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Nutrient Flux Associated with the Emergence of *Quesada gigas* Olivier (Hemiptera: Cicadidae) in an Urban Ecosystem

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Abstract

Large-bodied arthropods, such as cicadas, can be able to reallocate significant amounts of nutrients during adult emergence. Evidence suggests that Quesada gigas Olivier emergence constitutes an important nutrient flux from belowground to aboveground. The purpose of this study was to estimate the amount of nitrogen, proteins, and lipids resulting from the emergence of *O. gigas* in an urban ecosystem in Central Brazil. Adult specimens captured from September to November 2006 were weighed and submitted to biochemical analysis. Population density was approximately 4,200 individuals per hectare. Mean individual dry mass was 1.03 g and contained 12.6% proteins, 8.4% lipids, and 5% nitrogen. Total biomass input from the species was 4.3 kg ha⁻¹ y⁻¹, with a consequent annual reallocation of approximately 545 g of proteins, 363 g of lipids, and 216 g of nitrogen per hectare. The data obtained suggest that Q. gigas emergence can cause significant translocation of nutrients from belowground to aboveground, and is therefore an important biological event for ecosystem function.

Introduction

Arthropods constitute the majority of herbivores and detritivores in many ecosystems (Seastedt & Crossley 1984), influencing the amount of living and dead organic matter in a given system and consequently, the nutrient cycling (Seastedt 1984, 1988, Naiman 1988). Previous studies suggested that nutrients, mass and energy released by arthropods activities do not represent significant resource pulses in ecosystems (Schowalter & Crossley 1983, Seastedt & Crossley 1984). However, other studies have indicated that arthropods play a major role in plant productivity and nutrient cycling processes (Seastedt & Tate 1981, Brown & Gange 1989, Bardgett & Wardle 2003, Whiles & Charlton 2006).

Insect emergence has more often been associated with nutrient flux in terrestrial and aquatic ecosystems

(Nakano *et al* 1999, Collier *et al* 2002, Sabo & Power 2002, Ballinger & Lake 2006), but practically nothing is known about the importance of these animals with respect to nutrient and energy fluxes from belowground to aboveground in terrestrial ecosystems (Callaham Jr *et al* 2000).

In several cicada species emergence can be viewed as a resource pulse, available at high densities for a short period of time (Ostfeld & Keesing 2000). Due to their relatively large bodies, these arthropods are able of reallocate significant amounts of matter and energy during adult emergence (Callaham Jr *et al* 2000, Whiles *et al* 2001, Whiles & Charlton 2006). In addition, they cause an important translocation of nutrients from belowground to aboveground because cicada nymphs feed on plant root xylem whereas adults are aboveground feeders (Young 1980, Williams & Simon 1995).

Studies on the role of subterranean herbivorous insects, including cicadas, in tropical regions are scarce, and are virtually unknown in South American species (Blossey & Hunt-Joshi 2003). *Quesada gigas* Olivier is an annual cicada species with a wide geographic distribution (Aoki *et al* 2010). Adults have a large body size (~ 46 mm) and emerge at relatively high densities, suggesting that this species transfers nutrients and energy from belowground to aboveground.

The purpose of this study was to estimate the level of total nitrogen, total soluble proteins and total lipids associated to *Q. gigas* tissues which would correspond to their translocation from belowground to aboveground during species emergence in an urban ecosystem.

Material and Methods

Specimens were collected from September to November 2006 from six randomly selected 2,500 m² plots (1.5 ha in total). These plots were located in cultivated areas harboring native and introduced plant species, distributed among lawns and buildings at the campus of the Universidade Federal de Mato Grosso do Sul (UFMS), Campo Grande (MS), Brazil (20°29"59'S, 54°36"42'W). Climate in the region is Köppen's Aw (tropical savanna) subtype (Köppen 1948), with two well-defined seasons: dry and cold from May to September and rainy and hot from October to April. Average annual rainfall is 1,532 mm and relative air humidity is generally low, rarely reaching 80%. Average annual temperature range from 20°C to 22°C (Embrapa-CNPGC 1985).

To estimate species density, all *Q. gigas* exuviae found in the plots were manually collected every three days. Although other species of cicada emerge at the same time, morphological differences between exuviae ensure their correct identification. Adults were collected during the emergence period with the aid of an entomological net. These specimens were frozen, then weighed on a precision balance (0.01 g) and processed for biochemical analysis. Dry weight was determined after the specimens had remained in an incubator for four days at 50°C.

To estimate protein content, 0.4 g of dry material was homogenized in 20 ml of phosphate buffer (0.1 M, pH 7.0), centrifuged (1.850 g at 10°C for 30 min) and filtered through a 10 μ m nylon mesh. Soluble protein content was determined according to Bradford (1976), using bovine serum albumin as the standard. Extraction of total lipids was carried out in a Soxhlet apparatus for 1h at 105°C, using petroleum ether. The solvent was subsequently evaporated and cooled in a desiccator and the resulting material was weighed and used for total lipid determination (Pregnolatto & Pregnolatto 1985). Nitrogen content was measured using the Kjeldahl method

(Nelson & Sommers 1973), which involves digestion of the material followed by colorimetric analysis.

Results and Discussion

Emergence of *Q. gigas* on the UFMS Campus lasted for approximately 10 consecutive weeks, from early September to middle November. An average of 4,200 adults emerged per hectare (0.42 individuals/m²).

Mean adult fresh weight was 3.7 g (\pm 0.75), 72.3% (\pm 4.50) of which consisted of water. While seemingly a high percentage, this is within the 70-75% range typical for insects (Bell 1990). Mean dry weight was 1.0 g (\pm 0.25), comprising 12.6% (\pm 5.40) proteins, 8.4% (\pm 5.50) lipids, and 5% (\pm 0.90) nitrogen. Each adult consisted of approximately 129.8 mg of proteins, 86.5 mg of lipids, and 51.5 mg of nitrogen. The only previous investigation on cicadas which calculated protein and lipid content was the study by Brown & Chippendale (1973) on *Magicicada cassini* (Fisher). This species emerges at 17-year intervals and contains 51-110 mg of proteins and 19-58 mg of lipids per individual.

With regard to nitrogen content, Fagan et al (2002) conducted a comprehensive review of studies on a large number of orders of herbivorous insects and found a mean level of 9.6% (\pm 0.15%) of nitrogen per gram of dry weight, which falls within the 7-14% range reported earlier by Mattson (1980). While some of the reviewed studies were based on the Dumas combustion method for nitrogen quantification, the differences found in the concentrations of nitrogen were unlikely to have resulted from the choice of method (Dumas or Kjeldahl), since the results were sufficiently similar to allow for comparisons regardless the method adopted (Perez et al 2001, Watson & Galliher 2001). The relatively low level of nitrogen found in tissues of Q. gigas may stem from the low availability of nitrogen in their food source, since insects feeding on nitrogen-deficient matter usually have low nitrogen content in their tissues (Fagan et al 2002).

Despite the low percentage of nitrogen, adults of *Q. gigas* adults contribute with high amounts of nitrogen to the ecosystem due to their relatively large bodies. Each individual provides 14 folds more nitrogen than *Cicadetta calliope* (Walker) and twice as much as *Tibicen aurifera* Say and *M. cassini* (Callaham *et al* 2000, Whiles *et al* 2001), indicating the higher contribution neotropical cicadas may give aboveground by transferring nutrients from belowground as compared to other cicadas.

The estimated total dry mass associated with the emergence of Q. gigas was 4.3 kg ha^{-1} y^{-1} , with a consequent yearly reallocation of approximately 545 g of protein, 363 g of fat, and 216 g of nitrogen per hectare. Overall, the species reallocates up to 3% as much of the estimated nitrogen input for a neotropical savanna

area from precipitation, which is a major source for this nutrient in the ecosystem (Lima 1985).

Moreover, among soil-dwelling insects such as cicadas, emergence constitutes one of the few processes of nongaseous nitrogen flux from belowground to aboveground (Whiles et al 2001). In the same way that nitrogen translocation is effected by plants, nitrogen flux associated with the emergence of cicadas is not an input of "new" nitrogen into a system. Both nymphs and adults of Q. gigas obtain nutrients from the xylem of host plants. While plants are deprived of these nutrients for developing leaves, flowers, fruits, and seeds, all of which would be available aboveground, cicadas later serve as a food source for a wide variety of animals (e.g. spiders and insectivore birds) which do not feed directly on these plant structures. Furthermore, insect-derived proteins are preferred by birds because of their higher quality in relation to seed and fruit-derived proteins (Robbins et al 2005). These macronutrients are essential for growth and reproduction of most animals. Birds are key predators of cicadas in the study area, and a wide variety of species consume these cicadas (Aoki C., personal observation), similar to verified by Sazima (2009) at the urban Parque Ecológico Prof. Hermógenes F. Leitão Filho, in Campinas (SP). Lipid levels can affect clutch size (Ankney & Afton 1988), whereas the addition of proteins in the prereproductive diet may also augment the mass and size of eggs (Selman & Houston 1996, Williams 1996), which further influence on offspring survival (Bolton 1991, Magrath 1992, Blomqvist et al 1997). Given that most bird species breed in the rainy months (Matarazzo-Neuberger 1995, Sick 1997) and that Q. gigas emerges in the early period of this season, exhibiting scant anti predation behavior and producing no apparent toxic compounds, it may act as a key species during breeding season of local bird populations.

Quesada gigas emergence can also alter nutrient cycling ecosystem, since most nutrients released by this species reach at least one additional trophic level as they are consumed by a wide variety of predators, comprising both vertebrates (birds, mammals, and reptiles) and invertebrates (ants and spiders) (Aoki C., personal observation). Emergence can also alter the rate of nutrient cycling because decomposition of insect bodies usually occurs faster than does that of leaf litter, because degradation of complex organic matter such as plant cell walls are not involved (Schowalter et al 1986), allowing insects to be categorized as elements of the "fast cycle" of nutrient release (McNaughton et al 1988).

Although studies suggested that nutrients, mass and energy released by arthropods do not represent significant resource pulses in ecosystems (Schowalter & Crossley 1983, Seastedt & Crossley 1984, Wheeler *et al* 1992), our data show that *Q. gigas* represents an important pulse of resources from belowground

to aboveground, releasing essential nutrients for a wide range of secondary consumers and detritivores, and ultimately influencing nutrient cycling in the area investigated.

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References

- Ankney CD, Afton AD (1988) Bioenergetics of breeding Northern Shovelers: diet, nutrient reserves, clutch size, and incubation. Condor 90: 459-472.
- Aoki C, Lopes FS, Souza FL (2010) Insecta, Hemiptera, Cicadidae, *Quesada gigas* (Olivier, 1790), *Fidicina mannifera* (Fabricius, 1803), *Dorisiana viridis* (Olivier, 1790) and *Dorisiana drewseni* (Stål, 1854): first records for the state of Mato Grosso do Sul, Brazil. Check List 6: 162-163.
- Ballinger A, Lake PS (2006) Energy and nutrient fluxes from rivers and streams into terrestrial food webs. Mar Freshw Res 57: 15-28
- Bardgett RD, Wardle DA (2003) Herbivore mediated linkages between aboveground and belowground communities. Ecology 84: 2258-2268.
- Bell GP (1990) Birds and mammals on an insect diet: a primer on diet composition analysis in relation to ecological energetics. *Stud Avian Biol* 13: 416-422.
- Blomqvist D, Johansson OC, Götmark F (1997) Parental quality and egg size affect chick survival in a precocial bird, the Lapwing *Vanellus vanellus*. Oecologia 110: 18-24.
- Blossey B, Hunt-Joshi TR (2003) Belowground herbivory by insects: Influence on plants and aboveground herbivores. Annu Rev Entomol 48: 521-547.
- Bolton M (1991) Determinants of chick survival in the lesser black-backed gull: relative contributions of egg size and parental quality. *J Anim Ecol* 60: 949-960.
- Bradford MM (1976) A rapid e sensitive method for the quantitation of microgram quantities of protein utilizing the principle of protein-dye binding. Anal Biochem 72: 248-254.
- Brown JJ, Chippendale GM (1973) Nature and fate of the nutrient reserves of the periodical (17year) cicada. J Insect Physiol 19: 607-14.

- Brown VK, Gange AC (1989) Herbivory by soil-dwelling insects depresses plant species richness. Funct Ecol 3: 667-671.
- Callaham Jr MA, Whiles MR, Meyer CK, Brock BL, Charlton RE (2000) Feeding ecology and emergence production of annual cicadas (Homoptera: Cicadidae) in tallgrass prairie. Oecologia 123: 535-542.
- Collier KJ, Bury S, Gibbs M (2002) A stable isotope study of linkages between stream and terrestrial food webs through spider predation. Freshw Biol 47: 1651-659.
- Embrapa CNPGC (1985) Boletim agrometeorológico. Campo Grande. MS.
- Fagan W F, Siemann E, Mitter C, Denno R F, Huberty A F, Woods H A & Elser J J (2002) Nitrogen in insects: implications for trophic complexity and species diversification. Am Midl Nat 160: 784-802.
- Köppen W (1948) Climatologia. Fundo de Cultura Econômica. Buenos Aires. Trad de Guendriss du Klimakunde 1923p.
- Lima WP (1985) Ação das chuvas no ciclo biogeoquímico de nutrientes em plantações de pinheiros tropicais em cerradão. IPEF 30: 13-17.
- McNaughton SJ, Ruess RW, Seagle SW (1988) Large mammals and process dynamics in African ecosystems. Bioscience 38: 794-800.
- Magrath RD (1992) Roles of egg mass and incubation pattern in establishment of hatching hierarchies in the blackbird (*Turdus merula*). Auk 109: 474-487.
- Matarazzo-Neuberger WM (1995) Comunidade de cinco parques e praças da Grande São Paulo, estado de São Paulo. Ararajuba 3: 13-19.
- Mattson WJ (1980) Herbivory in relation to plant nitrogen content. Annu Rev Ecol Syst 11: 119-161.
- Naiman RJ (1988) Animal influences on ecosystem dynamics. BioScience 38: 750-752.
- Nakano S, Miyasaka H, Kuhara N (1999) Terrestrial-aquatic linkages: riparian arthropod inputs alter trophic cascades in a stream food web. Ecology 80: 2435-2441.
- Nelson DW, Sommers LE (1973) Determination of the total nitrogen in plant material. Agron J 65: 109-112.
- Ostfeld RS, Keesing F (2000) Pulsed resources and community dynamics of consumers in terrestrial ecosystems. Trends Ecol Evol 15: 232-237.
- Perez DV, Alcantara S, Arruda RJ, Meneghelli NDA (2001) Comparing two methods for soil carbon and nitrogen determination using selected Brazilian soils. Commun Soil Sci Plant Anal 32: 295-309.
- Pregnolatto W, Pregnolatto NP (1985) Métodos químicos e físicos para análises de alimentos. In Pregnolatto W, Pregnolatto NP (coord) Normas analíticas do Instituto Adolfo Lutz. 3ed., São Paulo, Instituto Adolfo Lutz, 533p.
- Robbins CT, Felicetti LA, Sponheimer M (2005) The effect of dietary protein quality on nitrogen isotope discrimination in mammals and birds. Oecologia 144: 534-540.

- Sabo JL, Power ME (2002) River–watershed exchange: effects of riverine subsidies on riparian lizards and their terrestrial prey. Ecology 83: 1860-1869.
- Sazima I (2009) Insect cornucopia: various bird types prey on the season's first giant cicadas in an urban park in southeastern Brazil. Biota Neotrop 9: 259-262.
- Schowalter TD, Crossley Jr DA (1983) Forest canopy arthropods as sodium, potassium, magnesium, and calcium pools in forests. For Ecol Manag 7: 143-148.
- Schowalter TD, Hargrove WW, Crossley Jr DA (1986) Herbivory in forested ecosystems. Annu Rev Entomol 31: 177-196.
- Seastedt TR (1984) The role of microarthropods in decomposition and mineralization processes. Annu Rev Entomol 29: 25-46.
- Seastedt TR (1988) Mass, nitrogen, and phosphorus dynamics in foliage and root detritus of tallgrass prairie. Ecology 69: 59-65.
- Seastedt TR, Crossley Jr DA (1984) The influence of arthropods on ecosystems. Bioscience 34: 157-161.
- Seastedt TR, Tate CM (1981) Decomposition rates and nutrient contents of arthropod remains in forest litter. Ecology 62: 13-19
- Selman RG, Houston DC (1996) The effect of prebreeding diet on reproductive output in zebra finches. Proc R Soc Lond B Biol Sci 263: 1585-1588.
- Sick H (1997) Ornitologia brasileira. Rio de Janeiro, Nova Fronteira, 912p.
- Watson ME, Galliher TL (2001) Comparison of Dumas and Kjeldahl methods with automatic analyzers on agricultural samples under routine rapid analysis conditions. Commun Soil Sci Plant Anal 32: 2007-2019.
- Wheeler GL, Williams KS, Smith KG (1992) Role of periodical cicadas (Homoptera: Cicadidae: Magicicada) in forest nutrient cycles. For Ecol Manag 51: 339-346.
- Whiles MR, Charlton RE (2006) The ecological significance of tallgrass prairie arthropods. Annu Rev Entomol 51: 387-412.
- Whiles MR, Callaham Jr MA, Meyer CK, Brock BL, Charlton RE (2001) Emergence of periodical Cicadas (*Magicicada cassini*) from a Kansas riparian forest: densities, biomass and nitrogen flux. Am Mid Nat 145: 176-187.
- Williams TD (1996) Variation in reproductive effort in female zebra finches (*Taeniopygia guttata*) in relation to nutrientspecific dietary supplements during egg laying. Physiol Zool 69: 1255-1275.
- Williams KS, Simon C (1995) The ecology, behavior and evolution of periodical cicadas. Annu Rev Entomol 40: 269-95.
- Young AM (1980) Habitat and seasonal relationship of some cicadas (Homoptera: Cicadidae) in Central Costa Rica. Am Mid Nat 103: 155-166.