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Oribatid mites (Acari: Oribatida) from the Sella massif (Dolomites, Trentino, Italy) with description of *Trichoribates valeriae* n. sp. (Ceratozetidae)

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Original research

ABSTRACT

Results from collections of oribatid mites in the cushion plant belt of Sass Pordoi and Piz Boè in the Italian Dolomites (Prov. Trento) are presented. A total of 10 oribatid species belonging to seven families were found. Most have a wide general distribution, *Anachipteria shtanchaevae* is restricted to the Alps. Three species are frequently or mainly found in the montane to alpine zone (*A. shtanchaevae, Camisia horrida, Oribatula interrupta*). Remarkable findings are *Camisia foveolata*, hitherto only known from the boreal climate zone, which indicates a preglacial relict distribution, and *Kunstidamaeus lengersdorfi* as highest recorded altitude for this species. *Trichoribates valeriae* n. sp., found in large numbers on both mountains, is described and compared with other congeners, and its systematic position is discussed. This species is characterized by a lamellar structure with small ridges and widely separated cusps, rounded rostrum with nose-like protuberance, 10 pairs of medium long notogastral setae with short bristles, notogastral porose area A1 divided in two parts, porose areas present on all tibiae and tarsi. Additional morphological information on the presence of porose areas on tarsi of *Trichoribates scilierensis* Bayartogtokh and Schatz, 2008 is given.

Keywords oribatid mites; faunistics; new records; high mountain region; preglacial relict; morphology; new species

Zoobank http://zoobank.org/52425B43-57A2-40C7-A407-9425E7606898

Introduction

The Sella group is a massif in the Dolomites, Northern Italy. The highest elevation is Piz Boè (3152 m a.s.l., Fig. 1A) at the intersection of South Tyrol, Trentino, and Veneto (Prov. Belluno). The Sass Pordoi (2950 m a.s.l., Fig. 1B), a plateau-shaped summit, is situated next to and west of Piz Boè. Both mountains are very popular touristic destinations, a funicular from the Pordoi Pass reaches the summit of Sass Pordoi. The isolated mountain ranges are fossil coral reefs of a Middle Triassic carbonate platform from the Raihl formation. It preserves fossils dating back to the Norian Middle Triassic sub-period during the Mesozoic Era (Reithofer 1928, Moroder 2008). Only small vegetation patches are scattered on karstic limestone.

During two excursions in 2017 soil and litter samples were taken on Sass Pordoi and the Piz Boè massif. A commented list of the found oribatid mite species is presented, and a new species to science is described. Results of sampling at the nearby Sella Pass on the edge of the Langkofel group (Gruppo di Sassolungo) are already published (Schatz 2017).
Materials and methods

Material examined

Figure 1

The material was collected during two excursions of the author and Irene Schatz on Sass Pordoi (2932 – 2940 m a.s.l.) and Piz Boè (2980 – 3150 m a.s.l.). A total of 10 soil and litter samples (each ca. 10 x 10 cm, volume ca. 0.5 liter) were taken in vegetation patches within the cushion plant zone, keeping the environmental impact as low as possible. The samples were extracted for 12 days with moderate heating with light, preservation fluid was 75% ethanol. Morphological investigations were carried out in temporary slides in lactic acid.

List of samples

- TN 157: 20. June 2017: Sass Pordoi, near cable car station, water logged with snowmelt, wet moss cushion (46°30.218′N, 11°48.270′E, 2935 m a.s.l., Fig. 1B).
- TN 159: 20. June 2017: Sass Pordoi, ibid., dry Cerastium uniflorum cushion from preceding year (46°30.218′N, 11°48.270′E, 2935 m a.s.l., Fig. 1F).
- TN 162: 19. July 2017: Piz Boè, southeast of summit, moist moss on rock (46°30.555′N, 11°49.697′E, 3149 m a.s.l., Fig. 1C).
- TN 163: 19. July 2017: Piz Boè, ibid., moist Cerastium uniflorum cushion (46°30.555′N, 11°49.697′E, 3150 m a.s.l., Fig. 1D).
- TN 164: 19. July 2017: Piz Boè, base of summit area, dry to moist Cerastium uniflorum cushion with humus and sand (46°30.383′N, 11°49.522′E, 2980 m a.s.l., Fig 1E).
- TN 165: 19. July 2017: Piz Boè, ibid., dry to moist Saxifraga oppositifolia cushion with moss and humus (46°30.383′N, 11°49.522′E, 2980 m a.s.l., Fig. 1E).
- TN 166: 19. July 2017: Sass Pordoi, near cable car station, below water logged (near TN157), moist to wet Carex grass cushion with humus (46°30.068′N, 11°48.455′E, 2932 m a.s.l., Fig. 1B).

Identification

Measurement of each parameter was done from the particular optimal parallax-free perspective. The total body length was measured in lateral view, from tip of the rostrum to the posterior edge of the notogaster, body width as maximal width of notogaster from dorsal view (without pteromorphs). Indicated are means, range in brackets. All measurements are given in micrometers (µm). The systematic arrangement follows Schatz et al. (2011), the terminology of morphological features that of F. Grandjean (summarized by Travé et al. 1996, Norton and Behan-Pelletier 2009), in the descriptive part of the Trichoribates species also that of Behan-Pelletier and Ermilov (2019).

Following abbreviations are used: L—length, W—width, no—nose-like protuberance on rostrum, ro, le, il, bo, ex—rostral, lamellar, interlamellar, bothridial, exobothridial seta, tu—tutorium, gt—genal tooth, pt I, pt II—pedotectum I, II, Ad—dorsal porose area, Am,
Figure 1 Sampling locality of *Trichoribates valeriae* n. sp. (all fotos Irene Schatz, Figs 1A – 1E: 19. July 2017, Fig. 1F: 20. June 2017). A – Piz Boè (3152 m a.s.l.) from opposing Sass Pordoi. B – Vegetation patches on Sass Pordoi (samples TN 157 and TN 166 were taken in this area). C – Piz Boè, moss (sample TN 162). D – Piz Boè, *Cerastium uniflorum* with moss (sample TN 163). E – Boz Boè, *Cerastium uniflorum* and *Saxifraga oppositifolia* (samples TN 164, 165). F – Sass Pordoi, dry *Cerastium uniflorum* cushion from preceding year (sample TN 159).
Ah, Al—anterior dorsolateral porose areas, Dp—dorsophragma, Pp—pleurophragma, len—lenticulus, c1-3, d1-3, e1-2, f1-3, ia, im, ip, dm, dp, h1-3, p1-3—notogastral seta, Aa, A1, A2, A3—notogastral porose areas, ap—postanal porose area, ia, im, ih, ip, ips—notogastral lyrifissures, gla—opisthontal gland opening, opisthontal gland region in immatures, cus—custodium, dis—discidium, cp—circumpedal carina, 1a-c, 2a, 3a-c, 4a-c—epimeral setae, adl-3—adanal setae, po—preanal organ, iad—adanal lyrifissure, ch, chb—cheliceral setae, σ, φ, φ1, φ2, ω, ω1, ω2—solenidia on genu, tibia, tarsus, pa—porose area on leg, PY—pygidial sclerite, ho—humeral organ.

Results

Species list

Family Brachychthoniidae Thor, 1934

Taxonomical notes — The three Liochthonius species found on Biz Boè are part of the “Lapponicus-group” (Moritz 1976) with common features as double-pointed sensillus, small velum on relatively short dorsal setae, and within this group they are separated from other members in having setae c1 – c1 slightly more distant than d1 – d1. They were distinguished by following characters:

• L. strenzkei: with small transverse crest anteriorly of setae le, notogastral setae fl, h1 on separate tubercles (not always easily visible).

• L. lapponicus: without tubercles on pygidium, without transverse crests near setae le.

• L. sellnicki: with transverse crests anteriorly and between setae le, with transverse crests posterior of setae fl, h1.

Liochthonius lapponicus (Trägårdh, 1910)


General distribution — Holarctic. Recorded also at nearby Sella Pass (Schatz 2017).

Liochthonius sellnicki (Thor, 1930)


Piz Boè — TN 165: numerous specimens.

General distribution — Holarctic, Oriental (China).

Liochthonius strenzkei Forsslund, 1963

Moritz 1976: p. 80, fig. 18 a, b. Weigmann 2006: p. 78, fig. 41 a, b.


Piz Boè — TN 162, TN 163, TN 164, TN 165: numerous specimens in each sample.

General distribution — Holarctic, Oriental (China). Recorded also at nearby Sella Pass (Schatz 2017).

Family Crotoniidae Thorell, 1876

Camisia foveolata Hammer, 1955


**Remarks** — The specimens from Sass Pordoi correspond widely to the redescription by Colloff (1993) and are considered conspecific. The juvenile instars correspond to the morphological description by Seniczak (1991b). Some variabilities are noteworthy:

- shape of setae *le*: frequently bent mediad in different intensity, but also almost straight in some specimens, length 30–40 (vs. Colloff 1993 “not curved into arch”, and Hammer 1955). Seniczak (1991b) illustrated setae *le* also bent mediad.

- thick and continuous transversal line between the apophyses of setae *le* as in Hammer (1955) (vs. weak or interrupted in Colloff 1993 and Seniczak 1991b).

- microsculpture between prodorsal ridges fine, punctate, forming small irregular lines (vs. areolate/scalloped in other descriptions).

- The asymmetrical number of genital setae as mentioned from Seniczak (1991b) could also be observed in some specimens of protonymphs, tritonymphs and adults.

This mosaic of differences of morphological characters in far distant populations do not justify the status of a separate species for the population in the Dolomites.

**General distribution** — Northern Holarctic (Alaska, Yukon, Greenland, Svalbard, Scandinavia), Chilenian Highland. New record for Central Europe, the Alps and Italy.

**Camisia horrida** (Hermann, 1804)


**Piz Boè** — TN 164: 2 adults, 1 protonymph, 2 deutonymphs, TN 165: 3 adult specimens.

**Taxonomical notes** — The juvenile instars correspond to the description by Seniczak (1991a) and were allocated to the respective instars according to body size and number of genital setae (protonymph — size 470 – 360, 1 pair of genital setae, deutonymph — size 540 – 550 x 310 – 330, 4 pairs of genital setae).

**General distribution** — Holarctic, Oriental, Ethiopian, Neotropical (Central America).

**Remarks** — Schweizer (1956) reported *C. horrida* frequently from the alpine zone in Switzerland up to 3109 m a.s.l. and considered it as a boreo-alpine species. In South Tyrol it was already found in the Dolomites in the nearby Schlern/Sciliar massif at 2200–2560 m a.s.l. (Schatz 2008a), and in the North-Tyrolean Central Alps (Austria) in the alpine grassland up to 2650 m a.s.l. (Schatz 1979). *Camisia horrida* was also reported from higher altitudes of Central America (Costa Rica: Volcán Irazú, 3400 m, Volcán Chirripó, 3800 m; Panama: Volcán Barú, 3400–3475 m a.s.l., Schatz 2006).

**Family Damaeidae Berlese, 1896**

**Kunstidamaeus lengersdorfi** (Willmann, 1932)


**Piz Boè** — TN 165: 3 adult specimens.

Remarks — The specimens from Piz Boè correspond to the redescription by Miko and Mourek (2008) and are considered conspecific. The relatively short distal parts of spinae adnatae vary. Willmann (1932, fig. 4) drew spinae adnatae with long distal part, apex slightly bent anteriad (“lange, geschweifte Spitze”). In Weigmann (2006, fig. 100a) the spinae adnatae also show a long distal part which is curved outward and directed more than 100° posteriad. In specimens from Harz (Miko and Mourek 2008, fig. 1A) the spinae adnatae are similarly bent as in the figure of Weigmann, in specimens from Slovak Karst) Miko and Mourek (op. cit.), fig. 2A) the spinae adnatae show a short distal part and are bent laterad to different degrees.

General distribution — Central Europe, frequently in caves. New record for the Dolomites and Italy.

Family Tectocepheidae Grandjean, 1954

Tectocepheus sarekensis (Trägårdh, 1910)


Sass Pordoi — TN160: 1 adult, TN 166: 1 protonymph.

Taxonomical and nomenclatorial notes — The specimens from the different samples on Sass Pordoi and Piz Boè show a mosaic of morphological characters between T. velatus and T. sarekensis. But they all are considered to be T. sarekensis based on characters given by Weigmann (2002, 2006) and Laumann et al. (2007) in having mainly broad and rounded cusp tip (with some exceptions), longitudinal striae on the interlamellar field (all specimens), and 2 – 4 pairs of notogastral depressions, the latter sometimes fainter (Table 1). The body length is slightly larger than in most other investigations (250 – 390 vs. 302 – 349 – Laumann et al. 2007, 312 – 374 – Pérez-Íñigo 1997, 295 – 362 – Weigmann 2002). The small granules of the specimen in TN 160 remind of T. alatus Berlese, 1913, but in that species the granules are larger, and small granules on the cutegument can be found in other Tectocepheus species too. Weigmann (2002, 2006), Subías (2004) and other authors list T. sarekensis as subspecies of T. velatus (Michael, 1880); I follow Laumann et al. (2007) in considering it a distinct species.

General distribution — cosmopolitan excl. Antarctic. Recorded also at nearby Sella Pass (Schatz 2017).

### Table 1 Tectocepheus sarekensis, variability of some morphological characters in adults within and between populations on Sass Pordoi and Piz Boè (Sella massif, Dolomites, Italy).

<table>
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<th>TN 163</th>
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<tr>
<td>(µm) (min – x̅ – max)</td>
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<td>235</td>
<td>226 (180–250)</td>
<td>224 (205–240)</td>
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<td>cuspis tip</td>
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<td>interlamellar field</td>
<td>broad, round longitudinal striae, body with small granules</td>
<td>smaller longitudinal striae present</td>
<td>most broad, round all with longitudinal striae</td>
<td>most broad, round all with longitudinal striae</td>
</tr>
<tr>
<td>pairs of notogastral depressions</td>
<td>4, well visible</td>
<td>3, 3rd weakly expressed</td>
<td>2, 3–4, mainly 3, all or 3rd sometimes weakly expressed</td>
<td>2–3, mainly 3, 3rd sometimes weakly expressed</td>
</tr>
</tbody>
</table>

Family Achipteriidae Thor, 1929

*Anachipteria shtanchaevae* Subías, 2009


- **Piz Boè** — TN 164: 26 adults, TN 165: 12 adults, juvenile instars in both samples.

**Nomenclatorial notes** — New name for *Oribata tecta alpina* Schweizer, 1922 nec Halbert.

**General distribution** — Alps, Central, South, Southeast Europe. Mainly in montane to alpine habitats.

Family Oribatulidae Thor, 1929

*Oribatula interrupta* (Willmann, 1939)


- **Piz Boè** — TN 164: 46 adults, TN 165: 340 adults, numerous juvenile instars in both samples.

**General distribution** — Holarctic, Ethiopian. In the Alps frequently in montane to alpine habitats. Recorded also at nearby Sella Pass (Schatz 2017).

Family Ceratozetidae Jacot, 1925

*Trichoribates valeriae* n. sp.

*Zoobank:* 5E466591-43EA-4CCA-806C-225A81E7608F


**Diagnosis**

*Trichoribates valeriae* n. sp. differs morphologically from other *Trichoribates* species by the following combination of characters: Total length of males 550–620 µm, females 595–670 µm, rostrum rounded with nose-like protuberance, lamella and translamella small ridges, cusps relatively wide apart, teeth on cusp minute or absent, lamellar and interlamellar setae long, setiform with small bristles, bothridial seta with small club-shaped head, tutorium large, anterior notogastral tectum present, lenticulus indistinct, triangular, 10 pairs of medium long notogastral setae, setiform with small bristles, four pairs of round or oval notogastral porose areas, *A1* divided in two parts, epimeral setal formula: 3–1–3–3. Small porose areas present on tibiae and tarsi I–IV. Nymphs with large pygidial sclerite, humeral organ absent in larva, present in nymphs. Larva with 12 pairs of gastronotal setae, nymphs with 15 pairs each.

**Description**

**Adult**

Figures 2–7


**Integument** — Body colour light brown. Dorsal and ventral part of body finely punctured.

**Prodorsum (Figs 2–5A)** — Rostrum broadly rounded in dorsal view, rarely with slightly anteriad projecting central part, with nose-like protuberance dorsally (no, about 18–22 elevated above prodorsal shield). Below front edge of rostral tectum a transverse trilobate thickening, lateral lobes with round or hornlike protuberances. Lamella small ridge, translamella small, about twice length (~50) as cusp (~25), lamellarp cusp shorter than lamella, small, teeth beside...
root of le minute or absent. Rostral (80–100), lamellar (120–135) and interlamellar (150–180) setae setiform with small bristles, bristles on antiaxial part of rostral seta slightly longer (up to 3), basal part of rostral seta covered by tutorium in dorsal view. Tutorium (tu) large, broad, anterior tip pointed (rarely with two small teeth), antiaxial side of tutorium with rough longitudinally striae. Bothridium partially or completely covered by anterior notogastral margin, without medial or lateral scales. Bothridial seta with short stalk (25–30) and small spoon-shaped head (length 25–30, width ~15), with pointed apex and short spines, directed anterolaterad. Exobothridial seta (25–30) setiform with small bristles. Dorsosejugal porose area Ad large,
medial to bothridium under anterior notogastral tectum. Internal apodemes dorsophragmata (Dp) and pleurophragmata (Pp) distinct, dorsophragmata small and short, close to each other, pleurophragmata larger, posterior to bothridia, muscle fibers visible through integument.

**Notogaster (Figs 2, 4, 5)** — Anterior margin bent anteriad forming broad notogastral tectum between pteromorphs. Lenticulus indistinct, triangular. Pteromorphs without hinge, immovable, broadly rounded laterally. Ten pairs of notogastral setae, all setiform with small bristles, series c, l, h longer (80–100), p1-3 slightly shorter (60–80). Four pairs of porose areas, round to oval in shape, Aa largest (15–17 x 20–23), A1–A3 smaller (6–8 x 10–12), A1 divided in two parts, lateral area larger, medial smaller. Lyrifissures (ia, im, ip, ih, ips) small, distinct,
best visible in lateral view. Opisthonotal gland (gla) opening located laterad to seta lp.


Lateral aspect of podosoma, epimeral region (Figs. 3, 4, 5A) — Pedotectum I (Pt I) large scale, dorsally convex, anteriorly rounded, with small longitudinal striae. Pedotectum II (Pt II) small shell-shaped scale. Horizontal folds present in integument dorsal of acetabula II and III. Discidium (dis) well developed, triangular, projecting laterally. Circumpedal carina (cp) merging into custodium. Custodium (cus) short, pointed anteriad. Humeral porose areas Am, Ah, Al indistinct (latter only visible in dorsolateral view, see Fig. 5A, or in dissected specimens). Epimeral setae setiform with small bristles, epimeral setal formula 3–1–3–3. length 1a, 2a, 3a, 4c (originating on base of discidium) 18–23, 1b, 3b, 4a, 4b 25–30, 1c, 3c (distal of custodium) 40–45. Apodemes blade-like, well sclerotized.

Anogenital region (Figs 3, 4, 5B, 6C, 6D) — All genito-anal setae setiform with small bristles, genital plates with 6 pairs (14–18), 2 anterior pairs situated side by side on anterior thickening of genital plate, 1 pair of aggenital (27–30), 2 pairs of anal (20–23), 3 pairs of adanal setae (ad1 35, ad2, ad3 20–23). Adanal lyrifissures (iad) short, adjacent and parallel to anal plates, level with their anterior half. Preanal organ (po) cup-shaped.
Figure 5 *Trichoribates valeriae* n. sp., adult: A – anterior part of the body, dorsolateral view, showing porose areas *Ad, Am, Ah, Al*. B – posterior view. Scale bar 100 µm.
**Legs (Fig. 7)** — Legs of moderate length (42 to 50 % of body size), leg I (including claws) ~260, leg II ~270, leg III ~260, leg IV ~310. All legs tridactylous, lateral claws thinner than medial claw. All femora with ventral thickening, trochanter and femur IV with transverse wrinkles and strong bladelike ventral keel each. Genua I and II distally with ventral spine. Solenidia ω1 and ω2 on tarsi I almost equal in length (95–100), φ1 on tibia I longest (140–160), φ2 shorter (30–40), σ on genu I about same length (35–40). Setae l” on genua and tibiae I and II considerably thickened (40–45). Almost all setae with short bristles, setae (ρ) and (υ) with very short bristles, seta v on trochanter IV smooth. Setal formula of legs (trochanter to tarsus, solenidia in parentheses): leg I 1 – 5 – 3(1) – 4(2) – 20(2), leg II 1 – 5 – 3(1) – 4(1) – 15(2), leg III 2 – 2 – 1(1) – 3(1) – 15, leg IV 1 – 2 – 2 – 3(1) – 12. Position and length of setae as in Fig. 7. Large porose areas (pa) on all femora and on trochanters III, IV. Porose areas on legs proximoventrally on tarsi I–IV and distoventrally on tibiae I–IV, all small and roundish to elliptical in shape.
**Immatures**
Figures 8 and 9

**Measurements** — Table 2.

**Integument** — Cuticle pale yellowish in larva, darker greyish-yellow in nymphs. Pygidial sclerite (PY) micropunctate in all nymphal instars. Opisthontal gland region dark brown. Dorsal part of gastronotum without wrinkles, laterally with small furrows, ventrally with some furrows leading from position of setae la and lm posterior around anal plates in all immature instars, and around genital plates in nymphs.

**Prodorsum** — Rostrum broadly rounded. Prodorsal setae setiform with short bristles. Interlamellar setae longest in all instars. Bothridial seta (ho) with club-shaped head and short spines in all instars, size according to the size of the instar (Table 2).

**Gastronomic region** — Generally weakly sclerotized. Setae of series c without small sclerites in larva, setae of series c, ps2, ps3 on small sclerites in nymphs. Pygidial sclerite in larva indicated by lateral furrows posterior to seta c3. Nymphs with a large pygidial sclerite bearing setae of l, d, h series and p1 (latter on edge of the sclerite), anterior region with c series and opisthontal regions separated. Humeral organ (ho) absent in larva, in nymphs present as protruding papilliform cupule. Larva with 12 pairs of gastronotal setae (c, l, d, h series, seta h3 lateral to medial part of anal plates), nymphs with 15 pairs each (including p series), all
setiform with short bristles, most setae of similar length except shorter $h_2$, $h_3$ in larva as well as $h_2$ and $p$ series shorter in nymphs. Lyrifissures expressed as small cupules, $ia$, $im$ visible on dorsal side, others on ventral side. Lateral sclerite surrounding opisthonotal gland opening dark colored.

**Ventral region** — Ontogenetic development of epimeral / anogenital setae: larva 3–1–2 / 0–0–0, protonymph 3–1–2–1 / 1–0–0–0, deutonymph 3–1–2–2 / 3–1–0–3, tritonymph 3–1–3–3 / 5–1–2–3. Setae setiform, short, slightly barbed. In larva large scale in place of Claparède’s organ instead of epimeral seta $lc$. Length of genital plate in different nymphal instars see Table 2.

**Legs** — All legs monodactylyous with strong claws. Setal formula of legs (trochanter to tarsus, solenidia in parentheses): larva I 0–2–2(1)–3(1)–16(1), II 0–2–2(1)–2(1)–13(1), III 0–2–1(1)–1(1)–13, protonymph I 0–2–2(1)–3(1)–16(2), II 0–2–2(1)–2(1)–13(1), III 0–2–1(1)–1(1)–13, IV 0–0–0–0–7, deutonymph I 0–4–2(1)–3(2)–16(2), II 0–4–2(1)–3(1)–13(2), III 1–2–1(1)–2(1)–13, IV 0–2–2–1(1)–12, tritonymph I 1–4–3(1)–4(2)–18(2), II 1–4–3(1)–4(1)–15(2), III 2–2–1(1)–3(1)–15, IV 1–2–2–3(1)–12. Lengths of solenidion $\varphi l$ in different nymphal instars see Table 2.

**Sexual dimorphism, eggs**
Adult females slightly larger than males, ranges overlapping. Genital plates slightly larger
Figure 9 Trichoribates valeriae n. sp., larva: A – dorsal view. B – ventral view (legs not shown). Scale bar 50 µm.

Table 2 Size and lengths of immatures and adults (males and females separated), of genital plates, and lengths of selected body setae in ontogenetic instars of Trichoribates valeriae n. sp. Abbreviations of setae see text. All measurements, except for total length and width of females (N=42) and males (N=24) were taken in five specimens of each stage and gender. 1 total length, mean, range in brackets. 2 maximal width of gastronotum in immatures resp. notogaster in adults, mean, range in brackets. Other measurements – range.

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<th>tritonymph</th>
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<th>adult males</th>
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<td>width ²</td>
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<td>189 (175–200)</td>
<td>206 (190–220)</td>
<td>268 (250–280)</td>
<td>379 (350–410)</td>
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<tr>
<td>length genital plate</td>
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in females (length 70-80) than in males (length 55-60). Apart from these size differences, no external sexual dimorphism could be observed. Most females bearing 1–5 eggs, frequently arranged diagonally or transverse in the posterior part of the body. Shape of eggs oval to elliptical, frequently kidney-shaped, surface pustulate. Dimensions \((n=12)\) length 165 (140–185), width 72 (60–80). Among the 42 investigated females 17 specimens (40%) do not bear eggs, 4 (10%) bear one egg, 5 (12%) two, 5 (12%) three, 10 (24%) four, and 1 (2%) bears five eggs.

**Variation**

Some specimens (including holotype) have seta 4a doubled on one side.

**Type deposition**

The holotype (adult male, TN-163) and six paratypes (TN-163), preserved in ethanol, are deposited in the collection of the Senckenberg Museum, Görlitz, Germany (SMNG). Additional material is deposited in the collection of the author which will finally also be placed in the Senckenberg Museum, Görlitz. Specimens are preserved in ethanol.

**Etymology**

This species is named in honour of my friend and colleague Dr. Valerie M. Behan-Pelletier, who has extensively contributed to our knowledge of oribatid mites. She supported my work in many ways.

**Remarks**

1. Systematic position. The broad tectum on the anterior margin of the notogaster of the new species and the small banded lamellar complex remind of *Jugatala* Ewing, 1913.

   The generic diagnosis of *Jugatala* (Ewing 1913) underlines the “broad shelf-like expansion” (anterior notogastral tectum) as the main trait distinguishing *Jugatala* species from other genera “except some of the species of the genus Pelops C. L. Koch” (Ewing 1913, p. 131). Sellnick (1923) noted that this character also occurs in other genera as *Punctoribates* or *Trichoribates*, and, according to him, the genus *Jugatala* is generally very close to *Trichoribates*. A main difference between *Trichoribates* and *Jugatala* is the shape of the pedotectum I which is concave dorsally in *Jugatala*, but convex in *Trichoribates* (Behan-Pelletier 2000, pers. comm.). In *Jugatala angulata* (C. L. Koch, 1839) the pedotectum I is straight to slightly concave dorsally (Bayartogtokh and Schatz 2008a, additional observations on own material).

   The revised generic diagnosis of *Trichoribates* given by Behan-Pelletier and Ermilov (2019) lists a number of specific apomorphies for the genus which also pertain to *T. valeriae* n. sp.

   This concerns mainly: anterior of notogaster forming tectum, usually covering at least base of bothridium; pedotectum I convex dorsally; pteromorphs without unsclerotized band; octotaxic system with four pairs of porose areas: A1 is divided in two parts of different size; humeral porose areas Ad, Am, Ah, Al present; axillary saccule on mentum present; lateral horizontal folds present dorsal ofacetabula I and II; femur III with 2 setae (seta \(\ell\) absent); humeral organ absent in larva, present in nymphs; gastronotal region of nymphs with large pygidial sclerite; larva with 12 pairs of gastronotal setae, nymphs with 15 pairs, adult with ten pairs of notogastral setae (in adult setae c1, c3, da, dm, dp lost).

2. *Trichoribates valeriae* n. sp. shows morphological similarities to *Trichoribates biarea* Gjelstrup and Solhøy, 1994 in having lamellar cusps relatively wide apart, bothridium without sharp projection, anterior notogastral tectum, 10 pairs of notogastral setae, porose area A1 divided in two parts. They differ in the shape of rostrum (two apices laterally, without nose-like protuberance in *T. biarea*), lamellar structure (lamella, cusp and translamella wider, well-developed teeth on cusp of *T. biarea*), tutorium (4–6 dentations in *T. biarea*), porose area Aa (larger in *T. biarea*). *Trichoribates biarea* was described from Iceland (Gjelstrup and Solhøy 1994) and was also found in montane and subalpine habitats of the Allgäu Alps in Southern Germany (Beck et al. 2018).

   The new species is also morphologically similar to *Trichoribates scilierensis* Bayartogtokh and Schatz, 2008 in having a nose-like protuberance on prodorsum, lamellar cusps relatively wide apart, bothridium without sharp projection, bothridial setae relatively small, anterior margin of notogaster arched anteriad, 10 pairs of notogastral setae, porose area Aa large, A1...
divided in two parts. The species differ in lamellar structure (lamella and cusp wider, well-developed teeth on cusp of *T. scilierensis*), shape of tutorium (part of *T. scilierensis* specimens with 4–5 small dentations at the dorsodistal end, others without teeth and broader end), epimeral setal formula (3–1–3–2 in *T. scilierensis*). *Trichoribates scilierensis* was described from the nearby Sciliar massif in the Italian Dolomites (Bayartogtokh and Schatz 2008a) and later found in further localities in the Italian and Austrian Alps (Fischer and Schatz 2013, Schatz 2018, 2020), mainly in alpine and high alpine altitudes.

Among all *Trichoribates* species only *T. rausensis* Aoki, 1982 has a comparable small lamellar structure. This species differs from *T. valeriae n. sp.* primarily in having longer cusp with developed outer tooth, shorter notogastral setae, and undivided porose area A1. *Trichoribates rausensis* was described from mountainous areas in Japan (Aoki 1982) and was also found at the foot of Himalaya in West Bengal, India (Mondal and Kundu 1999).

3 ● The fine muscle fibers on pleurophragmata are hitherto only reported in *T. biarea* (Fig. 3, Gjelstrup and Solhøy 1994) and commented by Beck et al. (2018, p. 161, fig. 17) (“A conspicuous Strahlenfigur [radiating pattern] exists inside behind the front of the notogaster, presumably consisting of fibrillar muscle attachments on apodemal arches”). By checking other species of *Trichoribates* from my collection I found these muscle fibers invisible or only very weakly visible (e.g. in *T. scilierensis*).

4 ● The frontal view of the retracted male spermatopositor (Fig. 6C) resembles that of *Damaeus omnustus* (Fig. 2 in Grandjean 1956). Only few studies on the male genital morphology of oribatid mites have been conducted (e.g. Alberti and Coons 1999, Grandjean 1955, 1956, Warren 1947, Woodring 1970), a comparison with other species was not possible in this context. The ovipositor (Fig. 6D) matches other oribatid species. Among the numerous specimens of this species I found only one female with an everted ovipositor, just releasing an egg.

5 ● The presence of porose areas on tibiae and tarsi has not been studied in most *Trichoribates* species. Beside *T. valeriae n. sp.* only *T. sidorchukae* Behan-Pelletier and Ermilov, 2019 is known to have porose areas distoventrally on all tibiae. Porose areas on tarsi are also known from *T. novus* Sellnick, 1928, *T. sidorchukae*, *T. striatus* (Behan-Pelletier, 1986) (in these species proximoventrally on tarsi I–IV), and *T. zingerlei* Bayartogtokh and Schatz, 2008 (on tarsi II, III, see Bayartogtokh and Schatz, 2008b). As Behan-Pelletier and Ermilov (2019) stated these characters may be more widespread.

A reexamination of *Trichoribates scilierensis* Bayartogtokh and Schatz, 2008 (specimens from the Dolomites and different places in the Alps) revealed the presence of porose areas on tarsi I–IV, all small and roundish, posterior to seta v′ on tarsus I, posterior on seta pv ′′ on tarsi I–IV. In that species no porose areas exist on tibiae I – IV.


7 ● The setation of legs of immature *Trichoribates valeriae n. sp.* coincides with *T. berlesei* (Seniczak 1980, sub *T. trimaculatus*) and *T. polaris* (Behan-Pelletier 1985). Differences with *Diapterobates brevidentatus*, a well-studied species regarding immature (Bayartogtokh and Ermilov 2016), are in tibia II of protonymph (3(1), including 1′”), tibia I of deutonymph (4(2), including v′”), tibia II of deutonymph (4(1), including v′”).

8 ● *Trichoribates valeriae n. sp.* is the most frequently recorded species on both mountains of this investigation. It was found in large numbers in *Cerastium uniflorum* cushions and in moist moss on the plateau of Sass Pordoi and the Piz Boè mountain up to the summit (3150 m a.s.l.). The latter site was chosen as type locality.

9 ● Several *Trichoribates* species and other Ceratozetoidea were found in the high mountains of Europe, Asia, Central and South America or in the Arctic zone. Many seem to be restricted to the harsh environment with short vegetation periods and show adaptations such as cold hardness (Schatz and Sømme 1981) and prolongation of life cycle (e.g. Schatz 1985, Grishina 1997).
Discussion

A total of ten oribatid species from seven families were found (Table 3). The samples were generally very poor in species, on Sass Pordoi four species were recorded, on Piz Boè nine species. The richest spot in species number was a vegetation patch at the foot of Piz Boè (TN 164, TN 165, 2980 m a.s.l.) with nine species, among them six species found only in this site (Anachipteria shtanchaevae, Camisia horrida, Kunstidamaeus lengersdorfi, Liochthonius lapponicus, L. sellnicki, Oribatula interrupta). Three species were recorded on both mountains (Trichoribates valeriae, Liochthonius strenzkei, Tectocepheus sarekensis), all in more than one sample and most in relatively high abundances. Camisia foveolata was only found in moss on Sass Pordoi.

The individual numbers were not estimated in all samples, especially not in the samples which were rich in individuals, but the results indicate an extreme aggregation of species and individuals wherever a suitable microhabitat exists. Especially the small section at the foot of Piz Boè with cushions of Saxifraga oppositifolia, Cerastium uniflorum, moist moss and a dry to moist underlying humus layer (samples TN 164, TN 165) shows a surprising diversity of species with high abundances. This reflects a new assessment of a possible minimum area for mesofaunal elements (Schatz and Schatz 1991).

The majority of the oribatid species from this study have a wide general distribution (holarctic or semicosmopolitan – 7 spp.), one species is known from the Alps, Central, South, Southeast Europe (Anachipteria shtanchaevae), one species is hitherto only known from some sites in Central Europe (Kunstidamaeus lengersdorfi). Trichoribates valeriae is new to science, and until now its known distribution is restricted to the investigation area. Four species were also recorded at the nearby Sella Pass (Liochthonius lapponicus, L. strenzkei, Tectocepheus sarekensis, Oribatula interrupta, Schatz 2017). Three species are new records for the Dolomites and Italy (Camisia foveolata, Kunstidamaeus lengersdorfi; the newly discovered Trichoribates valeriae enlarges the new records for Italy).

These data are snapshots of the investigated vegetation patches. Additional species might possibly occur in the area, but due to conservation considerations only few samples were taken in the scattered vegetation. Despite the small yield, the species spectrum gives an interesting insight into the faunal distribution of high summits. Besides some ubiquitous species three species are frequently or mainly found in the montane to alpine zone (Camisia horrida, Anachipteria shtanchaevae, Oribatula interrupta). Trichoribates valeriae might be added to this “alpine” species group.

Camisia foveolata was hitherto only known from the boreal climate zone in the Northern Holarctic (Subias 2004) and from the Chilean Andean highlands (Covarrubias 2004). The

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finding of this species in the Dolomites lines up with species of disjunct boreo-alpine or arcto-alpine distribution and restriction to high altitudes in the Alps (some remarkable examples are Ceratozetes spitsbergensis Thor, 1934 – cf. Fischer et al. 2016, Mycobates sarekensis (Trägårdh, 1910) – cf. Schatz 2020). These species are considered as preglacial relicts with a wider geographical extension previous to the last glaciation. They survived in certain habitats and retreats such as unglaciated alpine summits (nunataks), in subterranean niches, or in ice-free massifs de refuge along the margins of the Alps, with special adaptations to the extreme conditions in life cycle and development of cold-hardiness (Schatz 2008a, b, Fischer et al. 2016). A corresponding explanation is assumed for high alpine endemic species which could have evolved in recent isolation (e.g. Trichoribates valeriae – present study, Kunstidamaeus granulatus (Willmann, 1951), Mycobates alpinus (Willmann, 1951), Oppiella obscura (Mahunka and Mahunka-Papp, 2000), Trichoribates scilierensis Bayartogtokh and Schatz, 2008, T. zingerlei Bayartogtokh and Schatz, 2008).

Another remarkable finding is Kunstidamaeus lengersdorfi which marks the highest recorded altitude for this species. Kunstidamaeus lengersdorfi is known to be troglobiontic or troglophilic (Miko and Mourek 2008), most records were reported from caves or cave entrances in different parts of Central Europe (Austria, Belgium, Czechia, Germany, Hungary, Slovakia). The Piz Boé massif is a carbonate platform with many smaller and larger synclines, sinkholes, overhangs, and caves. The locality of K. lengersdorfi is situated on a small flat and plant covered section beside an overhanging rock with temporary trickle joining the rank of similar known habitats of that species.

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References


