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BIOGEOGRAPHICAL ANALYSIS WITHIN THE FAMILY PHYTOSEIIDAE BERLESE
(ACARI: MESOSTIGMATA): AN EXAMPLE FROM THE LARGE SUB-GENUS
TYPHLODROMUS (ANTHOSEIUS) DE LEON

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ABSTRACT — Despite their great interest for pest management, the biogeography of the Phytoseiidae is still poorly known. This study focuses on the spatial distribution of one of the largest taxa within the family: the sub-genus *Typhlodromus (Anthoseius)* currently comprising 322 species. It also provides a description of five discrete morphological characters used to discriminate between species and easily observed. This survey is the first analysis of the combined distribution of both species and associated morphological characters within the family Phytoseiidae. It reveals the great utility of taxonomic data compilation and associated databases for aspects beyond alpha-taxonomy. The highest number of species of *Typhlodromus (Anthoseius)* was reported from the East Palearctic region and the lowest from the Neotropical province. This was quite surprising as the highest species diversity of Phytoseiidae is reported from this latter region. Several hypotheses are discussed to explain this observation. In provinces showing the highest number of *Typhlodromus (Anthoseius)* species, we usually also observed the most diverse morphological forms for the five characters considered. High endemism levels were observed in all the provinces considered, even those where only few species were reported (especially the Neotropical province). Possible synonymies are thus discussed, as species could have been described more than once in different biogeographic provinces. Furthermore, some hypotheses are discussed to explain the high species and morphological diversity of the sub-genus *Typhlodromus (Anthoseius)* in the West and East Palearctic regions. Further phylogenetic analyses are required to test these different hypotheses.

KEYWORDS — biogeography; taxonomy; endemism; Gondwana; Laurasia

INTRODUCTION

The family Phytoseiidae Berlese (Acari: Mesostigmata) includes the most frequent predatory mites found on plants. Several species are used in the bio-control of mite and insect pests in several crops all over the world (McMurtry and Croft 1997; Gerson *et al.* 2003). It is also the family containing the highest number of known species within the or-

der Mesostigmata, with more than 2,250 species described worldwide (Moraes *et al.* 2004; Chant and McMurtry 2007; Krantz and Walter 2009).

Despite its great interest for pest management, many aspects of Phytoseiidae distribution still remain poorly analyzed. Recently, the computerization and compilation of those data have been initiated (Tixier *et al.* unpub. database). Tixier *et*

al. (2008a) analyzed the biogeographic distribution of the whole family, opening new perspectives for further studies on the distribution and occurrence of each genus. Indeed, understanding the distribution of taxa could help (i) in building a clearer picture of the evolutionary scenario of the considered group (Brown and Lomolino 1998; Humphries and Parenti 1999) and (ii) in more applied grounds, in searching for new predatory species to be employed in bio-control programs. This paper focuses on the biogeographic distribution of the known species of a specific sub-genus of the family Phytoseiidae, sub-family Typhlodrominae: *Typhlodromus* (*Anthoseius*) De Leon. Together, with its sister group [sub-genus *Typhlodromus* (*Typhlodromus*) Scheuten], it forms the genus *Typhlodromus* Scheuten, which is included in the tribe Typhlodromini with two other genera: *Neoseiulella* Muma and *Typhloseiulus* Chant and McMurtry (Chant and McMurtry 2007). The sub-genus *Typhlodromus* (*Anthoseius*), currently comprising 322 valid species and at least 20 suspected synonymies, is one of the largest within the family (Moraes *et al.* 2004; Chant and McMurtry 2007; Faraji *et al.* 2008, 2011; El-Banhawy *et al.* 2009; Rahmani *et al.* 2010; El-Banhawy and Knapp 2011; Jafari *et al.* 2011). Very few studies have provided revisions of this large group. Ueckermann and Loots (1988) and Ueckermann *et al.* (2008) revised the African species of *Typhlodromus* (*Anthoseius*). Denmark and Welbourn (2002) only listed 236 species in their world revision of the genus *Amblydromella* Muma, considered a junior synonym of *Typhlodromus* (*Anthoseius*) (Moraes *et al.* 2004, Chant and McMurtry 2007). We herein considered species of the sub-genus *Typhlodromus* (*Anthoseius*) as defined by Chant and McMurtry (2007): presence of the setae *z3*, *s6*, *S5* and absence of *Z1*, all dorsal setae thin, ventrianal shield not reduced (Chant and McMurtry 1994, 2007). Species of this sub-genus are delimited with some discrete characters (peritreme length, number of solenostomes on the dorsal and ventrianal shields, number of setae on the sternal and on ventrianal shields) but also with continuous characters as the dorsal and leg setal lengths (Denmark and Welbourn 2002; Ueckermann *et al.* 2008; Papadoulis *et al.* 2009; Ferragut *et al.* 2010).

The first objective of the present study was to provide an exhaustive survey of the geographic distribution of the known species of *Typhlodromus* (*Anthoseius*). The second objective was to determine whether the species found since now in each of the seven biogeographical provinces bear specific morphologically discrete features. Considering the low number of morphological characters available and the difficulty to encode them (especially for continuous characters), as well as the difficulty of examining all type specimens, we believe it would certainly be premature to carry out a proper phylogenetic analysis at this point. Thus, our objective was only to provide a description of the relationships between geographical distribution and morphological attributes in the aforementioned group.

MATERIALS AND METHODS

Species distribution

Data on the geographic distribution of species were obtained from the two world catalogues of the family Phytoseiidae (Moraes *et al.* 1986, 2004) and from subsequent papers (*e.g.* Bayan and Merheb 2006; Ueckermann *et al.* 2008, 2009; Faraji *et al.* 2008, 2011; El-Banhawy *et al.* 2009; Rahmani *et al.* 2010; El-Banhawy and Knapp 2011; Jafari *et al.* 2011). Since no geographical data was available for neither *Typhlodromus* (*Anthoseius*) *wichmanni* Hirschmann nor *T. (T.) chanti* Hirschmann those two species were not considered in the present analysis.

The zoogeographic provinces used in the analysis are those defined by Sclater (1858) and Wallace (1876). These areas, initially called realms, were based on the taxonomic or phylogenetic relationships of animals (especially vertebrates) and not on the adaptations to specific environments (Ebach and Humphries 2002). Despite the fact that the data used to delineate provinces were compiled long before advances in Palaeontology and the theory of Continental Drift, the basic concept of distribution and the names of the zoogeographic provinces are still in use today (Humphries and Parenti 1999; Cox 2001). The biogeographic provinces considered are as follows: Nearctic (North America excluding Florida), Neotropical (South and Central

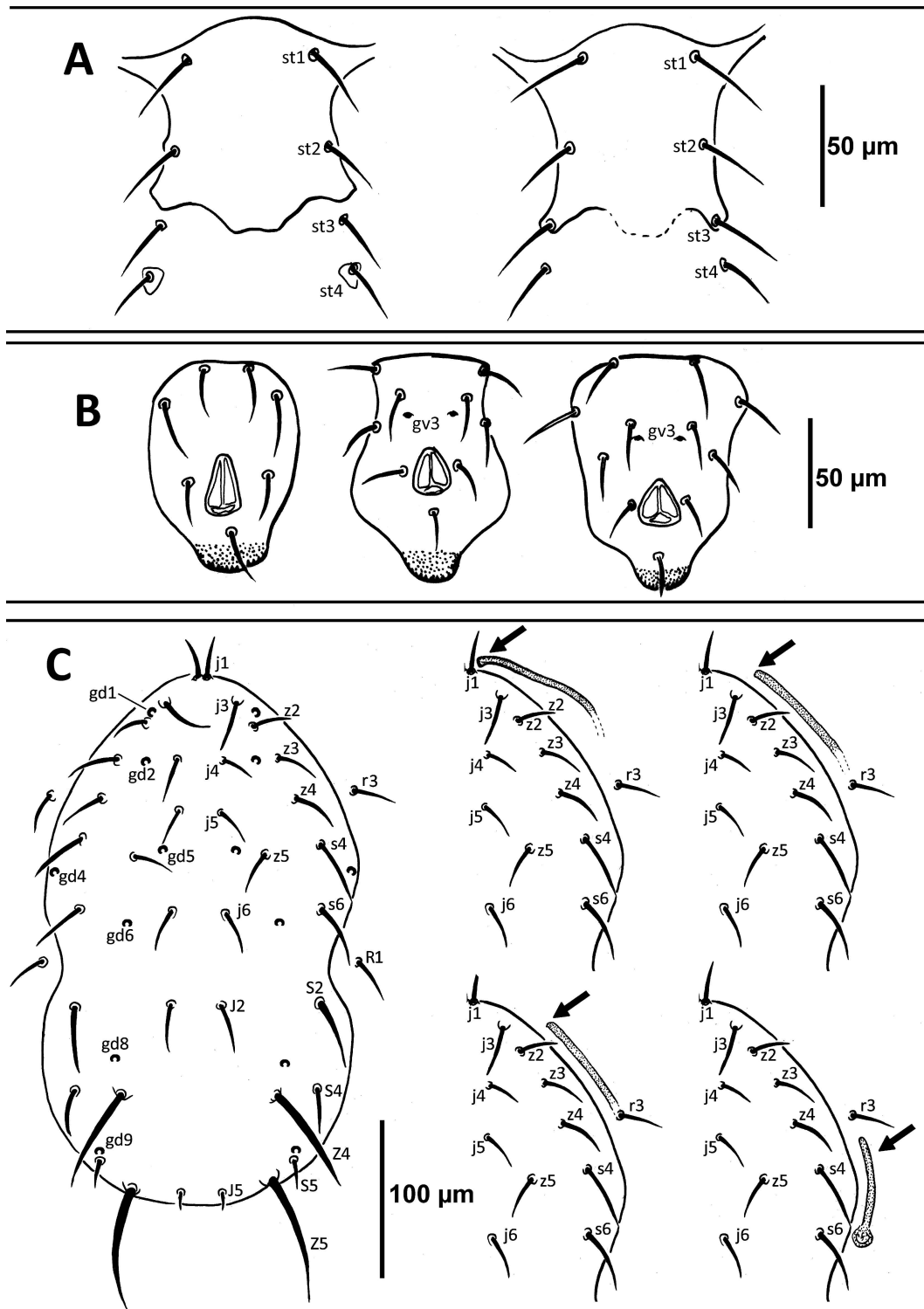


FIGURE 1: Morphological characters considered in the study of species of the sub-genus *Typhlodromus* (*Anthoseius*). A – the two sternal shield chaetotaxies (ST3 on or off the sternal shield), B – the three ventrianal shield chaetotaxies and position of the pair of pores (gv3) on the ventrianal shield, C – position and names of the dorsal setae (right-hand side of body) and solenostomes (left-hand side) and position of the peritreme endings.

America, Caribbean islands and Florida), Oriental (Taiwan, Malaysia, Philippines, India, Southern China, Okinawa, Japan, Thailand, South Korea), Australasian (Australia, New Zealand, New Caledonia, Papua New Guinea, Pacific islands), Ethiopian (sub-Saharan Africa including Madagascar), West Palaearctic (WP: Europe extending to Ural, Northern India and Northern Africa) and East Palaearctic (EP: Asia from Ural to Japan excluding Southern China and Okinawa). Because of its large surface, the Palaearctic region was divided in two parts, namely the West Palaearctic, and the East Palaearctic (*sensu lato*) (Gressitt 1974; Skuhrava and Skuhravy 2009). Furthermore, studies on the distribution of the Phytoseiidae species and genera revealed that different taxa are currently observed in these two areas (Tixier *et al.* 2008a; Tixier and Kreiter 2009). The Wallace Line (Darlington 1957) was used to separate the Oriental and Australasian provinces.

The database used for analyses includes the occurrence (presence/absence) of 320 species among the 322 presently described in the seven biogeographic regions (appendix 1). Species numbers and endemism levels were evaluated; endemism was considered when one species was reported only from a single biogeographic province.

Jaccard similarity indices were calculated between the areas as follows: $I = C / (N_1 + N_2 - C)$, where C is the number of species present in both areas, N_1 is the total number of species in area 1, and N_2 is the total number of species in area 2 (Brown and Lomolino 1998).

Spatial distribution of characters

Terminologies for chaetotaxy, poroidotaxy and adenotaxy used in this paper follow those proposed by Lindquist and Evans (1965) as adapted by Rowell *et al.* (1978) for the Phytoseiidae, and Athias-Henriot (1975), respectively.

The study of the spatial distribution of five qualitative discrete characters was carried out. We have chosen to analyze qualitative discrete characters in order to avoid interpretation errors (as it could be the case for continuous characters such as setae

lengths). Furthermore, they were quite easy to observe, and at the same time universally reported in description and re-descriptions. Finally, the five characters selected are currently used in species diagnostic (Ueckermann and Loots 1988; Denmark and Welbourn 2002; Ueckermann *et al.* 2008). Some rare intraspecific variations were observed but further experiments for testing if these variations really correspond to intraspecific morpholines would be clearly required (Rahmani *et al.* 2010, Jafari *et al.* 2011). The characters considered were as follows:

1. number of setae on the sternal shield, some species bear two pairs (*ST1* and *ST2*) whereas others bear three (*ST1*, *ST2*, *ST3*);
2. number of preanal setae on the ventrianal shield (*VAS*), some species bear two pairs, while others bear three or four;
3. presence of the pores *gv3* on the ventrianal shield;
4. number of solenostomes on the dorsal shield (ranging between one and seven: *gd1*, *gd2*, *gd4*, *gd5*, *gd6*, *gd8*, *gd9*);
5. peritreme length: 4 character states were considered, peritreme reaching at the levels of *j1*, between *j1-j3*, between *j3-z2* or between *z4-s4* (Figures 1 A-C).

To determine the state of these five characters, we used the original descriptions but also subsequent publications to check for the stability of the character state within a same species. However, although the overall literature on the sub-genus *Typhlodromus* (*Anthoseius*) was considered, information was not equally available for all the species and characters because of the poor quality and/or succinct descriptions. For some species, inconclusive characters were thus not considered. The most problematic feature was the number of solenostomes on the dorsal shield, because sometimes this was not mentioned in the description text, and drawings were often unclear. Furthermore, some authors (*e.g.* Gupta 1970) referred to solenostomes as "pairs of visible pores". In those cases, again, it is not conclusive to assume that those correspond

to solenostomes, or rather to other types of dorsal pores (e.g. poroids).

RESULTS

Species distribution

Species of the sub-genus *Typhlodromus* (*Anthoseius*) occur in the seven biogeographic provinces (Table 1). Distribution of species in these provinces are significantly different ($\text{Chi}^2_{(1,6)} = 210.16; P < 0.05$). Only seven species have been found in more than one biogeographic province [*T. (A.) bakeri* (Garnan), *T. (A.) caudiglans* Schuster, *T. (A.) persianus* McMurtry, *T. (A.) rhenanus* (Oudemans), *T. (A.) rhenanoides* Athias-Henriot, *T. (A.) richteri* Karg and *T. (A.) transvaalensis*] and a single species, *T. (A.) transvaalensis* (Nesbitt), was reported from all of them (Table 2). The East Palaearctic province shows the highest diversity (99 species), followed by the Ethiopian (87 species), the West Palaearctic (63 species) and the Oriental (59 species) provinces. Conversely, relatively few species are reported from the Neotropical (10 species), Nearctic (12 species) and Australasian (11 species) provinces (Table 1).

The great majority of species (313 species; 97.8%) are endemic to a single geographic province. Five provinces have endemism rates of at least 90 %

(Table 1). The highest endemism levels are observed in the four most species-rich areas (West Palaearctic, East Palaearctic, Ethiopian and Oriental) but also in the Neotropical province despite the low number of species reported in this latter area (10 species). The lowest endemism rate (50 %) is observed in the Nearctic province.

Jaccard indices were globally low, ranging from 0.93 % to 15 % (Table 3), which agrees with the high endemism rates previously discussed. The highest similarity rate is observed between the Nearctic and Australasian provinces (15 %) but this high rate is due to the occurrence of three species [*T. (A.) bakeri*, *T. (A.) caudiglans*, and *T. (A.) transvaalensis*] that are present in at least five provinces among the seven considered. The second highest similarity rate is observed between the Nearctic and West Palaearctic provinces (7.14 %). Again, among the five species common to these two provinces, four are worldwide distributed whereas a single one - *T. (A.) rhenanoides* - is only present in these two regions. Similarities around 5 % are noted between (i) Nearctic and Neotropical provinces (4.76 %), the single species in common, *T. (A.) transvaalensis*, is the most widespread species of the sub-genus; (ii) Australasian and Neotropical provinces [similarity of 5 %, one species in common: again, *T. (A.)*

TABLE 1: Number of species of the sub-genus *Typhlodromus* (*Anthoseius*) and endemism rates (%) in the seven biogeographic provinces considered

	Neotropical	Nearctic	Ethiopian	West Palaearctic	East Palaearctic	Australasian	Oriental
Number of species	10	12	87	63	99	11	59
Number of endemic species	9	6	83	57	95	8	55
% endemism / species of each zone	90%	50%	95%	90%	96%	73%	93%
% endemism / total of species	3%	2%	26%	18%	30%	3%	17%

TABLE 2: Species of the sub-genus *Typhlodromus* (*Anthoseius*) occurring in more than one biogeographic province.

	Neotropical	Nearctic	Ethiopian	West Palaearctic	East Palaearctic	Australasian	Oriental
<i>T.(A.) bakeri</i>		x		x	x	x	x
<i>T.(A.) caudiglans</i>		x	x	x	x	x	
<i>T.(A.) persianus</i>			x	x			
<i>T.(A.) rhenanoides</i>		x		x			
<i>T.(A.) rhenanus</i>		x	x	x			x
<i>T.(A.) rickeri</i>		x			x		x
<i>T.(A.) transvaalensis</i>	x	x	x	x	x	x	x

TABLE 3: Similarity indices of Jaccard (%) between the seven biogeographic provinces and species of the sub-genus *Typhlodromus* (*Anthoseius*).

	Neotropical	Nearctic	Ethiopian	West Palaearctic	East Palaearctic	Australasian
Nearctic	4.76					
Ethiopian	1.04	3.13				
West Palaearctic	1.38	7.14	2.74			
East Palaearctic	0.93	3.74	1.09	1.89		
Australasian	5.00	15.00	2.08	4.43	2.80	
Oriental	1.47	5.97	1.39	2.52	1.94	2.94

transvaalensis]. Finally, four species are simultaneously found in the Nearctic and Oriental provinces (similarity of 5.97%). However, again these species are the most worldwide distributed.

Spatial character distribution

Number of setae inserted on the sternal shield.

The number of species with two pairs (157 species) of setae on the sternal shield is slightly higher than the number of species bearing three pairs (139 species) (Figure 2). Species with two pairs are particularly numerous in the Ethiopian (52 species) and West Palaearctic provinces (45 species) where

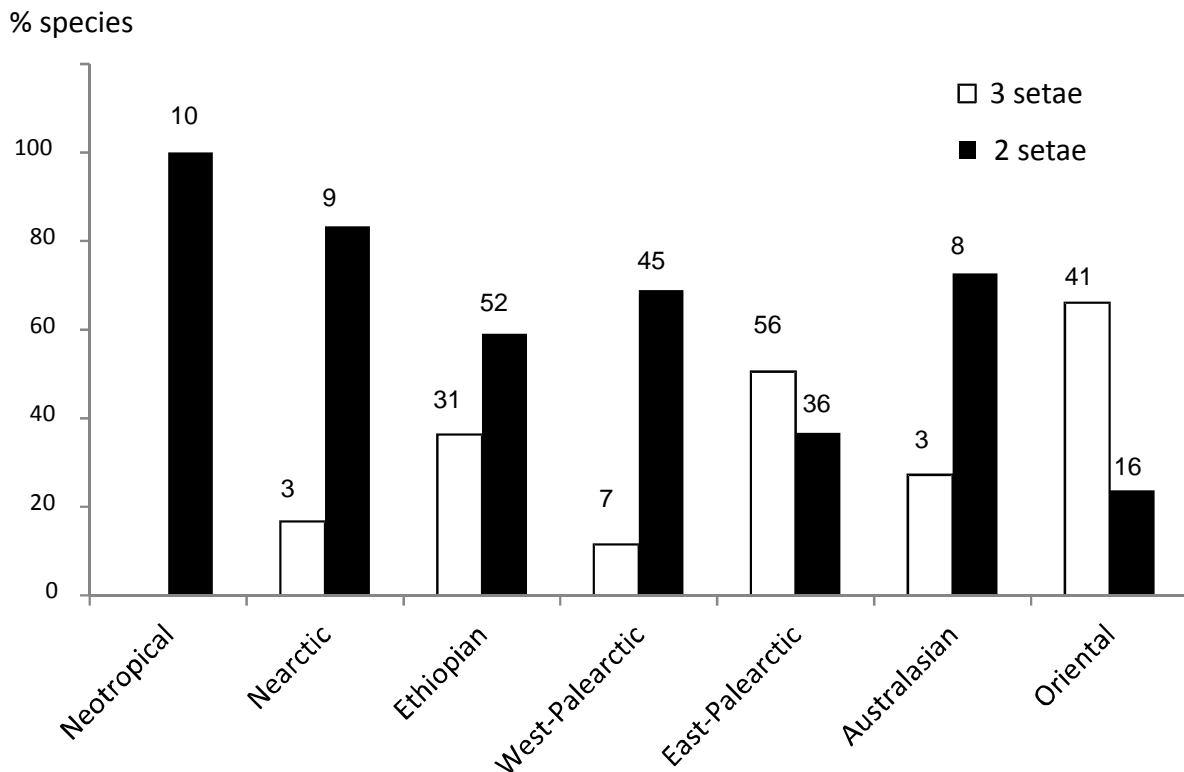


FIGURE 2: Number (above the bars) and percent of species of the sub-genus *Typhlodromus* (*Anthoseius*) with two or three pairs of setae on the female sternal shield in the seven biogeographic provinces considered.

they constitute the prevailing species. Species with two pairs of setae are also numerous in the East Palaearctic and Oriental provinces; however, in those areas they do not constitute the prevailing ones. No species with three setae on the sternal shield is reported from the Neotropical province and very few in the Nearctic and Australasian areas (Figure 2).

Number of setae inserted on the ventrianal shield (VAS).

Most species of *Typhlodromus (Anthoseius)* (280 species) have four pairs of pre-anal setae on the VAS; 31 species have three pairs whereas only four species have two (Figure 3). Species with four or three pairs are reported from all the seven provinces. However, they tend to be more numerous in the Palaearctic (both East and West), Ethiopian and Ethiopian provinces, which might be simply due to the higher number of species found in those provinces. Considering thus the percent of

species bearing these features in each province (Figure 3), the proportion of species with four setae on the VAS exceeds 80 % in all the provinces, except in the Neotropical and Nearctic provinces. Furthermore, in this latter province, 33.3 % of the endemic species have three setae whereas 66.6 % have four. One hundred percent and 77.8 % of endemic species have four setae on the VAS in the Australasian and Neotropical provinces, respectively. No clear delineation between geographical provinces regarding the presence of four or three setae on the VAS is thus visible. However, the four species with two setae on the VAS have only been recorded in the Palaearctic and Ethiopian provinces (Figure 3): *T. (A.) ilicis* Athias-Henriot in Morocco, France and Algeria (Athias-Henriot 1960, McMurtry and Bounfour 1989), *T. (A.) monosetus* Wang and Xu from China (Wang and Wu 1991), *T. (A.) poptari* (Kolodochka) in Russia (Kolodochka 1988), and *T. (A.) noonus* El-Banhawy and Knapp from Kenya (El-Banhawy and Knapp 2011).

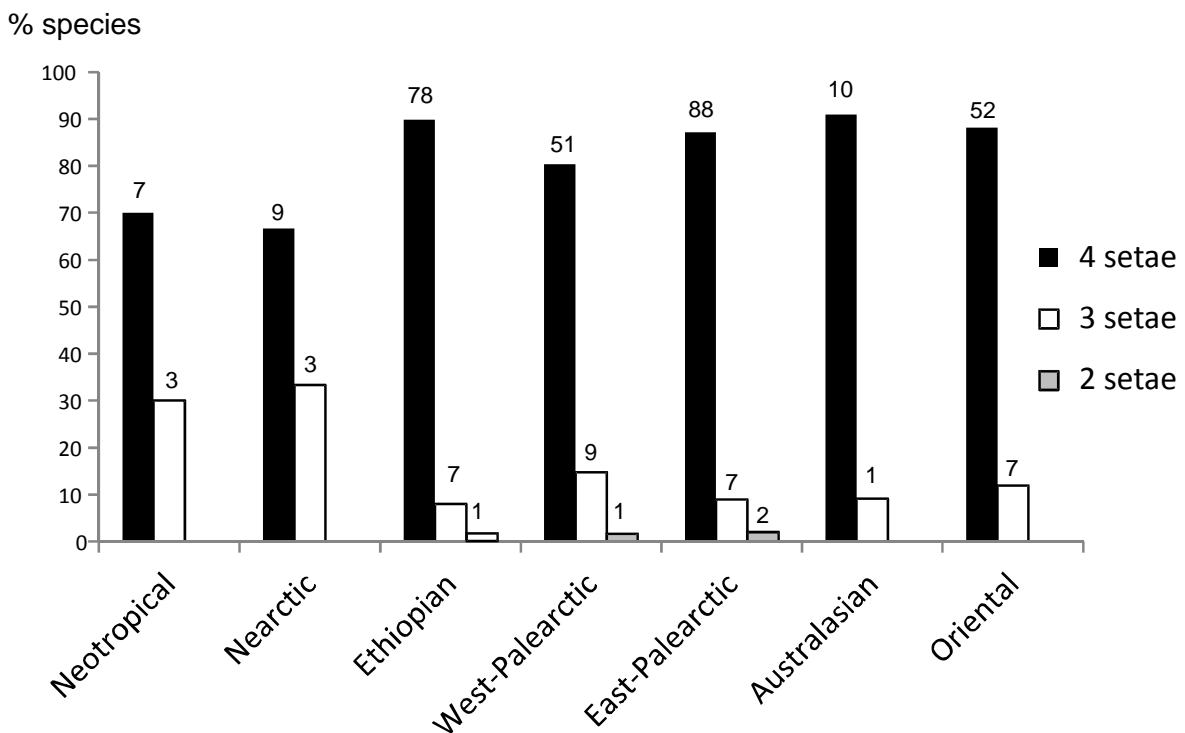


FIGURE 3: Number (above the bars) and percent of species of the sub-genus *Typhlodromus (Anthoseius)* with two, three or four pairs of preanal setae on the female ventrianal shield in the seven biogeographic provinces considered.

Presence of pores gv3 on the ventrianal shield.

gv3 Most species (262 species) of the sub-genus *Typhlodromus* (*Anthoseius*) have a pair of pores *gv3* on the ventrianal shield. These species are found mostly in the East Palaearctic (82 species), Ethiopian (80 species), Oriental (50 species), and West Palaearctic (40 species) provinces (Figure 4). Species reported from the Australasian and Nearctic provinces all bear a pore, as well as the majority of the Neotropical species.

Number of solenostomes on the dorsal shield.

The majority of the species of *Typhlodromus* (*Anthoseius*) for which this feature could be observed, has five solenostomes on the dorsal shield (126 species). They are found in all the seven geographic provinces, the highest number being reported from the Ethiopian (58 species) and the Palaearctic provinces (EP: 33 species, WP: 30 species) (Figure

5). Species with four solenostomes (24 species) are particularly numerous in the Palaearctic provinces (11 species in EP and 8 in WP). Species with three solenostomes are absent from the Neotropical zone, again these species are more commonly found in the Palaearctic provinces. Four species have only one solenostome (*gd2*), one is reported from the WP province [*Typhlodromus* (*Anthoseius*) *serratus* (Chaudhri)], two from EP [*T. (A.) balakotiensis* (Chaudhri, Akbar and Rasool) and *T. (A.) zafari* Chaudhri], and finally one from the Neotropics [*T. (A.) tecoma* (Denmark and Evans)]. Only two species have two solenostomes (*gd2* and *gd8*) [*T. (A.) luzonensis* Schicha and Corpuz-Raros and *T. (A.) manipurensis* (Gupta)]. However, for this latter species nothing was mentioned in the original description and this feature was inferred from the drawings. Finally, two species, *T. (A.) umbratus* (Chaudhri, Akbar and Rasool) and *T. (A.) noonus* El-Banhawy and

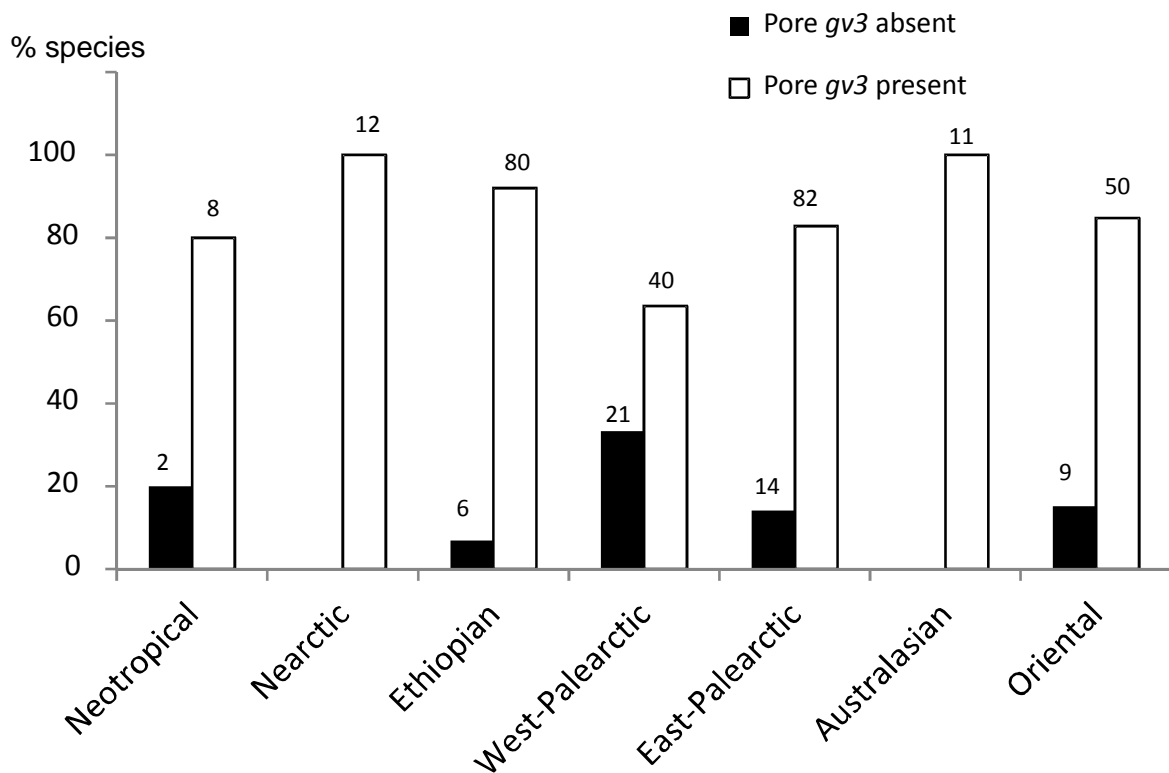


FIGURE 4: Number (above the bars) and percent of species of the sub-genus *Typhlodromus* (*Anthoseius*) with or without a pore (*gv3*) on the female ventrianal shield in the seven biogeographic provinces considered.

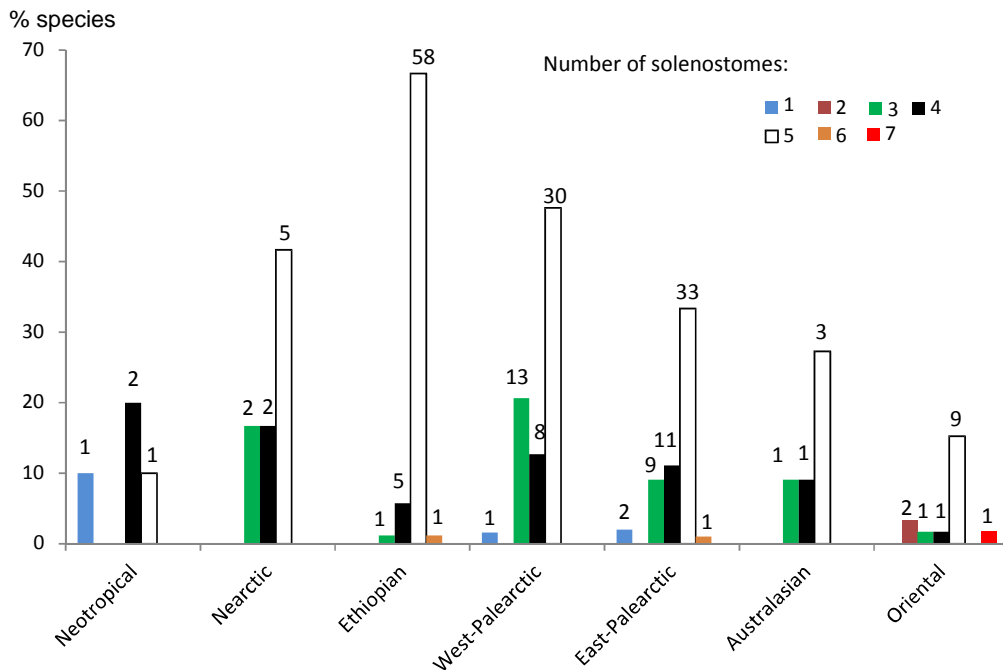


FIGURE 5: Number (above the bars) and percent of species of the sub-genus *Typhlodromus* (*Anthoseius*) with one to seven solenostomes on the female dorsal shield in the seven biogeographic provinces considered.

Knapp are reported to have 6 solenostomes, the former being endemic to the East Palaearctic province and the latter endemic to the Ethiopian region. Only one species, endemic to the Oriental province, has 7 solenostomes [*T. (A.) pruni* Gupta] (Figure 5).

Regions showing the highest diversity in the number of solenostomes are the Oriental, Ethiopian and the two Palaearctic provinces.

Peritreme length.

In most of the species (216 species), the peritreme reaches the level of the dorsal setae *j1*; only relatively few species show other positions. Species with this feature are numerous in all the provinces (Figure 6). Most of the species endemic to the Australasian and Neotropical provinces have the peritreme at *j1* position. Species with the shortest peritreme (15 species in total), *i.e.* reaching the level of dorsal setae *z4* to *s4*, are only present in the Palaearctic (8 species in WP, 3 species in EP), Oriental (3 species) and the Nearctic (1 species) provinces. Two types of peritreme are observed in species endemic to the Australasian province, and three in

the endemic species of the Nearctics and Neotropics (Figure 6).

DISCUSSION

This study focusing on the sub-genus *Typhlodromus* (*Anthoseius*) is the first one proposing combined analyses of the distribution of both species and morphological associated characters within the family Phytoseiidae. It clearly shows the utility of taxonomic data compilation and associated databases in broader aspects than only alpha-taxonomy (Meier and Dikow 2004).

This survey first emphasizes the worldwide distribution of *T. (A.) transvaalensis*. This species is found in many habitats including stored products (Amitai and Swirski 1978; Corpuz-Raros *et al.* 1988), and its wide distribution could be explained by the shipment of food from country to country.

The highest number of species of the sub-genus *Typhlodromus* (*Anthoseius*) was found in the whole Palaearctic region, whereas the lowest was reported

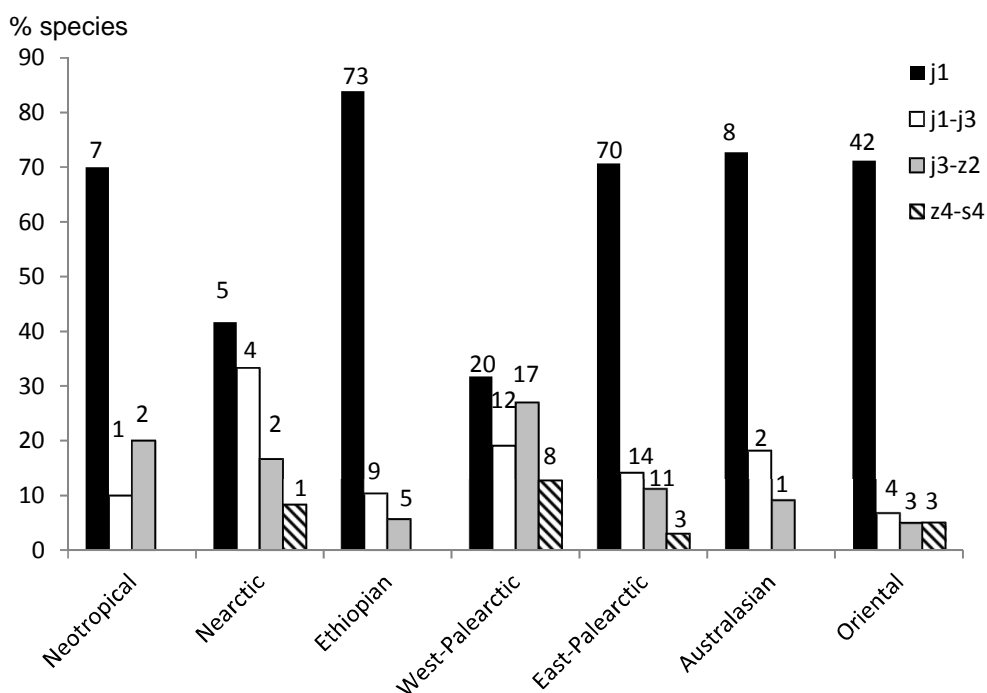


FIGURE 6: Number (above the bars) and percent of species of the sub-genus *Typhlodromus* (*Anthoseius*) having five different female peritreme lengths in the seven biogeographic provinces considered.

in the Neotropical province. This is quite surprising as this latter province contains the highest number of Phytoseiidae species reported to date (Tixier *et al.* 2008a). So one can wonder why a high diversity of *Typhlodromus* (*Anthoseius*) was not also observed in this area. Is it because species of this sub-genus have not been found yet in this province? In our opinion this hypothesis is weakly sustained, as the number of surveys carried out in the Neotropics is much higher than in other areas (Moraes *et al.* 2004; Tixier *et al.* 2008a; Tixier and Kreiter 2009). We could thus suggest other hypotheses as (i) species of this sub-genus did not colonize this province or (ii) have been extinct. Similarly, the number of species of *Typhlodromus* (*Anthoseius*) is also low in the Nearctics. The same hypotheses as those previously developed for the Neotropics could also be raised to explain the low number of species reported in the Nearctics.

The most diverse morphological forms of *Typhlodromus* (*Anthoseius*) have been observed in the two Palaearctic areas. This could appear quite logical.

Indeed, in provinces where the number of species of *Typhlodromus* (*Anthoseius*) was low, the number of morphological forms was also low (*e.g.* Neotropical and Australasian provinces) and *vice versa*. However, despite a great number of species reported in the Ethiopian province and in a less extent in the Oriental one, the number of morphological forms is much lower than in the Palaearctic provinces. This could suggest that only some morphological forms succeeded in colonizing these two former provinces, nevertheless with a great success in regards to the occurrence of many species. Other hypotheses could be that these forms have not been found yet or that they would have been extinct.

In the West and East Palaearctic provinces, we also observed rare character states (two setae on the ventrianal shield, short peritreme). The highest number of species and morphological diversity, as well as some rare characters in these latter provinces could suggest that these parts of the world are the centre of origin of the sub-genus. The distribution of the three other genera of the tribe

(Typhlodromini) [in which is included *Typhlodromus* (*Anthoseius*)] seems to concur to this hypothesis (Moraes *et al.* 2004; Tixier *et al.* 2008a). Indeed, all species of the genus *Typhloseiulus* Chant and McMurtry [same dorsal chaetotaxy as *Typhlodromus* (*Anthoseius*)] are reported from the West Palaearctic area; species of the sister sub-genus, *Typhlodromus* (*Typhlodromus*) Scheuten, are mainly reported from the Palaearctic province (60 % and 14 % of species from the West and East Palaearctic areas, respectively); and 38% of the species the genus *Neoseiulella* Muma are endemic to the West Palaearctic region. Furthermore, Kanouh (2010) questioned the polyphyly of this latter genus to explain that 50 % of species are endemic of the Australasian province. However, as high diversity and endemism in the Palaearctic provinces can be due to other factors, i.e. extinctions in the other provinces (Cain 1944; Brown and Lomolino 1998; Humphries and Parenti 1999), additional phylogenetic studies would be thus required to test the hypothesis presently discussed.

Finally, the present study emphasizes high endemism levels in all the provinces considered, even those where only few species were reported (*i.e.* Neotropics). Organisms can be endemic to a location because they originated in that place and never dispersed but also because they can be extinct in the original area and only survive in the secondary colonized zones (Brown and Lomolino 1998). Biology and history of species of *Typhlodromus* (*Anthoseius*) is poorly known, thus it would be highly speculative to try to explain the endemism rates. Our own experience suggests however that species of *Typhlodromus* (*Anthoseius*) are rare and when found the number of specimens is low. For instance, 269 species have been reported less than four times, whereas six species (the worldwide ones) have been reported more than 30 times. Finally, the high endemism rates at species level could also be due to an artefact. Indeed, the morphological features that really allow to discriminate between species within the family Phytoseiidae, and as a consequence in the sub-genus *Typhlodromus* (*Anthoseius*), are sometimes questionable (Tixier *et al.* 2003, 2006a, b, 2008b, 2010; Okassa *et al.* 2009, 2010). We can thus wonder if the number of synonymies which

is presently very low (6 %) would not be in reality much higher. Many species described or only reported from one province may have actually been described more than once from another province (under another name). Furthermore, many species have been described based on few specimens thus it is quite difficult to determine *a posteriori* if this is really a new species or a variant of a known one. Moreover, for some authors it could have been difficult to assess the huge taxonomic information (322 original descriptions in 58 countries and in nearly a dozen languages). To better assess the level of synonymy, different studies on this sub-genus are planned. At first, molecular experiments will be conducted to assess the reliability of some morphological characters presently used for diagnosis; a DNA sequence database will be built to assist diagnosis (La Salle *et al.* 2009). However, the main difficulty presently encountered is the collection of live material, given the rarity of the species. The second work consists in a compilation of the morphological characters of all the species of this sub-genus in order to construct an electronic polytomous key intended to be freely accessible on the web in order to facilitate the identification of the species of *Typhlodromus* (*Anthoseius*) and to avoid the description of species already known.

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APPENDIX 1

Species of *Typhlodromus* (*Anthoseius*) considered and their characteristics for the five morphological characters considered: (I) number of setae on the sternal shield; (II) number of setae on the ventrianal shield; (III) presence

of the pores *gv3* on the ventrianal shield (0 = absent; 1 = present); (IV) number of solenostomes on the dorsal shield; (V) peritreme extremity at the level of *j1* (1), between *j1* – *j3* (2), between *j3* – *z2* (3) or between *z4* and *s4* (4); U = unknown state or condition.

species	characters				
	I	II	III	IV	V
1 <i>T. (A.) abboudieius</i>	3	4	1	5	2
2 <i>T. (A.) acacia</i>	3	4	1	4	1
3 <i>T. (A.) acaciae</i>	3	4	1	5	1
4 <i>T. (A.) adenensis</i>	2	4	0	U	2
5 <i>T. (A.) admirabilis</i>	3	4	1	U	1
6 <i>T. (A.) aenaulus</i>	2	4	1	U	2
7 <i>T. (A.) aestivalis</i>	2	4	0	5	2
8 <i>T. (A.) agilis</i>	3	4	1	5	2
9 <i>T. (A.) ailanthi</i>	3	4	1	5	1
10 <i>T. (A.) aktherecus</i>	3	4	1	5	3
11 <i>T. (A.) algonquinensis</i>	2	3	1	3	2
12 <i>T. (A.) apoxys</i>	2	4	1	U	1
13 <i>T. (A.) applegum</i>	2	4	1	3	1
14 <i>T. (A.) argyronamus</i>	2	4	0	5	1
15 <i>T. (A.) arizonicus</i>	3	4	1	U	1
16 <i>T. (A.) arunachalensis</i>	3	4	1	U	1
17 <i>T. (A.) asperosetosus</i>	2	4	1	U	1
18 <i>T. (A.) astibus</i>	2	4	1	5	1
19 <i>T. (A.) asticus</i>	3	4	1	5	1
20 <i>T. (A.) athenas</i>	2	4	0	5	1
21 <i>T. (A.) atoosae</i>	2	4	1	5	3
22 <i>T. (A.) auratus</i>	2	4	1	5	1
23 <i>T. (A.) bagdasarjani</i>	2	4	0	5	3
24 <i>T. (A.) bakeri</i>	2	4	1	5	1
25 <i>T. (A.) balakotiensis</i>	U	4	1	1	1
26 <i>T. (A.) balanites</i>	3	4	1	5	1
27 <i>T. (A.) bambusae</i>	2	3	1	U	3
28 <i>T. (A.) bambusicolus</i>	3	3	1	U	1
29 <i>T. (A.) banahawensis</i>	3	4	1	U	1
30 <i>T. (A.) bergi</i>	3	4	1	5	1
31 <i>T. (A.) beskaravainyi</i>	2	4	0	5	3
32 <i>T. (A.) betulae</i>	3	4	1	U	1
33 <i>T. (A.) bifurcata</i>	2	4	1	U	1
34 <i>T. (A.) bondarenkoi</i>	U	4	1	5	2
35 <i>T. (A.) borealis</i>	2	4	1	U	1
36 <i>T. (A.) brevemedius</i>	2	4	1	U	1
37 <i>T. (A.) brisbanensis</i>	3	4	1	U	1
38 <i>T. (A.) buccalis</i>	3	4	1	5	1
39 <i>T. (A.) bullatus</i>	3	4	1	5	1
40 <i>T. (A.) cannabis</i>	3	4	1	5	1
41 <i>T. (A.) capparidis</i>	2	4	1	5	1
42 <i>T. (A.) caucasicus</i>	2	4	1	5	1
43 <i>T. (A.) caudiglans</i>	2	4	1	5	1
44 <i>T. (A.) celastrus</i>	2	4	1	3	1
45 <i>T. (A.) cephalochaitosus</i>	3	4	1	U	2
46 <i>T. (A.) cerasicolus</i>	2	4	0	5	2
47 <i>T. (A.) cervix</i>	3	4	1	U	1
48 <i>T. (A.) changi</i>	3	4	0	U	1
49 <i>T. (A.) channabasavannai</i>	3	4	1	U	1
50 <i>T. (A.) charactus</i>	2	3	1	U	2
51 <i>T. (A.) chazeaui</i>	3	4	1	U	1
52 <i>T. (A.) chinensis</i>	3	4	1	5	1
53 <i>T. (A.) chrysanthemi</i>	U	4	1	U	U
54 <i>T. (A.) clairathiasae</i>	U	4	0	U	2
55 <i>T. (A.) combretum</i>	2	4	1	4	1
56 <i>T. (A.) commenticus</i>	2	4	1	3	3
57 <i>T. (A.) communis</i>	3	4	1	U	U
58 <i>T. (A.) concavus</i>	2	4	1	U	2
59 <i>T. (A.) coniferculus</i>	2	4	1	5	1
60 <i>T. (A.) constrictus</i>	3	4	1	5	1
61 <i>T. (A.) coryli</i>	3	4	1	5	3

species	characters				
	I	II	III	IV	V
62 <i>T. (A.) coryphus</i>	U	4	0	U	1
63 <i>T. (A.) crassus</i>	3	4	1	5	1
64 <i>T. (A.) cuii</i>	3	4	1	U	2
65 <i>T. (A.) dactylifera</i>	3	4	1	4	1
66 <i>T. (A.) dalfardicus</i>	2	4	1	5	2
67 <i>T. (A.) daliu</i>	2	4	1	U	1
68 <i>T. (A.) daresalaami</i>	2	3	1	4	1
69 <i>T. (A.) darjeelingensis</i>	3	4	1	U	1
70 <i>T. (A.) dasiphorae</i>	2	3	0	U	3
71 <i>T. (A.) datongensis</i>	3	4	1	U	1
72 <i>T. (A.) deleari</i>	2	4	1	U	1
73 <i>T. (A.) denarus</i>	3	4	1	5	1
74 <i>T. (A.) denheyeri</i>	3	4	1	5	1
75 <i>T. (A.) denmarki</i>	2	4	1	U	4
76 <i>T. (A.) diumbokus</i>	3	3	0	U	3
77 <i>T. (A.) divergens</i>	3	4	1	4	1
78 <i>T. (A.) doreanae</i>	2	4	1	U	2
79 <i>T. (A.) dossei</i>	2	4	1	5	2
80 <i>T. (A.) drori</i>	2	4	0	4	3
81 <i>T. (A.) drymis</i>	2	4	1	5	1
82 <i>T. (A.) egypticus</i>	3	4	1	U	1
83 <i>T. (A.) elaeis</i>	3	4	1	5	1
84 <i>T. (A.) elmassri</i>	3	4	0	3	2
85 <i>T. (A.) eremicus</i>	3	3	1	5	3
86 <i>T. (A.) eremitidis</i>	3	4	1	U	2
87 <i>T. (A.) evectus</i>	2	4	1	U	1
88 <i>T. (A.) februs</i>	2	4	1	5	1
89 <i>T. (A.) fleschneri</i>	3	4	1	U	1
90 <i>T. (A.) foenilis</i>	2	4	0	5	2
91 <i>T. (A.) foraminosus</i>	2	3	1	U	3
92 <i>T. (A.) fujianensis</i>	2	4	1	U	1
93 <i>T. (A.) galpinii</i>	2	4	1	5	1
94 <i>T. (A.) galummatus</i>	3	4	1	3	1
95 <i>T. (A.) gameilus</i>	2	4	1	U	1
96 <i>T. (A.) gardeniae</i>	2	4	1	5	1
97 <i>T. (A.) garhwalicus</i>	3	4	1	U	1
98 <i>T. (A.) georgicus</i>	3	4	1	5	U
99 <i>T. (A.) ghanaensis</i>	U	4	1	5	1
100 <i>T. (A.) ghani</i>	2	4	1	U	1
101 <i>T. (A.) gopali</i>	3	4	1	5	1
102 <i>T. (A.) gouaniae</i>	3	4	1	5	1
103 <i>T. (A.) gracilentus</i>	3	4	1	U	1
104 <i>T. (A.) grastis</i>	2	4	1	5	1
105 <i>T. (A.) gressitti</i>	3	4	1	U	1
106 <i>T. (A.) grewiae</i>	U	3	1	5	1
107 <i>T. (A.) guangdongensis</i>	3	4	1	U	1
108 <i>T. (A.) guangxiensis</i>	3	4	1	U	1
109 <i>T. (A.) gulingensis</i>	3	4	0	U	1
110 <i>T. (A.) gutierrezii</i>	3	4	1	5	1
111 <i>T. (A.) hadii</i>	3	4	1	U	3
112 <i>T. (A.) hadzhievi</i>	U	4	0	5	2
113 <i>T. (A.) haiastanius</i>	2	3	1	U	4
114 <i>T. (A.) halinae</i>	2	4	1	5	2
115 <i>T. (A.) haramotoi</i>	3	4	1	U	1
116 <i>T. (A.) hartlandrowei</i>	2	3	1	5	2
117 <i>T. (A.) hebetis</i>	2	3	1	U	2
118 <i>T. (A.) hibernus</i>	3	4	1	5	2
119 <i>T. (A.) higoensis</i>	3	4	1	5	1
120 <i>T. (A.) himalayensis</i>	3	4	0	U	1
121 <i>T. (A.) hirashimai</i>	3	4	1	5	1
122 <i>T. (A.) homalii</i>	U	4	1	U	1

species	characters				
	I	II	III	IV	V
123 <i>T. (A.) hui</i>	3	4	1	U	1
124 <i>T. (A.) hungaricus</i>	U	4	0	U	3
125 <i>T. (A.) ilehoensis</i>	2	4	1	U	1
126 <i>T. (A.) ilicis</i>	2	2	0	U	3
127 <i>T. (A.) incasus</i>	3	4	1	3	1
128 <i>T. (A.) incertus</i>	U	U	U	5	3
129 <i>T. (A.) incisivus</i>	2	4	1	5	1
130 <i>T. (A.) inopinatus</i>	2	4	1	3	3
131 <i>T. (A.) inops</i>	2	4	0	U	1
132 <i>T. (A.) insularis</i>	3	4	1	U	1
133 <i>T. (A.) intercalaris</i>	2	3	1	3	4
134 <i>T. (A.) intermedius</i>	2	4	1	U	1
135 <i>T. (A.) invectus</i>	2	3	1	3	4
136 <i>T. (A.) involutus</i>	2	4	1	5	1
137 <i>T. (A.) iranensis</i>	2	4	0	3	4
138 <i>T. (A.) johannae</i>	2	4	1	5	1
139 <i>T. (A.) jordani</i>	2	4	0	5	3
140 <i>T. (A.) kadonoi</i>	2	4	1	5	3
141 <i>T. (A.) kazachstanicus</i>	2	4	1	3	2
142 <i>T. (A.) kazimiae</i>	2	4	1	5	1
143 <i>T. (A.) kenya</i>	U	4	1	5	1
144 <i>T. (A.) kerkirae</i>	2	4	1	5	1
145 <i>T. (A.) khosrovensis</i>	2	4	0	4	4
146 <i>T. (A.) kiambuensis</i>	3	4	1	U	1
147 <i>T. (A.) kikuyuensis</i>	3	4	1	5	2
148 <i>T. (A.) kirinyagae</i>	2	4	1	U	1
149 <i>T. (A.) kiso</i>	3	4	1	5	1
150 <i>T. (A.) kodaikanalensis</i>	3	4	1	U	1
151 <i>T. (A.) kolodochkai</i>	2	3	0	3	4
152 <i>T. (A.) kimbasi</i>	U	4	1	5	1
153 <i>T. (A.) kutabus</i>	3	3	1	5	2
154 <i>T. (A.) kuznetsovi</i>	2	4	1	4	1
155 <i>T. (A.) lazariensis</i>	3	4	1	5	2
156 <i>T. (A.) lanyuensis</i>	3	4	1	U	1
157 <i>T. (A.) lataniae</i>	3	4	0	U	1
158 <i>T. (A.) lateris</i>	3	4	U	U	1
159 <i>T. (A.) leilaseius</i>	2	4	1	U	1
160 <i>T. (A.) libitus</i>	3	4	1	5	1
161 <i>T. (A.) limitatus</i>	U	U	U	U	1
162 <i>T. (A.) limurus</i>	2	4	1	U	1
163 <i>T. (A.) linzhiensis</i>	2	4	1	U	1
164 <i>T. (A.) lobatus</i>	3	4	1	5	1
165 <i>T. (A.) longa</i>	2	4	1	U	1
166 <i>T. (A.) longicervix</i>	3	4	1	U	1
167 <i>T. (A.) lootsi</i>	2	4	0	5	1
168 <i>T. (A.) loralaiana</i>	2	4	1	U	1
169 <i>T. (A.) lushanensis</i>	3	4	1	3	1
170 <i>T. (A.) luzonensis</i>	2	4	1	2	1
171 <i>T. (A.) machaon</i>	2	4	1	U	1
172 <i>T. (A.) macroides</i>	3	4	1	5	1
173 <i>T. (A.) macrum</i>	U	U	U	U	U
174 <i>T. (A.) majumderi</i>	3	3	1	U	1
175 <i>T. (A.) makuyus</i>	2	4	1	5	1
176 <i>T. (A.) malawiensis</i>	2	4	1	U	1
177 <i>T. (A.) malicolus</i>	2	4	1	5	2
178 <i>T. (A.) mangiferus</i>	3	4	1	4	1
179 <i>T. (A.) manipurensis</i>	3	4	1	2	1
180 <i>T. (A.) maracus</i>	3	4	1	U	1
181 <i>T. (A.) marinus</i>	3	4	1	5	1
182 <i>T. (A.) masai</i>	2	4	1	U	1
183 <i>T. (A.) maspalomensis</i>	2	4	0	5	2

species	characters				
	I	II	III	IV	V
184 <i>T. (A.) mathyssei</i>	2	4	1	5	1
185 <i>T. (A.) meerutensis</i>	2	4	1	U	U
186 <i>T. (A.) meritus</i>	3	4	1	5	1
187 <i>T. (A.) mesasiaticus</i>	2	3	0	3	U
188 <i>T. (A.) michaeli</i>	3	4	1	5	1
189 <i>T. (A.) microbullatus</i>	2	4	1	5	1
190 <i>T. (A.) miyarai</i>	3	4	1	U	1
191 <i>T. (A.) monosetus</i>	3	2	0	U	1
192 <i>T. (A.) montanus</i>	2	4	1	4	4
193 <i>T. (A.) mori</i>	3	4	1	U	1
194 <i>T. (A.) muliebris</i>	3	4	1	5	1
195 <i>T. (A.) namaquaensis</i>	2	4	0	5	1
196 <i>T. (A.) ndibu</i>	3	4	1	4	1
197 <i>T. (A.) neobakeri</i>	2	4	1	U	1
198 <i>T. (A.) neoclastrus</i>	2	4	1	U	1
199 <i>T. (A.) neocrassus</i>	2	4	0	U	1
200 <i>T. (A.) neogutierrezii</i>	3	4	1	5	1
201 <i>T. (A.) neohartlandrowei</i>	2	3	1	5	2
202 <i>T. (A.) neomichaeli</i>	2	4	1	5	1
203 <i>T. (A.) neodibu</i>	2	4	1	U	1
204 <i>T. (A.) neorhenanus</i>	3	4	1	U	U
205 <i>T. (A.) neoterrulentis</i>	3	4	1	5	1
206 <i>T. (A.) neotransvaalensis</i>	3	4	1	U	1
207 <i>T. (A.) neyshabouris</i>	2	4	1	3	4
208 <i>T. (A.) nilgiriensis</i>	3	4	1	U	1
209 <i>T. (A.) njorogoeus</i>	3	4	1	U	1
210 <i>T. (A.) nobilis</i>	2	4	1	4	2
211 <i>T. (A.) noonus</i>	3	2	1	6	3
212 <i>T. (A.) nyahururus</i>	2	4	1	U	1
213 <i>T. (A.) oasis</i>	3	4	0	5	1
214 <i>T. (A.) obesus</i>	2	4	1	U	1
215 <i>T. (A.) occidius</i>	U	4	1	5	1
216 <i>T. (A.) octavus</i>	3	4	1	3	1
217 <i>T. (A.) ordinatus</i>	2	3	1	3	4
218 <i>T. (A.) orientalis</i>	3	4	1	U	1
219 <i>T. (A.) orissaensis</i>	3	4	0	U	U
220 <i>T. (A.) ornatulus</i>	3	4	1	U	1
221 <i>T. (A.) ornatus</i>	2	4	1	U	1
222 <i>T. (A.) paganus</i>	2	4	1	5	1
223 <i>T. (A.) paraevectus</i>	2	4	1	4	1
224 <i>T. (A.) parinopinatus</i>	U	4	1	3	3
225 <i>T. (A.) pegazzani</i>	2	4	1	5	3
226 <i>T. (A.) persianus</i>	2	4	1	5	1
227 <i>T. (A.) persicus</i>	3	4	1	U	U
228 <i>T. (A.) philippinensis</i>	3	4	1	5	1
229 <i>T. (A.) pineus</i>	3	4	1	U	1
230 <i>T. (A.) pirianykae</i>	2	4	1	5	1
231 <i>T. (A.) platycladus</i>	2	4	1	U	1
232 <i>T. (A.) ponticus</i>	2	4	1	5	1
233 <i>T. (A.) porathi</i>	3	4	0	4	2
234 <i>T. (A.) porus</i>	3	4	1	5	1
235 <i>T. (A.) povtari</i>	2	2	0	U	1
236 <i>T. (A.) praeacutus</i>	2	4	1	5	1
237 <i>T. (A.) pruni</i>	3	4	1	7	1
238 <i>T. (A.) pseudoserrulatus</i>	3	4	1	U	1
239 <i>T. (A.) psyllakisi</i>	2	4	1	4	U
240 <i>T. (A.) qianshanensis</i>	3	4	1	U	1
241 <i>T. (A.) quadratoides</i>	3	4	1	U	1
242 <i>T. (A.) quadratus</i>	2	4	1	U	1
243 <i>T. (A.) rapidus</i>	U	4	1	5	1
244 <i>T. (A.) rarus</i>	2	4	1	4	3

species	characters				
	I	II	III	IV	V
245 <i>T. (A.) rasilis</i>	2	4	1	5	1
246 <i>T. (A.) recki</i>	2	4	0	3	3
247 <i>T. (A.) religiosus</i>	3	4	1	5	1
248 <i>T. (A.) repens</i>	2	4	1	U	3
249 <i>T. (A.) rhenanoides</i>	2	4	1	5	2
250 <i>T. (A.) rhenanus</i>	2	4	1	5	3
251 <i>T. (A.) rhododendroni</i>	3	4	1	U	1
252 <i>T. (A.) ribei</i>	U	U	U	U	U
253 <i>T. (A.) richteri Karg</i>	U	4	1	4	1
254 <i>T. (A.) rickeri</i>	3	4	1	U	2
255 <i>T. (A.) rivulus</i>	U	4	1	U	1
256 <i>T. (A.) rodriguezii</i>	2	3	0	3	4
257 <i>T. (A.) rubetum</i>	U	U	U	U	U
258 <i>T. (A.) ruiro</i>	2	4	1	5	1
259 <i>T. (A.) ryukyuensis</i>	2	4	1	U	1
260 <i>T. (A.) saevus</i>	2	4	1	5	1
261 <i>T. (A.) salviae</i>	2	4	1	5	1
262 <i>T. (A.) samliensis</i>	3	4	1	U	1
263 <i>T. (A.) sapiens</i>	2	4	0	3	4
264 <i>T. (A.) serratosus</i>	3	3	1	5	1
265 <i>T. (A.) serratus</i>	2	4	1	1	1
266 <i>T. (A.) serrulatus</i>	3	4	1	5	1
267 <i>T. (A.) shibai</i>	2	4	0	5	2
268 <i>T. (A.) sica</i>	3	4	1	4	1
269 <i>T. (A.) sichirei</i>	3	4	1	U	1
270 <i>T. (A.) sijiensis</i>	3	4	0	U	1
271 <i>T. (A.) silvanus</i>	3	4	1	5	1
272 <i>T. (A.) singularis</i>	3	3	1	3	1
273 <i>T. (A.) sonprayagensis</i>	3	4	0	U	1
274 <i>T. (A.) spectatus</i>	2	4	1	U	3
275 <i>T. (A.) spiralis</i>	2	4	1	5	1
276 <i>T. (A.) subarcticus</i>	2	4	1	5	2
277 <i>T. (A.) subequalis</i>	2	4	0	U	2
278 <i>T. (A.) submarinus</i>	3	4	1	U	1
279 <i>T. (A.) subtilis</i>	3	4	1	5	2
280 <i>T. (A.) sudanicus</i>	3	4	1	U	1
281 <i>T. (A.) suecicus</i>	2	3	1	U	U
282 <i>T. (A.) sycomorus</i>	3	4	0	U	1
283 <i>T. (A.) taishanensis</i>	3	4	1	5	2
284 <i>T. (A.) tamaricis</i>	2	4	0	5	3
285 <i>T. (A.) tardus</i>	2	3	0	4	3
286 <i>T. (A.) tecoma</i>	2	4	0	1	1
287 <i>T. (A.) tenuis</i>	U	4	1	5	1
288 <i>T. (A.) ternatus</i>	2	3	1	3	3
289 <i>T. (A.) terrulentis</i>	2	4	1	5	1
290 <i>T. (A.) thailandicus</i>	3	4	1	5	1
291 <i>T. (A.) theroni</i>	2	4	1	5	1
292 <i>T. (A.) thesbites</i>	2	4	1	5	1
293 <i>T. (A.) torbatejamae</i>	2	4	0	3	4
294 <i>T. (A.) totifolianensis</i>	2	4	1	4	1
295 <i>T. (A.) transvaalensis</i>	2	3	1	4	3
296 <i>T. (A.) tridentiger</i>	3	4	1	U	1
297 <i>T. (A.) ulmi</i>	2	4	0	4	3
298 <i>T. (A.) umbraculus</i>	2	4	1	5	1
299 <i>T. (A.) umbratus</i>	3	4	1	6	1
300 <i>T. (A.) verbena</i>	3	3	0	4	1
301 <i>T. (A.) vascus</i>	3	4	1	5	1
302 <i>T. (A.) viniferae</i>	2	4	1	3	1
303 <i>T. (A.) votivus</i>	3	4	1	4	1
304 <i>T. (A.) vulgaris</i>	3	4	1	5	1
305 <i>T. (A.) wainsteini</i>	2	4	1	5	2

species	characters				
	I	II	III	IV	V
306 <i>T. (A.) werneri</i>	2	4	0	5	2
307 <i>T. (A.) wonkooi</i>	2	4	1	5	1
308 <i>T. (A.) wrenschae</i>	2	4	1	5	1
309 <i>T. (A.) xingchengensis</i>	2	4	1	U	U
310 <i>T. (A.) xini</i>	2	3	0	U	1
311 <i>T. (A.) xinjiangensis</i>	3	3	1	U	4
312 <i>T. (A.) xiufui</i>	3	4	1	U	1
313 <i>T. (A.) xizangensis</i>	3	4	1	U	1
314 <i>T. (A.) yamashitai</i>	3	4	1	3	1
315 <i>T. (A.) yasminae</i>	2	4	1	5	2
316 <i>T. (A.) yasumatsui</i>	2	4	1	U	1
317 <i>T. (A.) yinchuanensis</i>	2	4	1	U	1
318 <i>T. (A.) zafari</i>	U	4	1	1	1
319 <i>T. (A.) zhangyensis</i>	2	4	1	U	4
320 <i>T. (A.) zhaoi</i>	3	4	1	U	1

