Mandibles and labrum-epipharynx of tiger beetles: basic structure and evolution (Coleoptera, Carabidae, Cicindelitae)

George E. Ball¹, John H. Acorn², Danny Shpeley¹

¹ Department of Biological Sciences, University of Alberta, Edmonton, Alberta, Canada T6G 2E9 ² Department of Renewable Resources, University of Alberta, Edmonton, Alberta, Canada T6G 2H1

Corresponding author: Danny Shpeley (dshpeley@ualberta.ca)

Abstract

Using for comparison with, and as outgroups for, supertribe Cicindelitae, we describe and illustrate the mandibles and labrum-epipharynx of the basal geadephagans Trachypachus gibbii LeConte, 1861 (family Trachypachidae), and family Carabidae: Pelophila rudis (LeConte, 1863) (supertribe Nebriitae, tribe Pelophilini) and Ceroglossus chilensis (Eschscholtz, 1829) (supertribe Carabitae, tribe Ceroglossini). The range and pattern of variation in structure of mandibles and labrum-epipharynx within the supertribe Cicindelitae was assessed using scanning-electron (SEM) images of these structures in nine exemplar taxa: Amblycheila baroni (Rivers, 1890), Omus californicus (Eschscholtz, 1829) and Picnochile fallaciosa (Chevrolat, 1854) (representing the Amblycheilini); Manticora tuberculata (DeGeer, 1778) (representing the Manticorini); Tetracha carolina (Linnaeus, 1767) (representing the Megacephalini); Pogonostoma chalybeum (Klug, 1835) (representing the Collyridini); and Therates basalis Dejean, 1826, Oxycheila species, and Cicindela longilabris Say, 1824 (representing the Cicindelini). An evolutionary transformation series was postulated for the mandibles and labrum-epipharynx, based on a reconstructed phylogenetic sequence, which, in turn, was based on morphological and DNA evidence. Principal features of the transformation series for the mandibles included development of a densely setose basal face; wide quadridentate retinaculum; a lengthened incisor tooth; a multidentate terebra (one to five teeth; two-three most frequent), followed by subsequent loss of one or more such teeth; development of a diastema in the occlusal surface; development and subsequent loss of scrobal setae, and reduction and loss of the scrobe. Principal features of the transformation series for the labrum included evolution of form from transverse, sub-rectangular to elongate almost square, to triangular; position and number of setae evolved from dorsal to insertion on the apical margin, the number increased from 8–10 to as many as 36, and decreased to as few as four. The
epipharynx broadened evolutionarily, the pedium evolving in form from narrow, triangular and nearly flat, to broad, palatiform, and markedly convex; anterior parapedial setae both increased and decreased in number, and in orientation, from a row parallel to the parapedial ridge to a setal row extended forward at about a right angle to the latter.

Keywords
comparative morphology, mouthparts, pre-oral mill, evolution, Coleoptera, Trachypachidae, Carabidae, Carabitaee, Nebriitae, Cicindelitae

Introduction

In their review of variation in mandibular structure within the coleopteran suborder Adepha-ga, Acorn and Ball (1991) unfortunately did not include the distinctive but complex (and therefore difficult to interpret) mandibles of the tiger beetles (Carabidae: Cicindelitae). Since then, Ball et al (1995: 302–311) provided the basis for inclusion of the epipharynx in systematic studies, but did not include the tiger beetles or other basal geadephagan lineages. We (GEB, JHA) decided to fill in these gaps in knowledge of geadephagan structure, and to do so, enlisted in the cause our willing and able colleague, Danny Shpeley.

Our initial investigation of tiger beetle mandibles revealed for the Geadephaga a combination of unique features and in them, substantial variation. To understand these aspects, we sought an orienting principle in the relatively recent studies and postulates of tiger beetle evolution, admirably summarized by Pearson and Vogler (2001: 43–51). We turned to analyses (Maddison et al 1999) of related basal stocks of geadephagans to seek the antecedents of the tiger beetle mandibles and labrum-epipharynx. Tiger beetles have been placed as a supertribe (Cicindelitae) within the Carabidae (Erwin 1985: 467, Erwin 2007: 171, Erwin and Pearson 2008) or as a separate family, the Cicindelidae (Cassola 2001, Pearson and Vogler 2001, Deuve 2004: 31), or as tribe Cicindelini (Liebherr and Will 1998: 151). Recent phylogenetic analyses of the Geadephaga have consistently placed tiger beetles with the Carabidae, in some instances giving this group a basal position within the carabid phylogeny (Erwin and Pearson 2008, Deuve 1994, Maddison et al. 1999: 104, Fig. 1 (one alternative placement)) and in others placing the group higher in the tree (Arndt and Putchkov 1997, Beutel and Haas 1996: 201, Fig. 1; Liebherr and Will 1998: 142, Fig. 57A; Maddison et al. 1999 (another alternative placement)). Maddison et al. (1999: 115, Fig. 6) place Trachypachidae as the adelphotaxon for the remaining Geadephaga as do Dressler and Beutel (2010).

In this paper, we illustrate and describe the mandibles and labrum-epipharynx of tiger beetles representing the five tribes here recognized (Amblycheilini, Manticorini, Megacephalini, Collyridini, and Cicindelini). We arrange them in a phylogenetic sequence, based on a postulated evolutionary pattern (Pearson and Vogler 2001: 46), and relate this series to the form of the mouthparts in Trachypachidae, Nebriitae, and Carabitaee, the latter three taxa representing the three mouthpart configurations that may have preceded that of the tiger beetles in an evolutionary sense.
We are pleased to dedicate this study in comparative morphology to Ross and Joyce Bell, Department of Biology, University of Vermont, Burlington, Vermont, in recognition of their contributions to the study of tiger beetles, the use of mandibles as character systems, and more generally to the field of adephagan systematics (e.g., Bell 1966). Especially appropriate to note in the context of the present contribution are: Ross’ treatment of the North American Chlaeniini (Bell 1960), in which mandibular form and size were shown to be a useful diagnostic feature in classification; and his study of the mouthparts of rhyso dine carabids (Bell 1994), whose mandibles he showed to function as a sheath for the underlying maxillae, and to be non-biting. His interest in tiger beetles was demonstrated through co-authoring a field guide to cicindelids (Leonard & Bell 1991).

Material and methods

Material

We examined 12 specimens with SEM, and an additional 37 with light (Wild M5 and M3 stereoscope) microscopy, (Table 1, Appendix). These specimens are housed in the E. H. Strickland Entomological Museum, University of Alberta (UASM), Royal Alberta Museum, Edmonton, Alberta (RAMC), California Academy of Science, San Fran-

| Table 1. Names, sex, and classification of exemplar individuals and species with SEM-illustrated mandibles and labrum-epipharynx. |
|---------------------------------------------------------------|
| **Family TRACHYPACHIDAE**                                     |
| Tribe TRACHYPACHIDININI                                        |
| *Trachypachus* Motschulsky                                     |
| *T. gibbsi* LeConte, 1861 (male)                               |
| **Family CARABIDAE**                                           |
| Supertribe CARABITAE                                           |
| Tribe CEROGLOSSININI                                           |
| *Ceroglossus* Solier                                           |
| *C. chilensis* (Eschscholtz, 1829) (male)                      |
| Supertribe CICINDELITAE                                        |
| Tribe AMBLYCHEILININI                                          |
| *Amblycheila* Say                                              |
| *A. baroni* (Rivers, 1890) (female)                            |
| *Omus* Dejean                                                  |
| *O. californicus* (Eschscholtz, 1829) (male)                   |
| *Picnochile* Motschulsky                                       |
| *P. fallaciosa* (Chevrolat, 1835) (female)                     |
Tribe MANTICORINI Fabricius
   *Manticora* Fabricius
   *M. tuberculata* (DeGeer, 1778) (female)

Tribe MEGACEPHALINI
   *Tetracha* Hope
   *T. carolina* (Linnaeus, 1767) (male)

Tribe COLLYRIDINI
   *Pogonostoma* Klug
   *P. chalybeum* (Klug, 1835) (male)

Tribe CICINDELINI
   *Therates* Latreille
   *T. basalis* Dejean, 1826 (female)
   *Oxycheila* Dejean
   *O. species*, male
   *Cicindela* Linnaeus
   *C. longilabris* Say, 1824 (male)

Supertribe NEBRIITAE
Tribe PELOPHILINI
   *Pelophila* Dejean
   *P. rudis* (LeConte, 1863) (female)

Methods

**Taxon and specimen selection.** For the cicindelites, we chose exemplar specimens to represent the five currently recognized tribes: Amblycheilini, Manticorini, Megacephalini, Collyridini and Cicindelini. For comparative purposes, we chose representatives from basal geadephagan lineages: Trachypachidae (*Trachypachus gibbsii* (LeConte, 1861)), the putative adelphotaxon of the caraboid stock; and Carabidae—Carabinae—Carabidae-Ceroglossini (*Ceroglossus chilensis* (Eschscholtz, 1829)), and Nebriitae-Pelophilini (*Pelophila rudis* (LeConte, 1863)). These groups of Carabidae represent different feeding types (carabites, like cicindelites, primarily predatory fluid feeders; and nebrati, particulate feeders; see Evans and Forsythe (1985: 115).

**Specimen preparation.** Standard techniques were used to prepare specimens for examination with the SEM. Each specimen was relaxed and cleaned in warm water and the sclerites of interest were removed from the head capsule and sputter coated with gold before SEM images were prepared. For light microscopy, specimens were relaxed, cleaned, and the mandibles spread to their fullest extent. For some but not all taxa, the labrum epipharynx was excised and point-mounted.

**SEM image preparation.** Three aspects for both left and right mandibles are shown: dorsal, occlusal, and ventral; lateral aspects were noted, but not illustrated. For
Mandibles and labrum-epipharynx of tiger beetles

the labrum-epipharynx, the dorsal surface is the labrum, while the ventral surface is the epipharynx, keeping in mind that the epipharynx can also be thought of as the dorsal surface of the preoral cavity.

**Photographs.** The mouthparts of 12 taxa examined are also illustrated *in situ*, with the mandibles widely spread, hopefully providing better perspective on the relative coverage of the adducted mandibles by the labrum-epipharynx for each taxon, as well as some sense of potential occlusal relationships among mandible features, and bilateral asymmetry. Photographs were taken hand-held with a Nikon D300s camera set at ISO 800 and 1/200th of a second, equipped with an AF-S Micro-Nikkor 105mm lens set at f32, a TC-17EII (1.7X) teleconverter, and the Nikon R1C1 macro flash system with two flash heads, each equipped with hand made, double-layered, frosted Mylar light diffusers. These images were presented on two color plates.

**Identification of structural elements.** For tiger beetles (specifically *Cicindela hybrida* (Linnaeus, 1767)) Evans (1965) referred to the large dorsal anterior occlusal teeth as incisors, and the posterior and more ventral complex of large teeth as molars. Kritsky and Simon (1995) used similar terms in their study of sexual dimorphism in mandibles of a wide selection of North American *Cicindela* (*sensu latissime*) species. The major elements of the occlusal surfaces of adult geadephagan mandibles were identified by Acorn and Ball (1991: 639–641, Fig. 1) as terebra (with a distal incisor tooth, terebral ridge, and proximal terebral tooth) and retinaculum (with a distal anterior retinacular tooth, a double retinacular ridge, and proximal posterior retinacular tooth, or molar tooth. The retinaculum

![Figure 1. Measurements, plotted on photographic image of dorsal aspect of left mandible of *Amblycheila baroni* Rivers. Legend: B basal area; BL length of basal area; BW width of basal area; IL length of incisor tooth; T terebra; TEL length of terebra; TL total length.](image)
is posterior and ventral to the terebra. Here, based on similarity of position, we recognize the distal-most tooth as an incisor, the large dentiform projections along the occlusal margin as terebral teeth, and the posterior array of dentition as the retinaculum. See Table 2 for a complete list of the structural elements of mandibles and labrum-epipharynx. The abbreviations are used in the SEM figures to designate these structural elements.

**Table 2.** Terms and abbreviations for geadephagan mandibles (modified from Acorn and Ball 1991) and labrum-epipharynx (modified from Ball et al. 1995).

| Abbreviations | Terms                                      |
|---------------|--------------------------------------------|
| art           | anterior retinacular tooth                 |
| B             | base of mandible                           |
| bb            | basal brush                               |
| bfb           | basal face brush                           |
| irr           | inferior retinacular ridge                 |
| it            | incisor tooth                              |
| mss           | multiple scrobal setae                     |
| mt            | molar tooth                               |
| od            | occlusal diastema                          |
| prt           | posterior retinacular tooth                |
| rc 1          | retinacular tooth, cusp 1= art, in part    |
| rc 2          | retinacular tooth, cusp 2= art, in part    |
| rc 3          | retinacular tooth, cusp 3= prt, in part    |
| rc 4          | retinacular tooth, cusp 4= prt, in part    |
| ret           | retinaculum                                |
| rr            | retinacular ridge                          |
| s             | scrobe                                     |
| srr           | superior retinacular ridge                 |
| srt           | supplementary retinacular tooth            |
| ss            | single scrobal seta                        |
| T             | terebra                                    |
| tr            | terebral ridge                             |
| tt            | terebral tooth                             |
| tt 1          | terebral tooth 1                           |
| tt 1-1        | terebral tooth 1, cusp 1                   |
| tt 1-2        | terebral tooth 1, cusp 2                   |
| tt 1-3        | terebral tooth 1, cusp 3                   |
| tt 2          | terebral tooth 2                           |
| tt 3          | terebral tooth 3                           |
| tt 4          | terebral tooth 4                           |
| vg            | ventral groove                             |
| vm            | ventral microtrichia                       |
Measurements. To assist in characterizing mandibles, four measurements (Fig. 1) were taken and used to make ratios (Table 2). Being based on the figures of single specimens, differences in values of these ratios have no statistical significance. They are simply a means of standardizing descriptive statements.

Descriptions. Descriptions are brief, arranged in putative phylogenetic sequence, as reflected in the suprageneric taxa referred to in this paper.

Descriptions And Comparisons

Family Trachypachidae

Classification. Ranked as a family, this group of two genera and fewer than 10 species may be regarded as the adelphotaxon of the Carabidae (Kavanaugh, 1998: 337; Maddison et al., 1999: 116, Fig. 7; Dressler and Beutel, 2010: Fig. 22, p. 282 ), or as a group more closely related to the Hydradephaga (Acorn and Ball, 1991: 645; Beutel, 1998: 94, Fig. 1, and p. 101).

Exemplar taxon. Trachypachus gibbsii LeConte, 1861.

Figs 2–F, 5A–B

Structural features. Mandibles (Figs 2A-F). Trigonal in dorso-ventral aspect, robust, basal width one half total length; curved downward slightly (Figs 2C-D). Lateral surface basally with scrobe broad, asetose, delimited by a dorsolateral and ventrolateral ridge. Terebra (T) short (terebral length one half total length), in occlusal aspect broad, with short incisor tooth (iT) and with small terebral tooth (tT). Retinaculum slightly posteriad terebral tooth; short, broad, with two ridges (rR and irr), and two teeth (art and prT), anterior tooth near ventral margin (2C-D), posterior tooth dorsal. Basal face not evident. Basal brush (BB) small. Ventral surface smooth except for the ventral groove (Figs 2E-F, vG), extended anteriad anterior retinacular tooth, and with rather short and sparse microtrichia (VM).

Labrum (Fig. 5A). Articulated with, not immovably attached to, clypeus, labral-clypeal suture evident. Distinctly transverse, sub-rectangular, row of 14 tactile setae

| Abbreviations | Terms |
|---------------|-------|
| LABRUM-EPIPHARYNX |     |
| aps           | anterior parapedial setae |
| epd           | epipharynx, dorsal aspect |
| las           | labral apical seta |
| ped           | pedium |
| pp            | parapedial projection |
| pps           | posterior parapedial setae |
| pr            | parapedial ridge |
| sc            | sensillum coeloconicum |
Figure 2. SEM photographs of mandibles of *Trachypachus gibbsi* Le Conte. **A, C, E** left mandible, dorsal, occlusal, ventral aspects, respectively; **B, D, F** right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars = 0.2 mm.
(las) near anterior margin, on dorsal surface (Figure 5A). Form of anterior margin subtruncate, shallowly emarginate.

_Epipharynx_ (Figure 5B). Pedium (ped) trianguloid, apex posterior, slightly arched, with a short parapedial projection (pp), margined laterally each side by a thin parapedial ridge (pr); parapedial ridge anteriorly each side curved to lateral margin of labrum as a short lateral arm. Anterior parapedial setae (aps) in a row anterior and parallel to lateral arms of parapedial ridge. Posterior parapedial setae (pps) in a row.

**Family Carabidae**

Two major types of mandibles occur among the basal carabid lineages: the fluid-feeding _Ceroglossus_ type, and the particulate-feeding _Pelophila_ type, evidently depending upon manipulation of food (Evans and Forsythe 1985: 114). Associated with fluid-feeding, the labrum-epipharynx is immovably attached to the clypeus.

We identified two major types of epipharynx: the general one, shared with the Trachypachidae—pedium triangular in form, as described above; and the type confined to the tiger beetles—pedium broad, palatiform, markedly convex. For details, see below, under “Cicindelitae”.

**Supertribe Nebriitae, Tribe Pelophilini**

(Classification). This monogeneric group of two extant species is basal to the Supertribe Nebriitae, which in turn is a basal assemblage of the Carabidae (Kavanaugh 1998: 335, Fig. 3, Maddison et al. 1999: 104, Fig. 1).

_Exemplar taxon._ _Pelophila rudis_ (LeConte, 1863).

Figs 3A-F, 5C-D

**Structural features. Mandibles** (Figs 3A-F). Trigonal in dorso-ventral aspect, robust, basal width one third total length; curved downward (Figs 3C-D). Lateral surface basally with scrobe broad, delimited by a dorsolateral and ventrolateral ridge; scrobe triangular, moderately deep, with single seta (ss). Terebra (T) long (tuberal length two thirds total length), in occlusal aspect narrow, with short incisor tooth (it) and with small terebral tooth (tt). Retinaculum slightly posterior tuberal tooth, in dorso-ventral aspect (Figs 3A-B) narrow with single ridge (rr), and two teeth (art and prt), one at each end, in line with one another. Basal face not evident. Basal brush (bb) small. Ventral surface smooth except for the ventral groove (Figs 3E-F, vg), extended anterior retinacular tooth, and with rather short and dense microtrichia (vm).
Figure 3. SEM photographs of mandibles of *Pelophila rudis* LeConte. **A, C, E** left mandible, dorsal, occlusal, ventral aspects, respectively; **B, D, F** right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars = 0.5 mm.
Mandibles and labrum-epipharynx of tiger beetles...  49

Labrum (Fig. 5C). Articulated with, but not immovably attached to clypeus, labral-clypeal suture evident. Distinctly transverse, sub-rectangular, row of five tactile setae (las) near anterior margin, on dorsal surface (Fig. 5C). Form of anterior margin subtruncate.

Epipharynx (Fig. 5D). Pedium (ped) broadly trianguloid, apex posterior, slightly arched, with a short parapedial projection (pp); margined laterally each side by a thin parapedial ridge (pr); parapedial ridge anteriorly each side curved to lateral margin of labrum as a short lateral arm. Anterior parapedial setae (aps) in a row anterior and parallel to lateral arms of parapedial ridge. Posterior parapedial setae (pps) few, in row along parapedial ridge.

Comparisons. The prominent retinaculum seems to be a feature of the Nebriitae (see Kavanaugh 1978: 856, Figs 54–58; and Acorn and Ball 1991: 647, 11A-D).

Supertribe Carabitaee, Tribe Ceroglossini

Classification. The Supertribe Carabitaee is a basal lineage (Maddison et al. 1999: 104, Fig. 1; Kavanaugh 1998: 335, Fig. 3). Erwin and Pearson (2008: 19) cite Carabus Linnaeus (meaning supertribe Carabitaee, Erwin, personal communication, 2011) as the adelphotaxon of the Cicindelitae; Liebherr and Will (1998: 142, Fig. 57) include the Cicindelini as part of a quadritomy with three of the carabine tribes noted below.

The Carabitaee includes 14 genera, arranged in four tribes: Cychrini; Ceroglossini; Pamborini; and Carabini. The tribe Ceroglossini is monogeneric, including eight species.

Exemplar taxon. Ceroglossus chilensis (Eschscholtz, 1829).

Figures 4A-F, 5E-F, Plate 1A

Structural features. Mandibles (Figs 4A-F, Plate 1A). Trigonal in dorso-ventral aspect (Figures 4A-B), robust, basal width one quarter total length; curved downward (Figs 4C-D). Lateral surface basally with scrobe broad, triangular, moderately deep, asetose, delimited by a dorsolateral and ventrolateral ridge. Terebra (T) long (terebral length only one half total length, but basal area exceptionally long), in occlusal aspect (Figs 4C-D) broad; with short incisor tooth (it) and with small terebral tooth (tt). Retinaculum slightly posteriorid terebral tooth, short, oriented obliquely to long axis of mandible (Figs 4C-D), (broad single ridge (rr), and two teeth (art and prt), one at each end of terebral ridge, offset from one another. Basal face long, densely setose (bfb), setae seemingly continuous with microtrichia of ventral groove (Figs 4C-D). Basal brush (bb) small. Ventral surface smooth except for the ventral groove (Figures 4E-F, vg), extended anteriorly nearly to base of the incisor tooth, and with long and dense microtrichia (vm).

Labrum (Fig. 5E, Plate 1A). Immovably attached to clypeus, labral-clypeal suture evident. Distinctly transverse, sub-rectangular, transverse cluster of numerous tactile
setae (las) near anterior margin, on dorsal surface (Fig. 5E). Form of anterior margin moderately deeply emarginate.

*Epipharynx* (Fig. 5F, Plate 1A). Pedium (ped) rather narrowly trianguloid, apex posterior; slightly arched, with a long parapedial projection (pp); margined laterally each side by a thin parapedial ridge (pr); parapedial ridge anteriorly each side curved gradually to lateral margin of labrum. Anterior parapedial setae (aps) in a row anterior and parallel to lateral portion of parapedial ridge. Posterior parapedial setae (pps) rather numerous, in row along parapedial ridge.

**Comparisons.** The form of the mouthparts in *Ceroglossus* is in many ways reminiscent of that in cicindelites, and a more extensive survey of the Carabidae may well uncover additional shared features.

Another group of fluid feeders, the scaritines, was examined briefly, with inconclusive results. Some, (e.g., *Pasimachus* Bonelli, *Mouhotia* Laporte de Castelnau) appear not to possess a retinaculum, whereas in *Scarites* the retinaculum is either reduced and confluent with the base of the terebra, or the basal portion of the terebra is multidentate and vaguely reminiscent of what we interpret here as the retinaculum of cicindelites. Some evidence exists for a relationship between tiger beetles and scaritines (the “CPRS quartet” of Maddison et al. 1999, uniting tiger beetles, paussines, rhysoelines, and scaritines); even these authors suggest that convergence seems a more likely explanation for this morphologically incongruous assemblage (but see Bell and Bell 1962 and Bell, 1998 for evidence of a relationship between rhysoelines and scaritines). Further study of this issue seems warranted (see, for example, Makarov 2008, who, based on morphological features, proposes to place the rhysoelines and paussines in the suborder Archostemata).

**Supertribe Cicindelitae**

**Classification.** Based principally on the phylogenetic conclusions of Pearson and Vogler (2001: 46, Fig. 3.5) the tiger beetles are arranged here in five tribes: Amblycheilini; Manticorini; Megacephalini; Collyridini; and Cicindelini. Gálian et al. (2002: 1794, Fig. 1) indicate the Megacephalini as polyphyletic, based on their study of multiple sex chromosomes in the cicindelites, the problem taxa being the oxycheiline genera *Oxycheila* and *Cheilocyba*. Based on their 18sRNA evidence and the DNA evidence of Vogler and Barraclough (1998: 255, Fig. 1), these genera (and presumably *Pseudoxycheila*) belong in the tribe Cicindelini, and such a transfer obviates the taxonomic problem.

**Structural Features.** *Mandibles* (Figs 6A-F–8A-F, 10A-F–12A-F, and 14A-F–16A-F). Trigonal in dorso-ventral aspect, robust to slender, basal width one fifth to one third total length; planar to curved downward, ventral curvature simple to complex. Lateral surface basally with scrobe broad, delimited by a dorsolateral and ventrolateral ridge, or with ventrolateral ridge only distinct, or lateral surface convex, without a scrobe;
lateral surface asetose or multisetose (ss). Terebra (T) elongate (terebral length one half to three quarters total length), in occlusal aspect broad to slender, with long incisor tooth (it) (one third to one half total length), and with or without terebral teeth (tt) (in most taxa teeth two to three, variously prominent, unicuspidate or tricuspidate; in most taxa, sexually dimorphic). Retinaculum slightly posteriad basal terebral tooth or more widely separated by a distinct gap (the occlusal diastema, od; cf. Figs 12A, B, E, F). short, broad, with four or more cusps (rc 1–4); cusps of various sizes, in some taxa as large and prominent as terebral teeth. Basal face (bf) posteriad retinaculum, with long setae in form of a dense brush (bfb). Ventral surface smooth except for the ventral groove (vg) of various lengths, and with rather long and dense microtrichia (vm).

Labrum (Figs 9A, C; 13A, C, E; and 17A, C, E). Attached immovably to clypeus, although labral-clypeal suture evident. Of various proportions (Length/Width 0.20–1.02), i.e., distinctly transverse, sub-rectangular, to slightly longer than wide and trapezoid or triangularoid; row of tactile setae either near anterior margin, on dorsal surface (Fig. 9A), or on apical margin (Fig. 13E). Form of anterior margin various, from subtruncated (Fig. 13C) to simply projected medially (Fig. 9C), to markedly projected (Fig. 17C); projection one or several denticles, or broadly rectangular (Fig. 9C). See also Cazier (1954: 306–307, Figs 124–169).

Epipharynx (Figs 9B, D, F; 13B, D, F; and 17B, D, F). Pedium (ped) palatiform, markedly arched, ventral surface concave; posteriorly broadly rounded, without a parapedial projection, margined laterally each side by a thin parapedial ridge (pr); parapedial ridge anteriorly each side curved to lateral margin of labrum as a short lateral arm. Parapedial setae in a row or cluster anterior (aps) to or posterior (pps) to lateral arms of parapedal ridge.

### Table 3. Ratios for features of left mandible of exemplar specimens of Trachypachus, Pelophila, Ceroglossus, and nine genera of Cicindelitae, representing Tribes Amblycheilini, Manticorini, Megacephalini, Collyridini, and Cicindelini.

| Taxon                        | BW²/TL¹ | TeL³/TL | IL⁴/TL |
|------------------------------|---------|---------|--------|
| Trachypachus gibbsii LeConte | 0.47    | 0.53    | 0.12   |
| Pelophila rudis LeConte      | 0.32    | 0.76    | 0.10   |
| Ceroglossus chilensis Eschscholtz | 0.24   | 0.54    | 0.12   |
| Amblycheila baroni (Rivers)  | 0.27    | 0.77    | 0.46   |
| Omus californicus            | 0.27    | 0.59    | 0.57   |
| Picnochile fallaciosa (Chevrolat) | 0.32  | 0.71    | 0.43   |
| Manticora tuberculata (DeGeer) | 0.27   | 0.68    | 0.33   |
| Tetracha carolina (Linnaeus) | 0.30    | 0.70    | 0.34   |
| Pogonostoma chalybeum (Klug) | 0.20    | 0.69    | 0.42   |
| Therates basalis Dejean      | 0.19    | 0.66    | 0.37   |
| Oxycheila species            | 0.24    | 0.70    | 0.39   |
| Cicindela longilabris Say    | 0.17    | 0.68    | 0.45   |

² Total Length; ³ Basal width; ⁴ Terebral Length; ⁵ Incisor Length
Figure 4. SEM photographs of mandibles of *Ceroglossus chilensis* Eschscholtz. **A, C, E** left mandible, dorsal, occlusal, ventral aspects, respectively; **B, D, F** right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars = 1.0 mm.
Comments. Cicindelid mandibles are readily distinguishable from those of other geadephagans by a combination of complex and distinctive retinacular structure (essential to the “pre-oral mill” of Evans (1965: 64) and multi-toothed terebra. Among the tiger beetle tribes, the more plesiotypic manticorines and amblycheilines exhibit greater mandibular robustness compared to the more apotypic megacephalines, collyridines and cicindelines, and most taxa with either two or three terebral teeth, this number reduced to one in many collyridines (especially on the left mandible) and in some cicindelines (e.g., Therates), but increased to as many as four in other collyridine lineages and to five in some cicindeline lineages. The labrum-epipharynx is generally short in the amblycheilines, manticorines, and megacephalines, and sub rectangular or
elongate and dorsally convex (elongation presumably evolving several times) among the collyridines and cicindelines.

**Supertribe Cicindelitae: Tribe Amblycheilini**

**Synonymic note.** Commonly known as the Omini W. Horn (1907: 466) (for example, Ball and Bousquet 2001: 71; Pearson and Vogler: 2001: 48), in fact the correct name is Amblycheilinini Csiki (1903:124), based on the principle of priority. For details, see Madge (1989: 460 and 466).

**Classification.** This tribe includes three Western Hemisphere genera: the western Nearctic *Amblycheila* Say and *Omus* Dejean; and the southern Neotropical *Picnochile* Motschulsky (female). If the southern Afrotropical monobasic genus *Platychile* Maclay is placed in this group (e.g., Pearson and Vogler 2001: 48), the tribal name becomes Platychilinini W. Horn (1893: 325) (Madge, 1989: 460 and 466), but see below for our reasons for not choosing this arrangement.

**Exemplar taxa.** *Amblycheila baroni* (Rivers, 1890) (female); *Omus californicus* (Eschscholtz, 1829) (male); *Picnochile fallaciosa* (Chevrolat, 1854) (female).

Also examined but not treated in detail were two males of *Platychile pallida* (Fabricius, 1801).

**Structural features.** *Mandibles* (Figs 6A-F – 8A-F; Plate 1B).—With mandibular features of Cicindelitae, restricted as follows. Values for ratios BW/TL, T eL/TL, and IT/TL as in Table 3; planar (Figs 6C-D) to moderately curved ventrad (Figs 7C-D). Lateral surface basally with scrobe broad, delimited by a dorsolateral and ventrolateral ridge; scrobe multisetose (Figs 6A-F, ss) or glabrous. Diastema absent. Terebral teeth two, terebral tooth 1 tricuspidate (Figs 6A-F, tt 1-1, tt 1-2, tt 1-3). Retinacular cusps (Figs 6C-D) rc 1 and rc 2 directly opposite one another, also rc 3 and rc 4 directly opposite one another. Ventral groove (Figs 6E-F and 10E-F, vg) moderately long, extended about to middle of terebral tooth 1.

*Labrum* (Figs 9A, C, E; Plate 1B). Transverse (L/W 0.20–0.36), in form rectanguloid. Anterior margin distinctly projected medially, otherwise various: projection truncate (Fig. 9C) or narrowly notched (Fig. 9A); or anterior margin with two paramedial notches, anterior margin of medial projection broadly emarginate (Fig. 9E). Single preapical row of 8–10 setae (las).

*Epipharynx* (Figs 9B, D, F). Row of anterior parapedial setae (aps) extended laterally, parallel to lateral extensions of parapedial ridge (pr).

**Comments.** Gissler (1879: 234) postulated that the genus *Amblycheila* represented the “lowest.....and certainly the oldest line of descent...that probably diverged in the
Figure 6. SEM photographs of mandibles of *Amblycheila baroni* Rivers. **A, C, E** left mandible, dorsal, occlusal, ventral aspects, respectively; **B, D, F** right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars 1.0 mm.
Mesozoic age” - a remarkably prescient line of thought, considering that it was developed more than a century ago.

The genus *Platychile* deserves special comment here. We examined two males of *P. pallida* (Fabricius) using light microscopy. A number of mandibular features (planar, markedly curved; terebra bidentate; retinaculum quadridentate) place the genus among the more basal lineages. On the other hand, the mandibles are so flat as to virtually eliminate the scrobe (and are thus reminiscent of the more derived tiger beetles), and they possess a single cusp terebral tooth 1 instead of the three cusps characteristic of the Western Hemisphere genera Plate 1F). The rectangular labrum (Plate 1F) has an anterior margin with two paramedial dentiform projections, flanked each side by two more short and blunt projections (six, in all) and six setae on the anterior (not apical) surface. The labrum, in fact, in form and setation, is strongly reminiscent of that of the tribe Megacephalini. Further, the body size and form is not unlike that of what could be expected in the megacephaline genus *Phaeoxantha*. The color pattern is also megacephaline-like, and a careful reading of Pearson and Vogler (2001: 53–57) indicates that such a feature may have importance in tiger beetle evolution. The unusual habitus of *Platychile* may also derive in part from convergence or mimicry, since these nocturnal beetles show a consistent ecological association with the diurnal *Eurymorpha cyanipes* (Hope, 1838) (Werner, 2000), with which they share an oval dorsoventrally flattened appearance, without pronounced elytral humeri—a resemblance that seems unlikely to stem from mere coincidence. *Platychile* may be the adelphotaxon of the Western Hemisphere genera (Galian, et al. 2002: 1794, Fig. 1), or it could be treated as a monobasic group of uncertain affinity, our preferred arrangement here.

**Supertribe Cicindelitae: Tribe Manticorini**

**Classification.** This tribe includes the southern Afrotropical genera *Manticora* Fabricius, 1792 and *Mantica* Kolbe, 1896.

**Exemplar taxon.** *Manticora tuberculata* (DeGeer, 1778) (female).

Other specimens examined: see Appendix.

**Structural features.** *Mandibles* (Figs 10A-F). With mandibular features of Cicindelitae, restricted as follows. Values for ratios BW/TL, TeL/TL, and IT/TL as in Table 3. Robust; markedly curved ventrad (Figs 10C-D). Lateral surface basally with scrobe broad, delimited by a dorsolateral and ventrolateral ridge; scrobe multisetose (Figs 10A-F, ss). Diastema absent. Terebral teeth two (some female *Manticora* or three (male *Manticora*, some female *Mantica*, and both sexes of *Mantica*), terebral tooth 1 monocuspidate (Figs 10A-F, tt 1); or bicuspidate, tt 1-2 (appearing separate from tt 1-1 in genus *Mantica*. Retinacular cusps (Figs 10C-D) rc 1 and rc 2 directly opposite one another, also rc 3 and rc 4 directly opposite one another. Ventral groove (Figs 10E-F, vg) moderately long, extended about to middle of terebral tooth 1.
Figure 7. SEM photographs of mandibles of *Omus californicus* Eschscholtz. **A, C, E** left mandible, dorsal, occlusal, ventral aspects, respectively; **B, D, F** right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars = 1.0 mm.
Figure 8. SEM photographs of mandibles of *Picnochile fallaciosa* Chevrolat. A, C, E left mandible, dorsal, occlusal, ventral aspects, respectively; B, D, F right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars = 1.0 mm.
The mandibles of *Manticora* are pronouncedly sexually dimorphic, larger in males and asymmetric in form, the right mandible typically exhibiting greater elongation of the incisor region than the left. In *Mantica*, sexual dimorphism is slight, and some but not all males show larger left than right mandibles (Franzen and Heinz 2005: 299).
Labrum (Fig. 13A). Transverse (L/W 0.20–0.36), in form rectanguloid. Anterior margin distinctly crenate, Manticora with six teeth, Mantica with four teeth (Franzen and Heinz, 2005: 300), median projection short, broad, emarginate anteriorly. Single preapical row of 10 setae (las).

Epipharynx (Fig. 13B). Row of anterior parapedial setae (aps) extended laterally, parallel to lateral extensions of parapedial ridge (pr).

Comments. See also Plate 1C. In structure of mandibles and labrum-epipharynx, the Manticorini seems most similar to the Amblycheilini. However, the marked ventrad curvature of the mandibles is suggestive of the more derived megacephalines, cicindelines and collyrines. With the genera Amblycheila and Pogonostoma, members of Manticora share multisetose scrobes.

For details about way of life, classification and relationships of Manticora, see Oberprieler and Arndt (2000). Franzen and Heinz (2005) provide a valuable review, including illustrations of mandibles, of the monobasic genus Mantica (type species, M. horni Kolbe, 1896).

Supertribe Cicindelitae: Tribe Megacephalini

Classification. This tribe includes more than 100 species, arrayed in eight genera (Zerm et al. 2007, and Huber, 1994). Galián et. al. (2002: 1794) indicate that, based on 18sRNA analysis the Megacephalini is polyphyletic, with the genera Cheiloxya and Oxycheila sharing a closer relationship with the cicindelines than with the megacephalines. Vogler and Barraclough (1998: 256, Figure 2) had indicated that DNA evidence showed the same thing for Oxycheila. No doubt, Oxycheila, Cheiloxya and putative close relative Pseudoxycheila belong in the tribe Cicindelini.

Exemplar Taxon. Tetracha carolina (Linnaeus, 1767) (male).

Other specimens examined: see Appendix.

Structural Features. Mandibles (Figs 11A-F). With mandibular features of Cicindelitae, restricted as follows. Values for ratios BW/TL, TeL/TL, and IT/TL as in Table 3. Robust; markedly curved ventrad (Figs 11C-D). Lateral surface basally with scrobe broad, delimited by a dorsolateral and ventrolateral ridge; scrobe glabrous (Figs 11A-F). Diastema absent. Terebral teeth three, terebral tooth 1 monocuspidate (Figs 11A-F, tt 1). Retinaculum with supplementary tooth anteriorly (srt) in Tetracha and Megacephala, very small in Phaeoxantha. Retinacular cusps (Figs 11C-D) rc 1 and rc 2 directly opposite one another, also rc 3 and rc 4 directly opposite one another. Ventral groove (Figs 11E-F, vg) moderately long, extended about to base of terebral tooth 2.

Labrum (Fig. 13C). Transverse (L/W 0.28), in form rectanguloid. Anterior margin shallowly crenate, medially, median projection short, broad, emarginate anteriorly. Single preapical row of four setae (las).
Figure 10. SEM photographs of mandibles of *Manticona latipennis* Waterhouse. **A, C, E** left mandible, dorsal, occlusal, ventral aspects, respectively; **B, D, F** right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars = 5.0 mm.
Epipharynx (Fig. 13D). Row of anterior parapedial setae (aps) extended anteriorly, at about right angle to lateral extensions of parapedial ridge (pr).

Comments. See also Plate 1D and 1E. The long terebral tooth 3 illustrated for the specimen of *Tetracha carolina* seems to be characteristic of males of that genus, contrasting markedly with the shorter t3 of the corresponding females. Number of labral setae in Megacephalini ranges from four to seven.

**Supertribe Cicindelitae: Tribe Collyridini**

**Classification.** This group is the equivalent of and co-extensive with Walther Horn’s (1908) “Phylum” Alocosternalia. This tribe includes seven genera arranged in two subtribes: Ctenostomatina, and Collyridina (Lorenz 2005: 22).

**Exemplar taxon.** *Pogonostoma chalybeum* (Klug, 1835) (male).

Other collyridines examined include representatives of subtribes Ctenostomatina and Collyridina. See appendix.

**Structural features.** Mandibles (Figs 12A-F). With mandibular features of Cicindelitae, restricted as follows. Values for ratios BW/TL, TeL/TL, and IT/TL as in Table 3. Slender; markedly curved ventrad (Figs 12C-D). Lateral surface basally with scrobe broad, delimited by a dorsolateral and ventrolateral ridge; scrobe multisetose (Figs 12A-F, ss). Diastema (od) present. Terebral teeth various in size and number (see “Variation” below, for details). Retinaculum without or with (*Collyris* only) supplementary tooth anteriorly (srt). Retinacular complex large, cusps (Figs 12C-D) diagonally arranged, number of cusps various (see below for details) Ventral groove (Figs 12E-F, vg) moderately long, extended about to base of terebral tooth 2.

Labrum (Fig. 13E). Elongate (L/W 0.63), in form trapezoidal. Anterior margin shallowly crenate. Single apical row of nine setae (las), each seta inserted in base of crenulation. For details, see “Variation”, below.

Epipharynx (Fig. 13F). Pedium (ped) markedly concave. Row of anterior parapedial setae (aps) extended anteriorly, at about right angle to lateral extensions of parapedial ridge (pr). For details, see “Variation”, below.

**Variation.** Terebral teeth two (each mandible), tt 1 monocuspidate (Figs 12A-F) (*Pogonostoma*), or only one tooth on each mandible (*Collyris*); or terebral teeth asymmetric, with two on right mandible and one on left mandible (*Ctenostoma*), or one on left mandible, two on right mandible (*Tricondyla*). Retinacular cusps five each mandible (*Pogonostoma*), or seven on left mandible, five on right mandible (*Ctenostoma*), or five on left mandible, four on right mandible (*Tricondyla* and *Collyris*).

Although the labrum-epipharynx is consistently elongate and dorsally convex, the number of anterior marginal teeth varies from five to eight, some taxa with
Figure 11. SEM photographs of mandibles of *Tetracha carolina* Linnaeus. A, C, E left mandible, dorsal, occlusal, ventral aspects, respectively; B, D, F right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars = 1.0 mm.
medial crenulation and an odd number of teeth; some with median notch and an even number of teeth; lateral pair of teeth generally acute, median teeth in form of rounded crenulations; shallow grooves present, in some taxa on labrum and/or epipharynx, extended posteriad notch separating lateral and medial teeth. Number of labral setae various, from six to 14. Anterior parapedial setae extended almost to anterior margin in *Pogonostoma*, but less so in other collyridine taxa. It is not clear how medial teeth, notches, and/or setae have evolved from their paired bilateral homologues.

Comments. See Plate 2A for illustrations of the mandibles and labrum of *Ctenostoma ichneumoneum* Dejean, 1833. Clearly, although the collyridines present a diversity of mouthpart configurations, there is no obvious reason to doubt the use of *Pogonostoma* as an exemplar for the group, likely to exhibit a more or less plesiotypic structural condition. The ant-like body form of most collyridines may have constrained the head shape and therefore mouthpart structure of these beetles to some extent, but this is merely conjecture on our part.

Supertribe Cicindelitae: Tribe Cicindelini

Classification. This tribe includes more than 1500 species, arranged in five subtribes: Theratina; Oxycheilina; Iresina; Prothymina; and Cicindelina (Vogler and Barraclough, 1998).

Exemplar taxa. *Therates basalis* Dejean (Theratina); *Oxycheila* species (Oxycheilina); and *Cicindela longilabris* Say (Cicindelina).

Other cicindelines examined: see Appendix for names; plus numerous species of Nearctic *Cicindela*, principally for form, and for number of labral setae.

Structural features. Mandibles (Figs 14A-F; 16A-F; Plate 2F). With mandibular features of Cicindelitae, restricted as follows. Values for ratios BW/TL, TeL/TL, and IT/TL as in Table 3. Slender; markedly curved ventrad (Figs 14C-D). Lateral surface basally without scrobe, or scrobe very narrow, confined to lateroventral surface (*Dromica*), and without setae. Diastema present (Figs 14A-F and 16A-F, od) or absent (Figs 15A-F). Terebral teeth various in size and number: one, (Figs 14A-F, tt 2) to four (Figs 15A-F, tt 1 – tt 4), with a maximum of five, observed in *Oxygonia gloriola* Bates, 1872. Retinaculum without or with (*Therates*, Figs 14A-F) supplementary tooth anteriorly (srt). Retinacular complex large, cusps diagonally arranged, number of cusps four on each mandible (rc 1-rc 4). Ventral groove (Figs 15E-F, vg) moderately long, extended about to base of terebral tooth 2, or shorter, extended only to base of retinacular complex (Figs 16E-F).

Labrum (Figs 17A, C, E). Markedly varied. Illustrations as follows. Form transverse (L/W 0.41, Fig. 17E), as long as wide (L/W 1.00, Fig. 17A), or slightly longer than wide (L/W 1.02, Fig. 17C). Trapezoidal to trianguloid. Anterior margin nearly
Figure 12. SEM photographs of mandibles of Pogonostoma chalybeum Klug. **A, C, E** left mandible, dorsal, occlusal, ventral aspects, respectively; **B, D, F** right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars = 1.0 mm.
Figure 13. SEM photographs of labrum and epipharynx of: Manticora latipennis Waterhouse (A labrum, dorsal aspect; B epipharynx, ventral aspect); Tetracha carolina Linnaeus (C labrum, dorsal aspect; D epipharynx, ventral aspect); Pogonostoma chalybeum Klug (E labrum, dorsal aspect; F epipharynx, ventral aspect). Legend: see Table 2. Scale bars: 1.0 mm.

smooth or distinctly crenate. Labral setae (las, pls) four to 12, on dorsal or apical surface, inserted at base of crenations.

Epipharynx (Figs 17B, D, F). Anterior parapedial setae (aps) in row parallel to lateral arm of parapedial ridge (Fig. 17D), extended in short row anterior and
Figure 14. SEM photographs of mandibles of *Therates basalis* Dejean. **A, C, E** left mandible, dorsal, occlusal, ventral aspects, respectively; **B, D, F** right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars = 1.0 mm.
Figure 15. SEM photographs of mandibles of *Oxycheila* species. Eschscholtz. A, C, E left mandible, dorsal, occlusal, ventral aspects, respectively; B, D, F right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars = 1.0 mm.
Figure 16. SEM photographs of mandibles of *Cicindela longilabris* Say. **A, C, E** left mandible, dorsal, occlusal, ventral aspects, respectively; **B, D, F** right mandible, dorsal, occlusal, ventral aspects, respectively. Legend: see Table 2. Scale bars = 1.0 mm.
obliquely to lateral arm of parapedial ridge (Fig. 17E) or only very few setae laterally (Fig. 17B).

**Variation.** For illustrations of mandibles and labra of additional cicindeline taxa, see Plate 2C to 2E. The mandibles of three exemplar taxa described above, each representing a different cicindeline subtribe, differ strikingly from one another, seeming to indicate an appreciable level of divergence in this tribe. Within the subtribe Cicindelina, Kritsky and Simon (1995) showed that the mandibles of various Nearctic species of *Cicindela* exhibit more or less striking sexual dimorphism, the number of terebral teeth being constantly three, but differing in relative size. Similarly, Satoh and Hori (2004: 211) showed sexual dimorphism in the Palaearctic species, *Lophyridia angulata* (Fabricius, 1781), as did Oberprieler and Arndt (2000: 86) for *Manticora* adults, and Franzen and Heinz (2005: 299) for *Mantica* adults. Although not well studied, it appears that in all but a few aberrant individuals, the left mandible adducts above the right (“left-superior chirality”; Richardson, 2010).

In their remarkable study of geographical variation in *Cicindela dorsalis* Say, 1817, Boyd and Rust (1982: 225, 229) described a dentiform projection (the “submandibular tooth”) on the ventral terebral surface of the right mandible of males, only. This projection, of unknown function, was shown to vary in size (their paper, p. 228, Fig. 6) depending upon subspecies which, in turn, was correlated with overall body size.

Intensive investigation of mandibular length (Ganeshaiah and Belavadi, 1986; Mury Meyer, 1987; Niemelä and Ranta, 1993; Pearson, 1980; Pearson and Juliano, 1991; Pearson and Mury, 1979; Satoh et al 2003; and Satoh and Hori 2004) has shown that this factor is important in the structuring of tiger beetle communities, principally through resource partitioning (Pearson and Vogler 2001: 198–203).

Our three exemplar taxa differ markedly from one another in form and setation of the labrum. We note that the insertion of the labral setae on the dorsal surface of the labrum in *Cicindela longilabris* Say, 1824 (Fig. 17E) is a relatively basal condition. Cazier (1954: 2306–309, Figs 129–223), in his treatment of the Mexican species of *Cicindela*, illustrated striking differences in labra, particularly in form of the anterior margin of the labrum, proportions, and number of dorsal setae (from four to more than 30).

The epipharynges of the three exemplar taxa are basically similar to one another, but differ in the anterior parapedial setation. The small number of such setae exhibited by *Therates basalis* Dejean, 1826 (Fig. 17B) is the most derived, and is similar to that of the collyridine, *Ctenostoma metallicum* Laporte de Castelnau, 1834 (not illustrated).

**Evolution**

The following hypothesis of mandibular and labral-epipharyngeal evolution is illustrated in the reconstructed phylogeny (Fig. 18) based on Vogler and Barraclough (1998) and largely corroborated by other studies (Liebherr and Will, 1998; Gálian et al, 2002), although the latter study places the Manticorini as the adelphotaxon of
Figure 17. SEM photographs of labrum and epipharynx of: Therates basalis Dejean (A labrum, dorsal aspect; B epipharynx, ventral aspect); Oxycheila species (C labrum, dorsal aspect; D epipharynx, ventral aspect); Cicindela longilabris Say (E labrum, dorsal aspect; F epipharynx, ventral aspect). Legend: see Table 2. Scale bars: 1.0 mm.
the Amblycheilini. Reference points on this diagram are the lineages indicated by the capital letters A to P.

Lineage A represents the common ancestor of the Geadephaga, in which we assume the mandibles possessed a distinct terebra and retinaculum, of unknown form, but probably Pelophila-like. The labrum-epipharynx was likely movably articulated with the head capsule.

Lineage B represents the evolution of the Trachypachidae, with mandible (Figs 2A-F) and labral-epipharynx (Figs 5A-B) features as follows: mandibles, short, broad, slightly curved ventrally, terebra relatively short, with short incisor and short terebral teeth, a very short basal face, and the basal brush (bb) serving as the basal face setae. The retinaculum includes two long retinacular ridges (rr, srr), and is bicuspidate, teeth (art, prt) short.

The labrum-epipharynx is movably articulated with the head capsule: labrum transverse, rectanguloid, anterior margin subtruncate, dorsal surface with a row of numerous setae (14, more or less); epipharynx with short, trianguloid pedium, with few anterior and posterior parapedial setae, and a small parapedial projection. Diet yet to be determined: probably partially fluid and partially particulate matter.

Lineage C The stem of the Carabidae. Evolution of the mandibles includes only simplification of the retinaculum to a single broad ridge and possibly lengthening of the terebra. The diet was probably generalized, including both solid and fluid food (the “mixed feeders” of Forsythe, 1983: 371).

Lineage D Evolution of carabite-cicindelitide mandibles as the diet became fluid, only, the food principally soft-bodied invertebrates. Includes development of a somewhat enlarged and densely setose basal face for retention of the fluid component of prey tissues for extra-oral digestion, and the labrum-epipharynx becoming immovably attached to the head capsule.

Lineage O Evolution of mandibles (Figs 3A-F) of the nebriite- “rest of Carabidae” Lineage includes development of a scrobal seta (ss), and narrowing of the single ridged retinaculum (rr). (Not followed further here, but see Acorn and Ball, 1991).

Lineage E Evolution of mandibles of Carabidae (Figs 4A-F) includes marked lengthening of the basal face (bf) shortening of the retinaculum and its diagonal orientation (rr). The ventral groove (Figs 4E-F, vg) became markedly lengthened to nearly the base of the incisor tooth, and the ventral microtrichia (vm) became markedly lengthened. The anterior margin of the labrum (Fig. 5E) became markedly emarginate.

Lineage F Evolution of mandibles of Cicindelitae includes hypertrophy of the incisor tooth, development of a second terebral tooth, hypertrophy of the terebral teeth, and the scrobes becoming multisetose. But the most striking mandibular development is seen in the retinaculum, which becomes markedly enlarged and quadricuspidate – an important element of the preoral mill.

The labrum becomes lengthened, its anterior margin sinuously arched and sparsely setose, the number of anterior setae reduced to 8–10. The epipharynx is extensively modified, becoming palatiform, dorsally arched, and widened posteriorly, with loss of the para-
pedial projection. Evans and Forsythe (1985: 116) describe the mode of feeding, unique among fluid-feeding carabids, in which prey is held, punctured and sheared by the incisor and terebral teeth, then passed posteriorly by the maxillary lacinia to the crushing teeth of the retinaculum. Within the preoral mill (bordered dorsally by the epipharynx, ventrally
Plate 1. Digital images of head capsule, labrum, and mandibles dorso-frontal aspect, of: A Cercoglossus chilensis (Eschscholtz); B Amblycheila baroni (Rivers); C Manticora mygaloides Thomson; D Megacephala regalis Boheman; E Phaeoxantha tremolerasi (W. Horn); F Platychile pallida (Fabricius). Scale bars: A, B, E, F = 5 mm; C, D = 10 mm.
by the setose labium and laterally by the setose retinacular region of the mandibles), a food bolus is rotated posterio-dorsally and anterio-ventrally, bathed in midgut enzymes. Partially digested fluid is drawn through the mouth by a powerful pharyngeal pump until all but fragments of cuticle have been ingested, at which point the bolus is ejected.

Nodes and stems G-N map the evolution of the mandibles and labrum-epipharynx in the Cicindelitae.

Lineage G Evolution of the amblycheiline mandibles involves principally enlargement of the terebral tooth 1 (Figs 6C-D, tt 1-1), which becomes tricuspidate (tt 1-1 – tt 1-3). Within the Amblycheilini, the scrobal setae are retained in Amblycheila, but lost in the other genera (Figs 7A-B and 8A-B), and the slight down-curvature of the terebra is reduced (Figs 6C-D). The anterior margin of the labrum is variously modified, the median extension truncate (Fig. 9A), or narrowly notched (Fig. 9C), or broadly emarginate (Fig. 9E).

Node H Evolution of the remaining lineages (H-N) of the Cicindelitae. The basic mandibular and labral-epipharyngeal features are those of the Amblycheilini, outlined above, except terebral tooth 1 consists only of a single large cusp.

Lineage I Evolution of the manticorine mandibles (Lineage I) involves marked ventral curvature of the terebra (Figs 10C-D). The mandibles of Mantica and female Manticora are otherwise very amblycheiline-like, but those of males are remarkably hypertrophied, with an especially elongate incisor tooth. Mandibular sexual dimorphism developed in numerous lineages throughout the phylogenetic history of tiger beetles, but became most pronounced in the lineage leading to Manticora, in which the males developed tremendously elongate mandibles, especially in the incisor region. Mandibular dimorphism likely developed in concert with prolonged copulation, and mate-guarding and the fitting of the male mandible to the female metathoracic copulatory sulcus (Freitag, 1974). Oberprieler and Arndt (2000: 75–76) report that even the hypertrophied mandibles of these beetles function without any apparent awkwardness during tandem locomotion, and thus their allometric scaling appears appropriate for this purpose. The anterior margin of the labrum becomes shallowly emarginate, and develops four or six crenulations.

Lineage J Evolution of the Megacephalini and Collyridini + Cicindelini (Lineages K-N). Mandibles of this lineage develop a more or less extensive diastema between terebral tooth 1 and anterior margin of the retinaculum (Figs 11A-F – 13A-F, od), and retain the scrobal setae in a few genera (e.g., Megacephala, Pogonostoma). The epipharynx undergoes slight differentiation with the row of the anterior parapedial setae extended anteriorly (Figs 13D, F; 17F).

Lineage K Evolution of the mandibles of Megacephalini includes development of a supplementary retinacular tooth (Figs 11A-F, srt), and hence a longer retinaculum. Within this lineage, the terebral teeth differentiate in number (one to three) among taxa, and between sexes of the same taxon (cf. Pearson and Vogler, 2001:}
Plate 2. Digital images of head capsule, labrum, and mandibles dorso-frontal aspect, of: A *Ctenostoma ichneumoneum* Déjean; B *Therates erinnys* Bates; C *Cheiloxya binotata* Laporte de Castelnau; D *Pseudoxycheila* species?; E *Dromica junodi* Péringuey; F *Cicindela longilabris* Say. Scale bars: A = 3 mm; B, E = 4 mm; C, D, F = 5 mm.
Mandibles and labrum-epipharynx of tiger beetles

273, Fig. B-12). The number of labral setae is reduced from 10–12 to four. The anterior margin differentiates from slightly projected medially to virtually truncate (Fig. 13C). The epipharynx undergoes slight differentiation with the row of anterior parapedral setae extended anteriorly (Fig. 13D, \textit{aps}).

Lineage \textbf{L} The common ancestor of the tribes Collyridini and Cicindelini. Compared to those of its adelphotaxon (the Megacephalini), the mandibles (Figs 12A-F – 16A-F) become slender and labrum-epipharynx is extensively enlarged. The labral anterior margin becomes crenate, with the setae inserted apically rather than dorsally (Figs 13E and 17A, C). Since the arboreal members of the group evolved from ground-dwelling ancestors, tiger beetles serve as an example of the taxon pulse hypothesis (Erwin 1985) although the primarily terrestrial genus \textit{Cicindela} appears to have evolved from arboreal ancestors (Vogler and Barraclough 1998).

Lineage \textbf{M} The Collyridini. The mandibular retinaculum becomes more complex, with one or more additional cusps (Figs 12B, D, F). Within the tribe, the mandibles become markedly varied. We illustrate only what must be a relatively basal lineage—the genus \textit{Pogonostoma}, with only two terebral teeth on both left and right mandible, and only one additional retinacular cusp on only one (right) mandible. Scrobal setae are lost in all collyridine lineages examined with the exception of \textit{Pogonostoma}. Terebral teeth are reduced in number in, or lost from, the genera \textit{Ctenostoma} Klug, \textit{Tricondyla} Latreille, and \textit{Collyris} Fabricius.

Lineage \textbf{N} The Cicindelini. (Observations based principally on exemplar specimens representing the genus \textit{Therates} Latreille, \textit{Oxycheila} Dejean, and \textit{Cicindela} Linnaeus, arranged in evolutionary order of appearance, according to the reconstructed phylogeny of Vogler and Barraclough). Also examined: \textit{Oxygonia gloriola} Bates.

Within this tribe, and as in the Collyridini, the mandibles become markedly varied, as shown in Figs 14A-F, 15A-F and 16A-F. The scrobe, narrow in the more basal lineages, is lost from the more highly derived \textit{Cicindela}. The ancestral number of terebral teeth was probably three as in \textit{Cicindela} (Figs 16A-F), becoming four as in \textit{Oxycheila} (Figs 15A-F, \textit{tt 1 - tt 4}), and reducing to one, as in \textit{Therates} (Figs 14A-F, \textit{tt 2}). The supplementary retinacular tooth re-evolves in \textit{Therates} (Figs 14A-F, \textit{srt}). The ventral groove, normally quite long (Figs 15E-F, \textit{vg}), becomes shortened in \textit{Cicindela longilabris} (Figs 16E-F, \textit{vg}), and in other species of this genus (Pearson and Vogler, 2001: 198, Fig. 105). The position of the labral setae, though preapical in the more basal cicindelines, shifts back to the dorsal surface in the more recently evolved genus \textit{Cicindela}. Also, in this genus, the number of labral setae in some species is markedly increased, and in others decreased from the ancestral cicindelike 10–12, giving an overall range of 4 to 36. The labrum-epipharynx, quite long in the earlier-evolved cicindeline lineages (Figs 17A, C), becomes shortened in \textit{Cicindela} (Fig. 17E).

Vogler and Barraclough (1998) argue that rate of diversification (based on numbers of extant species) increased from the basal amblycheilines and manticorines to
the more derived megacephalines and collyridines, and then to an even greater extent in the cicindelines. They attributed this pattern to the broad geographic ranges of the pantropical collyridines and the cosmopolitan megacephalines and cicindelines, and to the role of collyridines and cicindelines as large-eyed, diurnal, visual predators. In our work, a similar pattern was observed, with a narrow range of mandible and labrum-epipharynx structure among the basal taxa, and a broad range, including increased complexity of the preoral mill, among the more derived taxa. It is tempting to suggest that mouthpart evolution also played a part in the diversification of the higher cicindelites, but we are also curious whether the currently restricted geographic ranges and low diversity of the amblycheilines and manticorines might be better interpreted as relictual.

Conclusions, and suggestions for further research

Building on increasingly sophisticated phylogenetic hypotheses for the Geadephaga, and the Cicindelitae (e.g., Maddison et al., 1999), as well as the functional morphology of Evans (1965) and Evans and Forsythe (1985), we are able here to propose a system of names and homologies for the structures of the tiger beetle mandibles and labrum-epipharynx, and to map hypothesized evolutionary changes in these structures on a generalized tree for the group. We propose relatively few multiple gains or losses of features, as follows: scrobal setae are lost in some but not all amblycheilines, megacephalines except Megacephala, and collyridines except Pogonostoma; the supplementary retinacular tooth evolves twice, in the megacephalines and the cicindeline genus Therates; terebral teeth are lost in many collyridines and the cicindeline genus Therates, terebral teeth increase in number in many cicindelines; and the labrum acquires the pleiotypic shortened condition in some cicindelines.

Further research would benefit from the examination of additional taxa, both within the Cicindelitae and among other relatively basal lineages of the Carabidae, in order to better address questions such as: is the Carabidae indeed the sister group of the Cicindelitae, and is Platychile more closely related to the amblycheilines or to the megacephalines? Incorporation of mouthpart features as character systems in phylogenetic analyses is recommended, as are further studies of the biomechanics of tiger beetle feeding and the use of mandibles during mating and mate-guarding, since the evolutionary changes we hypothesize are difficult to interpret without an appreciation of the functional consequences of changes in mouthpart configuration. In this light, further studies are also needed to adequately characterize sexual dimorphism, asymmetry, and chirality among tiger beetle mouthparts.
**Acknowledgements**

Through the good offices of Terry Thormin and Matthias Buck, The Royal Alberta Museum, Edmonton, Alberta, loaned to us the specimens used to launch our study of tiger beetle mouthparts. We are grateful to these colleagues for arranging this loan. We thank George Braybrook and De-ann Rollings (Department of Earth and Atmospheric Sciences, University of Alberta) for capturing the digital SEM images of our exemplar specimens. For a prompt response to our last-minute loan request for specimens of *Platychile pallida*, we thank David H. Kavanaugh (California Academy of Sciences, San Francisco, California); and for providing names for several Neotropical cicindeline species used in the present study, we thank Ronald L. Huber (Bloomington, Minnesota). Finally, we thank James K. Liebherr (Department of Entomology, Cornell University, Ithaca, New York) and David L. Pearson (Department of Zoology, Arizona State University, Tempe, Arizona) for insightful comments on earlier drafts of this paper. Publication was made possible by Natural Sciences and Engineering Research Council of Canada, NSERC Research Grant OGP 1399, to GEB.

**References**

Acorn JH, Ball GE (1991) The mandibles of some adult ground beetles: structure, function, and the evolution of herbivory (Coleoptera: Carabidae). Canadian Journal of Zoology 69: 638–650. doi: 10.1139/z91–094

Arndt E, Putchkov AV (1997) Phylogenetic investigation of Cicindelidae (Insecta, Coleoptera) using larval morphological characters. Zoologischer Anzeiger 235: 231–241.

Ball GE, Kavanaugh DH, Moore BP (1995) *Sugimotoa parallela* Habu (Coleoptera, Carabidae, Lebiini): redescription, geographical distribution, and relationships based on cladistic analysis of adult structural features. Japanese Journal of Coleopterology, Special Bulletin No. 4: 275–311.

Ball GE, Bousquet Y (2001) Family 6. Carabidae, in Arnett RH and Thomas, MC (eds.) American Beetles, Volume 1. CRC Press, Boca Raton,/ London/ New York/ Washington, DC. pp. 32–133,

Bell RT (1960) A revision of the genus *Chlaenius* Bonelli (Coleoptera: Carabidae) in North America. Miscellaneous Publications of the Entomological Society of America 1: 97–171.

Bell RT (1966) *Trachypachus* and the origin of the Hydradephaga (Coleoptera). Coleopterists Bulletin 20:107–112.

Bell RT (1994) Beetles that cannot bite: Functional morphology of the head of adult rhyssonines (Coleoptera: Carabidae or Rhysodidae. The Canadian Entomologist 126: 667–672. doi: 10.4039/Ent126667-3
Bell RT (1998) Where do the Rhysodini belong, in Ball GE, Casale A, Vigna Taglianti A (eds.) Phylogeny and Classification of Caraboidea. XX International Congress of Entomology (1996, Florence, Italy) Museo Regionali di Scienze Naturali, Torino, p. 261–272.
Bell RT, Bell JR (1962) The taxonomic position of the Rhysodinae. The Coleopterists Bulletin 15: 99–106.
Beutel RG (1998) Trachypachidae and the phylogeny of Adephaga (Coleoptera) In Ball GE, Casale A, Vigna-Taglianti A (eds), Phylogeny and classification of Caraboidea (Coleoptera: Adephaga). Atti, Museo Regionale di Scienze, Torino. pp. 81–106.
Beutel RG, Haas A (1996) Phylogenetic analysis of larval and adult characters of Adephaga (Coleoptera) using cladistic computer programs. Entomologica Scandinavica 27: 197–205. doi: 10.1163/187631296X00043
Boyd HP, Rust RW (1982) Intraspecific and geographic variations in Cicindela dorsalis Say (Coleoptera: Cicindelidae). Coleopterists Bulletin 36: 221–239.
Cassola F (2001) Studies of tiger beetles CXXIII Preliminary approach to the macrosystematics of the tiger beetles (Coleoptera: Cicindelidae). Russian Entomological Journal 10: 265–272.
Cazier MA (1954) A review of the Mexican tiger beetles of the genus Cicindela (Coleoptera, Cicindelidae). Bulletin of the American Museum of Natural History 103: 231–309.
Csiki E (1903) Die Cicindeliden Ungarns. Mathematische und Naturwissenschaftliche Berichte aus Ungarn 18: 121–144.
Deuve T (2004) Illustrated Catalogue of the Genus Carabus of the World. Pensoft Sofia-Moscow x + 461 pp.
Dressler C, Beutel RG (2010) The morphology and evolution of the adult head of Adephaga (Insecta: Coleoptera). Arthropod Systematics and Phylogeny 68 (2): 239–287.
Erwin TL (1985) The taxon pulse: a general pattern of lineage radiation and extinction among carabid beetles, in Ball GE (Ed.) Taxonomy, Phylogeny and Zoogeography of Beetles and Ants. A Volume Dedicated to the Memory of Philip Jackson Darlington, Jr. (1904–1983) Dr. W. Junk, Publishers Dordrecht/ Boston/ London pp. 437–472.
Erwin TL (2007) A Treatise on the Western Hemisphere Caraboidea (Coleoptera), Their Classification, Distribution, and Ways of Life. Volume I Trachypachidae, Carabidae--Nebriiformes 1). Pensoft, Sofia- Moscow, p. 171.
Erwin TL, Pearson DL (2008) A Treatise on the Western Hemisphere Caraboidea (Coleoptera), Their Classification, Distribution, and Ways of Life. Volume II (Carabidae--Nebriiformes 2--Cicindelitae). Pensoft, Sofia- Moscow, p. 18.
Evans MEG (1965) The feeding method of Cicindela hybrida L. (Coleoptera: Cicindelidae). Proceedings of the Royal Entomological Society (A) 40: 61–66. doi: 10.1111/j.1365-3032.1965.tb00314.x
Evans MEG, Forsythe TG (1985) Feeding mechanisms, and their variation in form, of some adult ground beetles (Coleoptera: Caraboidea). Journal of Zoology, London (A). 206: 113–143. doi: 10.1111/j.1469-7998.1985.tb05640.x
Forsythe TG (1983) Mouthparts and feeding of certain ground beetles (Coleoptera: Carabidae). Zoological Journal of the Linnean Society, 79: 319–376. doi: 10.1111/j.1096-3642.1983.tb01170.x
Mandibles and labrum-epipharynx of tiger beetles...

Franzen M, Heinz W (2005) Morphology, genitalia, and natural history notes on the enigmatic tiger beetle, Mantica horni Kolbe, 1896 (Coleoptera, Cicindelidae). Bonner zoologische Beiträge 53 (2004): 297–301.

Freitag R (1974) Selection for a non-genitalic mating structure in female tiger beetles of the genus Cicindela (Coleoptera: Cicindelidae). Canadian Entomologist 106: 561–568. doi: 10.4039/Ent106561-6

Galián J, Hogan JE, Vogler AP (2002) The origin of multiple sex chromosomes in tiger beetles. Molecular Biology and Evolution, 19 (10) 1792–1796.

Ganeshaiah KN, Belavadi VV (1986). Habitat segregation in four species of adult tiger beetles (Coleoptera: Cicindelidae). Ecological Entomology 11: 147–154. doi: 10.1111/j.1365-2311.1986.tb00289.x

Gissler CF (1879) The anatomy of Amblychila cylindriformis Say. Psyche 2: 233–244.

Horn W (1893) Bemerkungen und Nachträge zum “Catalogue systématique des Cicindelidae” par Fleutiaux (1893). Deutsche Entomologische Zeitschrift 1893: 321–347.

Horn W (1907) Brule’s “Odonotochila aus dem Baltischen Bernstein” und die Phylogene der Cicindeliden (Col.). Deutsche Entomologische Zeitschrift 1907: 461–466.

Horn W (1908) Coleoptera, Adepaha; Family Carabidae, Subfamily Cicindelinae. Genera Insectorum. P. Wytsman, Bruxelles. XIII, fasc. 82A, pp. 1–486.

Huber RL (1994) A new species of Tetracha from the west coast of Venezuela with comments on genus-level nomenclature (Coleoptera: Cicindelidae). Cicindela 26: 49–75.

Kavanaugh DH (1978) The Nearctic species of Nebria Latreille (Coleoptera: Carabidae: Nebriini): classification, phylogeny, zoogeography, and natural history. PhD thesis, University of Alberta, Edmonton, Alberta. 1041 pp.

Kavanaugh DH (1998) Investigations of phylogenetic relationships among some basal grade Carabidae (Coleoptera): a report upon work in progress. In Ball GE, Casale A, Vignataglianti A (eds), Phylogeny and classification of Caraboidea (Coleoptera: Adepaha). Atti, Museo Regionale di Scienze, Torino. pp. 329–342.

Kritsky G, Simon S (1995) Mandibular sexual dimorphism in Cicindela Linnaeus (Coleoptera: Cicindelidae). Coleopterists Bulletin 49: 143–148.

Leonard JG, Bell RT (1991) Northeastern Tiger Beetles: A Field Guide to Tiger Beetles of New England and Eastern Canada, CRC Press, Boca Raton/ London/ New York/ Washington, DC. xii + 176 pp.

Liebherr JK, Will KW (1998) Inferring phylogenetic relationships within Carabidae (Insecta, Coleoptera) from characters of the female reproductive tract. In Ball GE, Casale A, Vignataglianti A (eds), Phylogeny and classification of Caraboidea (Coleoptera: Adepaha). Atti, Museo Regionale di Scienze, Torino. pp. 107–170.

Lorenz W (2005) A Systematic List of extant ground beetles of the World ( Coleoptera : “Ge- adepaha”: Trachypachidae and Carabidae incl. Paussinae, Cicindelinae, Rhysodinae). 2nd edition (Tutzing: published by the author). 530 pp.

Maddison DR, Baker MD, Ober KA (1999) Phylogeny of carabid beetles as inferred from 18S ribosomal DNA (Coleoptera: Carabidae). Systematic Entomology 24: 103–138. doi: 10.1046/j.1365-3113.1999.00088.x
Makarov KV (2008) Larval chaetotaxy in the genus *Rhysodes* Dalman, 1823 and the position of Rhysodidae within the Adephaga (Coleoptera), in Penev L, Erwin TL, Assman T (eds), Back to the Roots and back to the Future, Towards a new Synthesis amongst Taxonomic, Ecological And Biogeographical Approaches to Carabidology. Proceedings of the XIII European Carabidologists Meeting, Blagovgrad, August 20–24, 2007. Pensoft Publishers, Sofia- Moscow. pp. 101–123.

Madge RB (1989) A catalogue of the family-group names in the Geodephaga, 1758–1985 (Coleoptera: Carabidae s. lat.). Entomologica Scandinavica 19: 459–474. doi: 10.1163/187631289X00573

Mury Meyer EJ (1987) Asymmetric resource use in two syntopic species of larval tiger beetles (Cicindelidae). Oikos +50: 167–175. doi: 10.2307/3565997

Niemelä J, Ranta E (1993) World-wide tiger beetle mandible length ratios: was something left unmentioned? Annales Zoologicae Fennici 30: 85–88.

Oberprieler RG, Arndt E (2000) On the biology of *Manticora* Fabricius (Coleoptera: Carabidae: Cicindelinae) with a description of the larva and taxonomic notes. Tijdschrift voor Entomologie 143: 71–89.

Pearson DL (1980) Patterns of limiting similarity in tropical forest tiger beetles (Coleoptera: Cicindelidae). Biotropica 12: 195–204. doi: 10.2307/2387971

Pearson DL, Blum MS, Jones TH, Fales HM, Gonda E, White BR (1988) Historical perspective and the interpretation of ecological patterns: defensive compounds of tiger beetles (Coleoptera: Cicindelidae). The American Naturalist 132: 404–416. doi: 10.1086/284860

Pearson DL, Mury EJ (1979) Character divergence and convergence among tiger beetles (Coleoptera: Cicindelidae). Ecology 60: 557–566. doi: 10.2307/1936076

Pearson, DL, Juliano SA (1991) Mandible length ratios as a mechanism for co-occurrence: evidence from a world-wide comparison of tiger beetle assemblages (Cicindelidae). Oikos. 61 (2): 223–233. doi: 10.2307/3545340

Pearson DL, Vogler AP (2001) Tiger Beetles: The Evolution, Ecology, and Diversity of the Cicindelids. Cornell University Press, Ithaca and London, xiii + 333 pp.

Richardson RK (2010) Mandibular chirality in tiger beetles (Carabidae: Cicindelidae). The Coleopterists Bulletin 64: 386–387. doi: 10.1649/0010-065X-64.4.386

Satoh A, Hori M (2004) Interpopulation differences in the mandible size of the coastal tiger beetle *Lophyridia angulata* associated with different sympatric species. Entomological Science 7: 211–217. doi: 10.1111/j.1479-8298.2004.00065.x

Satoh A, Uéda T, Enokido Y, Hori M (2003) Patterns of species assemblages and geographical distributions associated with mandible size differences in coastal tiger beetles in Japan. Population Ecology 45: 67–74. doi: 10.1007/s10144-003-0141-0

Vogler, AP, Barraclough TG (1998) Reconstructing shifts in diversification rate during the radiation of Cicindelidae (Coleoptera). In Ball GE, Casale A, Taglianti AV (Eds) Phylogeny and Classification of Caraboidea (Coleoptera: Adephaga), Museo Regionali di Scienze, Torino (Italia ), 251–260.

Werner K (2000) The tiger beetles of Africa. Volume 2 (Coleoptera: Cicindelidae). Taita Publishers, Hradec Kralove. 208 pp.
Appendix

List of names of tiger beetle taxa examined with Wild M3 and M5 stereomicroscopes for mandibular and/or labral-epipharyngeal features.

incertae sedis

*Platychile pallida* (Fabricius, 1801) (2 males)

Tribe Manticorini

*Manticora mygaloides* Thomson, 1859 (2 males, 2 females)
*Manticora latipennis* Waterhouse, 1837 (1 male, 1 female)
*Manticora tuberculata* (DeGeer, 1778) (3 males, 2 females)

Tribe Megacephalini

*Megacephala regalis* Boheman, 1848 (1 male)
*Phaeoxantha tremolerasi* (W. Horn, 1909) (1 female)
*Phaeoxantha wimmeri* Mandel, 1958 (1 male, 1 female)
*Tetracha annuligera* Lucas, 1857 (1 male, 1 female)
*Tetracha virginica* (Linnaeus, 1767) (1 male, 1 female)

Tribe Collyridini

*Ctenostoma metallicum* (Laporte de Castelnau, 1834) (1 male)
*Ctenostoma unifasciatum* Dejean, 1831 (1 male, 1 female)
*Ctenostoma ichneumoneum* Dejean, 1826 (1 female)
*Collyris dohrnii* Chaudoir. 1860 (1 female)
*Tricondyla* species? (3 males)

Tribe Cicindelini

*Therates erinnys* Bates, 1874 (1 male, 1 female)
*Pseudoxycheila ceratoma* Chaudoir, 1865 (1 male)
*Cheiloxya binotata* Laporte de Castelnau, 1833 (1 male)
*Dromica junodi* Péringuey, 1892 (2 males, 1 female)
*Oxygonia gloriola* Bates, 1872 (= *O. simplicipennis* Horn) (1 male)