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Drosophila Microbiota Modulates Host Metabolic Gene Expression via IMD/NF-κB Signaling



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Abstract

Most metazoans engage in mutualistic interactions with their intestinal microbiota. Despite recent progress the molecular mechanisms through which microbiota exerts its beneficial influences on host physiology are still largely uncharacterized. Here we use axenic *Drosophila melanogaster* adults associated with a standardized microbiota composed of a defined set of commensal bacterial strains to study the impact of microbiota association on its host transcriptome. Our results demonstrate that *Drosophila* microbiota has a marked impact on the midgut transcriptome and promotes the expression of genes involved in host digestive functions and primary metabolism. We identify the IMD/Relish signaling pathway as a central regulator of this microbiota-mediated transcriptional response and we reveal a marked transcriptional trade-off between the midgut response to its beneficial microbiota and to bacterial pathogens. Taken together our results indicate that microbiota association potentiates host nutrition and host metabolic state, two key physiological parameters influencing host fitness. Our work paves the way to subsequent mechanistic studies to reveal how these microbiota-dependent transcriptional signatures translate into host physiological benefits.

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Introduction

Metazoans establish functional interactions with their microbiota, the dynamic microbial communities that colonize their mucosal surfaces. These interactions contribute to many aspects of host physiology, notably metabolism and immunity [1]. Despite recent progress, the molecular mechanisms through which the microbiota exerts its beneficial influences on host physiology are still largely undefined.

Recently, *Drosophila melanogaster* has emerged as a powerful model to study host-microbiota interactions [2,3]. Compared to mammalian species, *Drosophila* carry microbial communities of low complexity, composed of only few dominant bacterial species (mostly of the *Acetobacteraceae* and *Lactobacillaceae* families). The ease to manipulate *Drosophila* commensal bacterial species and to cultivate Germ-Free (GF) animals, coupled to its powerful genetic tools makes *Drosophila* an ideal host model to study molecular mechanisms underlying microbiota-mediated physiological benefits.

Drosophila microbiota affects host biology throughout its life cycle [2–4]. In adults, Drosophila microbiota influences lifespan [5–7], shapes mating preference [8], increases host resistance to several

intestinal pathogens [9], modulates intestinal immune homeostasis [10–12] and promotes intestinal epithelium renewal [13,14]. During the juvenile (larval) phase the microbiota accelerates animal growth and maturation rate [14–16] when the host is under nutritional challenge. These observations point to an important role of *Drosophila* microbiota in shaping the biology of its host. However the molecular dialogue underlying these functional benefits remains elusive.

In this study, we used gnotobiotic *Drosophila* to reveal and study host-microbiota molecular dialogue. To this end, we performed a transcriptome analysis of germ-free and ex-germ-free animals reassociated with a standardized microbiota. Our results demonstrate that microbiota association sustains the expression of genes related to metabolism and digestion in the *Drosophila* midgut, partly via the activity of the IMD/Relish signaling cascade, a pathway previously associated to the regulation of processes related to immune responses. In addition, we further demonstrate that upon bacterial infection in the midgut, the expression of metabolic gene promoted by microbiota association is down-regulated, indicating the existence of host transcriptional trade-off between infection and normal physiology.

Results and Discussion

Drosophila microbiota impacts midgut genes expression

To gain insight into the molecular cross-talk between *Drosophila* microbiota and its host, we compared the transcriptomic changes between microbiota-associated adult flies and their Germ-Free (GF) siblings. Since the microbiota load and composition encountered in conventionally laboratory-reared flies (CONV) fluctuate highly [9,17] (and our unpublished observation), we chose to associate newly emerged GF adults with a standardized microbiota composed of four previously characterized *Drosophila* commensal bacterial strains (*Acetobacter pomorum*, *Commensalibacter intestini*, *Lactobacillus brevis* and *Lactobacillus plantarum*) [18] (Fig.1A). After data normalization and statistical analysis using Significance Analysis of Microarrays (SAM), using a false detection rate (FDR) of 0,2, we identified 105 transcripts whose expression level were significantly increased from 1.2 to 6 folds in the polyassociated flies compared to their GF siblings (Fig.1B and Fig.2).

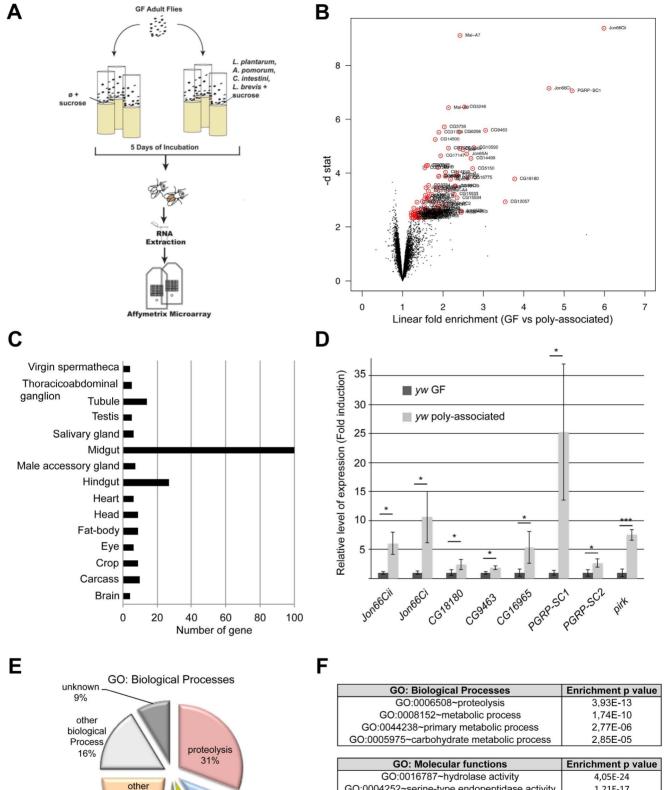
We classified the 105 selected genes according to their annotated tissue expression profile using the FlyMine tool [19] based on the FlyAtlas dataset [20]. Interestingly, despite the fact that the transcriptomic analysis was conducted on whole adult animals, we found that most (100/105) of the selected transcripts are expressed in the midgut of conventionally (CONV) raised adults (i.e with a microbiota) (Fig.1C). Recently, Buchon et al. and Marianes et al. reported that 60-65% of the Drosophila genes are detected in the midgut of CONV adults with specific patterns of expression in this tissue [21,22]. We therefore examined whether our list contains enrichment for genes sharing a given spatial pattern of expression in the midgut but we failed to detect any (data not shown). Nonetheless, the marked over-representation of midgut genes in our dataset indicates that the host transcriptional response to the microbiota is localized and occurs mainly in the Drosophila midgut. We then verified that microbiota-regulated transcripts were indeed enriched in the midgut upon microbiotaassociation. To do so, using RT-qPCR, we compared the expression levels of a set of the most strongly microbiota-regulated genes in dissected midguts from GF and microbiota-associated animals and found that the tested genes were all up-regulated in adult midguts associated with four commensal bacterial strains compared to GF controls (Fig.1D). These results therefore demonstrate that microbiota association impacts on genes expression in the midgut.

Drosophila microbiota sustains metabolic genes expression

We next used Gene Ontology (GO) clustering and enrichment analysis tools (Database for Annotation, Visualization, and Integrated Discovery or DAVID; [23]) to identify the functional categories within our gene list. We found that 74 out of 105 genes on the list are associated with metabolic processes (Fig.1E, Fig.2). Among the 74 genes, 33 are involved in proteolysis, 11 in carbohydrate metabolic process, six in lipid metabolic process and four in oxydo-reduction. From the same analysis, we found that the terms "metabolic process", "proteolysis", "primary metabolism" and "carbohydrate metabolism" are the most significantly enriched terms (Fig.1E-F, Fig.2). Among the remaining 31 genes, several are involved in processes that could either be connected to metabolic activities such as transport across membranes (4/31), chitin metabolism (4/31) and cellular ion homeostasis (2/31) or linked to host response to bacterial colonization (i.e, "response to biotic stimulus" (5/31) and "wound healing and related process" (2/31)) (Fig.1E and Fig.2). These results not only indicate that Drosophila presents an expected transcriptional response to bacterial colonization upon microbiota association, but also reveal a clear impact of microbiota association on host metabolic processes at least at the transcriptomic level.

Using the same GO terms tool, we further analyzed the molecular functions associated with the 105 genes (Fig.2). We identified a clear enrichment of genes encoding hydrolases and alkaline phosphatases (Fig.1F), which are two major classes of enzymes involved in digestive functions, suggesting that Drosophila microbiota enhances the host's enzymatic activities for macronutrients breakdown. Strikingly, among the 74 "metabolic genes", 60 genes encode hydrolases, including 33 peptidases (28 serineproteases including several members of the trypsin and Jonah proteases families, four metallo-proteases and the aspartic-protease Bace), 10 glycosyl-hydrolases (five alpha-glucosidases of the maltase family, two alpha-mannosidases, the alpha-amylase Amy-d, the beta-galactosidase Gal and one alpha, alpha-trehalase), six phosphatases (five alkaline and the acid-phosphatase Mipp1), five lipases (including magro, the Drosophila LipA homologue controlling cholesterol homeostasis, [24]), three non-lipase esterases (including EstQ), two nucleases and the biotinidase vanin-like (Fig.2). The other metabolic genes include five transferases (three acyl-transferases, one phospho-transferase and the beta-galactosyl-transferase Ugt86Dc), three Cytochrome P450 (Cyp6a17, Cyp6a21 and Cyp6d5), two Carbohydrate binding proteins, two genes related to Acyl-CoA metabolism, one Cytochrome-c oxidase and one 4-Coumarate-CoA ligase. These molecules are mostly involved in primary metabolic pathways, suggesting that microbiota association sustains these essential host metabolic activities. This finding is consistent with the observation that microbiota influences host energy homeostasis and carbohydrate allocation patterns in adults [15]. Among the 31 "non metabolic genes" regulated by microbiota association, we identified 11 genes that are still related to metabolic activities, which may be influenced by microbiota association. Five out of the 11 genes are involved in transport of micronutrients or xenobiotics including the vitamin-B12 transporter Amnionless, the sterol transporter Npc2d and the sodium transporter *Nha1*; four genes are associated to chitin metabolism, including the mucin Muc68E and the chitin binding molecule obst-A and two genes encode proteins involved in metal homeostasis (MtnB/C). In addition to these metabolic signatures, we identified seven genes that are clearly associated to host tissue response to bacterial challenges. Specifically, three of the seven genes are associated to "wound healing", such as one fibrinogen; five are innate immune genes whose expression is known to be controlled by the IMD signaling pathway such as the PGRP-LC/LE inhibitor pirk and the peptidoglycan amidases PGRP-LB, -SC1 and -SC2 which are all involved in dampening the IMD signaling strength to promote immune tolerance to indigenous microbiota [10-12]. This observation corroborates previous reports demonstrating that the gut microbiota modulates intestinal immune homeostasis and promotes intestinal epithelium renewal [4]. Finally, we identified the Zinc-finger transcription factor GATAe, which is required for the terminal differentiation of the Drosophila endoderm and maturation of the adult midgut [21,25,26]. Interestingly, among the 105 genes uncovered by our transcriptomic analysis, we could identify 31 genes which expression is altered upon GATAe genetic manipulation [21,27] (Fig.2). This observation reinforces the notion that microbiota may promote the maturation and the digestive functionalities of the midgut partly via GATAe-dependent regulation of digestive enzymes expression.

Taken together, our results clearly indicate that microbiota association influences the expression of host midgut genes encoding key actors involved in digestive functions, primary



GO:0016787~hydrolase activity 4,05E-24 GO:0004252~serine-type endopeptidase activity 1,21E-17	lue
GO:0004252~serine-type endopeptidase activity 1,21E-17	
GO:0004035~alkaline phosphatase activity 9,97E-05	

response to

biotic

stimulus

5%

metabolic processes 23%

carbohydrate metabolic

10%

lipid metabolic process

process 6%

Figure 1. *Drosophila* **microbiota impacts midgut genes expression.** (A) A schematic representation of the experimental set-up used for transcriptome analysis; GF: Germ-Free. (B) A volcano plot showing the absolute value of the computed statistics from the SAM analysis (d-stat) based on to the fold change in gene expression between the GF and poly-associated groups. The genes selected as differentially expressed by the SAM analysis are highlighted with a red circle around them. The selection was made using a value of Delta that corresponds to a FDR of 0.2. (C) Adult tissue enrichment classification of the 105 microbiota-regulated genes. (D) A bar graph representing the fold-enrichment of different microbiota-regulated transcripts in dissected midguts from GF vs poly-associated wild-type animals (the value of the relative $\Delta Ct^{gene}/\Delta Ct^{rp49}$ ratio was calculated for every sample and relativized to the ratio in the GF condition which was anchored to 1 to indicate fold induction). Statisticance of the result is represented (Student's t-test: ns \geq 0.05>* \geq 0.01>** \geq 0.001>***). Of note, *PGRP-SC1* mRNAs are detected to a 16 folds higher level than *PGRP-SC2* mRNA in young poly-associated animals ($\Delta Ct^{PGRP-SC1}/\Delta Ct^{Pp49} = 63$ and $\Delta Ct^{PGRP-SC2}/\Delta Ct^{P49} = 4$). (E) Relative percentage of Gene Ontology Terms: Biological processes in the list of microbiota-regulated gene (pie chart) and (F) Enrichment p-values (Benjamini-Hochberg corrected) for GO:Biological processes and GO:Molecular Functions in the list of microbiota-regulated genes.

metabolism and host tolerance to bacteria colonization and that *Drosophila* microbiota sustains these activities.

Correlation between microbiota and nutrients-mediated transcriptional signatures

Metabolic adaptation through metabolic gene regulation is essential for the host to respond to nutritional challenges. Now, having observed that microbiota association promotes the transcription of metabolic genes, we further compared our results with previous analysis on *Drosophila* transcriptome upon nutritional challenges [28,29]. Among the 105 microbiota-regulated genes, the expression of 30 genes was reported to fluctuate in response to sugar only diet [29] (Fig.2). Specifically, Zinke *et al.* reported that *sugarbabe* (*sug.*), a zinc-finger transcription factor that is strongly

activated upon sugar ingestion, represses the expression of several genes involved in dietary sugar and fat breakdown. We found in our list 16 "sug-regulated" genes among which four are Glycosylhydrolases (Amy-d, CG9463, Mal-A3 and Mal-A8) and four are lipases (CG15534, CG8093, mag and CG15533). In our experimental conditions, flies were reared on a sucrose-only diet prior and during the association. Therefore, the upregulation of sug-related genes upon microbiota association suggests that the repressive activity of Sug during sugar feeding is inhibited during host response to microbiota. Similarly, Li et al. identified the transcription factor Myc as one of the main regulators of metabolic genes expression in response to nutritional challenges. In this study we found 30 Myc-regulated genes in our list [28] (Fig.2). This correlation suggests that Myc is also a prime

Gene Name	Fold	GO:0006508~proteolysis	GATAe	Sugar only	Sug	Myc	Ecc15	Rel									
Jon66Cii	6.0	MF: serine-type endopeptidase activity	OMINO	Ougur Only	Oug	yes	up	inf	Gene Name	Fold	Other metabolic processes	GATAe	Sugar only	Sug	Myc	Ecc15	Rel
Jon66Ci	4.6	MF: serine-type endopeptidase activity		ves		ves	up	inf/basal	CG10592	2.8	MF: alkaline phosphatase activity		- again anny		yes		
CG18180	3.8	MF: serine-type endopeptidase activity	yes	,,,,		,,,,	op		CG5150	2,7	MF: alkaline phosphatase activity		yes	yes	,00	down	inf/basal
Jon65Ai	2.6	MF: serine-type endopeptidase activity	, , , ,	yes		yes			CG14499	2,7	MF: carbohydrate binding		yes	,			min basar
CG5246	2.5	MF: serine-type endopeptidase activity	yes	,,,,		yes			CG14219	2.1	MF: transferase activity, transferring acyl groups		,	ves		up	inf/basal
Jon25Bii	2.4	MF: serine-type endopeptidase activity	yes			ves	down	inf	CG14500	1.8	MF: carbohydrate binding			,00	ves	up	minousus
Jon99Ci	2.3	MF: serine-type endopeptidase activity	,,,,	ves		,,,,	down		Cyp6a21	1,8	GO:0055114~oxidation reduction				ves		
CG18179	2.3	MF: serine-type endopeptidase activity		yes	yes	yes	up	inf/basal	CG13813	1,7	MF: transferase activity, transferring phosphorus-containing groups				,00	down	inf/basal
CG7025	2,1	MF: metallocarboxypeptidase activity		yes	,,03	yes	down	inf/basal	Cyp6a17	1,7	GO:0055114~oxidation reduction		yes			down	IIII/Daaaii
Jon74E	2.1	MF: serine-type endopeptidase activity	yes				down	IIIIDUGGUI	CG4563	1.7	MF: 4-coumarate-CoA ligase activity		yes		yes		
CG3739	2	MF: serine-type endopeptidase activity	l yes				down	inf/basal	Uat86Dc	1.7	MF: 2-hydroxyacylsphingosine 1-beta-galactosyltransferase activity	,	, yes		yes	down	
CG8773	1,9	MF: aminopeptidase/metallopeptidase activity	ves	yes		yes	down	IIIIDaaaii	CG31104	1,6	MF: transferase activity, transferring phosphorus-containing groups					down	inf
Jon65Aii	1,9	MF: serine-type endopeptidase activity	yes	yes		yes	down	inf/basal	CG3841	1,6	MF: carboxylesterase activity				yes	down	inf/basal
CG31343	1.9	MF: aminopeptidase/metallopeptidase activity	,,,,	703			down	IIIIDaaaii	CG10827	1.6	MF: alkaline phosphatase activity				yes	uowii	IIII/Dadaii
CG7542	1,9	MF: serine-type endopeptidase activity				yes			CG8628	1,6	GO:0042049~cellular acvi-CoA homeostasis						
Bace	1.8	MF: aspartic-type endopeptidase activity		yes	yes	yes			CG14120	1,6	MF: metal ion binding/hydrolase activity/nucleic acid binding	yes	yes	yes		down	inf/basal
Jon44E	1.8	MF: serine-type endopeptidase activity		yes	yes	yes			CG3106	1.5	MF: transferase activity, transferring acyl groups	yes	yes	yes		down	IIII/Dasai
CG11911	1.8	MF: serine-type endopeptidase activity	yes	yes		yos			Cyp6d5	1,5	GO:0055114~oxidation reduction				yes	down	inf
CG10477	1.8	MF: serine-type endopeptidase activity	yes	yes		ves			CG42249	1.5	GO:0006139~nucleic acid metabolic process				yes	down	
CG3734	1,6	MF: serine-type endopeptidase activity MF: serine-type peptidase activity	ves			yes	down		CG1809	1,4	MF: alkaline phosphatase activity	ves				down	
CG31265	1,6	MF: serine-type endopeptidase activity	yes				down		vanin-like	1,4	GO:0006807~nitrogen compound metabolic process	yes					basal
CG4653	1,6	MF: serine-type endopeptidase activity				yes ves	up	inf/basal	CG3290	1,3	MF: alkaline phosphatase activity					up up	inf/basal
CG16749	1,6	MF: serine-type endopeptidase activity	yes			yes	down	inf	Est-Q	1,3	MF: GO:0004091~carboxylesterase activity		yes	yes		up	IIII/Dasai
Jon99Fi	1.6	MF: serine-type endopeptidase activity	yes			yes	down	1111	CG31644	1,3	GO:0055114~oxidation reduction	yes	yes	yes			
	1,6									1,3	MF: GO:0003993~acid phosphatase activity						inf/basal
zetaTry CG31269	1,6	MF: serine-type endopeptidase activity				yes			Mipp1	1,2	Mr. GO.0003995~acid priospriatase activity					down	IIII/Dasai
alphaTrv	1,5	MF: serine-type endopeptidase activity	yes														
CG15254	1,5	MF: serine-type endopeptidase activity MF: metalloendopeptidase activity	yes	yes													
Phae2	1,4				yes				Gene Name	Fold	Other biological process	GATAe	[C	C	Myc	Ecc15	Rel
Phae1	1,4	MF: serine-type endopeptidase activity MF: serine-type endopeptidase activity							Npc2d	2.2	GO:0015918~sterol transport	GATAE	Sugar only	Sug	wyc	na	na
Jon25Biii	1,4								CG17147	1,9	GO:0015918~sterol transport GO:0006030~chitin metabolic process						
		MF: serine-type endopeptidase activity		yes			down	inf/basal	CG30160							na na	na
CG17477	1,3	MF: serine-type endopeptidase activity	yes				aown	int/basai		1,9	InterPro: Tetraspanin						na
											ME, and a time and an establish in this is a						
thetaTry	1,3	MF: serine-type endopeptidase activity	yes				down		CG31704	1,9	MF: serine-type endopeptidase inhibitor activity					up	inf/basal
				I C	Cur	14		Dal	MtnB	1,9	GO:0006873~cellular ion homeostasis		yes			na	na
Gene Name	Fold	GO:0005975~carbohydrate metabolic process		Sugar only		Myc	Ecc15	Rel	MtnB MtnC	1,9 1,8	GO:0006873~cellular ion homeostasis GO:0006873~cellular ion homeostasis	yes	yes yes			na down	na na
Gene Name CG9463	Fold 3,1	GO:0005975~carbohydrate metabolic process GO:0006013~mannose metabolic process	GATAe	Sugar only yes	Sug	Мус		Rel inf/basal	MtnB MtnC Muc68E	1,9 1,8 1,6	GO:0006873∼cellular ion homeostasis GO:0006873∼cellular ion homeostasis GO:0006030∼chitin metabolic process	yes	yes			na down na	na na na
Gene Name CG9463 Mal-A7	Fold 3,1 2,4	GO:0005975~carbohydrate metabolic process GO:0006013~mannose metabolic process MF: alpha-glucosidase activity				Мус	Ecc15 down	inf/basal	MtnB MtnC Muc68E CG5550	1,9 1,8 1,6 1,6	GO:0006873~cellular ion homeostasis GO:0006873~cellular ion homeostasis GO:0006030~chitin metabolic process MF: fibrinogen / associated to wound healing					na down na up	na na na inf/basal
Gene Name CG9463 Mal-A7 CG9466	Fold 3,1 2,4 2,4	GO:0005975~carbohydrate metabolic process GO:0006013~mannose metabolic process MF: alpha-glucosidase activity GO:0006013-mannose metabolic process	GATAe yes	yes	yes		Ecc15		MtnB MtnC Muc68E CG5550 CG15170	1,9 1,8 1,6 1,6 1,6	GO:0006873~cellular ion homeostasis GO:0006873~cellular ion homeostasis GO:0006030~chitin metabolic process MF: fibrinogen / associated to wound healing GO:0042000~wound healing	yes	yes			na down na up na	na na na inf/basal na
Gene Name CG9463 Mal-A7 CG9466 Amy-d	Fold 3,1 2,4 2,4 2,3	GO:0005975~carbohydrate metabolic process GO:0006013~mannose metabolic process MF: alpha-glucosidase activity GO:0006013~mannose metabolic process MF: alpha-amylase activity	GATAe	yes	yes	yes	Ecc15 down down	inf/basal inf	MtnB MtnC Muc68E CG5550 CG15170 CG30272	1,9 1,8 1,6 1,6 1,6 1,6	GO:0006873-cellular ion homeostasis GO:0006873-cellular ion homeostasis GO:0006030-chilm metabolic process MF: fibrinogen / associated to wound healing GO:0042060-wound healing GO:0055085-transmembrane transport	yes yes	yes yes			na down na up na down	na na na inf/basal na inf/basal
Gene Name CG9463 Mal-A7 CG9466 Amy-d Mal-A3	Fold 3,1 2,4 2,4 2,3 2,3	GO 0005975-carbohydrate metabolic process GO:0006013-mannose metabolic process MF: alpha-glucosidase activity GO:0006013-mannose metabolic process MF: alpha-amylase activity MF: alpha-glucosidase activity	GATAe yes	yes	yes yes yes	yes yes	Ecc15 down down	inf/basal inf inf	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50	1,9 1,8 1,6 1,6 1,6 1,6 1,5	GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.008030-chitim metabolic process MF: fibrinogen i associated to wound healing GO.0042080-wound healing GO.0055085-transmembrane transport GO.0015898-drug transport	yes	yes	yes		na down na up na down down	na na na inf/basal na inf/basal inf
Gene Name CG9463 Mal-A7 CG9466 Amy-d Mal-A3 Mal-A8	Fold 3,1 2,4 2,4 2,3 2,3 2,1	GO:0005975-carbohydrate metabolic process GO:0006013-mannose metabolic process MF: alpha-glucosidase activity GO:0006013-mannose metabolic process MF: alpha-amylase activity MF: alpha-amylase activity MF: alpha-glucosidase activity	GATAe yes yes	yes yes yes	yes	yes yes yes	Ecc15 down down	inf/basal inf	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A	1,9 1,8 1,6 1,6 1,6 1,6 1,5 1,5	GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.006809-chitin metabolic process MF. fbrinogen / associated to wound healing GO.0042606-wound healing GO.0055065-transmetrane transport GO.006030-chitin metabolic process GO.006030-chitin metabolic process	yes yes	yes yes	yes		na down na up na down down na	na na na inf/basal na inf/basal inf
Gene Name CG9463 Mal-A7 CG9466 Amy-d Mal-A3 Mal-A8 Mal-A4	Fold 3,1 2,4 2,4 2,3 2,3 2,1 2,1	GC 0005975-carbohydrate metabolic process GC 0005013-mannose metabolic process MF: alpha-glucosidase activity GC 0006013-mannose metabolic process MF: alpha-amylase activity MF: alpha-glucosidase activity MF: alpha-glucosidase activity MF: alpha-glucosidase activity	GATAe yes	yes	yes yes yes	yes yes	Ecc15 down down down down	inf/basal inf inf	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A CG33523	1,9 1,8 1,6 1,6 1,6 1,6 1,5 1,5	GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.008030-chitin metabolic process MF: fibrinogen / associated to woud healing GO.004200-wound healing GO.005005-transmembrane transport GO.005005-transmembrane transport GO.0060030-chitin metabolic process Interpro: vesicule-associated membrane protein	yes yes	yes yes	yes		na down na up na down down na na	na na na inf/basal na inf/basal inf na
Gene Name CG9463 Mal-A7 CG9466 Amy-d Mal-A3 Mal-A8 Mal-A4 Mal-A6	Fold 3,1 2,4 2,4 2,3 2,3 2,1 2,1 1,9	GO 0005975-carbohydrate metabolic process GO 0006013-mannose metabolic process MF: ajbha-glucosidase activity GO 0006013-mannose metabolic process MF: ajbha-glucosidase activity	GATAe yes yes	yes yes yes	yes yes yes	yes yes yes	Ecc15 down down down down	inf/basal inf inf inf/basal	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A CG33523 GATAe	1,9 1,8 1,6 1,6 1,6 1,6 1,5 1,5 1,4	GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.006830-chitim metabolic process MF. fbrinogen / associated to wound healing GO.0042660-wound healing GO.0055068-fransmethrane transport GO.006030-chitim metabolic process Interpro: vesicule-associated membrane protein GO.007492-endodorem development	yes yes	yes yes	yes		na down na up na down down na na	na na na inf/basal na inf/basal inf na na
Gene Name CG9463 Mal-A7 CG9466 Amy-d Mal-A3 Mal-A8 Mal-A4 Mal-A6 CG16965	Fold 3,1 2,4 2,4 2,3 2,3 2,1 2,1 1,9 1,8	GO 0005975-carbohydrate metabolic process GO 0006913-mannose metabolic process MF. siphra-glucosidase activity GO 000013-mannose metabolic process MF. siphra-emylosse activity MF. siphra-emylosse activity MF. siphra-glucosidase activity	GATAe yes yes yes	yes yes yes	yes yes yes	yes yes yes	down down down down down up	inf/basal inf inf inf/basal inf/basal	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A CG33523 GATAe CG42397	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4	GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006803-chitim metabolic process MF: fibrinogen / associated to wound healing GO.004200-wound healing GO.005008-transmembrane transport GO.005008-transmembrane transport GO.005008-dept. metabolic process Interpro: vescule-associated membrane protein GO.0007492-endoderm development GO.0006030-chitim metabolic process	yes yes	yes yes	yes		na down na up na down down na na na	na na na inf/basal na inf/basal inf na na
Gene Name CG9463 Mal-A7 CG9466 Amy-d Mal-A3 Mal-A8 Mal-A4 Mal-A6 CG16965 Gal	Fold 3,1 2,4 2,4 2,3 2,3 2,1 2,1 1,9 1,8 1,4	GO 0005975-carbohydrate metabolic process GO 0006013-mannose metabolic process MF: ajbha-glucosidase activity GO 0006013-mannose metabolic process MF: ajbha-glucosidase activity MF: abba-glucosidase activity MF: abba-glucosidase activity MF: beta-glastosidase activity	GATAe yes yes	yes yes yes	yes yes yes	yes yes yes	down down down down down up down	inf/basal inf inf inf/basal inf/basal inf	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A CG33523 GATAe CG42397 Nha1	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,4	GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.006830-chitim metabolic process MF. fbrinogen / associated to wound healing GO.004260-wound healing GO.0055085-transmetrane transport GO.0065085-transmetrane transport GO.006030-chitim metabolic process Interpro: vesicule-associated membrane protein GO.007492-endodorem development GO.000630-chitim metabolic process GO.006830-chitim metabolic process GO.006830-chitim transport	yes yes	yes yes	yes		na down na up na down down na na na na	na na na inf/basal na inf/basal inf na na na
Gene Name CG9463 Mal-A7 CG9466 Amy-d Mal-A3 Mal-A8 Mal-A4 Mal-A6 CG16965	Fold 3,1 2,4 2,4 2,3 2,3 2,1 2,1 1,9 1,8	GO 0005975-carbohydrate metabolic process GO 0006913-mannose metabolic process MF. siphra-glucosidase activity GO 000013-mannose metabolic process MF. siphra-emylosse activity MF. siphra-emylosse activity MF. siphra-glucosidase activity	GATAe yes yes yes	yes yes yes	yes yes yes	yes yes yes	down down down down down up	inf/basal inf inf inf/basal inf/basal	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A CG33523 GATAe CG42397 Nha1 Amnionless	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,4	GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.000630-chitin metabolic process MF. fibringen/ associated to wound healing GO.000260-wound healing GO.005608-transmethrane transport GO.0005808-transmethrane transport GO.0005808-transmethrane transport GO.0005808-drain metabolic process GO.0007402-endoderm dewelopment GO.000760809-chitin metabolic process GO.00060809-chitin metabolic process GO.00060809-chitin metabolic process GO.0006081-ion transport	yes yes	yes yes	yes		na down na up na down down na na na na	na na na inf/basal inf/basal inf na inf na na na na na
Gene Name CG9483 Mal-A7 CG9486 Amy-d Mal-A3 Mal-A8 Mal-A4 Mal-A6 CG16965 Gal CG9394	Fold 3,1 2,4 2,4 2,3 2,3 2,1 1,9 1,8 1,4	GO 0005975-carbohydrate metabolic process GO 0006013-mannose metabolic process MF: ajbha-glucosidase activity GO 0006013-mannose metabolic process MF: ajbha-glucosidase activity MF: abba-glucosidase activity MF: abba-glucosidase activity GO 0006071-glycerol metabolic process	yes yes yes	yes yes yes yes	yes yes yes yes	yes yes yes yes	down down down down down up down down	inf/basal inf inf inf/basal inf/basal inf	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A CG33523 GATAe CG42397 Nha1	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,4	GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.006830-chitim metabolic process MF. fbrinogen / associated to wound healing GO.004260-wound healing GO.0055085-transmetrane transport GO.0065085-transmetrane transport GO.006030-chitim metabolic process Interpro: vesicule-associated membrane protein GO.007492-endodorem development GO.000630-chitim metabolic process GO.006830-chitim metabolic process GO.006830-chitim transport	yes yes	yes yes	yes		na down na up na down down na na na na	na na na inf/basal na inf/basal inf na na na
Gene Name CG9463 Mal-A7 CG9466 Amy-d Mal-A3 Mal-A4 Mal-A4 Mal-A6 CG16965 Gal CG9394 Gene Name	Fold 3,1 2,4 2,4 2,3 2,3 2,1 1,9 1,8 1,4 1,4	GO 0005975-carbohydrate metabolic process GO:0006013-mannose metabolic process MF: ajhra-glucosidase activity GO:0006013-mannose metabolic process MF: ajhra-glucosidase activity GO:0006071-glycerol metabolic process GO:0006023-glyd metabolic process	yes yes yes	yes yes yes yes	yes yes yes yes	yes yes yes	down down down down down up down	inf/basal inf inf inf/basal inf/basal inf inf	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A CG33523 GATAe CG42397 Nha1 Amnionless	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,4	GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.000630-chitin metabolic process MF. fibringen/ associated to wound healing GO.000260-wound healing GO.005608-transmethrane transport GO.0005808-transmethrane transport GO.0005808-transmethrane transport GO.0005808-drain metabolic process GO.0007402-endoderm dewelopment GO.000760809-chitin metabolic process GO.00060809-chitin metabolic process GO.00060809-chitin metabolic process GO.0006081-ion transport	yes yes	yes yes	yes		na down na up na down down na na na na	na na na inf/basal inf/basal inf na inf na na na na na
Gene Name' CG9463 Mal-A7 CG9466 Amy-d Mal-A8 Mal-A8 Mal-A4 CG16965 Gal CG9394 Gene Name' CG6296	Fold 3,1 2,4 2,4 2,3 2,3 2,1 2,1 1,9 1,8 1,4 1,4	GO 0005975-carbohydrate metabolic process GO 0006013-mannose metabolic process MF: ajbha-glucosidase activity GO 0006013-mannose metabolic process MF: ajbha-glucosidase activity MF: abpha-glucosidase activity MF: abpha-glucosidase activity GO 0006071-glycerol metabolic process GO 0006071-glycerol metabolic process MF: lipase activity	yes yes yes	yes yes yes yes	yes yes yes yes	yes yes yes yes	Ecc15 down down down down down up down down down	inf/basal inf inf inf/basal inf/basal inf inf	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A CG33523 GATAe CG42397 Nha1 Amnionless	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,4	GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.000630-chitin metabolic process MF. fibringen/ associated to wound healing GO.000260-wound healing GO.005608-transmethrane transport GO.0005808-transmethrane transport GO.0005808-transmethrane transport GO.0005808-drain metabolic process GO.0007402-endoderm dewelopment GO.000760809-chitin metabolic process GO.00060809-chitin metabolic process GO.00060809-chitin metabolic process GO.0006081-ion transport	yes yes	yes yes	yes		na down na up na down down na na na na	na na na inf/basal inf/basal inf na inf na na na na na
Gene Name CG9463 Mal-A7 CG9466 Amy-d Mal-A3 Mal-A4 Mal-A4 Mal-A6 CG16965 Gal CG9394 Gene Name CG6296 CG15534	Fold 3,1 2,4 2,4 2,3 2,3 2,1 2,1 1,9 1,8 1,4 1,4 1,4	GO 0005975-carbohydrate metabolic process GO:0006013-mannose metabolic process MF: ajhra-glucosidase activity GO:0006013-mannose metabolic process MF: ajhra-glucosidase activity MF: pathra-glucosidase activity GO:0006071-glycerol metabolic process MF: passe activity GO:0006063-philip metabolic process	yes yes yes GATAe	yes yes yes yes yes Sugar only	yes yes yes yes yes	yes yes yes yes	down down down down down up down down	inf/basal inf inf inf/basal inf/basal inf inf	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A CG33523 GATAe CG42397 Nha1 Amnionless Pask	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,4 1,3 1,3	GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006803-chitim metabolic process MF. fibringen/ associated to wound healing GO.0042060-wound healing GO.0055086-transmethrane transport GO.0016893-drug transport GO.000508-chitim metabolic process Interpro vesicule-associated membrane protein GO.007482-endoderm development GO.007482-endoderm development GO.0006811-ion transport Interpro Ammionless, Vil 812 transporter GO.0007165-signal transduction	yes yes yes	yes yes yes	ŕ		na down na up na down na na na na na na	na na na inf/basal na inf/basal inf na na na na na na na na
Gene Name CG9463 Mal-A7 CG9466 Amy-d Mal-A3 Mal-A6 GG15965 Gal CG9394 Gene Name CG6296 CG15534 CG15533	Fold 3,1 2,4 2,4 2,3 2,3 2,1 2,1 1,9 1,8 1,4 1,4 1,4 1,4	GO 0005975-carbohydrate metabolic process GO 0005013-mannose metabolic process MF: ajbha-glucosidase activity GO 0005013-mannose metabolic process MF: ajbha-glucosidase activity GO 0006071-glycerol metabolic process MF: lipase activity GO 0006655-sphingolipid metabolic process	yes yes yes GATAe yes yes	yes yes yes yes yes Sugar only	yes yes yes yes yes	yes yes yes yes	down down down up down down down	inf/basal inf inf inf/basal inf/basal inf inf	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A CG33523 GATAe CG42397 Nha1 Aminonless Pask	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,3 1,3 1,3	GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.0006873-cellular ion homeostasis GO.000630-chitin metabolic process MF. fibringen/ associated to wound healing GO.000260-wound healing GO.005608-transmethrane transport GO.0005808-transmethrane transport GO.0005808-transmethrane transport GO.0005808-drain metabolic process GO.0007402-endoderm dewelopment GO.000760809-chitin metabolic process GO.00060809-chitin metabolic process GO.00060809-chitin metabolic process GO.0006081-ion transport	yes yes	yes yes	yes	Мус	na down na up na down down na na na na na na	na na na inf/basal na inf/basal inf/basal inf na na na na na na na
Gene Name: CG9463 Mai-A7 CG9466 Amy-d Mai-A8 Mai-A8 Mai-A6 CG16965 Gal CG9394 Gene Name: CG6296 CG15534 CG15533 mag	Fold 3,1 2,4 2,3 2,3 2,1 1,9 1,8 1,4 1,4 Fold 2,4 2,3 2,1 2,1 2,1 2,1 2,1 2,1 2,1 2,1 2,1 2,1	GO 0005975-carbohydrate metabolic process GO:0006013-mannose metabolic process MF: ajhra-glucosidase activity GO:0006013-mannose metabolic process MF: ajhra-glucosidase activity GO:00060671-glycerol metabolic process MF: lipase activity GO:0006665-aphingolipid metabolic process GO:0006665-aphingolipid metabolic process GO:0006665-aphingolipid metabolic process MF: sterio eleterase activity	GATAe yes yes yes GATAe yes yes	yes yes yes yes Sugar only	yes yes yes yes yes	yes yes yes yes	Ecc15 down down down down down up down down down	inf/basal inf inf inf/basal inf/basal inf inf	MfnB MfnC Muc68E CG5550 CG15170 CG30272 Mfn50 obst-A G33523 GATAe CG42397 Nha1 Amnionless Pask	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,4 1,3 1,3 1,2	GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006803-chitim metabolic process MF. fibringen/ associated to wound healing GO.0042060-wound healing GO.0055086-transmethrane transport GO.0016893-drug transport GO.000508-chitim metabolic process Interpro vesicule-associated membrane protein GO.007482-endoderm development GO.007482-endoderm development GO.0006811-ion transport Interpro Ammionless, Vil 812 transporter GO.0007165-signal transduction	yes yes yes	yes yes yes Sugar only	ŕ	Myc yes	na down na up na down down na	na na na inf/basal na inf/basal inf na
Gene Name: CG9463 Mal-A7 CG9466 Amy-d Mal-A3 Mal-A8 Mal-A6 CG16965 Gal CG9394 Gene Name CG6296 CG15533 mag CG8093	Fold 3,1 2,4 2,3 2,3 2,1 1,9 1,8 1,4 1,4 1,4 Fold 2,4 2,3 2,1 1,9 1,9 1,9 1,9 1,9 1,9 1,9 1,9 1,9 1	GO 0005975-carbohydrate metabolic process GO 0006013-mannose metabolic process MF: ajbha-glucosidase activity GO 0006013-mannose metabolic process MF: ajbha-glucosidase activity MF: plaga-glactosidase activity GO 0006607-ajbqcm metabolic process MF: steroi estrase activity MF: lipase activity MF: lipase activity	yes yes yes GATAe yes yes	yes yes yes yes yes Sugar only	yes yes yes yes yes	yes yes yes yes	Ecc15 down down down down up down down down down down down down	inf/basal inf inf inf/basal inf/basal inf inf	MfnB MtnC Muc68E CG5550 CG15170 CG30272 Mdr50 obst-A CG33523 GA774e CG42397 Nmionless Pask	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,3 1,3 1,3 2,6	GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006803-chitim metabolic process MF. fibringen/ associated to wound healing GO.0042060-wound healing GO.0055086-transmethrane transport GO.0016893-drug transport GO.000508-chitim metabolic process Interpro vesicule-associated membrane protein GO.007482-endoderm development GO.007482-endoderm development GO.0006811-ion transport Interpro Ammionless, Vil 812 transporter GO.0007165-signal transduction	yes yes yes	yes yes yes Sugar only yes	ŕ	yes	na down na up na down down na	na na na inf/basal na inf/basal inf na
Gene Name: CG9463 Mai-A7 CG9466 Amy-d Mai-A8 Mai-A8 Mai-A6 CG16965 Gal CG9394 Gene Name: CG6296 CG15534 CG15533 mag	Fold 3,1 2,4 2,3 2,3 2,1 1,9 1,8 1,4 1,4 Fold 2,4 2,3 2,1 2,1 2,1 2,1 2,1 2,1 2,1 2,1 2,1 2,1	GO 0005975-carbohydrate metabolic process GO:0006013-mannose metabolic process MF: ajhra-glucosidase activity GO:0006013-mannose metabolic process MF: ajhra-glucosidase activity GO:00060671-glycerol metabolic process MF: lipase activity GO:0006665-aphingolipid metabolic process GO:0006665-aphingolipid metabolic process GO:0006665-aphingolipid metabolic process MF: sterio eleterase activity	GATAe yes yes yes GATAe yes yes	yes yes yes yes Sugar only	yes yes yes yes yes	yes yes yes yes	down down down up down down down	inf/basal inf inf inf/basal inf/basal inf inf	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mtn50 obst-A CG33523 GATAe CG42397 Nha1 Amnionless Pask	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,4 1,3 1,3 1,2	GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006803-chitim metabolic process MF. fibringen/ associated to wound healing GO.0042060-wound healing GO.0055086-transmethrane transport GO.0016893-drug transport GO.000508-chitim metabolic process Interpro vesicule-associated membrane protein GO.007482-endoderm development GO.007482-endoderm development GO.0006811-ion transport Interpro Ammionless, Vil 812 transporter GO.0007165-signal transduction	yes yes yes	yes yes yes Sugar only	ŕ		na down na up na down down na	na na na inf/basal na inf/basal inf na
Gene Name C9483 C9483 C9483 C9486 C9	Fold 3,1 2,4 2,4 2,3 2,3 2,1 1,9 1,8 1,4 Fold 2,4 2,3 2,3 2,1 1,9	GO 0005975-carbohydrate metabolic process GO 0006013-mannose metabolic process MF. sipha-glucosidase activity GO 0006013-mannose metabolic process MF. sipha-glucosidase activity MF. sipha-amylisias eactivity MF. sipha-glucosidase activity GO 0006631-fatty activi	GATAe yes yes yes yes GATAe yes yes yes	yes yes yes yes Sugar only yes yes	yes yes yes yes yes yes yes yes yes	yes yes yes yes	Ecc15 down down down down up down down down down down down down down	inf/basal inf inf inf/basal inf/basal inf inf Rel na inf/basal	MtnB MtnC Muc68E CG5550 CG15170 CG30272 Mtn50 obst-A CG33523 GATAe CG42397 Nha1 Amionless Pask	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,4 1,3 1,3 1,2	GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006803-chitim metabolic process MF. fibringen/ associated to wound healing GO.0042060-wound healing GO.0055086-transmethrane transport GO.0016893-drug transport GO.000508-chitim metabolic process Interpro vesicule-associated membrane protein GO.007482-endoderm development GO.007482-endoderm development GO.0006811-ion transport Interpro Ammionless, Vil 812 transporter GO.0007165-signal transduction	yes yes yes	yes yes yes Sugar only yes	ŕ	yes	na down na up na down down na	na na na inf/basal na na inf/basal inf na inf/basal na inf/basal na inf/basal na na inf/basal na inf
Gene Name CG9463 Mal-A7 CG9466 Amy-6d Mal-A3 Mal-A8 Mal-A8 Mal-A6 CG16965 Gal CG9394 Gene Name CG6296 CG15533 mag CG8093 CG9927 Gene Name	Fold 3,1 2,4 2,4 2,3 2,3 2,1 2,1 1,9 1,4 1,4 1,4 1,4 1,4 1,4 1,4 1,4 1,4 1,4	GO 0005975-carbohydrate metabolic process GO:0005013-mannose metabolic process MF: ajha-glucosidase activity GO:0006013-mannose metabolic process MF: ajha-glucosidase activity MF: ajha-amylase activity MF: ajha-glucosidase activity MF: alpha-glucosidase activity MF: place-glacistodisea activity GO:0006071-ajlectorial metabolic process MF: [spase activity GO:0006663-pinglogiligh metabolic process GO:0006665-pinglogiligh metabolic process GO:006665-pinglogiligh metabolic process MF: stere activity MF: [spase activity GO:0006651-fatty acid metabolic process GO:00066051-fatty acid metabolic process GO:00066051-fatty acid metabolic process GO:0006607-response to biolic stimulus	GATAe yes yes yes GATAe yes GATAe GATAe	yes yes yes yes Sugar only	yes yes yes yes yes yes yes yes yes	yes yes yes yes	Ecc15 down down down down up down down down down down down down	inf/basal inf inf inf inf/basal inf/basal inf Rel na inf/basal	MinB MinC Muc68E CG65590 CG15170 CG30272 Mtr50 Odst-A CG323523 GATAe CG42397 Nha1 Amioniless Pask	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,3 1,3 1,2 Fold 3,5 2,6 2,4 1,8	GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006803-chitim metabolic process MF. fibringen/ associated to wound healing GO.0042060-wound healing GO.0055086-transmethrane transport GO.0016893-drug transport GO.000508-chitim metabolic process Interpro vesicule-associated membrane protein GO.007482-endoderm development GO.007482-endoderm development GO.0006811-ion transport Interpro Ammionless, Vil 812 transporter GO.0007165-signal transduction	yes yes yes	yes yes yes Sugar only yes	ŕ	yes	na down na up na down na down up	na na na inf/basal na inf/basal inf na na na na na na na na na inf/basal inf/basal inf/basal inf/basal inf/basal inf/basal inf/basal inf/basal inf/basal
Gene Name CG9463 MgI-A7 CG9466 Amy-d MgI-A7 MGI-A8 MGI-A8 MGI-A8 MGI-A9	Fold 3.1 2.4 2.4 2.3 2.3 2.1 2.1 1.9 1.8 1.4 1.4 Fold 2.3 2.3 2.1 1.9 1.4 Fold 5.2	GO 0005975-carbohydrate metabolic process GO 0006013-mannose metabolic process MF. sipha-glucosidase activity GO 0006013-mannose metabolic process MF. sipha-glucosidase activity MF. sipha-amylisias eactivity MF. sipha-glucosidase activity GO 0006631-fatty activi	GATAe yes yes yes GATAe yes GATAe GATAe	yes yes yes yes Sugar only yes yes	yes yes yes yes yes yes yes yes yes	yes yes yes yes yes Myc yes	Ecc15 down down down down down down down down	inf/basal inf inf inf/basal inf/basal inf inf Rel na inf/basal	MtnB MtnC Muc68E CG590 CG590 CG590 OG550 OG550 OG3523 GATAe CG42397 Nha1 Amionless Pask	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,3 1,3 1,2 Fold 3,5 2,6 2,4 1,8 1,6	GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006803-chitim metabolic process MF. fibringen/ associated to wound healing GO.0042060-wound healing GO.0055086-transmethrane transport GO.0016893-drug transport GO.000508-chitim metabolic process Interpro vesicule-associated membrane protein GO.007482-endoderm development GO.007482-endoderm development GO.0006811-ion transport Interpro Ammionless, Vil 812 transporter GO.0007165-signal transduction	yes yes yes	yes yes yes Sugar only yes	ŕ	yes	na down na na na na na na na down up na down na	na na na inf/basal inf na inf/basal na inf/basal na inf/basal na inf/basal na
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Gene Name CG9463 Mal-A7 CG9468 Amy-d Mal-A7 CG9468 Amy-d Mal-A8 Mal-A8 Mal-A8 CG16965 CG19939 Gene Name CG6296 CG15533 may CG8093 CG957 Gene Name PRPP-SC1 pirk	Fold 3.1 2.4 2.4 2.3 2.1 2.1 1.9 1.8 1.4 1.4 1.4 Fold 2.4 2.3 2.1 1.9 1.4 Fold 5.2 2.5	GO 0005975-carbohydrate metabolic process GO:0005013-mannose metabolic process MF: ajha-glucosidase activity GO:0006013-mannose metabolic process MF: ajha-glucosidase activity MF: ajha-amylase activity MF: ajha-amylase activity MF: ajha-glucosidase activity GO:0006071-gl-glecosidase activity GO:0006071-gl-glecosidase activity MF: lipase activity GO:0006653-plingolipid metabolic process GO:0006651-fatro eletrase activity MF: lipase activity GO:0006651-fatro eletrase activity GO:0006651-fatro eletrase activity GO:0006671-response to biolic stimulus GO:0000270-peptidoglycan metabolic process system	GATAe yes yes yes GATAe yes yes GATAe yes yes yes	yes yes yes yes Sugar only yes yes	yes yes yes yes yes yes yes yes yes	yes yes yes yes yes Myc yes	Ecc15 down down down down down down down down	inf/basal	MtnB MtnC Muc68E CG5559 CG15170 CG30272 GG30272 GG30273 Mha1 Aminioness Pask	1,9 1,8 1,6 1,6 1,6 1,5 1,5 1,4 1,4 1,3 1,3 1,2 Fold 3,5 2,6 1,8 1,8 1,8 1,3	GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006873-cellular ion homeostasis GO.006803-chitim metabolic process MF. fibringen/ associated to wound healing GO.0042060-wound healing GO.0055086-transmethrane transport GO.0016893-drug transport GO.000508-chitim metabolic process Interpro vesicule-associated membrane protein GO.007482-endoderm development GO.007482-endoderm development GO.0006811-ion transport Interpro Ammionless, Vil 812 transporter GO.0007165-signal transduction	yes yes yes GATAe	yes yes yes Sugar only yes	ŕ	yes	na down na up na down na na na na na na na na down up na down down	na na na inf/basal na inf/basal na inf na na na na na na na inf/basal na inf/basal na inf/basal na inf/basal na basal

Figure 2. Microbiota-regulated genes. Microbiota-regulated genes are classified by Gene Ontology (GO):Biological function groups. Columns for each microbiota-regulated gene are designated as such: (1) Gene names from Flybase; (2) fold induction in microbiota associated animals vs GF; (3) information on gene function: lower hierarchy GO:Biological process or GO:Molecular function or insight on molecular function by mentioning an Interpro domain present in the protein; (4) Regulation by GATAe; (5) Gene deregulated upon sugar-only diet challenge; (6) Regulation by Sug; (7) Regulation by Myc; (8) Gene regulation in the midgut upon *Ecc15* infection (*Ecc15*) and (9) Relish dependence (Rel regulation) for gene expression in the midgut in basal condition (basal) or upon *Ecc15* infection (inf). For columns "GATAe" the data were collected from [21,27]; columns "Sugar only" and "Sug" data are from [29]; column "Myc" data are from [28]; columns "*Ecc15*" and "Rel" data are from [31].

candidate to mediate the transcriptional host response to microbiota association.

In summary, the host transcriptomic response to microbiota association includes the modulation of a significant number of genes required to adapt to nutritive challenges. This finding suggests that microbiota association potentiates nutrition via enhanced digestive enzyme expression mediated at least partly via Sug inhibition and Myc activation.

Trade-off between microbiota-mediated and infectionmediated midgut genes expression

Buchon et al. characterized the transcriptional signatures in dissected CONV adult midguts after an acute oral infection with the bacterial strain Erwinia carotovora carotovora 15 (Ecc15). Our current study shares many common features with that of Buchon et al., in that we adopted similar experimental protocols to study changes in transcriptomes after exposing the host to a specific set of bacteria and we both found a marked transcriptional response localized to the midgut. We therefore compared our dataset to that of Buchon et al. and found an evident overlap between the two lists: half of our microbiota-regulated genes are also modulated upon intestinal infection (52/105 genes, Fig.2). As expected, like in Buchon et al., we also found that the IMD pathway target genes (AttD, pirk, PGRP-LB/-SC1/-SC2) were up-regulated. However, most microbiota up-regulated genes (35/52) were in fact downregulated upon Ecc15 infection, suggesting the existence of a transcriptional trade-off between the response to indigenous bacteria (i.e the microbiota) and the response to infectious bacteria. We tested this hypothesis by infecting the flies associated with a standardized microbiota with Ecc15 and studying the expression of a selection of candidate genes from our list (Fig.3). As expected, we found that the immune-related genes pirk and AttD are up-regulated, and we confirmed that several microbiotaregulated genes such as digestive enzymes (CG3739, CG7025, CG9463), are markedly down-regulated upon Ecc15 infection. This result reinforces the notion that Ecc15 infection triggers a transcriptional trade-off to promote immune-related genes expression at the expense of the metabolic genes expression up-regulated by the microbiota association.

IMD/Relish pathway at least partly controls the expression of microbiota-regulated metabolic genes

Relish encodes the Drosophila orthologue of the mammalian NFkappaB factor p105, which functions downstream of the IMD signaling pathway and controls the induction of most IMD target genes [30]. Buchon et al. studied the midgut transcriptome of Relish mutants and found that changes in the expression of midgut genes upon Ecc15 infection are largely Relish-dependent [31]. Strikingly, from our list of genes, the vast majority of the 52 microbiotaregulated genes influenced by Ecc15 infection are also Relishdependent (41/52; Fig.2). Based on the dataset from Buchon et al., we derived the information of Relish dependence for the basal expression level of each midgut gene in CONV animals in their study, and compared this dataset with our list of microbiotaregulated genes. We found that 39 of the microbiota-regulated genes rely on Relish activity for their basal expression in the midgut of CONV animals (Fig.2). Strikingly, all these 39 genes were also regulated in a Relish dependent manner upon Ecc15 infection in CONV animals. These observations suggest that Relish, in addition to its known role to control the expression of immune-related genes, may also be an important transcriptional regulator of metabolic genes induced by the microbiota, which are likely independent of immune responses. To test this hypothesis,

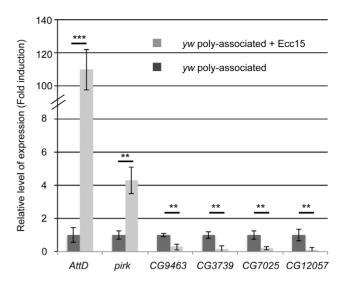


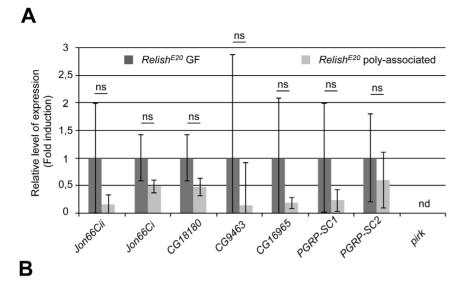
Figure 3. *Ecc15* infection promotes expression of microbiotaregulated immune genes while repressing microbiota-regulated metabolic genes in midguts. A bar graph representing the fold-enrichment of different microbiota-regulated transcripts in dissected midguts from poly-associated upon oral infection by *Ecc15* in WT animals (the value of the relative $\Delta Ct^{gene}/\Delta Ct^{rp49}$ ratios was calculated for every sample and relativized to the ratio in the GF condition which was anchored to 1 to indicate fold induction). Statistical significance of the result is represented (Student's t-test: $ns\geq0.05>*\geq0.01>***\geq0.001>***$).

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we studied the expression of a set of eight microbiota-regulated genes in the midguts of two mutants of the IMD signaling pathway: *Dredd* and *Relish. Dredd* encodes the *Drosophila* orthologue of Caspase-8 whose function is essential for IMD pathway signal transduction and Relish activation [32]. In *Dredd* or *Relish* mutant background, both immune-related (*pirk*, *PGRP-SC1/SC2*) and metabolism-related (*Jon66Cii*, *Jon66Ci*, *CG16965*, *CG18180* and *CG9463*) microbiota-regulated genes are no longer induced in the midgut upon standardized microbiota association (Fig.4). These results demonstrate that microbiota impacts metabolic gene expression partly via IMD/Relish activity.

Conclusion

The Drosophila indigenous microbiota modulates host physiology [2,3]. In this study, we have identified molecular signatures associated to these effects and pinpointed the central role of the IMD/Relish signaling pathway in controlling host transcriptional response to its microbiota. One striking result in our study is that the host transcriptome response to microbiota association is mostly restricted to the midgut, a major biological interface between the host and its environment and the primary site where host/ microbiota interactions occurs [4]. As described in previous studies, microbiota association triggers a transcriptional change related to host response to bacteria with similar molecular signatures (i.e, immune-related genes) to those elicited by pathogenic bacteria infection. However, microbiota association clearly favors a unique transcriptional response funneled towards promoting host metabolic capacities. Such response is severely dampened upon bacterial infection as a trade-off for the host to mount potent immune and tissue repair responses. Since the IMD/Relish pathway is instrumental to promote both the metabolic response to microbiota association and the response to infection, it is likely that the transcription factor Relish is at the



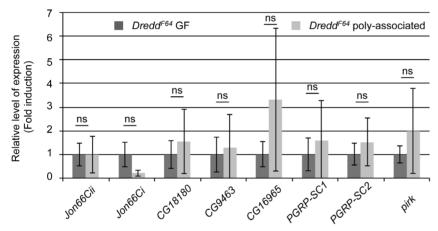


Figure 4. IMD/Relish signaling controls microbiota mediated transcript enrichment in midguts. Bar graphs representing the fold enrichment of different microbiota-regulated transcripts in dissected midguts from GF vs poly-associated $Relish^{E20}$ (A) or $Dredd^{F64}$ (B) mutant animals (the values of the relative $\Delta Ct^{gene}/\Delta Ct^{rp49}$ ratios were calculated for every sample and relativized to the ratio in the GF condition which was anchored to 1 to indicate fold induction). Statistical significance of the result is represented (Student's t-test: $ns\geq0.05>*\geq0.01>***\geq0.001>***$). nd: not detected.

doi:10.1371/journal.pone.0094729.g004

cornerstone of the transcriptional trade-off between the midgut response to beneficial microbiota and response to midgut pathogens. Other factors may also contribute to this trade-off, such as ATF3, which was recently reported to control immune and metabolic homeostasis in the *Drosophila* midgut [33].

Taken together, our results demonstrate that *Drosophila* microbiota has a marked impact on the expression of genes mainly involved in digestive functions and primary metabolism, suggesting that microbiota association potentiates host nutrition and host metabolic state, two key physiological parameters contributing to host fitness. Our results are in agreement with recent reports demonstrating that microbiota influence adult nutritional and metabolic phenotypes [15] and therefore pave the way to the subsequent mechanistic studies on how these microbiota-dependent transcriptional responses translate into host physiological benefits.

Materials and Methods

Drosophila lines and breeding

Drosophila were cultured at 25°C on a standard yeast/cornmeal medium containing 10 g.L⁻¹ agar (VWR, ref. #20768.361), 80 g.L⁻¹ cornmeal flour (Westhove, Farigel maize H1), 50 g.L⁻¹ inactivated dry yeast (Bio Springer, Springaline BA95/0-PW), 5.2 g.L⁻¹ Methylparaben sodium salt (MERCK, ref. #106756) and 4 ml.L⁻¹ of 99% propionic acid (CARLO ERBA, cref. #409553). Germ-free animals were obtained from bleached embryos cultured on autoclaved conventional medium. When needed GF stocks were maintained during few generations on antibiotic supplemented food (final concentration: 50 μg.L⁻¹ ampicilline, 50 μg.L⁻¹ kanamicine, 15 μg.L⁻¹ erythromycin, 50 μg.L⁻¹ tetracycline) to avoid bacterial contamination. Drosophila y,w flies were used as the reference strain in this work. The following mutant lines were used: y,w,Dredd^{F64} and y,w;;Relish^{E20} [34].

Bacterial strains and culture conditions

Erwinia carotovora carotovora¹⁵ (Ecc15), Lactobacillus plantarum ^{WJL}, Lactobacillus brevis ^{EW}, Commensalibacter intestini ^{A911T}, and Acetobacter pomorum strains were used in this study. All strains were described previously [18,35]. A. pomorum and C. intestini were cultivated in Mannitol medium (3 g.L⁻¹ Bactopeptone (Difco, cat. #0118-17), 5 g.L⁻¹ yeast extract (Difco, cat. #212750), 25 g.L⁻¹ D-Mannitol (Sigma, ref. M1902) at 30°C at least for 18 hours on agitation, L. brevis and L.plantarum were cultivated in Man, Rogosa and Sharpe (MRS) broth (Difco, ref. #288110) at 37°C standing. Ecc15 was cultivated in Luria-Bertani broth medium (Difco, ref. #244610) at 30°C for 24 hours on agitation.

Poly-association of GF adults with a standardized microbiota and Ecc15 infection

Association mixture: 150 µL of bacterial solution made of 75 µL of 5% sucrose solution sterilized by filtration through a 0.2 µm membrane (Pall Life Sciences, ref. #4652)+75 µL of a mix of equal amounts of four commensal bacterial cultures (Lactobacillus plantarum, Lactobacillus brevis, Commensalibacter intestini and Acetobacter pomorum) at an initial $OD_{600} = 1.0$ each. Control mixture: 150 µL of a control solution (75 μ L filter-sterilised sucrose +75 μ L of a 1:1 mixture of sterile MRS and sterile Mannitol). Either of these solutions was added to a disc of autoclaved paper (Whatman, ref. #3030917) covering standard fly culture media in standard fly tubes. For Ecc15 infection of poly-associated flies, Ecc15 culture at final $OD_{600} = 100$ resuspended in PBS was added to the mix of commensal bacterial strains and inoculated onto autoclaved paper disc. Germ-free adults derived from freshly made GF embryos were collected within the first 48 hours of adult emergence and placed in groups of 25-30 females and 10 males in experimental tubes. Flies were kept in such tubes for 2 days at 25°C and transferred into newly prepared vials with fresh bacterial cultures. 3 days after transfer, on day 5 post-inoculation flies were either sampled or infected with Ecc15 for 8 hrs. In each experiment we measured internal bacterial loads of representative experimental animals by plating fly homogenates on MRS and Mannitol agar plates to check the effectiveness of the re-association and infection processes or the axenic status of the flies. On average flies carried 10⁴ CFUs/animal upon re-association and >10⁶ CFUs/animals upon infection.

Microarray analysis

Three biological replicates of 20 females for each condition (GF or poly-associated) were homogenized in 500 μ l of Trizol (Invitrogen) and 100 μ l Choloroform using 0.75–1 mm glass beads and the Precellys24 Tissue Homogenizer (Bertin Technol-

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ogies, France). RNA pools were isolated and purified using NucleoSpin RNA kit (Macherey-Nagel). RNA was quantified on NanoDrop ND-1000 and RNA quality was controlled on Agilent 2100 Bioanalyzer chips. For each sample, 1 μg of total RNA was amplified and labeled using the GeneChip IVT Labeling Kit according to the protocol provided by the supplier. Affymetrix GeneChip Drosophila Genome 2.0 arrays were hybridized with 7.5 µg of labeled cRNA, washed, stained and scanned according to the Affymetrix' protocols. Raw data are deposited at NCBI GEO with the accession number: GSE56173. Raw data analysis was performed using R for Statistical Computing (R core team 2013) with Robust Multi-array Average (RMA) and Significance Analysis of Microarrays (SAM) from the package siggenes (http:// bioconductor.wustl.edu/bioc/html/siggenes.html). Gene selection was made using a value of Delta that correspond to a FDR of 0.2. Gene Ontology Term analysis and enrichments were made using DAVID [23].

RNA extraction and RT-qPCR analysis

Three biological replicates of a pool of 10 midguts dissected from females were used for RT-qPCR analysis. Foregut, hindgut, crop and malpighian tubes were carefully removed. Tissue homogenization, RNA extraction and purification were performed as described above for whole flies. Reverse transcription of 300 ng of RNA was performed using Superscript II enzyme (Invitrogen) and random primers (Invitrogen). Quantitative PCR was performed on a Biorad CFX96 apparatus (Biorad) using SYBR GreenER qPCR Supermix (Invitrogen), cDNA (1/160 dilution of the reverse transcription products) and gene specific primer sets (available upon request). Melting curves of the detected amplicons were analysed to ensure specific and unique amplification. PCR efficiency was calculated using serial dilution of cDNA. We used the $\Delta\Delta$ Ct method for data analysis and p49 as the reference gene. Results were expressed as a relative value of $\Delta Ct^{gene}/\Delta Ct^{rp49}$ ratios (fold induction).

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Author Contributions

Conceived and designed the experiments: BEC AD FL. Performed the experiments: BEC AD NB. Analyzed the data: BEC AD NB DP JR FL. Contributed reagents/materials/analysis tools: DP JR. Wrote the paper: BEC FL.

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