BRAZILIAN SPECIES OF LUMINESCENT ELATERIDS: BIOCHEMISTRY AND BIOLOGY

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The three main families of luminescent insects are widespread in Brazil, that is, Lampyridae (fireflies), Elateridae (firebeetles) and Phengodidae (railroad worms). In the past three decades, biological and biochemical research on luminescent insects have focused principally the lampirids, with very little effort being directed towards the other two families (Colepiccolo Neto et al., 1986). Taking advantage of the abundant distribution of elaterids (click-beetles) in Brazil, we decided to initiate a wide program of investigation on this family which includes the habitat, in vivo and in vitro bioluminescent spectra, luciferin/luciferase system, anti-oxidant enzymes, extra-corpooreal digestion, and the inquinism termite/Pyrearinus termitilluminans (Colepiccolo Neto, 1986; Bechara, 1988). Up to now, we have studied 14 species of luminescent Elateridae: Hapsoedrilini: Hapsoedrillus pyrotis, H. ignifer, H. sp.; Pyrophorini: Pyrophorus divergens, P. punctatissimus, P. sp., Pyrearinus candelarius, P. candens, P. micatius, P. janus, P. lineatus, P. termitilluminans, Opnelerter pyrophanus, Pyronates maculicollis.

In the larval stage (18-20 months), elaterids are predaceous and live into decaying logs, except P. termitilluminans which inhabits tunnels excavated into termite mounds found in the “cerrados” of Central Brazil (the “luminous termite hills”). Field and lab observations indicated that light emission by larval elaterids serves for prey attraction and, possibly, also defense purposes. The prothorax is the brightest larval segment. Adults of most luminescent elaterids (female, 3 months; male, 2 months) possess two light organs placed laterally on the dorsal prothorax, which emit green light continuously when at rest, walking or handled, and a larger ventral lantern (yellow light) hidden in a cleft in the first abdominal segment, which is seen only when forcibly exposed or during the nuptial flight (mating). Eggs and pupae of many elaterids described here emit a constant dim light. Adult and larval elaterids have been collected and successfully reared in laboratory.

Thin layer chromatography in five different elution conditions revealed that the luciferin extracted from 13 elaterid species, as well as that from Phengodes sp. and Phrixothrix sp. (both Phengodidae), at different stages of development and from either the thoracic or abdominal lanterns, is same as that of firefly luciferin (Colepiccolo Neto et al., 1986a; Colepiccolo Neto & Bechara, 1984). The biosynthesis of luciferin in larval P. termitilluminans was studied by injection of 14C-cystine—a putative luciferin precursor—in their abdominal segments followed by isolation and counting of labelled luciferin (Colepiccolo Neto et al., 1988). Dose and time response were observed; the radiochemical yield of luciferin reached a plateau at 3%, one day after the precursor incorporation. The observed temporal response to injected 14C-cystine is similar to that previously described for fireflies from a benzothiazol derivative, suggesting that the condensation of a second cystein moiety to the benzothiazol ring to form luciferin is the rate-limiting step.

In vivo and in vitro bioluminescence spectral data were recorded for 12 species of Elaterids in distinct stages of their life cycle. In adults, the emission color from the abdominal light organ is always shifted (20-40 nm) towards the red relative to that from the prothoracic lanterns (525-560 nm) (Colepiccolo Neto et al., 1986a). The peak wave length for either in vivo and in vitro spectrum is characteristic for a given species and depends on the metamorphic stage. This finding raises a potential use of the bioluminescence spectrum as a tool for taxonomy of luminescent elaterids, which can hardly be differentiated by morphological criteria when in the larval phase (Casari-Chen & Costa, 1986). The spectral data collected for these species, together with determination of Km for both ATP and luciferin using crude extracts, suggested the occurrence of luciferase isozymes in the abdominal and thoracic lanterns and in the four stage of their life cycle. The

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bioluminescence intensity from *P. termittillumina*ns eggs and larvae increases sharply upon increasing the temperature, with an apparent activation (Ea) energy of ca 25 kcal mol⁻¹ (in vitro, Ea ≈ 16 kcal mol⁻¹). Coincidently or not, this Ea value is within the range expected for dioxetane and dioxetanone thermal cleavage, which are putative intermediates of most chemical and bioluminescent reactions (Adams & Zinner, 1982).

Superoxide dismutase (SOD) activities were compared in crude extracts from larvae of luminous click-beetles (Colepicolo Neto et al., 1986b). The SOD levels were found to be 5- (P. divergens and an unidentified Pyrophorinae) and 15- (P. termittillumina*ns*) fold higher than in the non-luminous Elateridae (*Platycrepidus bicinctus*, *Chalcolepidus zonatus*, *Iscidiondus sp.*, and *Conoderus sp.*). It should be noted that the former two and the latter species live in decaying logs where oxygen tension drops to 2.5%, whereas larval *P. termittillumina*ns** dwell well-aerated galleries (20% oxygen) dug into termite mounds. It was also found that the brightest larval segments, that is, the prothorax and the 9-10th abdominal segments, exhibit two-fold higher SOD activities than the dim meso- and metathorax and lst-8th abdominal segments. SOD activities as high as those observed in luminous elaterids (8-30 U/mg protein) were also found in *Biceollonycha* sp. (Lampyride; 10 U/mg protein) and in *Phengodes* sp. (Phengodidae; 30 U/mg protein). These results were interpreted in terms of the need of SOD protection against deleterious effects of active oxygen species arising from storage of molecular oxygen to sustain the bioluminescent reaction.

Pre-oral digestion occurs in predaceous Elateridae larvae. When biting the prey, larval luminescent elaterids regurgitate a dark liquid buffered at pH ca 7.3, containing several hydrolases. Amylase, cellulase, beta-glucosidase and trypsin were found in major amounts in the midgut lumen and in regurgitated material of *P. termittillumina*ns larvae, whereas aminopeptidase, alpha-glucosidase and trehalase occur mainly in the midgut tissue (Colepicolo Neto et al., 1986c). The optimum pH of these enzymes varies in the range 6.0-8.0. About 40% of the cell bound enzymes are membrane associated. The properties of the pre-digestive enzymes from *P. termittillumina*ns and *P. divergens* determined by electrophoretic, isoelectric focusing, density-gradient centrifugation and kinetic procedures were found to be very similar (Colepicolo Neto et al., 1986c; 1987). These results compared with data obtained from *Rhynochosiara americana* and *Trichosia pubescens* (both Diptera), phylogenetically distant from the elaterids but living under decaying plants, suggest that the insect digestive enzymes reflect more phylogenetic traits than the adaptation to different habitats (Colepicolo Neto et al., 1987).

Regarding the "luminous termite mounds", first reported by German naturalists in the end of last century and recently described by Redford (1982), we have tentatively explained the inquilinism between termites and the larval *P. termittillumina*ns (Bechara, 1988). The "luminous mounds" are primarily built by *Cornitermes cumulans*, followed by their infestation by other termite species (mainly *Paracornitermes*, *Spinitermes* and *Embritermes*), ants, larval Diptera, larvae and adults of Coleoptera (*P. termittillumina*ns, *Tenebrionidae*, *Derestidae*, *Scarabaenidae*, *Cinclidaea*, *Cantharidae* and *Carabidae*), spiders, scorpions, centipedes, millipedes, and opilios. During the raining season (October through December), after sunset, the *P. termittillumina*ns larvae (ca 200-400 per mound) expose their head and prothorax at the mouth of tunnels and emit intense green light (peak intensity at 537 nm) to attract and catch alate prey, principally flying (adult) termites. Hundreds of luminous termite mounds illuminating the open fields can be admired specially under new moon. Field observations and polymer-injection molds of the larval galleries show that they dwell an intricate network of tunnels in the external layer (0.5-1 cm) of the mounds. Some tunnels branch perpendicularly toward the interior (8-10 cm), bend to the exterior turning U-shaped, and enlarge at the depth of ca 1.5 cm to form a small side chamber or atrium where the prey is consumed. In March through May, the mound basis is surrounded by a green halo due to thousands of first instar larvae eclosed from the eggs laid during late Spring and living on the ground/mound basis.

Work is in progress to enrich our collection of bioluminescence spectra, to isolate and characterize the luciferases, to elucidate the biosynthesis pathway of the luciferin, to better understand the oxyradical metabolism and its connection with bioluminescence, and to explore possible applications of the luciferin/luciferase system.
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REFERENCES


