# Research **Expression profiling of Drosophila imaginal discs** Ansgar Klebes\*, Brian Biehs\*, Francisco Cifuentes\*<sup>†</sup> and Thomas B Kornberg\*

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# Abstract

**Background:** In the *Drosophila* larva, imaginal discs are programmed to produce adult structures at metamorphosis. Although their fate is precisely determined, these organs remain largely undifferentiated in the larva. To identify genes that establish and express the different states of determination in discs and larval tissues, we used DNA microarrays to analyze mRNAs isolated from single imaginal discs.

**Results:** Linear amplification protocols were used to generate hybridization probes for microarray analysis from  $poly(A)^+$  RNA from single imaginal discs containing between 10,000 and 60,000 cells. Probe reproducibility and degree of representation were tested using microarrays with approximately 6,000 different cDNAs. Hybridizations with probes that had been prepared separately from the same starting RNA pool had a correlation coefficient of 0.97. Expression-profile comparisons of the left and right wing imaginal discs from the same larva correlated with a coefficient of 0.99, indicating a high degree of reproducibility of independent amplifications. Using this method, we identified genes with preferential expression in the different imaginal discs using pairwise comparisons of discs and larval organs. Whereas disc-to-disc comparisons revealed only moderate differences, profiles differed substantially between imaginal discs and larval tissues, such as larval endodermal midgut and mesodermal fat body.

**Conclusion:** The combination of linear RNA amplification and DNA microarray hybridization allowed us to determine the expression profiles of individual imaginal discs and larval tissues and to identify genes expressed in tissue-specific patterns. These methods should be widely applicable to comparisons of expression profiles for tissues or parts of tissues that are available only in small amounts.

#### Background

During the development of multicellular organisms, complexity builds sequentially in discrete steps as cells proliferate and their descendants choose between alternative developmental fates. Much of our understanding of these processes in *Drosophila* has come from mutants that have developmental defects. There are many examples. For instance, mutations in the homeodomain protein genes of the Antennapedia and Bithorax complexes provide evidence for the role of transcription factors in making developmental decisions. Flies with gain-of-function mutations in the *Antennapedia* (*Antp*) gene cannot grow normal antennae, but instead make extra legs with their antennal cells. Conversely, loss of Antp function causes transformation of the second leg to antenna-like structures [1]. Another example is the *glass* gene, which encodes a zinc-finger transcription factor that is expressed in many of the cell types in the eye imaginal disc and is required for photoreceptor development [2]. The common functions of these and many other genes in regulating gene expression suggest that developmental fates are manifested in part in the transcripts that different cell types produce.

Although the number of Drosophila mutants with interesting developmental phenotypes is large, we cannot assume that the genetic screens that have been carried out identified all the relevant genes. Many Drosophila genes have been refractory to genetic analysis, either because they are duplicated and may code for redundant functions, or because they have mutant phenotypes that are difficult to recognize. In addition, genetic and biochemical approaches have yielded few downstream genes that the key transcription factors regulate. The recent release of the Drosophila genome sequence [3] and the demonstration that cDNA microarrays can be used to catalog the transcriptome of both unicellular and multicellular organisms [4-20] opens up a new approach. Hybridization of mRNA pools to DNA microarrays can potentially identify the differences in gene activity that define every developmental state. Thus, transcriptional regulators and their downstream targets can be identified simultaneously.

The Drosophila larva contains two developmentally distinct sets of organs. One set comprises the vital organs of the larva - the epidermis and musculature, the central and peripheral nervous systems, and the organs of the digestive tract. Most of these tissues will be destroyed at metamorphosis and will be replaced by imaginal (adult) cells. The imaginal precursor cells have no functional role in the larva. For some adult structures such as the eyes, antennae, legs, wings, halteres and genitalia, the imaginal precursors are sequestered as physically distinct primordia - the imaginal discs. The different discs are determined as to disc type, but do not differentiate until the late larval and subsequent pupal stages when they form the adult structures. The distinct developmental phases of the imaginal discs, their easy accessibility, along with the numerous advantages of Drosophila as a model genetic and developmental organism make imaginal disc development an ideal system to explore the feasibility and usefulness of such an analysis at a genomic scale.

The principal technical difficulty such a study must surmount is that the quantity of probe needed for efficient hybridization to microarrays is large relative to the amount of RNA contained in a single *Drosophila* imaginal disc. Approximately 4  $\mu$ g poly(A)<sup>+</sup> RNA is required to make a probe, but a third instar wing imaginal disc has only about 1-4 ng (Figure 1). Thus, several thousand discs would need to



#### Figure I

Quality and amount of total RNA preparations from imaginal discs. Total RNA was prepared separately from 2, 4 and 12 third-instar larval wing imaginal discs using the Mini RNA Isolation Kit (Zymo Research). The total amount of RNA, based on the absorbance at 260 nm ( $A_{260}$ ), was 462, 540 and 1,530 ng, respectively. The A260/A280 ratios of 2.08, 2.2 and 2.27 for the separate preparations is indicative of high-quality RNA preparations. The amount of total RNA per disc was calculated to be 230, 140 and 130 ng, respectively. Assuming a poly(A)<sup>+</sup> RNA content of 1-2%, this amount of total RNA roughly corresponds to 1-4 ng poly(A)<sup>+</sup> RNA per wing disc (62,000 cells). The other discs are smaller and contain fewer cells. The equivalent of 1, 2 and 6 discs was separated on a denaturing agarose gel (lanes 1, 2 and 3, respectively). The two prominent bands represent the 18S and 28S ribosomal RNA populations (arrowheads). Poly(A)<sup>+</sup> RNA is detected as a smear. No obvious small-molecular-weight products were observed. Lane M contains molecular-weight markers, with numbers indicating the approximate lengths in nucleotides.

be dissected for each set of experiments. In previous reports of expression profiling, *Drosophila* embryos, adult flies and adult heads, and collections of animals were pooled after carefully timing or otherwise selecting individual animals [9,14,18,20]. However, any scheme to stage animals is inherently inexact. In addition, parameters such as genetic variability, nutritional state, pathogen exposure and effects of the isolation procedure cannot be easily controlled or measured. Pools of animals therefore yield only an average of their varied inputs, and depending on the extent of variability, critical differences between samples may be submerged. Furthermore, the level of resolution is relatively low when RNA preparations from whole animals are used. For many biological questions, it is necessary to detect differences in subsets of cells that might account for only a small fraction of the many thousands of cells in the animal. To circumvent these problems, we used techniques that allow us to carry out several hybridizations with probes derived from different tissues of a single animal.

We applied linear RNA amplification methods that were first introduced by Eberwine and co-workers [21,22] and have been refined by Baugh et al. [23]. We used two rounds of reverse transcription (RT) and in vitro transcription (IVT), and achieved as much as 1-5 x 105-fold amplification. The amplified RNA (aRNA) from different tissues of single larva was used to perform pairwise comparisons and to identify sets of genes with preferential expression. The preferential expression of many of the genes in these sets had been previously shown by genetic or molecular analysis. We also confirmed the preferential expression of ten genes that had not been previously characterized by in situ hybridization to imaginal discs. Several of these genes had particularly interesting patterns of expression. Our findings show that this method can be applied to determine the expression profile of individual tissues and to identify candidate genes with specific expression patterns.

# **Results and discussion Principle of amplification**

The amplification procedure we used is based on protocols described by Eberwine [21,22,24], Wang et al. [25], and Baugh et al. [23], and details are given in Materials and methods. Briefly, double-stranded cDNA was synthesized either from total RNA isolated from imaginal discs or from poly(A)+-purified embryonic RNA. The cDNA was used as a template for in vitro transcription, routinely generating an RNA product that represented an amplification of approximately 1,000 fold. This RNA was subjected to a second round of RT and IVT. The yield from the second round represented an approximately 100-fold amplification, resulting in an overall amplification of 1-5 x 105. Probes for hybridization were synthesized by RT and labeled with either Cy5 (red) or Cy3 (green). Arrays were hybridized simultaneously with Cy5- and Cy3-labeled probes, and the intensities and ratios of bound fluor were measured for each spot. Because of inherent variability in the hybridization efficiency of individual microarrays, only the ratios of the red and green fluorescence were interpreted to indicate relative transcript abundance in the starting pool of poly(A)<sup>+</sup> RNA.

A known drawback of linear amplification is that sequences at the 5' end of transcripts can be preferentially lost. This reduction in sequence complexity can probably be attributed to incomplete RT, to priming from subterminal locations, and to incomplete IVT. To overcome this problem, we supplemented the random hexanucleotides that were used to prime the second RT reaction with a template-switch (TS) primer [25]. This primer includes a guanosine triplet that is designed to pair with a 3' cytidine overhang created by the reverse transcriptase (Clontech [25]). Control amplifications with the TS primer alone or random hexanucleotides alone vielded similar quantities of aRNA. When these aRNA preparations were used for microarray hybridizations, many spots had significantly different signal intensities (data not shown). The 25 cDNAs with the most intense signals for the probe amplified with the TS primer alone had an average length of 1.6 kilobases (kb). In contrast, after amplification with hexanucleotides alone, the 25 cDNAs with the most intense signals had an average length of 1.1 kb. We interpret the bias towards long transcripts in TS-primer amplified probes as indicating better preservation of 5'-complexity.

#### **Representation of transcripts in amplified probes**

The most important feature of RNA pools that amplification must preserve is proportionality, the maintenance of the relative concentration of different RNAs. We established the quality of our amplification method in several ways. First, we isolated poly(A)+ RNA from embryos and subjected 100 pg, 1 ng and 10 ng aliquots to two rounds of linear amplification. The size distribution of the aRNA ranged from about 100 nucleotides to high-molecular-weight products, with the majority between 240 and 2,000 nucleotides (Figure 2a). aRNA from independent amplifications was then used to generate probes and to compare hybridization efficiencies of pairs of probes. Such separate amplifications of aliquots of 1 ng of the same embryonic poly(A)+ RNA produced signals with similar intensities. A scatterplot of the Cy3 and Cy5 signal intensities showed a tight correlation coefficient (CC) of 0.97, with a low standard deviations (SD) of the red/green (R/G) ratios for all spots (SD 0.25, log<sub>2</sub>transformed) (Figure 2b). Comparisons of probes made from 100 pg and 1 ng and from 1 ng and 10 ng poly(A)<sup>+</sup> RNA yielded essentially the same level of reproducibility (CC 0.95, SD 0.29; data not shown). We conclude that the amplifications were reproducible and that good proportionality of RNA was maintained during amplification even if the concentration of poly(A)+ RNA differed by tenfold.

Comparisons between amplified and unamplified RNA indicated that essentially no transcripts were lost during amplificaton. Of the spots that hybridized to probes generated from amplified and unamplified embryonic poly(A)<sup>+</sup> RNA 99% (5,514 out of 5,574) did so with both probes. Differences in intensities were, nevertheless, significant (CC 0.53 and SD 1.13, data not shown). We do not know whether these differences in intensities were a result of changes in relative abundance or in transcript length, both of which will affect hybridization signals. Nevertheless, the good correspondence of the hybridizing spots shows that the aRNA is an essentially complete representation of the population of



Linear amplification is highly reproducible. (a) Denaturing gel electrophoresis shows the size distribution of aRNA obtained after two rounds of linear amplification of 1 ng poly(A)<sup>+</sup> RNA. Lane 1, total RNA; lanes 2 and 3, products of independent amplifications; M, molecular-weight markers, with numbers indicating the approximate lengths in nucleotides. (b,c) Scatterplots of the Cy3 and Cy5 signal intensities from hybridizations of two probes derived from independent amplifications of embryonic or wing-disc RNA. Starting materials were (b) 1 ng embryonic poly(A)<sup>+</sup> RNA and (c) left and right wing imaginal discs of one wandering third-instar larva. CC, correlation coefficient; SD, standard deviation. SDs were calculated on the normalized  $log_2^{-}$  transformed R/G ratios.

transcripts present in the initial pool. We conclude that comparisons of two different probes on microarrays are valid if the RNA pools have been amplified in the same way.

# Pairwise comparisons of imaginal discs and larval organs

We carried out 43 separate hybridizations, comparing 13 different pairwise combinations of tissues (Table 1). This analysis identified many genes that are preferentially expressed in specific tissues; it also provided an independent and quantitative measure of the distinct nature of larval tissue and imaginal disc cells.

Hybridizations that were carried out were of two types: with probes generated from different imaginal discs; and with probes from imaginal discs and larval organs. Comparisons between imaginal disc probes revealed a high degree of similarity, but comparisons between imaginal disc and larval organ probes identified many differentially expressed genes. We now give a general description of these results, starting

Tab	le
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Overview	of the	comparisons
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Comparison	Number of experiments	
Wing : wing	I	
Wing : eye-antenna	П	
Wing : legl	2	
Leg1 : leg2	3	
Leg1 : leg3	2	
Leg2 : leg3	2	
Haltere : genital	5	
Wing : fat body	5	
LegI : fat body	2	
Haltere : fat body	I.	
Wing : brain/optic lobe	3	
Wing : salivary gland	3	
Wing : midgut	3	

Forty-three pairwise comparisons were carried out. Of these, 26 were between imaginal discs; 17 were between an imaginal disc and a larval organ (indicated in bold). The numbers indicate the times each type of comparison was repeated.

with those tissues that were most alike. It should be noted that these results were obtained with microarrays that represent a pool of cDNAs obtained from the Berkeley *Drosophila* Genome Project (BDGP) containing approximately 44% (6,000/13,600) of the transcription units predicted to be in the *Drosophila* genome. As this pool may not be a truly random subset, extrapolations to the whole genome may be of limited value. Indeed, a preliminary comparison of our cDNA arrays with microarrays that contain short cDNA sequences for all annotated genes suggests that the total number of genes that produce a signal with these probes is similar. It is possible that the set of 6,000 cDNAs distributed by the BDGP is biased toward genes expressed in the tissues we analyzed.

#### **Expression profiles of imaginal discs**

# Left and right wing discs

We first asked whether microarray hybridization could detect differences in expression profiles between left and right wing discs. Although few differences might be expected, previous studies have revealed a left/right difference in wing size, suggesting that the respective developmental programs are not identical [26]. When probes were prepared independently from the poly(A)<sup>+</sup> RNA isolated from left and right wing discs of a third instar larva, only minor differences in the intensities were observed (CC 0.99 and SD 0.14) (Figure 2c). This tight correlation again shows the reproducibility of the amplification procedure.

# Leg discs

We then asked if any of the genes represented on our arrays were differentially expressed in first, second or third leg discs. Among the several genes that have been shown to be differentially required or expressed in the different leg discs (for example, Ultrabithorax [27], Antp [28] and Sex combs reduced [29,30]), only Antp was among the set of cDNAs on our microarrays. Comparisons of probes from first, second and third leg discs yielded CC values in the 0.93 ranges. To identify differentially expressed genes in the leg discs, cluster analysis was used to group genes with similar expression profiles [31]. For this and all subsequent analyses a threshold setting of > 1.74 (=  $0.8 \log_2$ -transformed ratios) was applied to identify subclusters. We identified 2, 12 or 17 differentially expressed genes in the leg1:leg3, leg2:leg3 and leg1:leg2 disc arrays, respectively (Figure 3). The induced expression of *Antp* in the second leg discs was noteworthy. This finding is consistent with previous studies showing that although Antp is expressed in all three leg discs, it is expressed at a much greater level in second leg discs [28].

#### Leg and wing discs

Comparisons were also made between the first leg and wing imaginal discs (two experiments). Cluster analysis identified 23 genes that were preferentially expressed in the leg discs and 8 in the wing discs (Figure 4). Notable in the wing cluster are *broad*, which is expressed in imaginal cells and encodes a family of transcription factors [32], and *apterous*, which is expressed in the dorsal wing cells and is required for wing formation [33].

#### Wing and eye-antennal imaginal discs

Comparing probes from the wing and eye-antennal discs isolated from the same larva revealed a higher degree of divergence. Eleven independent wing:eye-antennal comparisons were made. Twenty-four genes were identified that had elevated expression in the wing discs, and 73 genes were identified with elevated expression in eye-antennal discs (Figure 5). Many of these genes are known from previous studies to be expressed specifically in these tissues. Among the genes with elevated expression in wing discs were *apterous* and engrailed. apterous is expressed in the dorsal compartment of the wing disc, but not in the eye-antennal disc; engrailed is expressed in the posterior compartment cells of both the wing and eye-antennal discs. However, inclusion of engrailed in the wing group is reasonable as the posterior compartment represents approximately half of the wing disc, but is less than a quarter of the eye-antennal disc. Among the genes with increased expression in eye-antennal discs were white, which encodes an eye pigment precursor transporter and the eye-specific transcription factor-encoding gene, glass. Consistent with the neuronal fate of many cells in the eve-antennal disc, ten genes known to be expressed and to function in neuronal or glial cells were grouped into the eyeantennal cluster: prospero, atonal, fasciclin I, longitudinal lacking, locomotion defects, Rapgap1, unc-13, sanpodo, Caps and beta-amyloid protein precursor-like.

In addition to the genes whose tissue-specific expression was confirmed by this array analysis, many have not been

Leg1:leg2	ID	Fold	Name and Function
91, 77, 142	Leg1           GH23165         GH23165           SD07683         SD07683           CG5210         LD21619           CG1772         CG1772           CG5113         LD34931           CG14476         GH04962           CG9415         GH09250           CG4124         LD47649           CG5431         LP01553           GH26692         GH26692	2.2 1.8 1.8 2.2 1.9 1.8 1.7 1.9 1.8 1.7	Unknown Unknown Chitinase-like, Chit, cuticle chitin catabolism dacapo, dap, cyclin-dependent kinase inhibitor Unknown, alpha-glucosidase II X box binding protein 1,Xbp1, transcription factor Unknown, Zn finger, conserved domain common to transcription factors TFIIS, elonginA. CRSP70 Tyrosine-ester sulfotransferase Unknown
91, 77, 142	Leg2           CG6824         LD47350           CG6139         GH16917           CG12880         LD29228           CG18812         GH03014           CG9747         GH07782           CG1028         GM05003           CG1607         GH27380	1.5 1.8 1.7 2.2 1.7 3.4 2	ovo, transcription factor, Zn finger Monoamine transporter Unknown A1pp domain Acyl-CoA delta(11)-desaturase Antennapedia, Antp, homeobox, transcription factor Amino-acid transporter
Leg1:leg3 94, 92	Leg1 CG15096 GH28013 CG4070 GH04518	3.6 2	High-affinity inorganic phosphate:sodium symporter Tis11 homolog, Tis11, Zn finger, DNA binding
Leg2:leg3			
106. 81	Leg2 CG8084 GH07389 CG4965 LD19391 CG1028 GM05003	2.6 2.1 3.1	anachronism, ana, expressed in larval glial cells and larval CNS twine, twe, protein phosphatase Antennapedia, Antp, homeobox, transcription factor
	Leg3           CG8502         LP07813           CG6416         GH19182           CG3505         LP10895           CG2471         LP11415           CG5171         LD21828           CG9124         GM14618           CG1213         GH03773           CG8549         GH01786           CG2718         LD47536	4.2 2 1.9 3.5 2.7 2.1 2 3.2 2.4	Structural protein of larval cuticle PDZ domian Monophenol monooxygenase activator Leucine-rich repeat, RNI-like Trehalose phosphatase eIF-3p40, translation initiation factor Glucose transporter Unknown Gs1, glutamate-ammonia ligase/glutamate synthase
Fold 5 2.5 1.5 0 1.5 2.5 5 3 absent	d		

Pairwise comparisons of prothoracic, mesothoracic and metathoracic leg discs (leg1, leg2 and leg3, respectively). Cluster analysis was carried out on the dataset with the requirement to show induction > 1.74 (0.8 of the  $log_2$ -transformed R/G ratio). In the color representation of the cluster results in this and the subsequent figures, the columns represent the different experiments and the rows indicate the genes. Seventeen genes were found to be differentially expressed between the first (ten genes, green cluster) and second leg discs (seven genes, red cluster) in three independently repeated experiments (numbers 91, 77 and 142). The comparison of first and third leg discs (two experiments, numbers 92 and 94) produced two genes in the leg1 cluster (green). Two experiments (numbers 106 and 81) revealed 12 genes that are differentially expressed between leg2 disc (three genes, green cluster) and her subclusters with consistent induction in both cases (see text). The columns indicate the subclusters with consistent induction in one channel, the gene identification numbers (ID), the average fold induction of the two or three comparisons (Fold), the gene name and function as published on Flybase [44]. The color code is indicate below with the numbers representing the fold induction.

characterized previously. One of these novel genes, CG9335, was notable for its 16-fold relative induction in eye-antennal discs (Figure 5). We also single out *Gliolectin (glec)* in the

eye-antennal cluster, as this is not consistent with previous studies. *glec* is expressed in midline glia cells, but previous *in situ* hybridization and immunohistochemistry failed to

Cluster	ID	Fold	Name	Function
	CG66921 LD14839 CG2555 GH23965 CG4311 GH22436 CG10578 SD08787 CG2858 LD36843 CG3479 LD14119 CG9709 LD22081 CG12789 GH23019 CG9261 GH13134 CG3441 GH23743 CG4070 GH04518 CG6906 LD26647 CG3168 GH13883 CG7777 LD27313 CG15085 LD15796 CG6467 LP10918 CG4192 GH12215 CG2923 LD39211 CG8280 GM14559 CG3050 GH07481 CG12505 LD41905	3.3 4.3 1.8 1.9 2.4 2.4 1.9 2.3 1.9 2.6 2.4 1.8 1.9 2.6 2.4 1.8 1.9 2 2.1 2.9 2.4 2.3 2.5 2.2 2.9 2.4	CG6921 CG2555 HMG Coenzyme A synthase, Hmgs DnaJ-like-1, DnaJ-1 CG2858 outspread, osp acyl-Coenzyme A oxidase at 57D distal, Acox57D-d Best:CK01577 Nervana 2, Nrv2 CG3441 Tis11 homolog, Tis11 CG6906 CG13907 CG3168 CG7777 modulator of the activity of Ets, mae CG6467 kekkon-3, kek3 CG2923 Dystrobrevin-like, Dyb Elongation factor 1alpha48D, Ef1alpha48D Cyp6d5 CG12505	Unknown Structural protein of larval cuticle Hydroxymethylglutaryl-coenzyme A synthase Chaperone NAD(P)-binding Rossman-fold domains PH-domain-like Acyl-CoA oxidase, palmitoyl CoA oxidase Scavenger receptor Sodium/potassium-exchanging ATPase Unknown DNA binding, zinc finger Carbonate dehydratase Monocarboxylate porter General substrate transporter domains Water transporter, MIP family Mediator of MAPK signaling Serine-type endopeptidase Immunoglobulin and major histocompatibility complex domain Unknown Cytoskeletal protein binding Translation elongation factor, protein-synthesizing GTPase Cytochrome P450 Zinc finger domain
	CG4914 LP11612 CG4766 GH11415 CG8376 SD05618 CG9008 GH14910 CG13574 RE56892 CG17278 SD04019 CG11407 GH20840 CG11491 LP10481	3.8 5.1 4.5 2.5 3.6 18.3 7.3 3	CG4914 CG4766 apterous, ap CG9008 CG13574 CG17278 CG17278 CG11407 broad, br	Serine-type endopeptidase, 'homeobox' antennapedia-type protein Unknown Homeobox domain transcription factor Unknown Unknown Kazal-type serine protease inhibitor family Long-chain fatty acid transporter BTB/POZ domain, zinc finger, transcription factor
Fol 5 2.5 1.5 0 1.5 2.5 2 8 2 8 0 1.5 2 2.5 8 2 8 0 1.5 2 5 0 1.5 1.5 0 1.5 0 1.5 0 1.5 0 1.5 0 1.5 0 1.5 0 1.5 0 1.5 0 1.5 0 1.5 0 1.5 0 1.5 0 1.5 1 1 1 1	ld 5 5 5 5 5 5 5 5			

A small number of genes are preferentially induced in the prothoracic leg disc. Two comparisons of the prothoracic leg disc (L1) to the wing disc (W) (numbers 83 and 93) are shown. The cluster analysis for these comparisons revealed 23 and 8 induced genes in the L1 (red) and W (green) subclusters, respectively. Note the expression of *apterous* in the wing disc cluster (see text).

detect it in discs [34]. We did not characterize *glec* further and cannot therefore distinguish whether this discrepancy should be attributed to greater sensitivity of the array analysis or to possible contamination by glial cell RNA in the eyeantennal RNA.

To gauge the validity of these sets of genes, eight genes with uncharacterized expression in imaginal discs - four from the wing-disc cluster and four from the eye-antennal-disc cluster - were chosen for examination by *in situ* hybridization. These genes were chosen to represent a spectrum of relative intensity of expression, from just above the threshold to the maximum observed. All of the patterns we obtained correlated with the array analysis and had the preferential expression as predicted for each cluster (Figure 6). Two out of eight genes were expressed specifically in either wing (CG10962; Figure 6c) or eye-antennal (CG9335; Figure 6g) discs. CG1607 was expressed specifically in the peripodial cells of wing discs (Figure 6b). CG11798 (Figure 6f) and CG9335 (Figure 6g) were expressed specifically by cells in or near the morphogenetic furrow of eye discs. CG9335 was expressed in R8 precursor cells that also stained with the neuronal-specific 22C10 antibody in a pattern reminiscent of atonal (data not shown); it was also expressed in the optic lobe, in Bolwig's organ (the light-sensing organ of the larva), and in a subset of neurons in the embryonic and larval central nervous system (CNS; data not shown). Another notable but unexpected finding was that Arrestin 2, the metarhodopsinbinding protein, was expressed in all imaginal discs we tested, not only in eye discs. Indeed, in situ hybridization indicated that it is expressed more abundantly in the wing than in the eye-antennal disc (Figure 6d). This distribution suggests that the Drosophila Arrestin 2 might function outside the visual system.

D         D	- ~ 2 t 66 65 33 4 ~ 5 3 3	Gene	Fold	Name	Function
CCC123         1.4         /         /         Honcom           CCC127         2.3         /         /         Honcom         Honcom           CCC128         2.3         /         /         Honcom         Honcom           CCC128         2.3         /         /         Honcom         Honcom           CCC128         2.3         /         /         Honcom         Honcom           CC128         1.3         /         /         Honcom         Honcom           CC128         1.3         /         /         Honcom         Honcom           CC128         1.4         /         Honcom         Honcom         Honcom           CC128         1.4         /         Honcom         Honcom         Honcom           CC128         1.4         /         Honcom         Honc	11 11 12 12 12 12 12 12 12 12 12 12 12 1	000400		,	Lister and
CC:SS2         1.1         /         /         /         /         /           CC:SS2         1.2         /         /         /         /         /         /         /           CC:SS2         2.4         /		CG3430 CG6139	1.4	/	Unknown Monoamine transporter
CG172         2.3         /           CG173         2.4         /           CG193         1.4         /		CG4526	1.4	/	Unknown
CC1140         1.1         //         Long-chain lay usit transporter           CC6400         1.0         2.0         //         Transpire           CC6400         1.0         //         Papin, Fgn         Anne add manotoria         Transpire           CC6400         1.0         //         Papin, Fgn         Anne add manotoria         Transpire         Transpire           CC6400         1.0         //         Papin, Fgn         Anne add manotoria         Transpire           CC6400         1.0         //         Papin, Fgn         Anne add manotoria         Transpire           CC6400         1.0         //         Papin, Fgn         Anne add manotoria         Transpire           CC6400         1.0         //         Papin, Fgn         Anne add manotoria         Transpire           CC6400         1.0         //         Papin, Fgn         Anneadd manotoria         Transpire		CG1727	2.3	1	Kazal-type S protease inhibitor family
Code#2/         2         /         /         Introduction           Code#2/         10         appaird(r) (n)         Z. https://bits         Transcription factor           Code#2/         2.0         //interview         Transcription factor         Transcription factor           Code#2/         2.0         //interview         Transcription factor         Transcription factor           Code#2/         2.0         //interview         Transcription factor         Transcription factor           Code#2         2.0         //interview         Transcription factor         Transcription factor           Code#2         2.0         //interview         Code#2         1.0         Headown           Code#2         1.0         //interview         Code#2         1.0         Headown           Code#2         1.0         //interview         Factorin f.Fast         Headown         Headown           Code#2		CG1140	1.8	1	Long-chain fatty acid transporter
Comparison         Compari		CG9427	2		I NIOREDOXIN-IIKE
Collision         2.         2.         7.         Transcription latter           Collision         1.7         //         Applin, Ppn         Applin, Ppn         Applin, Ppn           Collision         1.7         //         Applin, Ppn         Applin, Ppn         Applin, Ppn           Collision         1.7         //         Applin, Ppn         Applin, Ppn         Applin, Ppn           Collision         1.7         //         Applin, Ppn         Transcription latter         Ubinom           Collision         1.7         //         Collision         Applin, Ppn         Transcription latter           Collision         1.7         Collision         Applin, Ppn         Transcription latter         Applin, Ppn           Collision         1.7         Applin, Ppn         Applin,		CG9015	2.0	/ engrailed . en	Transcription factor
C01999         1.7         /<		CG1449	2.3	Zn finger homeodomain 2 , zfh2	Transcription factor
Constant of calcie Constant of calcie Consta		CG1999	1.7	/	Unknown
Constraint         Paper		CG8502	3.3	/	Structural protein of cuticle
CC68579         CC68579 <t< th=""><th></th><th>CG1843</th><th>1.7</th><th>Papilin, Ppn /</th><th>Amino acid transporter</th></t<>		CG1843	1.7	Papilin, Ppn /	Amino acid transporter
C01151         2:5         broad, br'         Transcription factor           C01068         1;7         C)p310a T         C)p310a T         C)p310a T           C01068         1;7         C)p310a T         C)p310a T         C)p310a T           C01068         1;8         //         Historia         C)p310a T           C01068         1;8         //         Historia         C)p310a T           C01068         1;8         //         Historia         Historia           C0107         1;8         //         Historia         Historia           C0108         1;8         /         Historia         Historia           C0108         1;5         /         Historia         Historia           C0108         1;5         /         Historia         Historia           C0108         1;7         /         Historia         Historia <tr< th=""><th></th><th>CG8376</th><th>9.1</th><th>apterous, ap</th><th>Transcription factor. LIM homeodomain</th></tr<>		CG8376	9.1	apterous, ap	Transcription factor. LIM homeodomain
LP103         C)         C/p370 at /         C/p370 at / <thc <="" at="" p370="" th=""> <thc <="" at="" p370="" th=""> <thc <="" at="" p370="" th=""><th></th><th>CG1151</th><th>2.5</th><th>broad , br</th><th>Transcription factor</th></thc></thc></thc>		CG1151	2.5	broad , br	Transcription factor
Contrast         a)         Cyperiod         Cyperiod           Contrast         a)         Cyperiod         Contrast         Contrast           Contrast         1.5         / mestrix 2, An2         Unknown           Contrast         1.5         / mestrix 2, An2         / mestrix 2, An2           Cont		LP1093	1.7		Unknown
CG897 1:5 CG897 1:6 CG897 1:6 CG898 1:6		CG1039	3	Cyp310a1	Cytochrome P450 Oxidoreductase
CG6693 1.5 / mestan 2. Arc2 Methodopain inactivation CG7142 1.5 / / CG7142 1.5 / / CG7142 1.5 / / CG7142 1.5 / / CG7142 1.5 // CG7142 1.5 // CG714		CG1057	2.7	/	Unknown
CG1971 15 Arealm 2, Ar2 CG1970 15 / Arealm 2, Ar2 CG1970 15 / Arealm 2, Ar2 CG1970 15 / Arealm 2, Ar2 CG1970 16 / Arealm 2, Ar2 CG1970 17 / Fast Hermitian Construction C		CG8664	1.5	1	Unknown
Cold Hall       1.3       /       Production in formation of the description of the desc		GH2350	1.5	Arrestin 2, Arr2	Metarhodopsin inactivation
OC16770         1.6         /         Monophenol monocorgania activator           C05688         1.7         Flackbirt, Flast         Neuronal cell adhesion (gamma cells)         Neuronal cell adhesion (gamma cells)           C05688         1.4         Joe         Neuronal cell adhesion (gamma cells)         Neuronal cell adhesion (gamma cells)           C05688         1.4         Joe         Effector cappase         Neuronal cell adhesion (gamma cells)           C05688         1.7         /         Scolumphosphate schwarts         Neuronal cell adhesion (gamma cells)           C05689         1.3         /         Globechin, ginc         Scolumphosphate schwarts           C05189         2.4         sprdyor, sprd         Scolumphosphate schwarts         Scolumphosphate schwarts           C05189         2.4         sprdyor, sprd         Scolumphosphate schwarts         Scolumphosphate schwarts           C05189         2.4         Scolumphosphate schwarts         Scolumphosphate schwarts         Scolumphosphate schwarts           C05189         2.4         Scolumphosphate schwarts         Scolumphosphate schwarts         Neuronal cell adhesite schwarts           C05189         1.7         Scolumphosphate schwarts         Scolumphosphate schwarts         Neuronal cell adhesite schwarts           C05182         2.4		CG/144	1.9		Saccharopine dehydrogenase
CG5588 1.7 CG5588 1.6 CG5678 1.6 CG5678 1.6 CG1697 1.8 CG1697 1.8 CG5678 1.4 CG1697 1.8 CG5678 1.4 CG1697 1.3 CG5678 1.4 CG5678 1.4 CG568 1.4		CG1670	1.6	/	Monophenol monooxygenase activator
C6588         1.7         Fascial (I, Fast         Neuroni ola admission           000729         1.3         /         Ka         Lincown           000729         1.3         /         Ka         Lincown           000729         1.3         /         Ka         Lincown           000729         1.3         /         Sodum/phosphase         Codial field fiel			-		
Cusasa         1.0         /         Image: Cusasa         Image:		CG6588	1.7	Fasciclin1, Fas1	Neuronal cell adhesion
Norzyse       1.54       / 6s       Utter manual stress         Vietner       1.56       /       Solum Phosphate sympother         CG3036       1.7       /       Solum Phosphate sympother         CG3036       1.8       /       Solum Phosphate sympother         CG3036       1.8       /       Solum Phosphate sympother         CG3036       2.4       Solum Phosphate sympother       Utter manual sympother         CG3037       2.4       Solum Phosphate sympother       Utter manual sympother         CG3038       2.4       Solum Phosphate sympother       Utter manual sympother         CG3038       1.8       /       Solum Phosphate sympother         CG3047       1.7       Isolum Sympother       Utter manual sympother         CG3058       1.6       Solum Sympother       Utter manual sympother         CG3058       1.6       BG DSO180.10       DN replication factor family         CG3059       1.6       DC20809.11       Isolum Sympother       DN replication factor family         CG3059       1.6       DC20809.11       Isolum Sympother       DN replication factor family         CG3050       1.6       DC20809.11       Isolum Sympother       DN replication factor sympother         CG3051 <th></th> <th>CG8434</th> <th>1.6</th> <th></th> <th>ig and MHC domain</th>		CG8434	1.6		ig and MHC domain
CG1067       1.6       -         CG3035       -       -         CG3035       -       -         CG4067       2       -         CG4067       2       -         CG4067       2       -         CG4057       1.7       E         CG4058       1.8       E         CG4057       2       E         CG4057       2       E         CG4158       2       E         CG4159       2.4       F         CG4151       2.4       F <th></th> <th>CG7788</th> <td>1.5</td> <td>, Ice</td> <td>Effector caspase</td>		CG7788	1.5	, Ice	Effector caspase
CG3035       1.7       /       Sodiumphosphate sympother         Unknown       CG4067       1.8       //         CG4067       1.8       //       CG4067         CG4067       1.9       //       CG407         CG4068       2.1       //       CG407         CG4069       1.7       CG407       CG407         CG4069       1.8       CG408       CG408         CG4069       1.8       CG408       CG408         CG4069       1.8       CG408       CG408         CG4069       1.8       CG408       CG408         CG407       1.8       CG408       CG408         CG4081       2.4       //////>CG408       CG408         CG407       1.8       /////>CG407       CG408         CG4172       2.5       //////>CG407       CG4172         CG4172       2.5       ///////>CG		CG1067	1.8	1	Unknown
CG4967       2       /       Uhinoom         CG1957       1.9       /       Control       Bar Zontain, Ph domain-like         CG1958       2.4       anadop, sp0       Bar Zontain, Ph domain-like       Bar Zontain, Ph domain-like         CG1958       2.1       PFTAHE-interaction factor: 1A or 1B, PH JAN       Bar Zontain, Ph domain-like         CG1959       1.7       Bc/XA GH7485       Bar Zontain, Ph domain-like         CG1959       1.7       Bc/XA GH7485       Bar Zontain, Ph domain-like         CG1950       1.7       Bc/XA GH7485       Bar Zontain, Ph domain-like         CG1950       1.7       Bc/XA GH7485       Bar Zontain, Ph domain-like         CG1950       1.8       Bar Zontain, Ph domain-like       Bar Zontain, Ph domain-like         CG1950       1.8       Bar Zontain, Ph domain-like       Bar Zontain, Ph domain-like         CG1950       1.8       Bar Zontain, Ph domain-like       Bar Zontain, Ph domain-like         CG1950       2.4       Interview       Bar Zontain, Ph domain-like		CG3036	1.7	7	Sodium:phosphate symporter
<ul> <li>Jossev J. 10.1</li> &lt;</ul>		CG5467	2	/	Unknown
121       Circlewin gloc         Construction       Sandop, spin         Construction       Construction         Construction       PFTAHE:Interaction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE         Construction       Tist 11 construction factor 1A or 1B, PHTAHE		CG1057	1.8 1.8	/	Sec7 domain. PH domain-like
CG36908 2.4 CG36909 2.4 CG36909 2.7 CG36909 2.7 CG36909 1.7 BcDNACH745G CG3690 1.7 BcDNACH745G CG3690 1.7 BcDNACH745G CG3690 1.7 BcDNACH745G CG3690 1.7 BcDNACH745G CG3690 1.7 BcDNACH745G CG3690 1.7 BcDNACH745G DAN bright harding data lifeternitation CG3690 1.7 BcDNACH745G DAN bright harding data lifeternitation CG390 1.7 BcDNACH745G CG390 1.7 BcDNACH745G CG390 1.7 BcDNACH745G CG390 1.7 BcDNACH745G CG390 1.7 BcDNACH745G CG390 1.7 BcDNACH745G CG390 1.7 CG3977 2.7 / CG397 2.7 / CG397 2.7 / CG397 2.7 / CG399 2.4 / CG399 1.7 BcDNACH752G CG390 1.7 BcDNACH752G CG390 1.7 BcDNACH752G CG390 1.7 BcDNACH752G CG390 1.7 CG390 2.4 / CG390 1.7 CG390 2.4 / CG390 1.7 CG390 2.4 / CG390 2.4 / CG390 2.4 / CG390 1.7 CG390 2.4 / CG390 2.4 / CG390 2.4 / CG390 2.4 / CG390 2.4 / CG390 1.7 CG390 2.4 / CG390 2.4 / CG390 1.7 CG390 2.4 / CG390 1.6 / CG390 2.4 / CG390 1.6 / CG390 2.4 / CG390 2.4 / CG390 1.6 / CG390 1.7 / CG390 1.7 / CG390 1.7 / CG390 1.7 / CG390 1.7 / CG390 1.7 / CG390		CG6575	2.2	Gliolectin, glec	Carbohydrate binding protein
CG9808         2.1         PF7/IRE-interaction factor 1A or 18, P11/14/15         D2R replication factor, chromatin binding disc prolifeation factor, the original discretion factor factor, chromatin binding D3R replication factor, chromatin binding Paintity/CA oxidase           D40 F71         BD/AN CH07285         Paintity/CA oxidase         Paintity/CA oxidase           D40 F71         BD/AN CH07285         Paintity/CA oxidase         Paintity/CA oxidase           D40 F71         BD/AN CH07285         Paintity/CA oxidase         Paintity/CA oxidase           D41 F71         BD/AN CH07285         Paintity/CA oxidase         Paintity/CA oxidase           D42 F71         BD/AN CH07285         Paintity/CA oxidase         Paintity/CA oxidase           D42 F71         BD/AN CH07285         Paintity/CA oxidase         Paintity/CA oxidase           D43 F71         BD/AN CH07285         Paintity/CA oxidase         Paintity/CA oxidase           D43 F71         BD/AN CH07285         Paintity/CA oxidase         Paintity/CA oxidase           D43 F71         BD/AN CH07285		CG1539	2.4	sandopo, spdo	Actin and tropomyosin binding
V-11010         C-1100         Calc promeasure allowation and upper section acceleration and upper section acceleration		CG9808	2.1	PFTAIRE-interaction factor 1A or 1B, Pif1A/1B	bZIP transcription factor family
Text I homious_Tust I       Text I homious_Tust I         RG:BS20180.10       RG:BS20180.10         CG8342       1.9         Jocomolon defects, loco       RG:BS20180.10         CG8342       1.9         Jocomolon defects, loco       RG:BS20180.10         CG122       1.8         /       CG122         RG:BS20180.10       RG:BS20180.10         CG122       1.8         /       //         CG122       1.8         CG122       1.8         CG122       1.9         CG122       1.9         CG122       1.1         CG122       1.1         CG122       1.1         CG122       1.1         CG122       1.1         CG122       1.1         CG123       1.1     <		CG5009	2 17	BcDNA:GH07485	PalmitovI-CoA oxidase
CG8942       1.7       BG:DS0160.10       Wit-protein Spinling, "Income Division," Spinlar, "Income Division," S		CG4070	1.5	Tis11 homolog, Tis11	DNA binding, zinc finger
CG5284 1.9 bccombino defects, loco CG5282 1.8 c/DA/CH/03529 CG1292 1.8 c/DA/CH/03529 CG1292 1.3 brg/tudhais lacking, loia, LD28033 brg/tudhais lacking, loia, LD28034 brg/tudhais lacking, loia, LD28044 brg/tudhais lacking, loia		CG8942	1.7	BG:DS00180.10	Wnt-protein binding
Current         Discussion         Carrent		CG5248	1.9	locomotion defects, loco	G-protein signaling, glia cell differentiation
Fractions       Factbacking, lola, LD2903       Coli 197         CG1937       2.3       Joinghudinals lacking, lola, LD29033       Transcription factor, axon guidance         CG1937       2.3       Joinghudinals lacking, lola, LD29033       Transcription factor, axon guidance         CG1937       2.3       Joinghudinals lacking, lola, LD29033       Transcription factor, axon guidance         CG1937       2.4       /       With semicore         CG1937       2.7       /       With semicore         CG1959       2.4       /       With semicore         CG1959       2.4       /       Chith binding domain, Igan-binding domain         CG1972       2.5       /       Transcription factor, glia cell differentiation         CG1971       1.6       extra macrochaetae, enc       Statimin         CG1971       1.6       /       Coli 100       Coli 100         CG1971       1.6       /       Coli 100       Coli 100       Coli 100         CG1971       1.6       /       Coli 100       Coli 100       Coli 100       Coli 100         CG1971       1.6       /       Coli 100       Coli		CG5869 CG1202	1.5 1.8	DG:D302740.9 /	Acum binaing, aepolymerizing proteins Unknown
C01187       2.5       Iongitudinals lacking, Iola, LD28033       Transcription factor, zono guidance         Virkow       yunc-13       yunc-13       Synaptic vesicle exccytosis         C01155       2.4       /       Unknown         C01777       2.7       /       Unknown         C01772       2.7       /       Chithing         C01772       2.3       prospero, pros       Transcription factor, sill cell differentiation         C01172       2.5       /       Chithinase, cubic chinnic       Chithinase, Cubic chinnic         C01172       2.5       /       Chithinase, Cubic chinnic       Transcription factor, sill cell differentiation         C01172       2.5       /       Chithinase, Cubic chinnic catabolism       Transcription factor, sill cell chinic catabolism         C01172       2.6       Chithinase, Cubic chini catabolism       Transcription factor, sill cell chinic catabolism         C01173       1       /       Colisias       1.6       Transcription factor, sill cell chinic catabolism         C01184       2       E(pi) region transcript mbeta, HLHmbeta       Solium dependent multivitamin transporter       Transcription factor, sill cell chinic chinic cells, solic chinic conscription         C01184       2       Colisia       1.6       /       Chinhicom		CG2086	1.5	/ BcDNA:GH03529	Cell adhesion. EGF/Laminin
CG4716       1.8       //       //         CG299       CG1515       2.8       //       RN-like         CG3756       2.2       /       Water transporter, MIP family       Unknown         CG3757       2.7       /       Water transporter, MIP family       Control binding domain, igand-binding domain         CG4756       2.4       /       macrobalata, and       Artin binding       Transcription factor, gia cell differentiation         CG1720       2.3       /       macrobalata, and       Transcription factor, gia cell differentiation         CG1721       2.4       /       Transcription factor, gia cell differentiation       Transcription factor, gia cell differentiation         CG1722       2.3       /       Transcription factor, gia cell differentiation       Transcription factor, gia cell differentiation         CG1729       2.5       /       Transcription factor, gia cell differentiation       Transcription factor, gia cell differentiation         CG1729       1.6       /       Transcription factor, gia cell differentiation       Transcription factor, gia cell differentiation         CG1730       2.6       Cpip fegin transcript mbeta, HL/Himbeta       Clinkinow       Sodium dependent multivitamin transporter         CG6682       2.8       Rapapa       /       Clinkinow		CG1837	2.5	longitudinals lacking, lola, LD28033	Transcription factor, axon guidance
CC2999       1.9       Inc. 13       Symphic Vesicle exceptosis         CC31515       2.8       /       Hikine         CC31512       2.8       /       Hikine         CC31512       2.3       prospero, pros       Tarascription factor, gla cel differentiation         Hikine       Hikine       Hikine       Hikine         CC31512       2.3       Chinasse-like, Chi       Tarascription co-repressor         Minecipies       CG3143       CG1444       Progent transcript mbeta, HLHmbeta       Tarascription factor, myc-type         CC31515       1.6       /       E(pi) region transcript mbeta, HLHmbeta       Sofum dependent multivitamin transporter         CG31535       1.8       Tarascription factor, myc-type       Unknown         CG3553       1.8       Faggap       Hikinin factor         CG3553       1.6       /       Hikinin transporter         CG3553       1.6       /       Hikinin transporter         CG3553       1.6       /		CG4716	1.8	/	Unknown
Citilitie2.2///CG38312.3//UnknownCG37772.7/Water transporter, MIP family Chitin binding domain, ligand-binding domain Actin bindingTranscription factor, glia cell differentiation Histamine-gated chioride channelCG1722.5/Transcription factor, glia cell differentiationCG1722.5/Transcription factor, glia cell differentiationCG1721.6extra macrochaetae, emc staftminTranscription co-repressoreCG39311.6staftminMicrotubule binding Chitinase-like, ChitChitinase-like, ChitCG1721.5/Flop) region transcript mbeta, HLHmbetaTranscription factor, myc-type UhroomCG31311.5/Flop) region transcript mbeta, HLHmbetaUhroomCG31311.6/Flop) region transcript mbeta, HLMmbetaUhroomCG31311.7/Claikin activated protein for secretion, CapsFlop region factorCG31322.6Claikin activated protein for secretion, CapsFlop region factorCG31347/UhroomCG11346/Plass, glCG31367/Signaphilin, Spr; E62CG31367/Signaphilin, Spr; E62CG31367/Signaphilin, Spr; E62CG31367/Signaphilin, Spr; E62CG31367/Signaphilin, Spr; E62CG31367/Signaphilin, Spr; E62CG31367<		CG2999	1.9	unc-13	Synaptic vesicle exocytosis
CG7777       2.7       /       Viater magodity MP family         CG756       2.4       /       Viater magodity MP family         CG756       2.4       /       Cith binding         CG172       2.3       prospero, pros       Tanacription factor, gla cell differentiation         CG172       2.5       /       Tanacription factor, gla cell differentiation         CG172       2.5       /       Tanacription tack, grapheses       Tanacription tack, grapheses         CG1750       2.7       Chimase-like, Chit       Tianscription co-regressor       Microbuble binding         CG1751       /       CG1453       E[spi) region transcript mbeta, HLHmbeta       Tianscription tackr, my-cype         CG1838       1.6       /       Vinknown       Signaling, Noth antagonist         CG3836       1.8       Twin of n4. Torn       Signaling, Noth antagonist       Fase activator         CG49542       Caticium activated protein for secretion, Cass       Signaling or carrier, lectin       Homeodomain-like         CG1956       1.6       /       Adomain       Ligad binding or carrier, lectin         CG1844       1.6       /       Padomain       Signaling or carrier, lectin         CG1845       1.6       Let amyloid protein precursor-like, April       <		CG3831	∠.8 2,2	/	Unknown
CG1566 2.4 / CG1572 2.3 prospero, pros CG1572 2.5 / CG1007 1.6 extra macrochaetae, erac statmin CG5571 2.2 Chilinase-like, Chit CG1732 1.6 extra macrochaetae, erac SG5781 1.6 statmin CG5781 1.6 f. CG1383 1.6 / CG1383 1.6 / CG1383 1.6 / CG1383 1.6 / CG1383 1.6 / CG5781 2.3 BcDNA:CH07826 CG5860 1.6 / CG5860 1.6 / CG7872 4.1 glass, gl CG1184 6.9 / CG1184 6.9 / CG1184 6.9 / CG7727 4.6 beta amyloid protein precursor-like, Appl CG1868 1.7 / CG1868 1.7 / CG1868 1.7 / CG1868 2.7 Raggap CG5860 1.6 / CG5860 1.6 / CG5860 1.6 / CG5860 1.6 / CG5860 1.6 / CG5860 1.6 / CG5860 1.6 / CG6868 3.3 / CG6868 4.3 / CG6		CG7777	2.7	1	Water transporter, MIP family
CG1509       2.4       / prospero, pros         CG1722       2.5       /         CG1722       2.5       /         CG1722       2.5       /         CG1722       2.5       /         CG1072       2.5       /         CG1072       2.5       /         CG1071       1.6       extra macrochaetae, enc         CG1072       2.7       ////////////////////////////////////		CG8756	2.4	1	Chitin binding domain, ligand-binding domain
CG1172 2.5 prosperior pros CG172 2.5 prosperior pros extra macrochaetae, error statmin e-gated oblinitie charmal CG500 1.6 / CG510 2 2 Chilinase-like, Chil CG510 2 2 Chilinase-like, Chil CG510 2 2 Chilinase-like, Chil CG510 1.6 / CG510 2 2 Classifier (CG1454 2 CG1333 1.6 / CG515 1.8 T/win of m4, Tom CG515 1.8 T/win of m4, Tom CG550 2.2 Rapgap CG5600 1.6 / CG5600 1.6 / CG5700 2.2 Calcium activated protein for secretion, Caps atonal ato CG570 2.1 Spinophilin, Spn; E62 CG7906 1.7 / CG7906 1.7 / CG		CG1509	2.4		Actin binding
CG:1007       1.6       exta macrochaetae, enc       Transcription co-repressor         CG:200       2       Chifnase-tike, Chit       Chifnase, cuited china catabolism         CG:4150       2       Chifnase-tike, Chit       Chifnase, cuited china catabolism         CG:4151       1.6       /       Chifnase, cuited china catabolism         CG:4154       2       E(spl) region transcript mbeta, HLHmbeta       Chifnase, cuited china catabolism         CG:4154       2       E(spl) region transcript mbeta, HLHmbeta       Chifnase, cuited china catabolism         CG:4154       2       E(spl) region transcript mbeta, HLHmbeta       Chifnase         CG:4152       2.8       EGMAGHO7626       Fatyapa         CG:4150       1.8       Transcription factor, myc-type       Unknown         Signaling, Notch antagonist       Fatyapa       Fatyapa         CG:4152       Calcium activated protein for secretion, Caps       Fatyapa         CG:4154       2.7       /       Calcium activated protein precursor-like, Appl         CG:4154       2.7       /       Calcium activated protein precursor-like, Appl         CG:4154       2.7       /       Calcium activated protein precursor-like, Appl         CG:4154       2.9       /       Calcium activated protein precursor-like,		CG1472	2.5	/	Histamine-gated chloride channel
CGS891 1.6 statmin Chilinase-like, Chit CG1179 1.5 CG1174 2. CG1184 2. CG1184 2. CG1184 2. CG1185 1.6 / CG1185 1.6 / CG1185 1.6 / CG1185 1.6 / CG1185 1.6 / CG1185 1.6 / CG1185 1.6 / CG1185 1.6 / CG1185 1.6 / CG1185 1.6 / CG1195 1.7 / CG1195 2.1 / CG1195 2.3 / CG1996 1.7 / CG1195 2.3 / CG1996 1.7 / CG1196 1.7 / CG119		CG1007	1.6	extra macrochaetae, emc	Transcription co-repressor
CG6210 CG6451 CG6454 CG6454 CG6454 CG6454 CG6454 CG6454 CG6454 CG6454 CG6454 CG6454 CG6458 1.8 CG6353 1.8 CG6585 1.8 CG6585 2.3 BcDN4.5H07826 CG6682 2 Rapgap CG6680 CG6686 2.2 Rapgap CG6680 CG6686 2.2 Rapgap CG6680 CG6686 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6985 3.4 CG6986 3.4 CG6985 3.4 CG7906 1.7 CG1727 4.5 beta amyloid protein precursor-like, Appl CG7906 1.7 CG1727 4.5 beta amyloid protein precursor-like, Appl CG1727 4.5 beta amyloid protein precursor transporter Transcription factor, fork head domain RNA binding, riboruclepridase Protein prosphataset binding RNA binding, riboruclepridase Protein prosphataset binding RNA binding, riboruclepridase Protein proteophataset binding RNA binding, riboruclepridase Protein prosphataset binding RNA binding, riboruclepridase Protein proteophataset binding RNA binding, riboruclepridase Protein proteophataset binding RNA binding, riboruclepridase Protein pro		CG5981	1.6	stathmin	Microtubule binding
CG4451       1.6       /         CG4451       1.6       /         CG4451       2       E(spl) region transcript mbeta, HLHmbeta         CG4516       1.8       Twin of m4, Tom         CG4762       Rapgap         CG4956       1.6         CG4956       2         CG4956       3.3         CG4956       3.4         CG4957       1.6         CG4956       1.7         CG4957       1.9         CG4956       2.3         CG4777       1.6         Dilkaru genet, hig         CG3562       2.3         CG4767       1.5         Solum MDeptidase       Protein phosphatase1 binding, PDZ domain         CG4767       1.5         Solup Bilin,		CG5210	2	Chitinase-like, Chit	Chitinase, cuticle chitin catabolism
CG1454       2       E(spl) region transcript mbeta, HLHmbeta       Transcription factor, myc-type         CG1383       1.6       /       F(spl) region transcript mbeta, HLHmbeta       Transcription factor, myc-type         CG1383       1.6       /       /       Signaling, Notch antagonist         CG3523       2.3       BcDNA:GH07S26       Ragap       RAS GTPase activator         CG1900       1.6       /       /       Unknown         CG1901       2       Calcium activated protein for secretion, Capa       RAS GTPase activator         CG1902       2       Calcium activated protein procursor-like, Appl       RA Gmain       Unknown         CG1903       3.4       atonal, ato       RA domain       Ligand binding or carrier, lectin         CG1904       2.3       hikaru genki, hig       glass, gl       Transcription factor, photoreceptor determining         CG1906       1.7       /       white, w       Stopal particle sciptor for termining         CG1906       1.7       /       white, w       Stopal part procursor transporter         CG1906       1.7       /       White, w       Stopal part procursor transporter         CG1906       1.7       /       Stopal part procursor transporter       Transcription factor, fork head domain <th></th> <th>CG8451</th> <th>1.5</th> <th>/ /</th> <th>Zine iniger domain Sodium dependent multivitamin transporter</th>		CG8451	1.5	/ /	Zine iniger domain Sodium dependent multivitamin transporter
CG13831.6Twin of m4, Tom Tun of m4, Tom B200A:GH02526Unknown Signaling, Notch antagonist Fatty-acid synthaseCG51832.3BcDNA:GH075266Fatty-acid synthaseCG66822RaggapGAS GTPase activatorCG67002Calcium activated protein for secretion, CapsSynaptic vesicle exocytosisCG75083.4atonal, atoFatsynaptic vesicle exocytosisCG75083.4/HomeodomainCG75083.4/Homeodomain-likeCG75083.4/Homeodomain-likeCG75083.4/Homeodomain-likeCG75083.4/Homeodomain-likeCG75083.4/Homeodomain-likeCG75071.6/Homeodomain-likeCG78351.6/Homeodomain-likeCG78081.7/GG7908CG79081.7/GadesCG79081.7/GadesCG79081.7/CG16752.3Cyp 6d5CG30502.3Cyp 6d5CG16051.7/CG16051.7/CG16051.7/CG17361.8Sloppy paired 1, slp1CG11341.4/CG11341.4CG11341.4CG11341.4CG11341.4CG11341.4CG11341.4CG11341.4CG11341.4CG11351.8CG1136 <th></th> <th>CG1454</th> <th>2</th> <th>, E(spl) region transcript mbeta, HLHmbeta</th> <th>Transcription factor, myc-type</th>		CG1454	2	, E(spl) region transcript mbeta, HLHmbeta	Transcription factor, myc-type
CG5185 CG523 CG5682 CG6682 CG6780 CG16 CG798 CG798 S4 atonal, ato CG6965 CG798 S4 atonal, ato CG6965 CG798 CG772 CG1184 CG914 CG772 CG1184 CG772 CG172 CG172 CG172 CG173		CG1383	1.6	/	Unknown
CG332-3       23       ECLIVA:CHUT02C0       Fatty-3cdI Synthas9         CG6662       2       Ragap       RASG TPass activator         CG5600       1.6       /       Unknown         CG1802       2       Calcium activated protein for secretion, Caps       Synaptic vesicle exocytosis         CG7508       3.4       atonal, ato       RA domain         CG9134       2.7       /       Homeodomain-like         CG9135       16       /       Unknown         CG9136       1.6       /       Unknown         CG9134       2.7       /       Homeodomain-like         CG9134       2.7       /       Homeodomain-like         CG9134       2.7       /       Homeodomain-like         CG9135       16       /       Unknown         CG7727       4.6       beta amyloid protein precursor-like, Appl       Amyloid protein         CG7672       1.1       glass, gl       Celitathesion molecule, selectin       Transcription factor, photoreceptor determining         CG1675       2.1       Spinophilin, Spn; E62       Cyrotrome P450       Unknown         CG1676       1.7       Vhile, w       Sloppy paired 1, sip1       Retinal binding         CG1676		CG5185	1.8	Twin of m4, Tom	Signaling, Notch antagonist
CG5500 GH0977 1.9 CG5500 GH0977 1.9 CG7508 2 CG67508 3.4 CG9965 3.3 / CG9965 3.3 / CG9965 3.3 / CG9965 3.4 CG9965 3.3 / CG9965 3.3 / CG9965 1.7 CG7727 4.6 beta amyloid protein precursor-like, Appl Micro and a comparison of the second of		CG6682	2.3	DCDIVA:GHU/626 Bangan	Fally-acid synthase BAS GTPase activator
GH0977 CG1802       1.9       /       Juknown         CG1802       2       Calcium activated protein for secretion, Caps       Synaptic vesicle exocytosis         CG706       3.4       atonal, ato       Synaptic vesicle exocytosis         CG8965       3.3       /       Inscription factor         CG9184       2.7       /       RA domain         CG9184       6.9       /       Homeodomain-like         CG7727       4.6       beta amyloid protein precursor-like, Appl       Amoliding or carrier, lectin         CG708       1.7       /       Gell adhesion molecule, selectin         CG7090       1.7       /       Metalloendopeptidase         CG1675       2.1       Spinophilin, Spr; E62       Cyp 6d5         CG1675       1.5       sloppy paired 1, slp1       Transcription factor, fork head domain         CG1675       1.7       /       Vechrome 7450         Unknown       CG1673       smooth, sm       Retinal lording         CG3136       2.3       //       Vechrome 7450         Unknown       CG1673       smooth, sm       Retinal lording, ribonucleoprotein         CG1673       1.6       smooth, sm       Retinal lording, ribonucleoprotein         CG3136		CG5600	1.6	/	Unknown
CG1802       2       Calcium activated protein for secretion, Caps atonal, ato       Synaptic vesicle exocytosis         Transcription factor       RA domain         CG9134       2.7       /         CG1184       6.9       /         CG3335       16       /         CG7727       4.6       beta amyloid protein precursor-like, Appl       Homeodomain-like         CG7727       4.6       beta amyloid protein precursor-like, Appl       Calcium activated protein precursor-like, Appl         CG7672       4.1       glass, gl       Cell adhesion molecule, selectin         CG1675       2.1       Spinophilin, Spn; E62       Cytochrome P450         CG2269       1.6       //       Unknown         CG1673       1.5       sloppy paired 1, slp1       Transcription factor, fork head domain         CG1673       1.5       sloppy paired 1, slp1       Kazal-type serine protease inhibitor family         CG7870       1.9       /       Dolichyl-phosphatase1 beta-glucosyltransferase		GH0977	1.9	1	Unknown
CG1790e       3-4       atonal, ato       Instruction factor         CG9134       2.7       /       R4 domain         CG9134       2.7       /       Homeodomain-like         CG9134       6.9       /       Homeodomain-like         CG9135       16       /       Homeodomain-like         CG7727       4.6       beta amyloid protein precursor-like, Appl       Amyloid protein         CG7672       4.1       glass, gl       Cell adhesion molecule, selectin         CG7672       1.3       Spinophilin, Spn; E62       Protein phosphatase1 binding, PDZ domain         CG3050       2.3       Cyp 6d5       Unknown         CG1675       2.1       Spinophilin, Spn; E62       Protein phosphatase1 binding, PDZ domain         CG3050       1.7       white, w       G61673       Sclopp paired 1, slp1         CG1675       1.5       sloppy paired 1, slp1       Transcription factor, fork head domain         CG1965       1.7       /       Dickin/-phosphate beta-glucosyltransferase         CG1970       1.9       /       Dickin/-phosphate beta-glucosyltransferase         CG1970       1.5       BcDNA:GH02976       Structural protein of peritrophic membrane         CG3136       2.3       //       Di		CG1802	2	Calcium activated protein for secretion, Caps	Synaptic vesicle exocytosis
CG9134 C.7 / Ligand binding or carrier, lectin CG9134 6.9 / CG9135 16 / CG9727 4.6 beta amyloid protein precursor-like, Appl Nikaru genki, hig CG7727 4.6 beta amyloid protein precursor-like, Appl CG7727 4.7 / CG906 1.7 / CG3050 2.3 Cyp 6d5 CG2269 1.6 / CG2269 1.6 / CG2269 1.6 / CG1673 1.5 sloppy paired 1, slp1 CG7906 1.7 / CG7906 1.7 / CG9218 1.6 smooth, sm CG4468 1.3 / CG4114 1.4 / CG9218 1.6 smooth, sm CG4468 1.3 / CG4119 2.3 // CG4119 2.3 // CG4119 2.3 // CG4119 2.9 / CG3136 2.3 / CG3048 2.3 TNF-receptor-associated factor 1, Tra11 CG1990 3.4 Ecdyson-inducible gene L2, ImpL2 CG9991 1.6 / CG9991 1.6 / CG9992 2.2 onecut CG1904 2.2 SP555 SD025 Smoth of Statu-inbihitters		CG8965	3.4	awnai, aw /	RA domain
CG1184       6.9       /         CG9335       16       //         CG7727       4.6       beta amyloid protein precursor-like, Appl       Honeodomain-like         CG7727       4.6       beta amyloid protein precursor-like, Appl       Amount         CG7727       4.6       beta amyloid protein precursor-like, Appl       Transcription factor, photoreceptor determining         CG7906       1.7       /       Spinophilin, Spn; E62       Cg1675         CG1673       1.5       sloppy paired 1, slp1       Cychorme P450       Unknown         CG1673       1.7       /       CG1675       sloppy paired 1, slp1         CG1673       1.6       smooth, sm       Transcription factor, fork head domain         CG9269       1.7       /       Dolichyl-phosphate beta-glucosyltransferase         Unknown       CG1673       1.6       smooth, sm       RNA binding, ribonucleoprotein         CG1874       1.4       /       Dolichyl-phosphate beta-glucosyltransferase       Unknown         CG3136       2.3       /       RNA binding, ribonucleoprotein         Unknown       CG4778       1.5       ScDNA:GH02976       Structural protein of peritrophic membrane         CG3136       2.3       /       ATP binding, Thioredoxin-like </td <th></th> <th>CG9134</th> <td>2.7</td> <td>1</td> <td>Ligand binding or carrier, lectin</td>		CG9134	2.7	1	Ligand binding or carrier, lectin
CG3335       16       /       Unknown         CG7727       4.6       beta amyloid protein precursor-like, Appl       Mikaru genki, hig         CG7672       4.1       glass, gl       Cell adhesion molecule, selectin         CG7672       4.1       glass, gl       Matalloendopeptidase         CG1675       2.1       Spinophilin, Spn; E62       Protein phosphatase1 binding, PDZ domain         CG3050       2.3       Cyp 6d5       Unknown         CG1675       1.5       sloppy paired 1, slp1       Transcription factor, fork head domain         CG16675       1.7       /       Kailonding, ribonucleoprotein         CG1676       1.7       /       Kailonding, ribonucleoprotein         CG7870       1.9       /       Volknown         CG7870       1.9       /       Dolichyl-phosphataset binding, ribonucleoprotein         CG4134       1.4       /       Dolichyl-phosphataset beta-glucosyltransferase         Unknown       CG47870       1.9       /       Dolichyl-phosphataset beta-glucosyltransferase         Unknown       CG47870       1.9       /       Dolichyl-phosphataset beta-glucosyltransferase         Unknown       CG47878       1.5       BcDNA:GH02976       Structural protein of peritrophic membrane <th></th> <th>CG1184</th> <td>6.9</td> <td>/</td> <td>Homeodomain-like</td>		CG1184	6.9	/	Homeodomain-like
CG1727 4.0 Deta amycion protein precursor-like, Appl CG7672 4.1 GG1675 2.1 CG1675 2.1 CG1675 2.1 CG1675 2.1 CG1675 2.1 CG1675 2.1 CG1675 1.2 CG2269 1.7 CG1757 1.5 Spinophilin, Spn; E62 CG2269 1.7 CG1757 1.5 Soloppy paired 1, slp1 CG1673 1.5 Soloppy paired 1, slp1 CG1665 1.7 CG1767 1.5 Soloppy paired 1, slp1 CG1965 1.7 CG1965 1.7 CG1965 1.7 CG1964 1.4 CG1970 1.9 CG1970 1.9 CG1970 1.9 CG1970 1.9 CG1970 1.5 Solophilin, Spn; E62 CG1970 1.7 CG1921 1.5 Solophy paired 1, slp1 CG1965 1.7 CG1964 1.4 CG1970 1.9 CG1970 1.9 CG1970 1.9 CG1970 1.9 CG1970 1.9 CG1974 1.5 BcDNA:GH02976 CG3136 2.3 CG3048 2.3 TNF-receptor-associated factor 1, Traf1 CG1920 2.2 CG3991 1.6 CG1921 2.2 CG1921 2.2 Spinophilin, Spn; E62 CG3048 2.3 TNF-receptor-associated factor 1, Traf1 CG1920 2.2 Spi555 Solopy calcoluble gene L2, ImpL2 Spice Solop and the spice of the spice o		CG9335	16	/ hoto amulaid protein measurer "	Unknown
CG7672 4.1 glass, gl CG7908 1.7 / CG7908 1.7 / CG7908 1.7 / CG7908 1.7 / CG7908 1.7 / CG3050 2.3 Cyp 6d5 Cytochrome P450 Unknown CG2269 1.6 / CG2269 1.6 / CG2759 1.7 white, w CG1673 1.5 sloppy paired 1, slp1 CG7906 1.7 / CG7906 1.7 / CG7906 1.7 / CG7906 1.7 / CG7870 1.9 / CG7870 1.9 / CG7870 1.9 / CG7870 1.9 / CG7870 1.9 / CG1134 1.4 / CG1134 1.4 / CG1134 1.4 / CG1135 1.5 BcDNA:GH02976 Structural protein factor, hore down in mozoplase CG3136 2.3 / CG3136 2.3 / C		CG2040	4.b 2.3	beta amytoto protein precursor-like, Appl hikaru genki, hig	Cell adhesion molecule selectin
CG7908       1.7       /       /       Metalloendopeptidase         CG1675       2.1       Spinophilin, Spn; E62       Protein phosphatase1 binding, PDZ domain         CG3050       C.3       Cyp 6d5       Cytochrome P450         CG1673       1.5       sloppy paired 1, slp1       Transcription factor, fork head domain         CG1673       1.7       /       Kazal-lype serine protease inhibitor family         CG1065       1.7       /       Kazal-lype serine protease inhibitor family         CG1073       1.9       /       Dolichyl-phosphate beta-glucosyltransferase         CG1134       1.4       /       Dolichyl-phosphate beta-glucosyltransferase         CG3136       2.3       /       RNA binding, ribonucleoprotein         CG4178       1.5       BcDNA:GH02976       Structural protein of peritrophic membrane         CG3136       2.3       /       ATP binding, Thioredoxin-like         CG3195       1.8       SP1029       ATimopeptidase, NOT aminoacyclase         CG39691       1.6       /       Uhknown         CG39691       1.6       /       Uhknown         CG39691       1.6       /       CG39691         CG1692       2.2       onecut       Uhknown		CG7672	4.1	glass, gl	Transcription factor, photoreceptor determining
CG1675       2.1       Spinophilin, Spr; E62       Protein phosphatase1 binding, PDZ domain         CG3050       2.3       Cyp 6d5       Uhknown         CG2759       1.7       white, w       Eye pigment precursor transporter         CG1675       1.7       slopy paired 1, slp1       Transcription factor, fork head domain         CG1665       1.7       /       Dolichyl-phosphatase1 binding, PDZ domain         CG1065       1.7       /       Transcription factor, fork head domain         CG1067       1.9       /       Dolichyl-phosphate beta-glucosyltransferase         Unknown       CG7870       1.9       /       Dolichyl-phosphate beta-glucosyltransferase         CG1134       1.4       /       RNA binding, ribonucleoprotein       Unknown         CG4288       1.6       smooth, sm       RNA binding, ribonucleoprotein       Unknown         CG3136       2.3       /       Structural protein of peritrophic membrane       bZIP transcription factor family         CG3136       2.3       /       Structural protein of peritrophic membrane       bZIP transcription factor, homeobox domain         CG3136       2.3       /       ZIP transcription factor, homeobox domain       Cell         CG3136       2.3       /       ZIP transcription factor		CG7908	1.7	7	Metalloendopeptidase
CG309U 2.3 Cyp bab Cyp bab Cg2269 1.6 / CG2759 1.7 white, w CG1673 1.5 sloppy paired 1, slp1 Eve pigment precursor transporter CG1065 1.7 / CG7900 1.9 / CG7900 1.9 / CG1741 1.4 / CG12759 1.5 sloppy paired 1, slp1 Retinal binding CG7906 1.7 / CG12759 1.5 Smooth, sm CG1275 1.5 Smooth, sm CG4468 1.3 / CG4761 2.9 / CG4511 2.9 / CG3046 2.3 TNF-receptor-associated factor 1, Traf1 CG4501 3.4 Ecdyson-inducible gene L2, ImpL2 CG1922 2.2 onecut CG1924 2.2 SP555 Smooth 1.7 CHEMICAL State 1.5 Smooth, sm CG4775 1.5 Smooth, sm CG4776 1.5 Smooth, sm CG4776 CG3046 2.3 / CG4511 2.9 / CG3048 2.3 TNF-receptor-associated factor 1, Traf1 CG450 3.4 Ecdyson-inducible gene L2, ImpL2 CG3046 2.3 Chemical State 1.5 Smooth, sm CG4776 CG3046 2.3 Chemical State 1.5 Smooth, sm CG1927 CG3047 Chemical State 1.5 Smooth, sm CG1927 CG3048 2.3 Chemical State 1.5 Smooth, sm CG4776 CG3048 2.3 Chemical State 1.5 Smooth, sm CG47777 CHemical State 1.5 Smooth, sm CG47777		CG1675	2.1	Spinophilin, Spn; E62	Protein phosphatase1 binding, PDZ domain
CG2759 1.7 // White, w CG2759 1.7 // White, w CG1673 1.5 sloppy paired 1, slp1 CG1906 1.7 / CG7906 1.7 / CG7906 1.7 / CG7906 1.7 / CG7870 1.9 / CG7870 1.9 / CG7870 1.9 / CG7871 1.4 / CG9218 1.6 smooth, sm CG4468 1.3 / CG4468 1.3 / CG459 1.8 / CG4		CG3050	2.3	Сур 6d5 /	Cytochrome P450
CG1673       1.5       sloppy paired 1, slp1       Transcription factor, fork head domain         CG1065       1.7       /       Retinal binding         CG7060       1.7       /       Kazal-type serine protease inhibitor family         CG7870       1.9       /       Unknown         CG91134       1.4       /       Unknown         CG9218       1.6       smooth, sm       RNA binding, ribonucleoprotein         Unknown       CG4468       1.3       /       Unknown         CG4136       2.3       /       Structural protein of peritrophic membrane         CG3136       2.3       /       ATP binding, Thioredoxin-like         CG3048       2.3       TNF-receptor-associated factor 1, Tra11       Defense response, signal transduction         CG3195       1.6       oneut       Unknown       Celeson         CG3048       2.3       TNF-receptor-associated factor 1, Tra11       Defense response, signal transduction         CG3691       1.6       Unknown       Celeson       Celeson         CG3048       2.3       TNF-receptor-associated factor 1, Tra11       Defense response, signal transduction         CG3691       1.6       Unknown       Celeson       Celeson         CG3691		CG2759	1.7	, white, w	Eye pigment precursor transporter
CG1065       1.7       /       Retinal binding         CG7906       1.7       /       Kazal-type serine protease inhibitor family         CG7870       1.9       /       Dolicht/i-phosphate beta-glucosyltransferase         Unknown       CG134       1.4       /       Dolicht/i-phosphate beta-glucosyltransferase         Unknown       CG4787       1.5       smooth, sm       RNA binding, ribonucleoprotein         Unknown       CG4134       2.3       /       Structural protein of peritrophic membrane         CG4186       1.3       /       Structural protein of peritrophic membrane       Structural protein of peritrophic membrane         CG4178       1.5       BcDNA:GH02976       Structural protein of peritrophic membrane       Structural protein of peritrophic membrane         CG3136       2.3       /       Defense response, signal transduction       CG3048         CG3195       1.6       SP1029       Aminopeptidase, NOT aminoacyclase       Defense response, signal transduction         CG1950       3.4       Ecdyson-inducible gene L2, ImpL2       Unknown       Celf1922       Onecut         CG1952       2.2       opecut       SOCS domain, Cettrophic actor, homeobox domain       CG3641         CG1922       2.2       SP555       SOCS domain, Cettrophic		CG1673	1.5	sloppy paired 1, slp1	Transcription factor, fork head domain
CG7906       1.7       /       Kazal-type serine protease inhibitor family         CG7870       1.9       /       Dolichyl-phosphate beta-glucosyltransferase         CG1134       1.4       /       Unknown         CG4786       1.6       smooth, sm       RNA binding, ribonucleoprotein         CG4778       1.5       BcDNA:GH02976       Structural protein of peritrophic membrane         CG4778       2.3       /       ATP binding, Tiboredoxin-like         CG1195       1.8       SP1029       ATP binding, Tiboredoxin-like         CG1950       3.4       Ecdyson-inducible gene L2, ImpL2       Cel adhesion         CG6991       1.6       /       Unknown         CG1922       2.2       onecut       CG1922       Cancut         CG4964       1.6       /       Unknown       Cefseen exponse, signal transduction         CG1921       2.2       onecut       CG1922       Cel adhesion       Unknown		CG1065	1.7	1	Retinal binding
CG1134       1.3       /       Unknown         CG1134       1.4       /       Unknown         CG9218       1.6       smooth, sm       RNA binding, ribonucleoprotein         CG4778       1.5       BcDNA:GH02976       Structural protein of peritrophic membrane         CG4778       2.3       /       ATP binding, Thioredoxin-like         CG3136       2.3       /       ATP binding, Thioredoxin-like         CG3145       1.8       SP1029       Aminopeptidaes, NOT aminoacyclase         CG3048       2.3       TNF-receptor-associated factor 1, Traf1       Defense response, signal transduction         CG1500       3.4       Ecdyson-inducible gene L2, ImpL2       Cell adhesion       Unknown         CG1922       2.2       onecut       Transcription factor, homeobox domain         CG1924       2.2       SP555       SOCS domain, C-terminus of STAT-inbinititres		CG7906	1.7	/	nazal-type serine protease inhibitor family
CG9218       1.6       smooth, sm       RNA binding, ribonucleoprotein         CG4268       1.3       /       Unknown         CG4468       1.3       /       Structural protein of peritrophic membrane         CG3136       2.3       /       Structural protein of peritrophic membrane         CG4511       2.9       /       ATP binding, Tiloredoin-like         CG3136       2.3       /       ATP binding, Tiloredoin-like         CG4511       2.9       /       ATP binding, Tiloredoin-like         CG3048       2.3       TNF-receptor-associated factor 1, Traf1       Defense response, signal transduction         CG3050       3.4       Ecdyson-inducible gene L2, ImpL2       Uhknown         CG3691       1.6       Uhknown       Cel adhesion         CG3192       2.2       onecut       Transcription factor, homeobox domain         CG3192       2.2       SP555       SOCS domain, C-terminus of STAT-inhithtres		CG1134	1.9	, ,	Unknown
CG4468     1.3     /     Unknown       CG4778     1.5     BcDNA:GH02976     Unknown       CG3136     2.3     /     bZIP transcription factor family       CG4511     2.9     /     bZIP transcription factor family       CG4511     2.9     /     ATP binding, Thioredoxin-like       CG3048     2.3     TNF-receptor-associated factor 1, Traf1     Defense response, signal transduction       CG39691     1.6     /     Unknown       CG1922     2.2     onecut     Transcription factor, homeobox domain       CG1904     2.2     SP555     SOCS domain, Clefringius of STAT_inbibititres		CG9218	1.6	smooth, sm	RNA binding, ribonucleoprotein
CG4778     1.5     BcDNA:GH02976     Structural protein of peritrophic membrane bZIP transcription factor family       CG4511     2.9     /     ATP binding, Thioredoxin-like       CG4511     2.9     /     ATP binding, Thioredoxin-like       CG4514     2.9     /     ATP binding, Thioredoxin-like       CG4515     1.8     SP1029     ATP binding, Thioredoxin-like       CG3048     2.3     TNF-receptor-associated factor 1, Traf1     Defense response, signal transduction       CG50691     1.6     /     Unknown     Unknown       CG1922     2.2     onecut     Transcription factor, homeobox domain       CG1044     2.2     SP555     SOCS domain, C-terminus of STAT-inhibitors		CG4468	1.3	/	Unknown
CG3136       2.3       /       bZIP transcription factor family         CG4511       2.9       /       ATP binding, Thioredoxin-like         CG195       1.8       SP1029       Aminopeptidase, NOT aminoacyclase         CG3048       2.3       TNF-receptor-associated factor 1, Traf1       Defense response, signal transduction         CG1950       3.4       Ecdyson-inducible gene L2, ImpL2       Cell adhesion         CG9691       1.6       /       Unknown         CG1922       2.2       onecut       Transcription factor, homeobox domain         CG1904       2.2       SP555       SOCS domain, C-terminus of STAT-inhibitors		CG4778	1.5	BcDNA:GH02976	Structural protein of peritrophic membrane
CG1195 C.3 / SP1029 Animopptidase, NOT animoacyclase CG3048 2.3 TNF-receptor-associated factor 1, Traf1 CG1500 3.4 Ecdyson-inducible gene L2, ImpL2 Defense response, signal transduction CG3691 1.6 / Unknown CG1922 2.2 onecut Transcription factor, homeobox domain CG1904 2.2 SP555 SOCS domain. C-terminus of STAT-inhibitors		CG4511	2.3		ATP binding Thioredoxin-like
CG3048       2.3       TNF-receptor-associated factor 1, Traf1       Defense response, signal transduction         CG1500       3.4       Ecdyson-inducible gene L2, ImpL2       Cell adhesion         CG9691       1.6       /       Unknown         CG1902       2.2       onecut       Transcription factor, homeobox domain         CG1044       2.2       SP555       SOCS domain. C-terminus of STAT-inhibitors		CG1195	1.8	, SP1029	Aminopeptidase, NOT aminoacvclase
CG1500         3.4         Ecdyson-inducible gene L2, ImpL2         Cell adhesion           CG9691         1.6         /         Unknown           CG1922         2.2         onecut         Transcription factor, homeobox domain           CG1404         2.2         SP555         SOCS domain. C-terminus of STAT-inhibitors		CG3048	2.3	TNF-receptor-associated factor 1, Traf1	Defense response, signal transduction
CG192 2.2 SP55 Unc		CG1500	3.4	Ecdyson-inducible gene L2, ImpL2	Cell adhesion
CG1404 2.2 SP55 SC domain. Of STAT-inhibitors		CG9691	1.6	/ opecut	Unknown
		CG1404	2.2	SP555	SOCS domain, C-terminus of STAT-inhibitors
CG3556 1.7 / Terpenoid cyclase/protein prenyltransferase		CG3556	1.7	/	Terpenoid cyclase/protein prenyltransferase

Figure 5 (see the legend on the next page)

Haltere and genital imaginal discs

In five direct comparisons of haltere and genital imaginal discs, 130 genes were found to be preferentially expressed in the haltere disc and 54 in the genital disc (see Additional data files). As with the wing/eye-antennal analysis, known genes clustered as predicted from previous studies. For example, *caudal*, a homeobox domain transcription factor, partitioned into the genital-disc cluster. *caudal* is expressed in the genital disc but not in more anterior tissues such as the haltere disc [35]. *abdominal B*, which would be predicted to show preferential expression in the genital disc, was not represented on the arrays. These 184 differentially expressed genes correspond to 3% of the entire set of 6,000 and represent the largest cluster we obtained in all pairwise comparisons of imaginal discs.

### Expression profiles of larval organs

In order to compare and contrast the expression profiles of larval organs and imaginal discs, we made probes from the poly(A)<sup>+</sup> RNA isolated from several larval organs. We directly compared probes from the wing imaginal disc with: the salivary gland, an anterior fraction of the midgut, parts of the fat body and the brain hemisphere, including the optic lobe primordium. For each experiment, probes prepared from a single larva were compared. With the exception of the comparisons between the first leg disc and fat body which was carried out once, each of the other comparisons was repeated at least three times.

Visual inspection of the array scans of imaginal disc to larval tissue comparisons revealed a large number of red and green spots, indicative of a high degree of divergence. A representative block from one wing disc-to-fat body scan is shown in Figure 7b. These results contrast with disc to disc comparisons which produced mostly yellow and only few red or green spots on the arrays. For purposes of illustration, the same block of spots from a leg1 disc-to-leg3 disc scan is shown in Figure 7a. When the intensity of Cy3 and Cy5 signals for all cDNAs was plotted on a scatterplot, only a small number of data points are positioned away from the bisector on which signals of equal intensity in the two channels fall (Figure 7c). This is indicative of strong similarity in

expression profiles and is reflected in a high CC (0.93). In contrast, the wing disc-to-fat body comparison had many data points located on each side of the bisector (CC 0.37) (Figure 7d). This disparity between leg disc-to-leg disc and wing disc-to-fat body comparison was also observed for all other disc-to-disc and disc-to-larval organ comparisons we carried out. Figure 8a illustrates the numbers of cDNAs that were expressed differentially in the various pairwise comparisons as a bar diagram. All disc-to-disc comparisons showed small numbers of differentially expressed cDNAs, whereas the disc-to-nondisc comparisons revealed many. The only exceptions were the comparisons of the wing imaginal disc to the brain/optic lobe (see below). To score for the differences in expression for all 6,000 genes, the standard deviations of the R/G ratios (log<sub>2</sub>-transformed) for each gene were calculated for all 42 experiments and represented in a scatterplot (Figure 8b). Standard deviations from the datasets derived from disc-to-disc comparisons (25 experiments) were all < 1, indicating the high level of similarity among the different imaginal discs. In contrast, the disc-tonondisc comparisons (17 experiments) were, with a single exception (see below) > 1, reflecting a high degree of divergence (Figure 8b). These observations are consistent with the distinct developmental programs of imaginal and larval cells.

The only exceptions to emerge in this analysis were the cells of the optic lobe/brain hemisphere. Comparisons of these neuronal tissues with wing discs revealed only a small number of differentially expressed genes (Figure 8a) and had low standard deviations that were in the range of discto-disc comparisons (Figure 8b). The optic lobe primordium and brain of late third instar larvae have only limited functions in larval photoreception and contain only a small number of mature neurons. The larval brain is populated with clusters of immature neurons, neuroblasts and ganglion mother cells whose descendants will generate the neuronal and glial populations of the adult optic neuropil and the brain hemisphere. The brain hemisphere also differs from other larval tissues by not degenerating during metamorphosis. Instead, most neurons of the larva join with newly formed adult-specific neurons to build the CNS of the adult

Figure 5 (see the figure on the previous page)

Genes preferentially expressed in wing and eye-antennal discs. Amplified RNA from single discs of five individual larvae was used to carry out direct comparisons between a wing imaginal disc and an eye-antenna imaginal disc of the same larva (experiments 51, 88, 146, 155, 156, 157, 173 and 175). Whenever samples from one larva were used for more than one experiment, different left to right combinations of the individual discs were made. In two experiments, combinations of discs from two larvae (experiment 74) or pools of five larvae (ten wing and ten eye-antenna discs, experiment 98) were used. Cluster analysis was performed on the dataset with the requirement to show induction > 1.74 (0.8 of the log<sub>2</sub>-transformed R/G ratio) in at least five experiments. One hundred and forty genes grouped into different subclusters. After removal of double hits or genes with inconsistent induction, 97 genes remained. Twenty-four genes were induced in the wing imaginal disc (red) and 73 genes in the eye-antennal disc (green). Black indicates lack of induction. Fold induction was calculated as an average induction in all experiments (column 3). The name, description and molecular or biological function is indicated as in Flybase [44] (columns 4 and 5). Some genes with known expression in the respective tissue are included in the clusters, such as *apterous*, *engrailed* and *glass* (see text). Of this set, 22 genes are uncharacterized and do not code for known protein domains. Note the high induction of CG9335 (16 fold) and CG11849 (6.9 fold) in the eye-antennal cluster. Genes marked in red were chosen for *in situ* experiments (see Figure 6).



Expression patterns of genes in the wing and eye-antennal clusters. *In situ* hybridizations of genes from (a-d) the wing and (f-i) eye-antennal disc clusters show the disc-specific patterns of expression. Confirming the mircoarray data, the signal intensities are higher in the wing discs in (a-d) (red frame) and the eye-antennal discs in (f-i) (green frame). Note the refined expression pattern in the wing disc for CG10962 in (c). In two cases, CG10962 (c) and CG9335 (g), signal could only be detected in the predicted disc. The arrowhead in (g) indicates the morphogenetic furrow. The number indicates the average fold induction in the 11 experiments. Arrestin2 (d) and CG11798 (f) were included in the clusters because their relative induction was > 1.74 in more than five experiments. (e) CG6680 and (j) LD11162 failed the threshold criteria for the cluster depicted in Figure 4, but were part of a larger cluster with lower threshold settings. For both genes, the *in situ* patterns confirm the predicted expression. Discs are oriented anterior to the left, and dorsal uppermost.



Imaginal discs share a similar expression profile but differ from differentiated larval tissues. (a) Comparison between first and third leg discs. An enlargement of a representative block out of 32 blocks on the microarray is shown. It produced mostly yellow spots on the superimposed red (Cy5 labeling) and green (Cy3 labeling) images, indicative of a high degree of similarity in the respective expression profiles. (b) In a comparison of wing disc and fat body, the same block contained mostly red and green spots, indicating a high degree of divergence. (c,d) Scatterplots of Cy3 and Cy5 intensities in comparisons (c) between leg discs and (d) between wing disc and larval fat body. In the leg comparison the spots are in close proximity to the bisector (CC 0.93, SD 0.38) with only a small number of genes induced in either the leg1 or leg3 disc. In contrast, spots are spread widely to both sides of the bisector for the wing-to-fat body comparison (CC 0.37, SD 1.17), indicating a large number of differentially expressed genes. The data points are color coded such that spots that are induced > 1.74 fold in the Cy5 channel are colored red and those induced > 1.74 in the Cy3 channel in green. Ratios within a threshold of 1.74 are represented in black.

([36] and references therein). We interpret the high degree of similarity in the expression profiles of imaginal discs and the brain/optic lobe preparations as a manifestation of the apparent commonalities of their developmental programs.

The number of genes that were found to be induced in the larval organs is significantly greater than the number of genes in the sets defined by the disc-to-disc comparisons. Most of the genes in these sets have not been characterized previously and therefore have no known function, but the known genes in these clusters illustrate the predictive power of the method; we mention three examples.

Many of the genes that are preferentially expressed in the optic lobe/brain hemisphere have predicted or described expression and/or function in neurons or glial cells (data not shown). One of these is *lola*, which encodes a transcription factor involved in axon guidance and is expressed in the embryonic optic lobe placodes as well as other tissues [37]. Another is *beta amyloid protein precursor-like (Appl)*, with known expression in the larval optic lobe [38].

The expression profile of the cyclin-dependent kinase 1 (*Cdk-1/cdc2*), is also noteworthy. Cdk-1 is believed to regulate the G2 checkpoint of the cell cycle and is required to maintain diploidy. Mutations in *Cdk-1* drive cells that would normally remain diploid into endoreplication [39]. We found *Cdk-1* (CG5363) to be repressed in larval tissues when compared to imaginal discs (Figure 9). As most of the cells in the larval fat body, salivary gland and midgut are polyploid (reviewed in [40]), the array hybridizations are in good agreement with the known function of *Cdk-1*.



The mRNA populations in larval and imaginal tissues are distinct. (a) Graphical representation of the number of spots with intensity differences > 1.74 when imaginal and larval tissues were compared. The numbers of spots are derived from cluster analysis of the repeated experiments. All disc-to-disc comparisons are represented by red bars, whereas the disc-to-nondisc comparisons are shown in blue. EA, eye-antennal; FB, fat body; G, genital; GUT, anterior part of the midgut; H, haltere; L1, leg 1; L2, leg 2; L3, leg 3, OL, optic lobe/brain; SG, salivary gland; W, wing. (b) A plot of the level of divergence as measured by the SD of the log<sub>2</sub>-transformed R/G ratios for all genes in 42 experiments (leaving out the wing-to-wing experiment). Of these, 25 experiments were comparisons of one imaginal disc to another (see text and Table 1) and the remaining 17 compared imaginal discs to larval tissue, that is, salivary gland, midgut, fat body and optic lobe/brain hemisphere. The disc-to-disc comparisons were placed on the lower line, whereas disc-to-nondisc comparisons group to the right, with SD > 1. A group of three experiments of the experiments of disc-to-nondisc comparisons. These are the wing disc-to-optic lobe/brain hemisphere comparisons. These are the wing disc-to-optic lobe/brain hemisphere comparisons (arrow).



Cyclin-dependent kinase I (Cdk I/cdc2) is expressed in imaginal but not in larval tissues. The cluster analysis for Cdk I is shown for 41 pairwise comparisons with each box representing one experiment. All the disc-to-disc comparisons (see text) group to the left and display ratios of the intensities close to I, as indicated by the dark shades of red and green (color coding as in Figure 3). The *in situ* hybridization to the various imaginal discs confirmed that Cdk I is expressed in all discs. Cluster analysis of the comparisons of discs (labeled in green) and larval tissues (red) showed a strong induction in imaginal discs (represented by the intense green staining). This finding is confirmed by previous descriptions that Cdk I is downregulated in endoreplicated tissues and by the lack of signal after *in situ* hybridization (FB, GUT and SG). The wing disc (W) to brain/optic lobe (OL) comparisons revealed less induction of Cdk I in the wing disc as indicated by the darker shades of green. This is in good agreement with *in situ* hybridization data that showed some expression, particularly in the proliferation zones of the optic lobe, but no signal in most parts of the brain. Abbreviations as in Figure 8.

### Experimental design and data analysis

Two methods for comparative hybridizations to DNA microarrays have been used in previous studies. Most commonly, experimental samples have been compared to a common reference sample that contains an appropriate diversity of cell types and states. An alternative method, which we have used in this study, is to directly compare probes generated from two experimental samples. This second method should be the more sensitive. The amplification technique we used was such that product generated from a single imaginal disc was sufficient to make several probes and to carry out multiple array hybridizations. By comparing samples that had been isolated from a single animal, this method minimized differences due to biological variability. By repeating experiments several times and by reversing the label (dye-flip) for experimental samples in repeat hybridizations, we also attempted to minimize experimental noise.

In addition to these design aspects of the experimental protocols, we applied a number of filters to the data analysis. We selected only those genes with hybridization signals over background in 80% of all experiments and with a 1.74-fold difference between the two channels in at least 45%. Applying this method to 11 arrays that compared wing and eyeantennal discs, we found two subclusters representing genes preferentially expressed in either disc. These lists of genes are largely validated by the previously described expression pattern in either tissue and by *in situ* hybridization analysis. We understand that the use of any such threshold settings is likely to filter out biologically relevant genes. For example, when less stringent settings were used (for example, a 1.6fold intensity difference increased the gene sets by 20%), additional genes such as twin of eyeless were included in the eye-antennal cluster and Antp and CG6680 (Figure 6e) in the wing cluster. Relaxing the requirement to produce a ratio in at least 80% of all experiments to 70% identifies yet another group of genes, and includes LD11162 (Figure 6j). Conversely, setting a more stringent threshold to 1.9-fold intensity difference decreased the gene sets by 25% and removed genes such as Arrestin2 and CG11798 (Figure 6d,f).

Setting a threshold level for the clusters is arbitrary, and as some candidate genes may be expressed by only a small subset of cells in the experimental sample, each analysis must be evaluated separately. Nevertheless, we interpret the good correlation between the array analysis and expression patterns we have described as indicating that the method is both efficient and reliable. Subsequent to the analyses described above, we carried out single-disc comparisons with microarrays that represent a nearly complete set of the approximately 14,000 annotated Drosophila genes. The short cDNAs spotted on these arrays were produced with specific primer pairs obtained from Incyte Genomics, Inc. Use of these arrays for disc-to-disc comparisons also identified only a relatively small number of differentially expressed genes (data not shown). The list of induced genes was apparently more complete, however, and included, for example, vestigal in the wing-to-leg disc comparisons (A.K., G. Schubiger and T.B.K., unpublished observations).

Although the absolute number of genes included in the lists of tissue-specific genes is not meaningful, the relatively small number of genes that differ between imaginal tissues and the significantly greater number of genes that differ between imaginal and larval tissues certainly is.

# Conclusions

We have shown that linear RNA amplification is reproducible and can be used to generate probes for microarray experiments with limited quantities of starting material. We used this technique to determine the expression profiles of imaginal and larval tissues by pairwise comparisons of tissues dissected from single larvae. Our approach was validated by the presence of characterized genes in the appropriate cluster and by confirmation of expression patterns by direct analysis. Applying stringent threshold criteria, we found that less than 3% of the genes represented on the arrays were preferentially expressed when discs were compared to each other. Many of these genes have either not been characterized previously or lack obvious similarity to known coding domains. Those that have been previously characterized or share sequence similarities represent a broad array of cellular functions. These include basic metabolic and transporting functions, structural, cell-adhesive and signaling functions, as well as transcriptional regulation. We lack sufficient understanding of the specific processes in most cell types to interpret the significance of the genes that they specifically express. Future studies will be required to identify the functional networks that the identified candidate genes describe and to establish how they contribute to the execution of specific developmental programs.

We anticipate that the methods we describe here can be applied to determine and compare the expression profiles of small cell groups in *Drosophila* tissues. We look forward to a further enhancement in resolution that can be achieved by combining cell-sorting techniques with this type of array analysis. We found that starting with as little as 10 pg poly(A)<sup>+</sup> RNA yields enough aRNA for one hybridization. This amount roughly corresponds to 300 cells of the third instar wing imaginal disc.

# Materials and methods Microarray production and labeling

Spotted cDNA microarrays were produced essentially as described at [41]. In brief, the Drosophila Gene Collection (DGC 1.0, kindly provided by the Berkeley Drosophila Genome Project) of 5,849 nonredundant cDNAs was amplified using universal primers essentially as described at [42]. The amplification products were purified with 96-well format Qiagen PCR purification columns. All PCR products were analyzed on agarose gels, and reactions with no detectable product, multiple bands or bands of unexpected size were repeated using Herculase (Amersham) or Expand polymerase (Roche). One hundred and seventy-five cDNAs from our lab collection were amplified and added to the set, resulting in a total of 6,024 cDNAs. Reactions were arrayed into 384-well plates and printed on poly-L-lysine-coated glass slides using a linear servo arrayer and ArrayMaker Version 2 control software. To minimize background caused by oxidation of the polylysine coating, slides were pretreated by a protocol suggested by Paul Ebert. Before post-processing, slides were incubated for 5 min in 3x SSC; 0.2% SDS at 65°C, washed first in water and then in 95% ethanol, followed by centrifugation to dry the slides. Slides were then treated, and the hybridization reaction was carried out as described at [42]. We indirectly labeled the hybridization probes by incorporation of amino-allyl modified nucleotides in a first-strand cDNA RT reaction. Monofunctional Cy5 or Cy3 dye (Amersham) was subsequently coupled to the reactive residues. Multiple hybridizations were carried out for most experiments, and dye labeling was reversed to avoid systematic bias.

# Larval dissections, RNA isolation and *in situ* hybridization

Third instar wandering larvae were dissected and washed several times in Ringer solution (130 mM NaCl, 5 mM KCl, 2 mM  $Na_2HPO_4$ , 1.5 mM  $CaCl_2$ , 0.37 mM  $KH_2PO_4$ ). Fatbody fragments were mainly anterior plates. Midgut was the anterior portion just posterior to the gastric caeca. Brain/optic lobe consisted of one brain hemisphere including the optic lobe primordium.

Total RNA from larval tissues was extracted using the Mini RNA Isolation Kit (Zymo Research, Orange, CA) and was eluted with RNase-free water. Total RNA from embryos was extracted with Trizol reagent (Invitrogen). Embryonic poly(A)<sup>+</sup> RNA was purified using the Oligotex mRNA Mini Kit (Qiagen). Poly(A)<sup>+</sup> RNA was not purified before amplification of disc RNA because of the small amounts (see Figure 1).

*In situ* hybridizations were carried out as described in O'Neill and Bier [43]. The plasmids used to generate *in situ* probes were derived by subcloning an aliquot of the PCR product that was printed onto the microarray with Topo TA cloning (Invitrogen). All cloned products were sequenced and separate hybridizations were carried out for sense and antisense probes.

#### **RNA** amplification

Methods for amplification were adapted from Wang et al. [25] and Baugh et al. [23]. Total RNA isolated from dissected larval tissues or poly(A)+ RNA purified from embryos was incubated with 100 ng oligo(dT)24 -T7 primer (GGCCAGTGAATTGTAATACGACTCACTATAGGGAGGCG-(AAGCAGTGGTAACAACGCAGAGTACGCGGG) in 5 µl at 65°C for 10 min and cooled on ice. Six microliters first-strand premix were added and incubated at 42°C for 2 h (6 µl mix was made up with 2 µl 5x first-strand buffer, 1 µl 0.1 M DTT, dNTPs (10 mM each), 0.3 µl T4gp32 (USB,  $1 \, \mu l$ 13.82 mg/ml), 0.5 µl RNasin (Promega), 1 µl Superscript II (Invitrogen)). The reaction was incubated at 65°C for 10 min and cooled on ice. Cold second-strand premix (64.5 µl prepared on ice: 45 µl RNase-free water, 15 µl 5x second-strand buffer (Invitrogen), 1.5 µl dNTPs (10 mM each), 0.5 µl E. coli ligase (10 U/ $\mu$ l), 2  $\mu$ l *E. coli* polymerase (10 U/ $\mu$ l), 0.5  $\mu$ l E. coli RNaseH (2 U/µl)). Enzymes for the second-strand

premix (Invitrogen) were added and incubated at 16°C for 2 h, followed by the addition of 2 units T4 DNA polymerase (Promega) and incubation at 16°C for 15 min before heat inactivation at 70°C for 10 min. Clean-up was performed with DNA clean and concentrator-5 (Zymo Research), eluting twice with 8 µl RNase-free water. The total volume was adjusted to 8 µl in a Speed Vac. The first IVT was performed with Megascript T7 (Ambion) in a 20 µl volume for 5-6 h, followed by DNA digestion. aRNA was purified using Mini RNA Isolation Kit or RNA Clean-up kit (Zymo Research), eluting in 2x 8 µl RNase-free water. Random hexanucleotides (250 ng) and TS primer  $(1 \mu g)$  were added and the volume adjusted to 5 µl. The mix was incubated at 65°C for 10 min before 5 µl first-strand premix was added followed by incubation at 42°C for 2 h. Before the second-strand synthesis, 200-500 ng oligo(dT)24 -T7 primer was added and denatured at 65°C for 10 min. The second-round second-strand synthesis, clean-up and IVT were carried out as for the first round, except that no ligase was added to the second-strand premix. One-quarter to one-third of the total aRNA was applied to the clean-up columns after the second IVT to avoid overloading. All reactions were carried out in a thermocycler with heated lid or air incubator to avoid evaporation.

#### Data analysis

Hybridized microarrays were scanned with a GenePix 4000A Microarray Scanner (Axon Instruments, Union City, CA). Data were analyzed and displayed with Cluster and Treeview [31], AMAD [41], Genepix PRO (Axon Instruments), and Microsoft Excel. Normalization for cluster analysis and calculation of standard deviations were done with AMAD. Only genes that qualified with a combined median intensity > 300 above background in both channels in at least 80% of the repeated experiments were included in the analysis. A threshold of > 1.74 (= 0.8 of the  $log_2$ -transformed ratios) was chosen for all comparisons. A requirement to show induction above threshold in 100% of the arrays was applied for experiments that were repeated three times or less. For experiments that were repeated more than three times this requirement was relaxed to 45%. Standard deviations were calculated on the log<sub>2</sub>-transformed normalized R/G ratios using the 'nonbiased' method. To calculate the correlation coefficients, the intensities of both channels were normalized to equalize the sums of the intensities.

# Additional data files

A comparison of expression profiles of genital and haltere discs is available with the online version of this paper. Five comparisons between the haltere (H) and genital (G) disc were carried out. Four of the G discs were female (nos 48, 149, 150, 224) and one was male (no. 148). To control for contaminating fat body (FB) cells in the disc preparations, one direct comparison between haltere disc and FB (no. 129) was included. The columns indicate the subclusters with consistent induction in one channel (H, green; G, red), the

gene identification numbers (ID), the average fold induction of the five H:G comparisons, the gene name and function as published on [44]. 130 and 54 genes were induced in the H and G subclusters, respectively, using the same threshold settings as in Figure 3. Note the 2.5-fold induction of *caudal* (CG1759) in the G subcluster.

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