |
|---------------|-----|-----|-----|-----|-----|-------|
| Amphibia      |     |     |     |     |     |       |
| Anura         |     |     |     |     |     |       |
| Arachnida     |     |     |     |     |     |       |
| Araneae       |     |     |     |     |     |       |
| Scorpiones    |     |     |     |     |     |       |
| Magnoliopsida |     |     |     |     |     |       |
| Asterales     |     |     |     |     |     |       |
| Insecta       |     |     |     |     |     |       |
| Coleoptera    | 4 (5.63) |     | 33 (46.47) | 8 (11.26) | 26 (36.61) | 71 |
| Ephemeroptera |     |     | 7 (35.0) | 1 (5.0) | 12 (60.0) | 20 |
| Hemiptera     |     |     | 1 (20.0) | 4 (80.0) |     | 5 |
| Odonata       |     |     | 3 (37.5) | 5 (62.5) |     | 8 |
| Orthoptera    |     |     | 3 (33.3) |     | 6 (66.6) | 9 |
| Phasmida      |     |     |     |     | 1 (100.0) | 1 |
| Plecoptera    |     |     | 2 (8.7) |     | 21 (91.3) | 23 |
| Reptilia      |     |     |     |     |     |       |
| Squamata      |     |     | 6 (75.0) |     | 2 (25.0) | 8 |
| Testudines    |     |     | 1 (100.0) |     |     | 1 |
| Total         | 4 (2.26) | 2 (1.12) | 72 (40.67) | 19 (10.73) | 80 (45.2) | 177 |

Note. EG: Endemic Gondwanic; Ne: Nearctic; NG: Neotropical Gondwanic; Pa: Pangeic; PG: Patagonian Gondwanic.
the Patagonian steppe as forming part of a Neotropical pattern linked to the Andes uplift and to Quaternary glaciations (Katinas et al., 1999).

Our results show not only that the Patagonian Steppe is composed of a similar proportion of BEs of different origins, noticeably Patagonian and Neotropical Gondwanic elements, but also that the proportion of these elements varies among the subprovinces of the Patagonian Steppe. This is the case for Ephemeroptera, where 12 genera were found to belong to the Patagonian Gondwanic element (see Table 1) and nine of them are distributed only across the Sub-Andean subprovince. The three remaining genera occupy the Western, Sub-Andean and Central steppe, that is 84.9% of the Patagonian Gondwanic genera of Ephemeroptera occur in regions of the steppe that is close to Nothofagus forests. On the other hand, all Ephemeroptera genera of Neotropical Gondwanic origin, except for two (Apobaetis and Callibaetis), are mainly distributed in the central steppe. This pattern repeats itself in other taxa, especially in Plecoptera, where the Neotropical Gondwanic BEs analysed are mainly distributed in the central region of the steppe, whereas the Patagonian Gondwanic BEs are distributed in natural areas of the steppe close to Nothofagus forests, such as the Sub-Andean and Western subprovinces. This pattern shows that, although the different subprovinces of the Patagonian Steppe share common botanical features, the proportion of BEs present in each of them, varies according to their geographical location and proximity to the Andean region as well as to the STZ.

Furthermore, we found that, within the Patagonian Steppe, the proportion of Neotropical Gondwanic and Patagonian Gondwanic elements may also differ among taxonomic groups. Amphibia, Asterales and Araneae have an equal or similar proportion of Patagonian Gondwanic and Neotropical Gondwanic BEs. Other taxa present a predominance of Neotropical Gondwanic BEs, such as reptiles and Coleoptera, and the four analysed genera of scorpions which belong to the Neotropical BE, even the genus Urophonius Pocock that is Patagonian Gondwanic and related to the genus Orobothriatus Maury with whom it forms the sister group of a large Neotropical clade (Ojanguren-Affilastro & Ramírez, 2008). Taxa showing predominance of Patagonian BEs are Orthoptera and Ephemeroptera. With the exception of reptiles, scorpions and the order Plecoptera, which are herein represented by few genera, no taxonomic group surpasses 70% of BEs of Patagonian or Neotropical Gondwanic origin, not even those groups such as Coleoptera, which are represented by the highest number of genera, show this proportion.

Considering the Coleoptera, the taxa with the highest number of genera analysed, there are different families...
with diverse origins. For example, one-third of their analysed genera are phylogenetically related to taxa inhabiting the Nothofagus forests or circum-Antarctic areas of South America, or other austral regions of the world such as Australia, New Zealand and South Africa. This is the case of Brosocini (Roig-Juñent, 2000), Antarctini (Straneo, 1951) (Coleoptera: Carabidae) and Listeroderini (Morrone, 2013) (Coleoptera: Curculionidae). On the other hand, a high proportion (46%) has a neotropical origin such as the tribe Nycteliini that is one of the most conspicuous elements of Tenebrionidae (Coleoptera) in the Patagonian Steppe (Kuschel, 1969). Patagonian genera of Nycteliini show phylogenetic relationships with taxa of neotropical origin that inhabit other arid biogeographic provinces of the STZ such as Monte, Puna, and Prepuna, and the Neotropical Chacoan, Caatinga and Cerrado (Flores, 1997, 2000; Flores & Triplehorn, 2002).

This study of the biogeographic elements of the Patagonian Steppe has allowed us to propose its inclusion in the STZ based on a quantitative approach, and future research on these elements will allow a better understanding of the biogeographic elements represented in the Patagonian Steppe’s current diversity. Transition areas constitute the borders between regions and present sectors where their biota has more or less biogeographic affinity with that of their bordering regions. This pattern is also found within each of the biogeographic provinces that constitute the transition area, as shown in this study by the heterogeneity found in the different subprovinces of the Patagonian Steppe.

Finally, the shapefile of the Patagonian Steppe is based mainly on Roig (1998) who delimited natural areas based on vegetation. The map depicts floral districts proposed by Soriano (1956) and subprovinces according to endemism areas determined on the basis of insect distribution data proposed by Domínguez et al. (2006). In addition, this map also includes ecolonal areas of the Patagonian Steppe, that is smaller areas of biotic overlap between neighbouring subprovinces or provinces.

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