

Hemipteran - host plant interactions: focus on some model insect saliva

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Introduction

The interactions between herbivorous insects and the plants they consume have resulted in the evolution of a fascinating and complex web of chemical signals, behavioral responses, and genetic changes.

Insect salivary components play important roles in plant-insect interactions. A variety of enzymes and organic components in saliva of herbivory insects can induce series of biochemical responses in damaged plants, which could be very specific.

It has been demonstrated that the oral secretions of several chewing insects contain elicitors that either stimulate the plant defense, or promote infestation by manipulating plant metabolism/physiology. The effects of salivary compounds injected by piercing-sucking insects into the plant are much less understood.

Objectives

In our lab, several studies have been screening proteins in different aphid saliva (Harmel et al., 2008, for *Myzus persicae* ; Vandermoten et al., 2013, for *M. persicae*, *Megoura viciae* and *Acyrtosiphon pisum*).

The purpose of our current and future works is to widen that field of study to other piercing-sucking pests and host plant models. We focus on pure saliva and salivary glands extract.

Three models will be compared: the pea aphid (feeds on Fabaceae), the invasive Asian brown marmorated stink bug (tree fruit), and the invasive South America green stink bug (Fabaceae, Solanaceae...).

Brown marmorated stink bug (*Halyomorpha halys* Stål)



Broad bean
(*Vicia faba* L.)



Bean
(*Phaseolus vulgaris* L.)

Saliva proteome comparison
(gel-free liquid chromatography)
(Experiment in process)

Southern green stink bug (*Nezara viridula* L.)



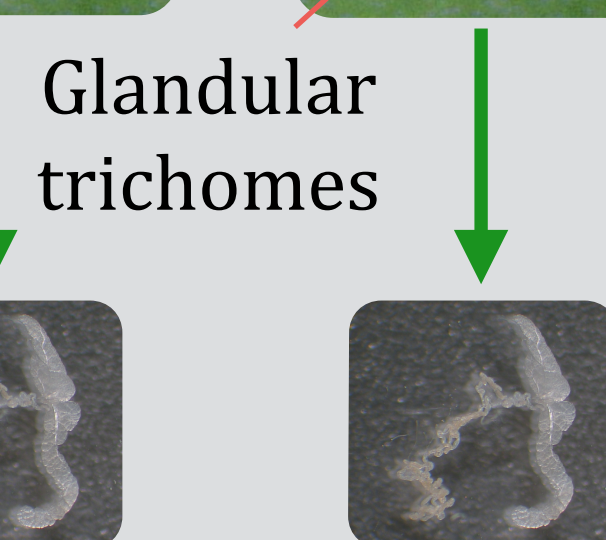
Broad bean



Tomato
(*Solanum lycopersicum* L.)

Salivary glands proteome comparison
(2D-DIGE-MALDI-ToF/MS)

(Analysis in process)



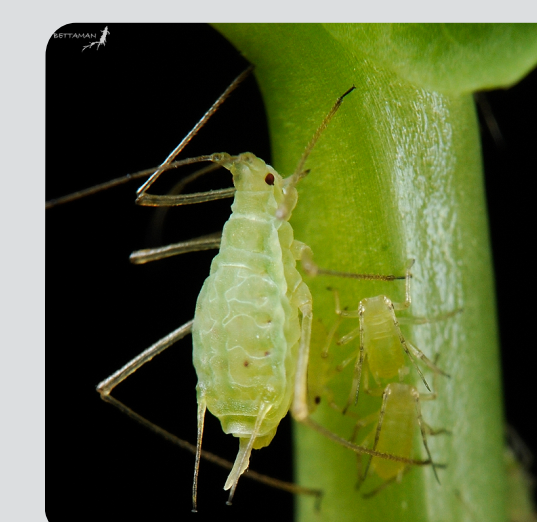
2D-DIGE

N°	sp	Accession	Normalized Wildtype	Normalized Mutant	Fold	Title	Accession	Mass Score	MS Coverage	pI (DB)	pI (Exp)	Query coverage	E value	Source organism	Database
Energy metabolism															
762	1.184	1.086	1.1			ATP synthase subunit alpha	gi55162848	122	24	55116	5.6			Rhodospirillum rubrum	Bacteria
767	1.035	0.829	1.2			ATP synthase subunit alpha	gi55162848	95	22	55116	5.6			Rhodospirillum rubrum	Bacteria
931	0.790	1.112	1.4			Polysphosphate kinase	gi55162848	80	26	37908	5.3			Caulobacter crescentus	Bacteria
DNA replication															
995	0.935	0.882	1.4			Primase	gi55162848	80	23	51133	8.6			Rhodospirillum rubrum	Bacteria
Translation															
1129	0.930	1.497	1.6			Tyrosyl-tRNA synthetase	gi55162848	85	11	43891	5.3			Gemmatimonas	Bacteria
Cell wall															
1812	1.202	1.043	1.2			Pentapeptide repeat-containing protein	gi55162848	71	15	48391	7.7	70%	3E-84	Xanthomonas maltophilia	Bacteria
Protein export															
1860	1.194	1.052	1.1			Cold-shock domain-containing protein 132	gi55162848	62	4	112133	5.6	88%	0	Bacteroides caccae	Cytoplast
Acetification															
1902	0.700	1.444	1.8			Zn-dependent hydrolase including glyoxylase	gi55162848	69	16	24119	5.9	99%	4E-166	Lactobacillus fermentum	Bacteria
Phosphatidyl kinase															
2040	1.223	1.018	1.2			AtvE	gi55162848	74	14	105580	6.8			Pseudomonas fluorescens	Bacteria
Signal transduction															
2102	1.183	0.890	1.3			Putative large ATPase-rich protein	gi55162848	85	19	151434	4.9	90%	0	Streptococcus dysgalactiae	Bacteria
Lipases															
2186	0.884	1.221	1.4			Membrane protein	gi55162848	74	16	22140	5.6			Agrobacterium tumefaciens	Bacteria
1899	1.285	0.761	1.7			Hypothetical protein	gi55162848	76	12	29712	5.9			Acetivibrio butyraceus	Bacteria
1862	0.569	0.964	1.7			Hypothetical protein	gi55162848	67	22	8419	4.1			Bacillus thuringiensis	Bacteria

Comparison between 2 stink bugs with different feeding preferences (herbaceous plant for *N. viridula* and tree fruits for *H. halys*)

Differential expressions of bacterial proteins in salivary glands when *N. viridula* were exposed to tomato with or without glandular trichomes.

Pea aphid (*Acyrtosiphon pisum* Harris)



Different symbiont profiles

Saliva proteome comparison
(Experiment in process)

Impacts of endosymbionts on salivary proteins

Discussion

This study takes place as a continuation of a first description of *H. halys* salivary proteins (Peiffer & Felton, 2014). Salivary glands and gut of stink bugs may be colonized by a bacterial community, which could explain the presence of bacterial proteins in these glands. This is in concordance with other proteomic studies on aphid saliva (Vandermoten et al., 2013 ; Chaudhary et al., 2014). Then it will be relevant to analyze the impact of aphid secondary symbionts on salivary proteins in order to assess their role in aphid performance on its host plant. Some of the identified proteins might indeed play a role in induction or repression of plant defence mechanisms.

The perspective would be applying saliva, salivary gland extracts and/or purified proteins on plant in order to screen its defensive responses by complementary “omic” approaches. Thanks to this comparative study of these three insect models, we would be able to move forward in the understanding of general and specific interactions between plants and Hemipteran pests.

Chaudhary R. et al., 2014. GroEL from the endosymbiont Buchnera aphidicola betrays the aphid by triggering plant defense. *PNAS*, **111**(24): 8919-8924.
Harmel et al., 2008. Identification of aphid salivary proteins: a proteomic investigation of Myzus persicae. *Insect Molecular Biology* **17**(2): 165-174.
Peiffer M. & Felton G.W., 2014. Insights into the saliva of the brown marmorated stink bug Halyomorpha halys (Hemiptera: Pentatomidae). *PLoS One*, **9**.
Vandermoten et al., 2013. Comparative analysis of salivary proteins from three aphid species. *Insect Molecular Biology*, **32**.