Cytokinin-mediated leaf manipulation by a leafminer caterpillar

David Giron1,*, Wilfried Kaiser1, Nadine Imbault2 and Jérome Casas1

1Institut de Recherche sur la Biologie de l’Insecte, UMR-CNRS-6035, and 2EA2106 Biomolécules et Biotechnologies Ve´ga´tales, UFR Sciences et Techniques, Université Francois Rabelais, 37200 Tours, France
*Author for correspondence (david.giron@insect-tours.fr).

A large number of hypotheses have been proposed to explain the adaptive significance and evolution of the endophagous-feeding mode, nutritional benefits being considered to be one of the main advantages. Leaf-mining insects should feed on most nutritional tissues and avoid tissues with high structural and/or biochemical plant defences. This selective feeding behaviour could furthermore be reinforced by manipulating the plant physiology, as suggested by the autumnal formation of ‘green islands’ around mining caterpillars in yellow leaves. The question we address here is how such metabolic manipulation occurs and what the nutritional consequences for the insect are. We report a large accumulation of cytokinins in the mined tissues which is responsible for the preservation of functional nutrient-rich green tissues at a time when leaves are otherwise turning yellow. The analogy with other plant manipulating organisms, in particular gall-forming insects, is striking.

Keywords: leafminer; cytokinins; green island; nutrient acquisition; endophytic organisms

1. INTRODUCTION

Leaf mining is a means by which some plant-eating insects consume live foliage while simultaneously dwelling inside it. A large number of hypotheses have been proposed to explain the adaptive significance and the evolution of this endophagous-feeding mode. The mine presumably functions as a shelter from the detrimental effects of the physical environment, as a protection from attack by natural enemies, enables an increased development rate due to higher temperatures and is therefore the total surface available to later stages. The two following instars are selective tissue feeders.

Samples were collected early in autumn 2005 on 12-year-old apple (Malus domestica Borkh.) cv. ‘Allstar’s trees grown in a biologically managed orchard. Collected leaves were directly dissected on site, frozen on dry ice and stored at −80°C until analysis. Samples were controlled for leaf age and natural exposition in the orchard (Sun versus shade—position in the rows). Mined age (only one mine per leaf) was also controlled and corresponds to L4 tissue-feeding larvae.

(a) Cytokinin quantification in mined and non-mined tissues according to leaf senescence

Leaf samples were lyophilized and pulverized. Similar amounts of control, mined, ipsilateral and contralateral tissues were extracted overnight in aqueous methanol containing butylated hydroxytoluene as an antioxidant.

(i) Purification

Purification was performed using a nitrocellulose prefiler (4.5 μm, Sartorius, Germany) connected to an Oasis cartridge (Waters, USA) with a Teflon filter (0.2 μm, Sartorius, Germany) at the outlet.

(ii) Separation

Eluates were reduced by rotary evaporation, taken up with acidified water and injected into a reverse-phase HPLC column (Water, USA). After sample injection, the column was eluted at 1 ml min−1 with a convex gradient of acidified water and pure acetonitrile (HPLC grade) until the eluant was 100% acetonitrile. Retention time of cytokinins was determined by separate and pooled injections of pure zeatin (Z), ribosyl zeatin (ZR), isopentenyladenine (iP) and isopentenyl adenosine (iPA) (Sigma, USA) as standards. Fractions centred on the retention time of each cytokinin were collected and evaporated to dryness in a speed-vac concentrator (Savant, USA).
Quantification

Aliquots of each fraction corresponding to the retention time of each hormone standard were subjected to enzyme-linked immunosorbent assay (ELISA) as previously published (Jourdain et al. 1997). ELISA quantification could be done owing to a strict separation of each compound by HPLC fractionation. Optical densities were measured at 405 nm. See electronic supplementary materials for details.

(b) Nutrient quantification in mined and non-mined tissues according to leaf senescence

Quantification of the amount of nutrients in leaf samples was carried out using colorimetric techniques successfully used for plant material (Yemm & Willis 1954; Jones et al. 1989). Briefly, samples were lyophilized, weighed and ground. Nutrients were measured using a spectrophotometer after reaction with Anthrone reagent for sugars and Bradford reagent for proteins. See electronic supplementary materials for details.

(c) Statistical analysis

Statistical analyses were performed using R v. 2.3.0 software. Data were analysed by non-parametric Kruskal–Wallis tests followed by Behrens–Fisher post hoc tests.

3. RESULTS

(a) Cytokinin content of mined area exceeds that of non-infected or control areas

To evaluate the role of phytohormones on ‘green island’ formation, we compared cytokinin contents in mined and non-mined tissues (figure 2). As expected, cytokinin content decreases with leaf senescence, some compounds such as isopentenyladenine, falling below the detection level (green versus yellow controls). However, the global cytokinin content of mined areas far exceeded that of uninfected areas (ipsi- and contralateral) and controls (green and yellow uninfected leaves). This phenomenon is particularly true for isopentenyl adenine and isopentenyladenine, both compounds being at the beginning of the synthesis pathway of cytokinins. High levels of cytokinins are also observed in the near vicinity of the mine, ipsilateral tissues showing an amount of iP higher than contralateral tissues.

Figure 1. An apple tree leaf infected with the tentiform leafminer Phyllonorycter blancardella larva in autumn. The ‘Green island’ (feeding area) exhibits intact green chlorophyll-containing tissues, while the remaining leaf tissues undergo leaf senescence. The white spots on the mine are feeding windows, where all but the epidermis has been consumed by the caterpillar. Letters show areas used for cytokinin and nutrient content analysis: ‘a’ mined, ‘b’ ipsilateral and ‘c’ contralateral plant tissues.

(b) Nutrient content of mined tissues does not decrease in senescing leaves

We next conducted experiments investigating the nutritional benefits of the hormonal manipulation. Both food quality parameters decreased in senescing leaves falling from 135.56 ± 25.35 to 53.25 ± 4.27 µg mg⁻¹ for sugars (n=15) and from 29.10 ± 0.91 to 16.22 ± 1.67 µg mg⁻¹ for proteins (n=15; figure 3). Mined areas showed similar content of carbohydrates and proteins as uninfected green tissues. Carbohydrate and protein amounts in distant
insects themselves could potentially synthesize cytokinins. Large quantities of cytokinins have already been measured in labial glands of leafminers (Engelbrecht et al. 1969) and the ability to produce cytokinins has been shown in several plant-associated organisms like fungi and bacteria (Barry et al. 1984; Stevens & Berry 1988; Morris 1995).

During the senescence of apple tree leaves, the levels of proteins and carbohydrates decrease, but mined tissues displayed a similar level of both proteins and sugars as non-senescing uninfected green leaves. Owing to the maintenance of the photosynthetic apparatus within the mine—as a direct consequence of high levels of phytohormones—natural production of sugars and proteins ‘on site’ cannot be excluded. However, modifications of the water flow and accumulation of nutrient-attracting cytokinins in the mined area could favour the translocation of nutrients from the leaf to the mine as well. In any case, both processes result in a net accumulation of nutrients in the mined tissues with potential competition between the plant and the mine for nutrients originating from senescing cells (change in the source–sink relationships—Balibrea et al. 2004; Wingler et al. 2006).

The maintenance of functional green tissues is, of course, of considerable ecological value to the development of the larvae as it allows the insect to maintain a favourable nutritional environment in an otherwise degenerating context. The ‘green island’, enriched in cytokinins, enables the larvae to retain nutrients within the mine and to undergo a long development later in the season. Interestingly, remote parts of infected leaves have higher concentration of cytokinins than uninfested yellow leaves (except for iP). Yellow leaves also totally lack iPA, while this class of cytokinin is present in all tissues of infected leaves. Both results are a clear proof of manipulation of the plant physiology by the herbivore and over a much larger scale than the mine itself. The observed metabolic changes taking place in the mine and the manipulation of plant physiology induced by the insect strengthens the hypothesis that mines behave independently from the hosting leaf, operating a metabolic machinery of their own. Similar conclusions have been drawn with galling insects also known to produce galling tissues with high levels of cytokinins (e.g. Leicht 1994; Carol & Davies 2001). Despite great morphological differences between galls and mines, our results suggest a probable generalized process associated with the colonization of plants by endophagous organisms. Such intimate associations probably facilitated biochemical and hormonal cross-talk between insects and plants, setting the ground for host-plant manipulation by insects.

This work was supported by the ANR grant no. ANR-05-JCJC-0203-01. We also thank two anonymous reviewers for their comments on the manuscript and L. H. for full access to his orchard and helpful discussions.


