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***Wheat streak mosaic virus* alters the transcriptome of its vector, wheat curl mite (*Aceria tosichella* Keifer), to enhance mite development and population expansion**

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Wheat streak mosaic virus alters the transcriptome of its vector, wheat curl mite (*Aceria tosichella* Keifer), to enhance mite development and population expansion

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Abstract

Wheat streak mosaic virus (WSMV; genus *Tritimovirus*; family *Potyviridae*) is an economically important wheat virus that is transmitted by the wheat curl mite (WCM; *Aceria tosichella* Keifer) in a persistent manner. Virus–vector coevolution may potentially influence vector gene expression to prolong viral association and thus increase virus transmission efficiency and spread. To understand the transcriptomic responses of WCM to WSMV, RNA sequencing was performed to assemble and analyse transcriptomes of WSMV viruliferous and aviruliferous mites. Among 7291 *de novo*-assembled unigenes, 1020 were differentially expressed between viruliferous and aviruliferous WCMs using edgeR at a false discovery rate ≤ 0.05 . Differentially expressed unigenes were enriched for 108 gene ontology terms, with the majority of the unigenes showing downregulation in viruliferous mites in comparison to only a few unigenes that were upregulated. Protein family and metabolic pathway enrichment analyses revealed that most downregulated unigenes encoded enzymes and proteins linked to stress response, immunity and development. Mechanistically, these predicted changes in mite physiology induced by viral association could be suggestive of pathways needed for promoting virus–vector interactions. Overall, our data suggest that transcriptional changes in viruliferous mites facilitate prolonged viral association and alter WCM development to expedite population expansion, both of which could enhance viral transmission.

INTRODUCTION

The wheat curl mite (WCM; *Aceria tosichella* Keifer; family *Eriophyidae*) causes damage to wheat plants by feeding on leaf sap, which ultimately results in dehydration and the formation of characteristic leaf curling. Cumulatively, the loss of sap and leaf surface area due to curling hamper the photosynthetic ability of the plant. However, the most detrimental effect of WCM infestation is its ability to transmit several wheat viruses, including *Wheat streak mosaic tritimovirus* (WSMV) [1], *High Plains wheat mosaic emaravirus* [2], *Brome streak mosaic rymovirus* [3] and *Triticum mosaic poacevirus* [4]. Two distinct genotypes of WCM, type 1 and type 2, have been identified in the USA [5]. Both mite genotypes often occur as mixed populations within wheat fields;

however, these genotypes vary in their ability to transmit wheat viruses [6–8].

Among WCM-transmitted viruses, WSMV is the most economically important virus in the Great Plains of the USA [9, 10]. Although WSMV on average causes 3–5 % annual yield loss in wheat [11], yield losses of as high as 100 % have been observed in infected winter wheat [12]. WSMV is the type species of the genus *Tritimovirus* in the family *Potyviridae* [13]. The 9384-nucleotide single-stranded, positive-sense genomic RNA of WSMV is encapsidated in flexuous filamentous virions of 690–700 × 11–15 nm. WSMV genomic RNA contains a single large open reading frame (ORF) encoding a polyprotein of ~350 kDa that is processed into at least 10 mature proteins by 3 virus-encoded proteinases: P1, HC-Pro and NIa-Pro [13]. WCMs efficiently transmit

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Keywords: Wheat curl mites; *Wheat streak mosaic virus*; transcriptome; differential gene expression; population expansion; behavioral changes. The NCBI BioProject PRJNA489675 contains all raw reads of wheat curl mite transcriptome data. SRR7796148, SRR7796152 and SRR7796153 contain reads derived from viruliferous mites, and SRR7796149, SRR7796150 and SRR7796151 contain reads derived from aviruliferous control mite samples.

Four supplementary tables and one supplementary figure are available with the online version of this article.

WSMV at transmission rates of approximately 50 and 100% with single and 5 to 10 viruliferous mites per test plant, respectively [1, 8]. HC-Pro and coat protein (CP) are required for WCM transmission of WSMV [14–16]. The mode of WCM transmission of WSMV appears to be persistent and circulative, as virus-like particles and inclusion bodies have been found in the digestive tracts of viruliferous mites [17, 18]. Juveniles acquire the virus while feeding on infected wheat and mites remain viruliferous through moulting [17, 19, 20].

The plant cell wall and cuticle offer primary lines of defence against biotic and abiotic factors, including viruses. To circumvent these physical barriers, many plant viruses have evolved precise vector-specific interactions with herbivorous insects [21]. In some cases, viruses are capable of manipulating vector behaviour and physiology to enable more efficient viral transmission and spread, a phenomenon termed ‘adaptive manipulation’ [22]. These behavioural changes can be quite dramatic and include enhanced feeding and increased preference of viruliferous vectors for noninfected host [22–25]. In addition to behavioural modifications, viruses have evolved mechanisms to facilitate acquisition, retention and, occasionally, replication in vectors, often governed by signature motifs in the viral proteome. Often, these molecular determinants facilitate specific interactions with receptors of the vector [26]. In some cases, initial binding of viruses with vector receptors triggers transcriptional responses that directly alter the behaviour of vectors [27].

Studying virus–vector interactions is particularly important because vector-borne plant viruses cause severe economic damage to crops, and vectors can turn an apparently non-significant disease into an epidemic by enhancing virus transmission. Understanding how viruses modulate the metabolism of the vector can provide important information on the biology of these complex interactions and potentially lead to development of improved strategies for integrated pest management. Although virus–vector interactions can be queried using different methodologies, high-throughput RNA sequencing has become a powerful tool to study the global transcriptomic response of vectors influenced by viruses. A number of studies comparing aviruliferous and viruliferous arthropod vectors have been conducted in recent years and have provided insights into how viruses interact with vectors. For example, in viruliferous *Graminella nigrifrons* (black-faced leafhopper), which transmits *Maize chlorotic dwarf waikavirus* and *Maize fine streak rhabdovirus*, viral acquisition was associated with higher expression of genes related to energy production and innate immunity pathways and significant downregulation of peptidoglycan recognition genes [28, 29]. Similarly, viruliferous *Frankliniella occidentalis* (western flower thrips), which transmits *Tomato spotted wilt virus*, showed elevated expression levels of genes associated with insecticide resistance and immune response pathways [30, 31]. *Southern rice black-streaked dwarf fijivirus*, which is transmitted by *Sogatella furcifera* (white-backed plant-hopper), induced

higher stress responses in vectors as compared to aviruliferous insects [32], whereas the acquisition of *Pea enation mosaic luteovirus* was associated with the upregulation of genes involved in transcytosis in the vector, *Acyrtosiphon pisum* (pea aphid) [33]. Additionally, acquisition of *Tomato yellow leaf curl China begamovirus* was associated with the downregulation of genes linked to immune responses and autophagy in whiteflies [34].

The microscopic nature (~200 µm) of WCMs and the lack of adequate genomic and transcriptomic resources for this important vector have hampered research to discern the impact of virus acquisition on vector physiology and delineate the virus–vector dynamics in this economically important pathosystem. In this study, high-throughput RNA sequencing and complementary bioinformatic analyses were used to assemble the *de novo* transcriptome of the WCM and evaluate the transcriptomic signatures of WCMs upon acquisition of WSMV.

METHODS

Preparation of WSMV viruliferous and aviruliferous control WCMs

Type 2 mites reared on the WCM-susceptible wheat cultivar Settler CL (NH03614 CL) grown in caged pots maintained at the University of Nebraska-Lincoln [5] were used for this study. Single seeds of wheat cultivar Settler CL were sown in 4 cm diameter cone-tainers (Stuewe and Sons, Inc., Tangent, OR, USA) filled with standard steam-sterilized greenhouse soil. The cone-tainers were covered with plastic tubular cages of 60 cm in height with two circular air vents of 6 cm diameter and the top was guarded by Nitex Bolting cloth (BioQuip Products, Inc. Compton, CA, USA). The cone-tainers were maintained in a growth chamber at 25 ± 2 °C with 16 h light. Wheat seedlings were infested with a single type 2 WCM at the two-leaf stage using an eyelash attached to a wooden dowel. The successful transfer of each mite was confirmed via observation under a stereo microscope. At 21 days post-infestation, curled top leaves were microscopically observed to confirm the establishment of mite infestations. WCMs from these plants were used to infest WSMV-infected and buffer-inoculated wheat seedlings (see below).

Wheat cv. Settler CL was sown in two sets of five 15 cm diameter pots filled with standard greenhouse soil. Both sets of pots were maintained in separate growth chambers at 25 ± 2 °C with 16 h light. Wheat seedlings at the two-leaf stage were thinned to 15 seedlings per pot. Wheat seedlings in one set of pots designated for viruliferous mites were mechanically inoculated with freshly prepared crude sap from wheat leaves infected with *in vitro* transcripts of WSMV isolate Sidney 81 at a 1:20 dilution in 20 mM sodium phosphate buffer, pH 7.0 (inoculation buffer) [35]. The second set of wheat seedlings were inoculated with inoculation buffer as a control. At 4 days post-inoculation, both sets of plants were infested with WCMs by placing 1 cm-long pieces of infested leaves from the source plants

between the leaf bases of the virus- and buffer-inoculated plants. Three independent infestations of viruliferous and aviruliferous mites were prepared to generate three biological replicates to be used for RNA-seq and RT-qPCR studies.

Mites were collected separately from infected (viruliferous) and uninfected control plants (aviruliferous) at 28 days after infestation. Infested plants were processed in batches of five to six plants, which were cut into 5 cm-long pieces and vortexed in 30 ml of phosphate-buffered saline (PBS) with 0.2% polysorbate 20. Mites were filtered from the washings using a cell strainer with a pore size of 40 µm (EASYstrainer Cell Sieves) and were transferred into a clean Petri dish. WCMs were inspected visually under a stereo light microscope and selectively aspirated using a micropipette to separate the mites from any contaminating plant debris. The mites were snap frozen in liquid nitrogen in 200 µl aliquots in PBS and stored at -80°C until further processing for RNA extraction.

RNA extraction and Illumina sequencing

Total RNA (rRNA+mRNA) was extracted from three biological replicates each of frozen viruliferous and aviruliferous mites by pulverizing the samples in liquid nitrogen using a mortar and pestle and adding 1 ml of TriPure isolation reagent (Sigma-Aldrich, St Louis, MO, USA). RNA was processed following the manufacturer's directions and treated with RT enhancer to eliminate genomic DNA contamination. The quantity and integrity of each RNA sample were validated on an Agilent 2100 Bioanalyzer (Agilent Technologies, Santa Clara, CA, USA) (data not shown). The total RNA ($\sim 1.0\ \mu\text{g}$) from each sample was subjected to mRNA enrichment using oligo dT magnetic beads and subsequently each RNA sample was prepared for reverse transcription and sequencing using the TruSeq Stranded mRNA Library Preparation Kit (Illumina, Inc., San Diego, CA, USA) per the manufacturer's instructions. The six barcoded libraries were combined into a single library pool and sequenced on an Illumina HiSeq 2500 platform to a depth of approximately 40 million 2×50 bp paired-end reads (40 Gb) per sample at the University of Minnesota Genomics Center (Minneapolis, MN, USA). All raw reads were submitted to the NCBI Sequence Read Archive under BioProject PRJNA489675. SRR7796148, SRR7796152 and SRR7796153 contain reads derived from viruliferous mites, and SRR7796149, SRR7796150 and SRR7796151 contain reads derived from aviruliferous control mite samples.

De novo transcriptome assembly, filtering and annotation

Reads from all biological replicates from both treatments were quality-checked using the program FastQC (<https://www.bioinformatics.babraham.ac.uk/projects/fastqc/>) and pooled together to build a *de novo* transcriptome assembly with Trinity (version 2.0.6) [36]. Reads were normalized *in silico* prior to assembly to a maximum coverage of $50\times$ to reduce the frequency of erroneous kmers in reads used to build the assembly [37], and the normalized reads were

used for assembly with the following parameters: `min_kmer_cov 1`, `min_contig_length 200`, and `SS_lib_type RF` for strand specificity. All other parameters were set to default. Transcriptome assemblies are inherently noisy and often contain multiple allelic variants derived from the same parent unigenes, which is caused by pooling multiple unrelated individuals together to obtain sufficient levels of RNA for extraction and library preparation. Additionally, they can also contain fusion transcripts and/or misassembled transcripts [38, 39]. In order to reduce the abundance of these spurious transcripts and unigenes in the assembly, it was filtered to retain only high-quality transcripts containing full-length or near full-length coding regions. To accomplish this, several steps were undertaken. First, all reads were mapped back to the assembly using Bowtie2 [40] with the `align_and_estimate_abundance.pl` script packaged with Trinity and abundance was computed on per isoform and per unigene levels using RSEM [41]. Next, coding regions were predicted using Transdecoder (<https://github.com/TransDecoder/TransDecoder/>). Default parameters were used with the addition of HMMER (version 3.0) [42] to identify Pfam-A domains and facilitate the detection of ORFs. Additionally, transcripts derived from potential microbial contaminants were identified by BLASTP and BLASTN comparisons (version 2.4.0+) to the non-redundant protein and nucleotide databases (NR and NT; downloaded 17 February 2018). The top five matches with *e*-values $\leq 1e-5$ were retained for each transcript and used for taxonomic classification with MEGAN [43]. Because lower-quality and misassembled transcripts are usually low in abundance, we removed transcripts and unigenes with TPM values below 0.5 and transcripts that represented less than 5% of the abundance of the dominant isoform of each unigene from the assembly. Transcripts and unigenes that did not have BLASTP or BLASTN matches to arthropod-derived sequences in the nucleotide database (nt; downloaded 17 February 2018) were also removed from the assembly.

Functional annotations of protein-coding unigenes were performed using Trinotate (<https://trinotate.github.io/>). In brief, the top BLASTP and BLASTX matches to the UniProtKB/Swiss-Prot 2018_02 database (downloaded 17 February 2018) for each protein coding unigene were retrieved, Pfam-A domains were identified using HMMER (version 3.0) [42], transmembrane domains were identified using TMHMM [44], and signal peptides were identified using signalP [45]. GO, eggNOG and KEGG [46] terms were retrieved from the Trinotate.sqlite database using the top BLASTP matches to the Swiss-Prot/Uniprot database as queries. Transcripts were screened for any remaining adapter sequence using BLASTN searches against the UniVec database (downloaded on 15 May 2018) and fully annotated transcripts were submitted to the NCBI's Transcriptome Shotgun Assembly (TSA) database (accession number GGY000000000). The programs Transvestigator [47] and Annie [48] were used to facilitate NCBI submission of the annotated transcripts. Other supporting information, such as Trinotate annotations and predicted protein translations for all unigenes

Table 1. Summary of Trinity *de novo* assembly of read mapping and contig read counts for wheat curl mite

Before length-filtering	No. of bases	29.96 Mb
	No. of transcripts	26989
	No. of unigenes	25678
	Contig N50 (bp)	1857
	Median contig length (bp)	656
After length-filtering*	No. of bases	15.87 Mb
	No. of transcripts	7785
	No. of unigenes	7291
	Contig N50 (bp)	2644
	Median contig length (bp)	1718

*Transcripts with lengths less than 200 bp (minimum cutoff), transcripts putatively derived from microbial taxa and transcripts with transcripts per million (TPM) values below 0.5 were removed from the assembly prior to differential expression analysis.

(including the unfiltered unigene set), can be found at <https://data.nal.usda.gov/dataset/de-novo-transcriptome-assembly-and-annotations-wheat-curl-mite-aceria-tosichella>.

Differential expression analysis

To compare global unigene expression patterns between viruliferous and aviruliferous mites, reads from each of the three biological replicates from the two treatments were separately mapped back to the filtered transcriptome assembly using Bowtie2 [40], and transcript abundances were determined using RSEM [41]. Differential expression analysis between transcriptomes from viruliferous and aviruliferous mites was performed at the unigene level using EdgeR [49]. Read counts were normalized using the trimmed mean of M-values (TMM) approach and transcripts with counts ≤ 10 across at least four samples were removed. Transcripts with \log_2 -fold changes ≥ 0.5 and FDR-corrected P -values ≤ 0.05 were defined as differentially expressed. A \log_2 -fold change of 0.5 was chosen because even small fold changes in

Table 2. Annotation metrics for unigenes from the filtered wheat curl mite transcriptome assembly. Values indicate numbers of unigenes

	No. of unigenes
BLASTX matches to Sprot/Uniprot	6749
BLASTP matches to Sprot/Uniprot	6798
Pfam domains	6592
Signal peptides	516
Transmembrane domains	1691
Gene ontology (GO) assignments	5019
KEGG assignments	5412
BLASTP matches to NR	7788
Matches to hypothetical proteins (NR)	1204
Full-length ORFs	5636
5' partial ORFs	1530
3' partial ORFs	622

Table 3. MEGAN taxonomy classification for highest scoring BLASTP matches of protein-coding unigenes from filtered wheat curl mite transcriptome assembly against the non-redundant protein database

Order	No. of unigenes
Anura	99
Coleoptera	407
Diplostraca	381
Diptera	878
Echinoida	92
Enteropneusta	141
Hemiptera	258
Hymenoptera	836
Ixodida	1010
Neoptera	289
Phthiraptera	229
Rhabditida	160
Other	3011
Could not be classified to order	26

expression levels can have biological and phenotypic significance and FDR is well controlled by EdgeR when \log_2 -fold change thresholds >0.3 and $FDR \leq 0.05$ are used [50].

GO enrichment analyses for up- and downregulated genes in viruliferous mites were performed using GoSeq [51]. Basal terms for all GO assignments were retrieved using the `extract_GO_assignments_from_Trinotate_xls.pl` script included in the Trinotate package. For both up- and downregulated genes, the entire list of transcripts/unigenes tested for differential expression was used as a reference to determine enrichment. All genes were weighted by gene length and enriched terms were identified using the Wallenius approximation ('pwf' option) with Benjamini–Hochberg adjusted P -values ≤ 0.05 .

KEGG and Pfam enrichment analysis

Pfam, KO and KEGG pathway enrichment analyses for up- and downregulated genes were performed using the GeneOverlap package in R, which calculates enrichment significance using Fisher's exact test on individual gene sets [52], and the Trinotate Pfam and KO annotations [53]. Benjamini–Hochberg adjustment was used on the resulting P -values to correct for multiple testing [54].

Assessment of transcriptome completeness and comparison to other mite transcriptomes

To assess the completeness of the transcriptome, Benchmarking Using Single Copy Orthologs (BUSCO) analysis [55] was performed using the Arthropod odb9 orthologue set available at https://busco.ezlab.org/datasets/arthropoda_odb9.tar.gz and representation of core metabolic pathways was validated using the KEGG pathway assignments computed in the previous section. Further, comparisons were made to the genomes and transcriptomes of other mites, focusing primarily on phytophagous species belonging to the same superorder as WCM (Acariformes). Because

protein-coding sequences were not available for the mite species found in TSA, transcriptomes from the raw sequence read archive reads were reassembled and reannotated using Trinity and Trinotate as described above for WCM. TBLASTN searches were performed using predicted protein coding sequences derived from the fully annotated *T. urticae* genome assembly [56] as queries against the three mite transcriptome assemblies [*P. ulmi* [57], *P. citri* [58] and *A. tosichella* (this study)] to identify homologous coding regions that may have been missed by Trinotate. To facilitate comparisons, orthology assignments were computed using bidirectional BLASTP searches and the program OrthoFinder (2.1.2) [59] to identify unigenes that were conserved among these mites, unigenes that were unique to *A. tosichella* and unigenes whose copy number may be expanded in *A. tosichella* relative to the other mite species. To ensure that transcript isoforms derived from the same unigene did not contribute to copy number inflation, only the most highly expressed isoform per unigene from transcriptome assemblies was included in the orthology analysis.

RT-qPCR

Total RNA (1.0 µg) was treated with RT enhancer to eliminate genomic DNA contamination and was reverse-transcribed using random hexamers and AMV RT (Roche, Indianapolis, IN, USA) in a 10 µl reaction. RT-qPCR was performed on technical duplicates with cDNA prepared from the same three biological replicates of total RNA that were used for the RNA-seq experiment. SsoAdvanced SYBR Green Supermix (Bio-Rad, Hercules, CA, USA) was used per the manufacturer's instructions and analysis was performed using the Bio-Rad CFX Connect Real-Time PCR System. The primers used for RT-qPCR are listed in Table S1 (available in the online version of this article) with

the transcript identifier, annotation of templates along with orientation and corresponding nucleotides. The amplification conditions used for qPCR were 95 °C for 2 min, followed by 40 cycles of 95 °C for 10 s, 50 °C for 30 s and 72 °C for 1 min. No template and no RT reactions were included as negative controls. The relative expression levels of WCM unigenes in viruliferous and aviruliferous mites were calculated with the $\Delta\Delta C_t$ method [60] using actin and glucose-6-phosphate dehydrogenase as references for normalization.

RESULTS

WCM transcriptome assembly and annotation

To study the transcriptomic response of WCMs exposed to WSMV, RNA-seq analysis was performed on three biological replicates of mites fed on WSMV-infected and buffer-inoculated wheat plants. A total of approximately 270 million 2×50 bp paired reads totalling over 27 Gb were generated from the 6 transcriptomic libraries with an average coverage of approximately 45.7 million reads (4.6 Gb) per library. Few PCR duplicates were identified in the raw reads (<0.3%) and the average PHRED quality scores (Illumina 1.9) exceeded 32 along the entire lengths of the forward and reverse read. Approximately 14.5 million high-quality normalized read pairs (~1.5 Gb) were used to generate the *de novo* transcriptome assembly, which led to the initial assembly of 26 989 transcripts derived from 25 678 unigenes (Table 1); although the reads were not quality-filtered prior to assembly, 99.9% of the raw reads matched the consensus sequence for each transcript, indicating that the raw data used to build the assemblies were of extremely high quality. From this initial assembly, the majority of the transcripts had matches to proteins derived from metazoa; however, approximately 2800 unigenes had the highest scoring

Table 4. Summary of RNA-seq-derived raw reads and mapped sequences from *Wheat streak mosaic virus* (WSMV) viruliferous and control wheat curl mites (WCMs)

Sample description ^a	No. of raw reads	No. of reads mapped to the transcriptome assembly in proper pairs	% of reads mapped in proper pairs	% of reads mapped to other genomes		
				Wheat	WSMV	Fungus
Aviruliferous WCMs-1	46 848 229	26 045 552	55.60	0.14	0	1.08
Aviruliferous WCMs-2	43 317 338	24 217 625	55.91	0.12	0	1.09
Aviruliferous WCMs-3	45 938 305	24 974 195	54.36	0.2	0	0.25
Mean	45 367 957	25 079 124	55.29	0.15	0	0.81
Viruliferous WCMs-1	41 971 907	21 188 142	50.48	0.12	7.49	0.34
Viruliferous WCMs-2	43 694 956	23 160 099	53.00	0.1	4.46	0.23
Viruliferous WCMs-3	48 296 451	25 506 536	52.81	0.09	3.81	0.25
Mean	45 038 641	23 284 926	52.10	0.11	5.25	0.27

a, 1–3 represent RNA from biological replicates prepared from three independent experiments. Aviruliferous WCMs and viruliferous WCMs represent wheat curl mites reared on buffer-inoculated and WSMV-infected wheat plants, respectively.

BLASTP matches to fungal taxa, while 44 and 3 unigenes had matches to bacteria and archaea, respectively. Overall, the majority of the unigenes (~2500) derived from microbes had the highest scoring BLASTP matches to the fungal class Sordariomycetes, with the majority of these unigenes having matches to the Nectriaceae (*Fusarium*, *Gibberella* and *Nectria*). A small percentage of reads (<1%) were derived from fungi in both the aviruliferous and viruliferous mites (Table 4). In addition, approximately 29% of the unigenes (4651) that coded for putative proteins showed no significant matches to any annotated GenBank protein sequence, suggesting significant divergence from previously annotated proteins. Although many of these proteins could represent erroneous protein predictions, 404 of these proteins (8.7%) had predicted Pfam domains, 1049 had predicted transmembrane domains (22.6%) and 964 had predicted signal peptides (20.7%), suggesting that at least some of these unigenes encode functional proteins.

After filtering to remove non-coding sequences, low-abundance transcripts with transcripts per million (TPM) values ≤ 0.5 and sequences derived from microbes, 7785 transcripts derived from 7291 unigenes remained. The contig N50 was 2644 nt, the median contig length was 1718 nt and the total assembly length was approximately 15.9 Mb (GC content: 46.07%) (Table 1). When only the longest isoform per unigene was considered, the contig N50 decreased slightly to 2626 nt, median contig length remained approximately the

same at 1709 nt, and total assembly length decreased to approximately 15.87 Mb (Table 1). Of the 7291 unigenes that coded for proteins, 6749 had BLASTX matches to the Swiss-Prot/UniProt database, 6798 had BLASTP matches to the Swiss-Prot/UniProt database, approximately 7700 had BLASTP matches to NR, 6592 had PfamA domains, 516 had predicted signal peptides and 1691 had predicted transmembrane domains (Table 2). The vast majority of the unigenes identified had the highest scoring BLASTP matches to proteins derived from phylum Arthropoda (~70.1% of the protein-coding unigenes). Because few genomic resources exist for taxa derived from the subclass Acari, which includes mites, the most abundant BLASTP matches to the NR database were actually to proteins derived from the class Insecta (~46.1% of the protein coding unigenes), with Arachnida representing the class with the second most frequent matches (~15.5%, of which most matched to members of the subclass Acari). This finding indicates that the nucleotide and amino acid sequences derived from the WCM transcriptome are divergent from potential orthologues and homologues from other mite species represented in the NR database. At finer taxonomic resolutions, the highest scoring BLASTP matches were broadly distributed over more than 50 different orders; however, the most frequent orders observed included Ixodida, Diptera and Hymenoptera (Table 3).

Additionally, one unigene that was approximately 6160 nt in length and corresponded to the WSMV polyprotein was

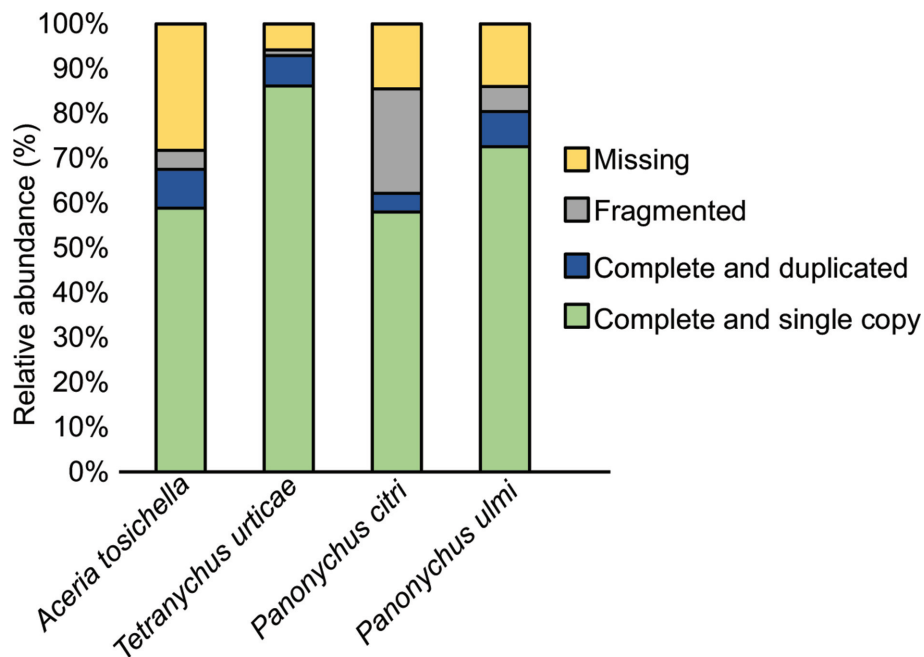


Fig. 1. Benchmarking Universal Single-Copy orthologs (BUSCO) analysis of WCM transcriptome and transcriptomes and genomes of other mites belonging to the superorder Acariformes. One isoform from each WCM unigene was searched against the arthropoda odb 9 database. WCM results were compared to results from protein-coding unigenes derived from the *Panonychus citri* [97] and *Panonychus ulmi* [95] transcriptome assemblies and protein-coding genes derived from the *Tetranychus urticae* [96] genome assembly.

Table 5. Top 30 enriched orthogroups in the wheat curl mite compared to other mite genomes and transcriptomes

Orthogroup	Annotation	Representative unigene	<i>P. ulmi</i>	<i>P. citri</i>	WCM	<i>T. urticae</i>
OG0000243	C2H2-type zinc finger	TR1204 c0_g1	1	2	4	3
OG0000200	Troponin	TR133 c0_g1	1	2	6	2
OG0000246	Cyclophilin-type peptidyl-prolyl cis-trans isomerase/CLD	TR2015 c0_g1	3	1	4	2
OG0000186	Ion transport protein	TR2139 c0_g1	1	3	4	3
OG0000178	DEAD/DEAH box helicase	TR2337 c0_g1	2	2	4	3
OG0000072	Cathepsin propeptide inhibitor domain (I29)	TR254 c0_g1	1	0	15	0
OG0000009	ABC transporter	TR3201 c1_g1	5	10	18	9
OG0000202	Trypsin	TR3233 c0_g1	0	2	6	3
OG0000137	RyR domain	TR3639 c0_g1	2	3	6	1
OG0000173	Hypothetical	TR3639 c3_g1	1	3	5	2
OG0000149	Cytochrome P450	TR3694 c0_g1	0	4	5	3
OG0000073	Zinc carboxypeptidase	TR3749 c0_g1	1	2	9	4
OG0000066	Fatty acid desaturase	TR3970 c1_g1	5	2	6	3
OG0000195	Tropomyosin	TR4510 c0_g1	2	2	5	2
OG0000097	Eukaryotic elongation factor 1 beta central acidic region	TR479 c0_g1	3	1	8	2
OG0000067	Acytransferase family	TR4951 c0_g1	2	3	7	4
OG0000099	Hypothetical	TR4951 c4_g1	3	4	6	1
OG0000059	Aldo/keto reductase family	TR5435 c0_g1	4	2	7	4
OG0000114	C2 domain	TR5758 c0_g1	2	3	5	3
OG0000227	RasGEF N-terminal motif	TR5959 c12_g2	1	3	5	1
OG0000171	Adenylate and guanylate cyclase catalytic domain	TR5974 c21_g1	3	2	4	2
OG0000237	Hypothetical	TR5991 c32_g1	1	2	6	1
OG0000140	VWA N-terminal	TR6023 c1_g1	3	2	5	2
OG0000110	Hypothetical	TR6040 c11_g1	2	2	5	4
OG0000179	Protein kinase domain	TR6042 c0_g1	2	1	6	2
OG0000235	C2 domain	TR6044 c16_g1	3	1	4	2
OG0000241	Pyruvate phosphate dikinase, PEP/pyruvate-binding domain	TR6118 c0_g1	3	1	5	1
OG0000098	Hypothetical	TR6144 c0_g1	2	4	7	1
OG0000182	Phorbol esters/diacylglycerol-binding domain (C1 domain)	TR6149 c11_g1	2	2	4	3
OG0000068	Low-density lipoprotein receptor repeat class B	TR6207 c9_g2	4	3	8	1

All-versus-all BLASTP searches in combination with the program OrthoFinder were used to identify gene families whose copy number was potentially expanded in WCM compared to other mites. The transcriptomes for *P. ulmi* and *P. citri* were assembled and annotated as described in the Methods section, and the proteome of *T. urticae* was downloaded from NCBI. Single isoforms for each gene or unigene were used in the all-versus-all BLASTP searches to ensure that different isoforms did not contribute to any copy number inflations.

recovered from the assembly. BLASTP and BLASTN matches were 100 and 99 % identical to the WSMV Sidney 81 strain (GenBank: AF057533.1), respectively, and the nucleotide sequence for the unigene aligned with positions 3225 to 9384 of its closest match. A small percentage of reads from each of the libraries derived from viruliferous WCMs also mapped to the WSMV transcriptome, confirming that the virus had been successfully transmitted to the mites (Table 4). No reads from the control libraries mapped to WSMV and only a small percentage of reads in all samples mapped to the wheat genome (*Triticum aestivum* v2.2, phytozome.jgi.doe.gov) (Table 4).

Overview of WCM transcriptome annotations and orthology analysis

Overall, the recovery of Benchmarking Universal Single-Copy Orthologs (BUSCOs) from the WCM was similar to

recovery of BUSCOs from the transcriptomes and genomes of other phytophagous mites, suggesting that the transcriptome assembly represents a fairly complete reconstruction relative to other mites. Approximately 59 % of the arthropod BUSCOs were identified as complete and single copy, 8.6 % were identified as complete and duplicated, and 28.2 % were identified as missing (Fig. 1). The percentages of recovered and missing BUSCOs may seem low and high, respectively, especially in comparison to metrics that are typically reported for insect genomes and transcriptomes. However, these values are similar to the number of BUSCOs recovered from transcriptomes of two mites belonging to the genus *Panonychus* (58 % complete/single copy and 14.5 % missing in *P. citri*, and 72.6 % complete/single copy and 14.0 % missing in *Panonychus ulmi*). Additionally, 5.8 % of arthropod BUSCOs were missing from the genome assembly of two-spotted spider mite (*Tetranychus urticae*) (Fig. 1). This

suggests that it may be difficult to obtain a complete gene space inventory from a transcriptome assembly in some mite species or that the BUSCO arthropod database might not perform well with species of Arachnida. Additionally, the percentage of complete and duplicated BUSCOs was lower in the filtered assembly compared to the raw transcriptome assembly (8.6 vs 11.6%, respectively) and the recovery of complete and single copy BUSCOs and fragmented BUSCOs did not differ (data not shown). This finding indicates that removing low-abundance transcripts and isoforms from the data potentially improved our assembly by removing duplications caused by allelic variation in the sequence data.

Orthology analyses among the 4 different mite species revealed the presence of over 3000 different orthologue groups that included representative proteins from all 4 mites, and of these 903 consisted exclusively of single-copy orthologues. Approximately 500 different orthogroups contained higher numbers of assigned unigenes in the WCM compared to other mite transcriptomes and genomes, which is potentially indicative of copy number expansion in this species (Table 5). The majority of these unigenes encoded proteins such as trypsin, CYP450, ABC transporters, aldoketo reductases (AKRs), catalases, cuticular proteins, glutathione S-transferases (GSTs), juvenile hormone-binding proteins (JHBPs), heat-shock proteins (HSPs), leucine-rich repeat proteins (LRRs), sugar transporters, chemoreceptors and insulinases, which tend to be amplified in taxon-specific manners in the genomes of many other arthropod species. Notably, proteins containing cathepsin propeptide inhibitor

domains (I29) were uniquely expanded in the transcriptome of the WCM compared to the other 3 mites, and were assigned to an orthogroup containing 16 of the copies found in the WCM compared to 1 copy found in *P. citri* and no copies found in the other 2 mites (Table 5). Other notable expansions included orthogroups of ABC transporters (18 unigenes in the WCM compared to <10 copies in the other mites), zinc carboxypeptidases (9 unigenes in the WCM compared to <5 copies in the other mites), lipoproteins (8 unigenes in the WCM compared to ≤ 5 copies in the other mites), sodium channel proteins (7 unigenes in the WCM compared to ≤ 4 copies in the other mites) and conserved hypothetical proteins (6 unigenes in the WCM compared to ≤ 4 copies in the other mites) (Table 5). Although these expansions are of potential importance for the biology of the WCM, genome assembly will be required to validate the expansion of these paralogues observed in the transcriptome assembly.

In general, unigenes encoding proteins involved in signalling, structural proteins, proteinases and immunity were among the most abundant protein families observed in the WCM transcriptome (Fig. 2). Specifically, protein kinases were the most abundant Pfams, followed by WD domains, trypsins, homeobox proteins, immunoglobulins, ubiquitin-conjugating enzymes, RNA-recognition motifs, DEAD/DEAH box helicases and cuticle proteins (Fig. 2). Unigenes encoding trypsins (86 unigenes) were the most common types of proteinases identified in the assembly, followed by papain family cysteine proteinases (27) and aspartyl

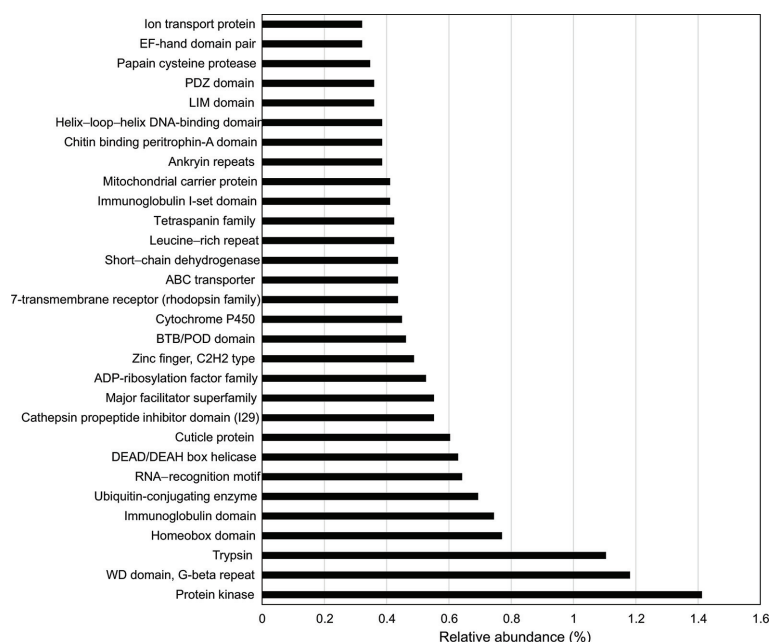


Fig. 2. Relative abundances of the most dominant Pfam domains in the wheat curl mite transcriptome assembly. Pfam annotations were retrieved using Trinotate as described in the Methods section. Occurrences of Pfam domains were counted only once for each protein and abundance was expressed relative to the total number of proteins in the transcriptome that had detectable Pfam domains.

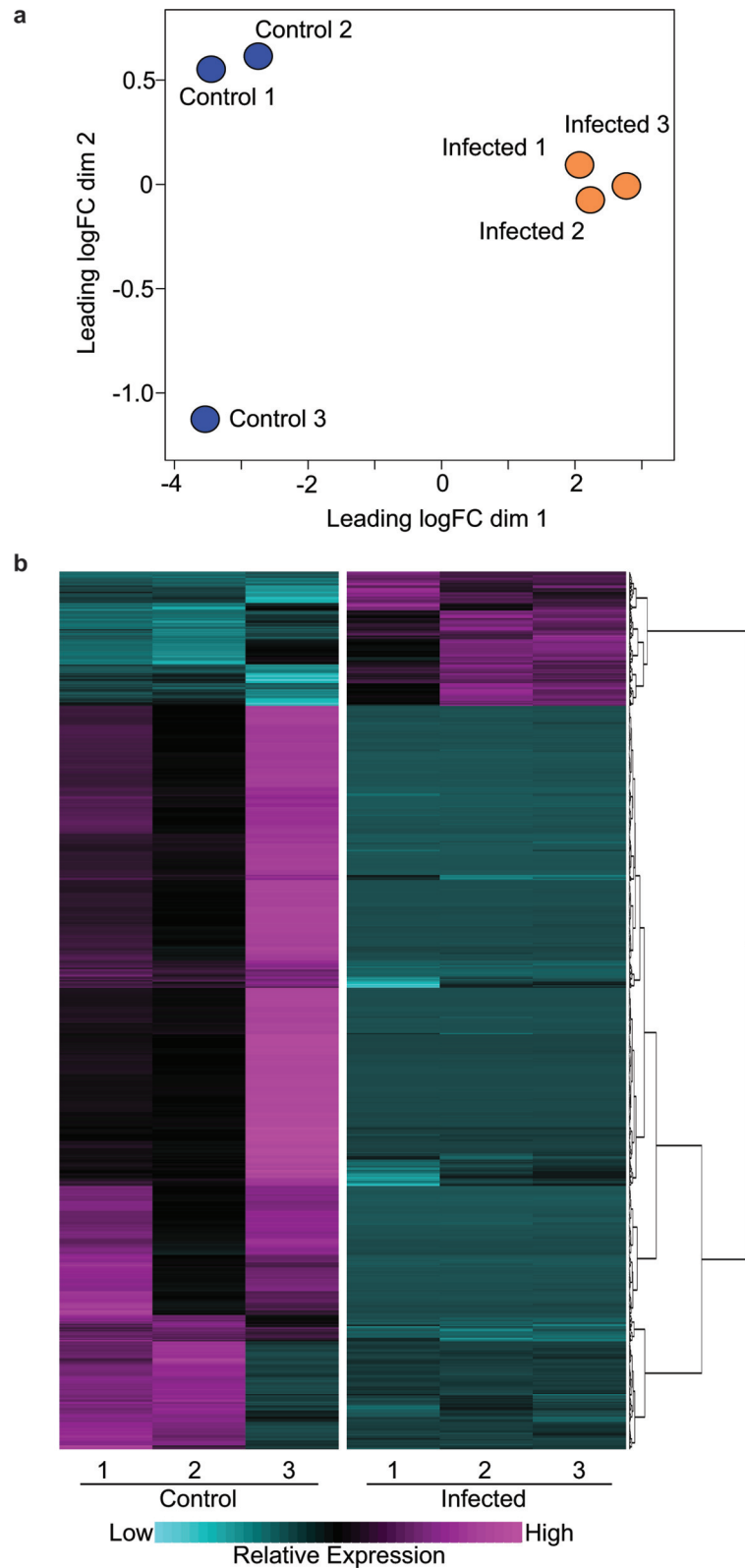


Fig. 3. Differentially expressed genes in *Wheat streak mosaic virus* viruliferous wheat curl mites. (a) Non-metric multidimensional scaling (NMDS) analysis of the expression profiles of viruliferous and aviruliferous WCMs. The NMDS analysis was based on Euclidean dissimilarities between the sample pairs and was performed using the plotMDS function in edgeR. Aviruliferous control samples are shown in blue and viruliferous samples are shown in orange. (b) Heatmap and clustering analysis of differentially expressed unigenes

identified in viruliferous WCMs. Differentially expressed unigenes were identified using edgeR with an FDR cutoff of ≤ 0.05 and a \log_2FC of ≥ 0.5 , the heatmap was prepared using JMP (v12) and clustering was performed using Ward's method. The colour key depicts the magnitude of the expression levels for each unigene ranging, from very low expression (cyan) through very high expression (magenta) to moderate expression (black).

proteases (15). Other protein families linked to immunity and stress response that were found in the assembly included cytochrome P450s (35 unigenes), short-chain dehydrogenases (34), LRRs (33), lipases (21), GSTs (15), peroxidases (15), GMC oxidoreductases (8), carboxylesterases (7) and UDP-glucuronosyltransferases (UGTs; 2). Similar to other phloem-feeding arthropod taxa, unigenes encoding enzymes capable of degrading plant cell walls were not particularly abundant; however, one unigene predicted to encode a glycoside hydrolase (GH) family 45 β -1,4-endoglucanase was identified. Additional unigenes encoding GH family proteins included GH 18 chitinases (10), GH 1 β -glucosidases (6), GH 15 phosphorylase kinases (glycogen breakdown; 2), GH 31 lysosomal α -glucosidases (4), GH 30 glucosylceramidases (5), GH 38 lysosomal α -glucosidases (2) and GH 47 α -mannosidase-like proteins involved in degrading misfolded glycoproteins (5). A unigene encoding a pectin acetylesterase also was identified, which could be important for digesting homogalacturonan polysaccharides found in plant primary cell walls.

Influence of WSMV on WCM transcriptome

Overall, a total of 1020 differentially expressed genes were detected in WSMV viruliferous mites relative to the aviruliferous control mites at a false discovery rate (FDR)-corrected *P*-value of 0.05 and a \log_2 -fold change of 0.5 (Fig. S1). The number of reads mapped to the reference assembly did not differ among the viruliferous and aviruliferous mites, which ranged from 50–52 % and 54–55 % of the raw reads, respectively (Table 4). Of these, 155 unigenes were upregulated in viruliferous mites relative to the controls and 865 were downregulated (Fig. 3a, b). Additionally, the two treatments were clearly separable from one another via non-metric multidimensional scaling, and correlation values among the three biological replicates within each treatment were ≥ 0.95 (Fig. 3a), suggesting consistent responses among replicates within each treatment. The downregulated unigenes were associated with the enrichment of 112 different gene ontology terms, of which 85, 111 and 16 were assigned to the biological processes (BP), molecular function (MF) and cellular component (CC) parent categories, respectively (Fig. 4a; Tables 6 and S2). BP gene ontology (GO) terms were mostly related to purine nucleoside metabolic processes, hydrogen ion transmembrane transport, glycosyl compound metabolic processes, translational elongation, energy coupled proton transport and oxidation-reduction, while CC was enriched for terms related to ribosomes and MF was enriched for the structural constituents of ribosomes, hydrolase activity, peptidase activity, magnesium chelatase activity and ligase activity (Fig. 4a; Tables 6 and S2). Among the upregulated unigenes, two MF terms showed enrichment, including structural constituents of cuticle and iron ion binding

(Table 7, Fig. 4a). Although not significantly enriched, several terms were highly represented among upregulated unigenes, including steroid metabolic process (4 out of 15 annotated unigenes) and fatty-acyl-CoA (alcohol-forming) activity (4 out of 12 annotated unigenes) (Table S2).

Pfam, Kyoto Encyclopedia of Genes and Genomes (KEGG) orthology (KO) and KEGG pathway enrichment analyses of differentially expressed unigenes provided some additional insights into the impacts of viral acquisition on the WCM transcriptome. Overall, a total of 115 different Pfam domains were associated with 1020 differentially expressed unigenes detected in this study. Over 40 Pfam domains (~35 %) were enriched among unigenes that were downregulated in response to WSMV and only one Pfam domain was enriched among upregulated unigenes (Table 8; Fig. 4b). Enriched Pfam domains from unigenes that were downregulated in viruliferous mites included aldo-keto reductases (AKRs), α -amylases, insect cuticle proteins (22 downregulated unigenes), GSTs, trypsin, cathepsin propeptide inhibitor domain I29 proteins, HSPs, JHBPs, lipocalins, ferritins and ubiquitins (Table 8). Among upregulated unigenes, enriched Pfam domains also included insect cuticle proteins (10 upregulated unigenes) (Fig. 3b). Although not enriched, other prominent Pfam domains included CYP450s, male sterility proteins, animal haem peroxidases, HSP20s and lipases (Table S3; Fig. 4b), most of which were associated with the downregulated unigenes. Overall, Pfam enrichments and pathway analyses suggest that WSMV influenced the development, reproduction, apoptosis, immunity, oxidative stress, and digestive and metabolic processes of WCM.

KO analysis showed an enrichment of 34 different terms among unigenes that were downregulated in viruliferous WCMs, which included myosin heavy chain; aldehyde reductase; apolipoprotein D and lipocalin family protein; cathepsins L, B and D; elongation factor 1- α ; HSP 70; transmembrane serine protease; and glutathione peroxidase (Table 9; Fig. 5a). No significantly enriched KO terms were found in upregulated unigenes; however, high numbers of unigenes were assigned to terms related to CYP450s, dystonin and calcium/calmodulin-dependent protein kinases were upregulated (Fig. 5a). Several KEGG pathways were enriched among differentially expressed unigenes, including 22 that were enriched in downregulated unigenes (Table 10; Fig. 5b). These pathways were mostly related to ribosome, oxidative phosphorylation, apoptosis, glutathione metabolism, metabolism of xenobiotics by CYP450, translation, innate immunity, galactose metabolism and the pentose phosphate pathway (Table 10; Fig. 5b). Although not significantly enriched, several unigenes encoding proteins involved in steroid hormone biosynthesis were upregulated (Fig. 5b).

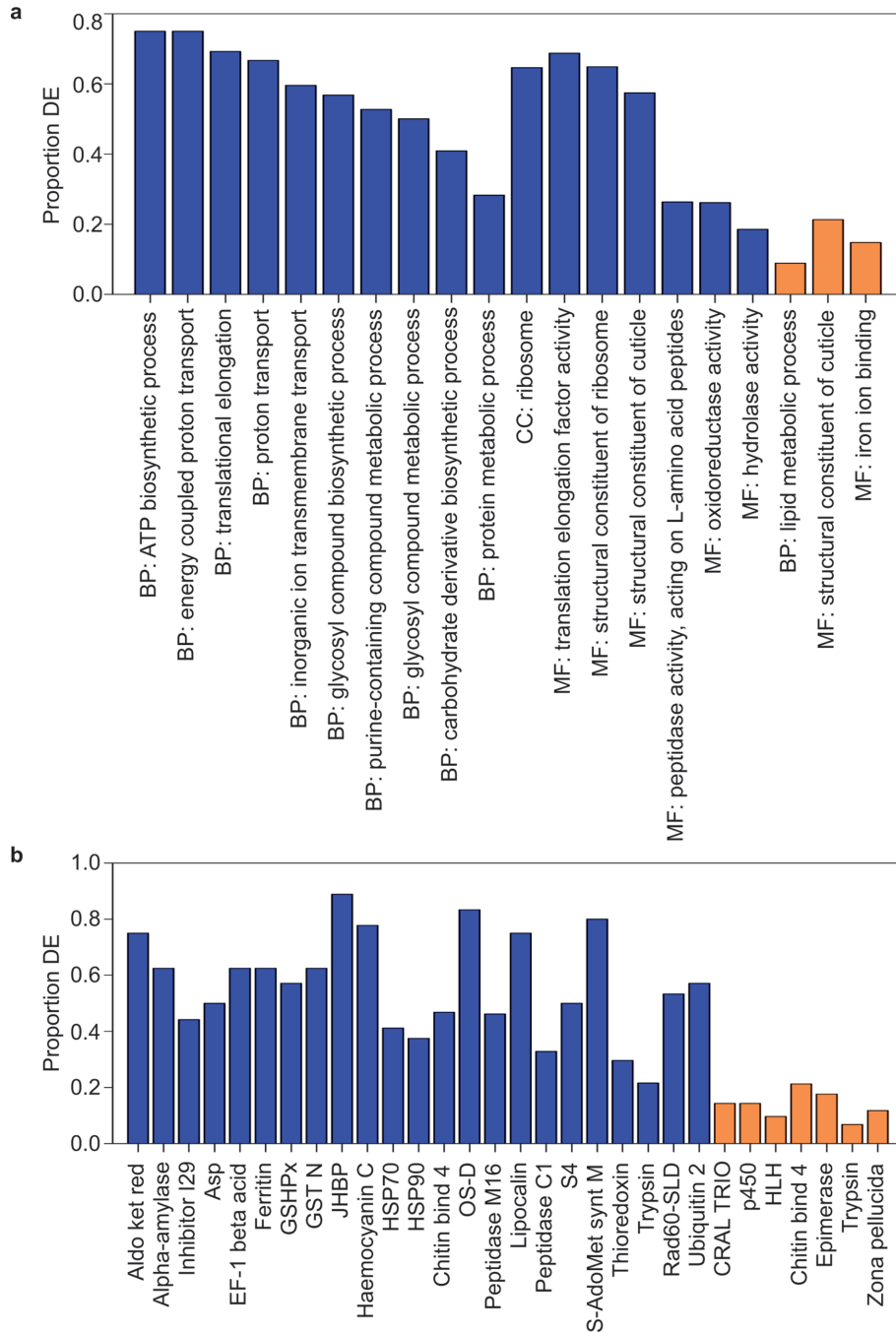


Fig. 4. Select enriched gene ontology (GO) terms and Pfam domains in differentially expressed wheat curl mite unigenes under the influence of *Wheat streak mosaic virus*. (a) Graphical representation of select significantly enriched (FDR <0.05) GO categories in differentially expressed unigenes after WSMV acquisition. (b) Graphical representation of select significantly enriched (P -value<0.05) Pfam domains in differentially expressed unigenes. In both (a) and (b), the bars depict the percentage of unigenes assigned to each category that were downregulated (blue) and upregulated (orange) in viruliferous mites relative to the total number of unigenes in the transcriptome.

Additional influences of WSMV on unigenes linked to stress response, development and digestion

Overall, the strongest transcriptional impacts on WCMs that had acquired WSMV were to unigenes encoding

ribosomal proteins, the majority of which were downregulated in viruliferous mites. In addition to those unigenes, other unigenes related to translation, protein biosynthesis and processing, folding and transport were downregulated

Table 6. Top 20 enriched gene ontology (GO) terms observed in unigenes that were downregulated in *Wheat streak mosaic virus*-viruliferous wheat curl mites

Term	Ontology	No. of downregulated unigenes	No. of assigned unigenes	FDR corrected P-value	Description
GO:0008152	BP	469	1962	4.30E-14	Metabolic process
GO:0005198	MF	173	287	7.66E-14	Structural molecule activity
GO:0003735	MF	146	225	3.12E-13	Structural constituent of ribosome
GO:0005840	CC	146	226	3.49E-13	Ribosome
GO:0006412	BP	139	216	1.29E-12	Translation
GO:0030529	CC	146	240	1.43E-11	Intracellular ribonucleoprotein complex
GO:0043228	CC	146	253	5.29E-11	Non-membrane-bounded organelle
GO:0043232	CC	146	253	5.29E-11	Intracellular non-membrane-bounded organelle
GO:0044444	CC	193	405	1.64E-10	Cytoplasmic part
GO:1901576	BP	197	479	4.26E-10	Organic substance biosynthetic process
GO:0034645	BP	158	322	8.86E-10	Cellular macromolecule biosynthetic process
GO:0071704	BP	374	1550	8.86E-10	Organic substance metabolic process
GO:0009058	BP	202	524	2.03E-09	Biosynthetic process
GO:0044249	BP	191	463	2.36E-09	Cellular biosynthetic process
GO:0044238	BP	352	1473	1.31E-08	Primary metabolic process
GO:0032991	CC	210	534	1.86E-08	Macromolecular complex
GO:0009059	BP	158	347	3.76E-08	Macromolecule biosynthetic process
GO:0019538	BP	241	853	1.27E-05	Protein metabolic process
GO:0072521	BP	39	74	0.000214	Purine-containing compound metabolic process

GoSeq was used to identify enriched GO terms among the downregulated unigenes. All genes were weighted by length and enrichment was determined using an FDR-corrected *P*-value of 0.05. BP, biological processes; MF, molecular function; CC, cellular complex. All GO enrichment results can be found in Table S2.

in viruliferous mites, including unigenes encoding proteins of the following pathways: RNA transport via the nuclear export complex, translation initiation elongation factor proteins, protein processing in the endoplasmic reticulum, ubiquitin-mediated proteolysis, and protein digestion and absorption (Table S4). These data suggest that the majority of unigenes encoding enzymes and structural proteins involved in protein biosynthesis were downregulated in viruliferous mites. In tandem, unigenes encoding the majority of enzymes linked to the citric acid cycle (12) and pyruvate metabolism (10) were downregulated, along with unigenes linked to oxidative phosphorylation (62), glycolysis (18) and the pentose phosphate pathway (10) (Table S4), suggesting that energy production was likely reduced in viruliferous

mites. Downregulated unigenes derived from other KEGG pathways included those linked to lysosome (42 unigenes), the cell cycle (4), starch and sucrose metabolism (8), regulation of actin cytoskeleton (8), glutathione metabolism (17), calcium signalling (14), peroxisome (10), fatty acid degradation (5), fatty acid elongation (2) and endocytosis (8) (Table S4). Few pathways were impacted with upregulated unigenes; however, three unigenes encoding enzymes involved in the synthesis of glycosphingolipids were upregulated in viruliferous mites, along with two unigenes encoding GNS1/SUR4 proteins (Table S4). GNS1/SUR4 enzymes can produce precursors for sphingolipid biosynthesis.

Acquisition of WSMV was associated with a large-scale downregulation of unigenes encoding enzymes and proteins

Table 7. Enriched gene ontology (GO) terms observed in unigenes that were upregulated in *Wheat streak mosaic virus* viruliferous wheat curl mites

Term	Ontology	No. of downregulated unigenes	No. of assigned unigenes	FDR corrected P-value	Description
GO:0042302	MF	10	47	1.94E-06	Structural constituent of cuticle
GO:0005506	MF	9	61	0.001677	Iron ion binding
GO:0020037	MF	8	60	0.00963	Heme binding
GO:0046906	MF	8	60	0.00963	Tetrapyrrole binding
GO:0006629	BP	11	124	0.012473	Lipid metabolic process

GoSeq was used to identify enriched GO terms among the downregulated unigenes. All genes were weighted by length and enrichment was determined using an FDR-corrected *P*-value of 0.05. BP, biological processes; MF, molecular function.

Table 8. Enriched Pfam domains in downregulated unigenes

Accession	PFAM_name	PFAM description	Total no. of unigenes	Total no. of downregulated unigenes	FDR
PF00379.18	Chitin_bind_4	Insect cuticle protein	47	22	3.93E-06
PF08246.7	Inhibitor_I29	Cathepsin propeptide inhibitor domain (I29)	43	19	6.73E-05
PF06585.6	JHBP	Haemolymph juvenile hormone-binding protein (JHBP)	9	8	8.99E-05
PF00061.18	Lipocalin	Lipocalin/cytosolic fatty acid-binding protein family	12	9	0.000128
PF00248.16	Aldo_ket_red	Aldo/keto reductase family	12	9	0.000128
PF02798.15	GST_N	Glutathione S-transferase, N-terminal domain	16	10	0.00022
PF00467.24	KOW	KOW motif	16	10	0.00022
PF00112.18	Peptidase_C1	Papain family cysteine protease	67	22	0.000559
PF03723.9	Hemocyanin_C	Haemocyanin, ig-like domain	9	7	0.00065
PF00240.18	Ubiquitin	Ubiquitin family	24	11	0.002232
PF00736.14	EF1_GNE	EF-1 guanine nucleotide exchange domain	8	6	0.002803
PF14560.1	Ubiquitin_2	Ubiquitin-like domain	14	8	0.002803
PF03392.8	OS-D	Insect pheromone-binding family, A10/OS-D	6	5	0.004589
PF11976.3	Rad60-SLD	Ubiquitin-2-like Rad60 SUMO-like	15	8	0.004589
PF05193.16	Peptidase_M16_C	Peptidase M16 inactive domain	12	7	0.005549
PF00026.18	Asp	Eukaryotic aspartyl protease	16	8	0.007471
PF00372.14	Hemocyanin_M	Haemocyanin, copper-containing domain	7	5	0.011572
PF01849.13	NAC	NAC domain	7	5	0.011572
PF00578.16	AhpC-TSA	AhpC/TSA family	14	7	0.015625
PF03953.12	Tubulin_C	Tubulin C-terminal domain	11	6	0.017497
PF00213.13	OSCP	ATP synthase delta (OSCP) subunit	5	4	0.017497
PF02772.11	S-AdoMet_synt_M	S-adenosylmethionine synthetase, central domain	5	4	0.017497
PF00956.13	NAP	Nucleosome assembly protein (NAP)	5	4	0.017497
PF00128.19	Alpha-amylase	Alpha amylase, catalytic domain	8	5	0.018002
PF00210.19	Ferritin	Ferritin-like domain	8	5	0.018002
PF10587.4	EF-1_beta_acid	Eukaryotic elongation factor 1 beta central acidic region	8	5	0.018002
PF00012.15	HSP70	Hsp70 protein	17	7	0.038901
PF13881.1	Rad60-SLD_2	Ubiquitin-2 like Rad60 SUMO-like	13	6	0.038901
PF00675.15	Peptidase_M16	Insulinase (peptidase family M16)	13	6	0.038901
PF00464.14	SHMT	Serine hydroxymethyltransferase	6	4	0.038901

Only one enriched term was associated with the upregulated unigenes (PF00379.18 insect cuticle protein; P -value=5.77E-7).

linked to immunity and stress responses, including several proteins involved in canonical detoxification reactions (Table S4). These included proteins containing AhpC/TSA domains (7 unigenes), which can act as antioxidants, aldehyde dehydrogenases (2), antifungal peptide (1), aspartyl proteases (8), catalases (2), CYP450s (4), superoxide dismutases (2), glutaredoxins (2), glutathione peroxidases (4), immunoglobulin domain protein (1), LRRs (1), lipases (5), lipocalins (9), papain cysteine proteinases (22) and thioredoxins (8). In contrast, only a handful of unigenes encoding stress-responsive proteins were upregulated, which included carboxylesterase (1), CYP450s (5), carbonic anhydrase (1), HSP20s (2), immunoglobulin domain protein (1), lipases (3), animal haem peroxidases (3) and trypsins (6) (Table S4). The expression levels of several unigenes encoding enzymes related to digestion and nutrient acquisition were affected in viruliferous mites. Downregulated unigenes encoding proteins involved in these processes were annotated as ABC transporters (2), α -amylase (5), ferritin-like proteins involved in iron storage (5), major facilitator

superfamily transporters (4), melibiase (1) and uricase (1) (Table S4). As was the case with unigenes linked to stress response, fewer unigenes associated with digestion and nutrient acquisition were upregulated in viruliferous mites. Such unigenes included amino acid permease (1) and major facilitator superfamily transporter (1). Additionally, some of the most strongly upregulated unigenes in WSMV-exposed mites included unigenes encoding CYP450s, cuticle proteins, dual oxidase maturation factors and a putative defence protein (Fig. 6). In contrast, some of the most strongly downregulated unigenes included those encoding apolipoprotein, trypsin, lysosomal aspartic protease, trypsins, JHBPs, cathepsins, cytochrome C oxidase and GSTs (Fig. 6).

Validation of differential expression by RT-qPCR

Differential expression of five each of up- and downregulated unigenes identified through RNA-seq were validated. RT-qPCR reactions were performed on the same pools of RNA samples that were used for the RNA-seq study. Log₂ fold change values ($\Delta\Delta C_q$, viruliferous–aviruliferous)

Table 9. Enriched KEGG orthology (KO) terms in downregulated unigenes

Term	Description	FDR
K17751	MYH6_7; myosin heavy chain 6/7	0.00046
K00011	AKR1B; aldehyde reductase [EC:1.1.1.21]	0.00046
K03098	APOD; apolipoprotein D and lipocalin family protein	0.00105
K04097	HPGDS; prostaglandin-H2 D-isomerase/glutathione transferase [EC:5.3.99.2 2.5.1.18]	0.00105
K01365	CTSL; cathepsin L [EC:3.4.22.15]	0.00227
K17732	PMPCB, MAS1; mitochondrial-processing peptidase subunit beta [EC:3.4.24.64]	0.00227
K02132	ATPeF1A, ATP5A1, ATP1; F-type H ⁺ -transporting ATPase subunit alpha	0.00227
K02898	RP-L26e, RPL26; large subunit ribosomal protein L26e	0.00227
K02973	RP-S23e, RPS23; small subunit ribosomal protein S23e	0.00227
K02989	RP-S5e, RPS5; small subunit ribosomal protein S5e	0.00227
K01379	CTSD; cathepsin D [EC:3.4.23.5]	0.0037
K13247	CRYL1; L-gulonate 3-dehydrogenase [EC:1.1.1.45]	0.00594
K03231	EEF1A; elongation factor 1-alpha	0.0073
K05863	SLC25A4S, ANT; solute carrier family 25 (mitochondrial adenine nucleotide translocator), member 4/5/6/31	0.0073
K14753	RACK1; guanine nucleotide-binding protein subunit beta-2-like 1 protein	0.0073
K00789	metK; S-adenosylmethionine synthetase [EC:2.5.1.6]	0.0073
K02137	ATPeFOO, ATP5O, ATP5; F-type H ⁺ -transporting ATPase subunit O	0.0073
K02889	RP-L21e, RPL21; large subunit ribosomal protein L21e	0.0073
K02920	RP-L36e, RPL36; large subunit ribosomal protein L36e	0.0073
K02934	RP-L6e, RPL6; large subunit ribosomal protein L6e	0.0073
K02966	RP-S19e, RPS19; small subunit ribosomal protein S19e	0.0073
K03283	HSPA1_8; heat shock 70 kDa protein 1/8	0.00741
K00615	E2.2.1.1, tktA, tktB; transketolase [EC:2.2.1.1]	0.00741
K07374	TUBA; tubulin alpha	0.01429
K09640	TMPRSS9; transmembrane protease serine 9 [EC:3.4.21.-]	0.01429
K00600	glyA, SHMT; glycine hydroxymethyltransferase [EC:2.1.2.1]	0.01429
K01363	CTSB; cathepsin B [EC:3.4.22.1]	0.01429
K02896	RP-L24e, RPL24; large subunit ribosomal protein L24e	0.01429
K09580	PDIA1, P4HB; protein disulfide isomerase A1 [EC:5.3.4.1]	0.01429
K00432	gpx; glutathione peroxidase [EC:1.11.1.9]	0.02907
K03257	EIF4A; translation initiation factor 4A	0.02928
K01623	ALDO; fructose biphosphate aldolase, class I [EC:4.1.2.13]	0.02928
K02865	RP-L10Ae, RPL10A; large subunit ribosomal protein L10Ae	0.02928
K02998	RP-SAe, RPSA; small subunit ribosomal protein SAe	0.02928

No enriched terms with FDR <0.05 were identified in the upregulated unigenes.

calculated from relative expression determined by RT-qPCR were plotted along with Pearson's correlation between the RNA-seq expression values and the RT-qPCR ΔCq values for each sample (Fig. 7). With the exception of data derived from the unigene encoding the ryanodine receptor, the RT-qPCR and RNA-seq data were highly correlated.

DISCUSSION

Viruses can modulate a variety of biochemical pathways and behaviours to promote vector colonization and prolong association with vectors. This phenomenon, termed adaptive manipulation, often results in rapid vector population expansions and provide viruses with obvious selective advantages [22–25, 61–63]. To study the impact of WSMV on the WCM, we generated a *de novo*-assembled

transcriptome of WCM and profiled global changes in gene expression that occurred after the acquisition of WSMV. A majority of unigenes were downregulated in viruliferous mites, with only a few unigenes being upregulated (Table S4). Protein family and metabolic pathway enrichment analyses revealed that the majority of downregulated unigenes coded for enzymes and proteins linked to stress response, immunity and development (such as JHBP), HSPs, GSTs, trypsin, aspartyl proteases, cuticle proteins and cathepsin propeptide inhibitors. Overall, the downregulation of these unigenes may inhibit WCM immune response to prolong viral association and/or alter WCM development to expedite population expansion, both of which could enhance viral transmission or vector spread, or both.

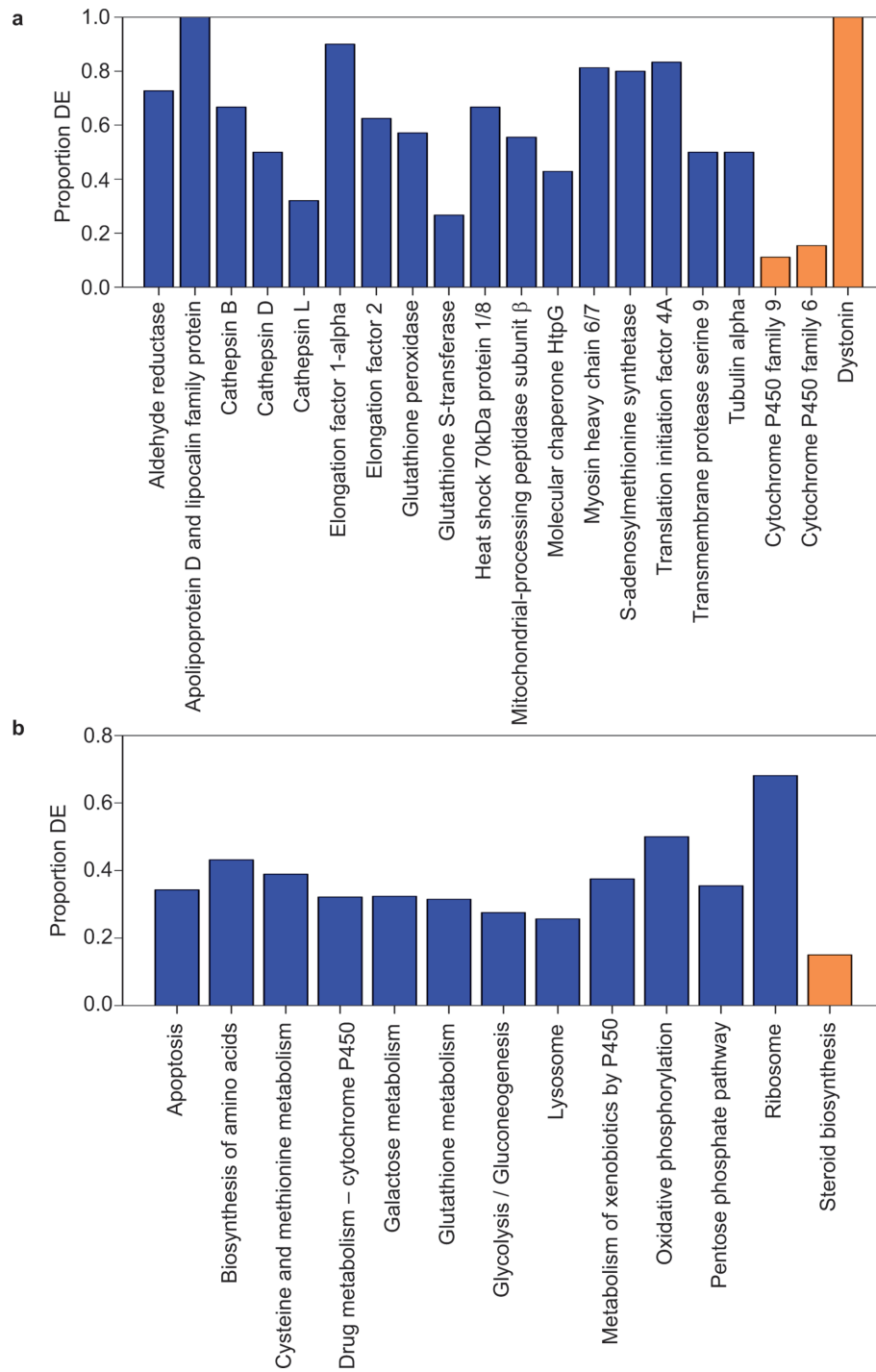


Fig. 5. Select enriched KEGG orthology (KO) terms and KEGG pathways in differentially expressed wheat curl mite unigenes under the influence of *Wheat streak mosaic virus*. (a) Graphical representation of select significantly enriched (P -value <0.05) KO categories in differentially expressed unigenes after WSMV acquisition. (b) Graphical representation of select significantly enriched (P -value <0.05) KEGG pathways in differentially expressed unigenes after WSMV acquisition. In both (a) and (b), the bars depict the percentage of unigenes assigned to each category that were downregulated (blue) and upregulated (orange) in viruliferous mites relative to the total number of unigenes in the transcriptome.

Table 10. Enriched KEGG pathways in downregulated unigenes

Pathway	FDR
Ribosome	0.0000
Oxidative phosphorylation	0.0000
Biosynthesis of amino acids	0.0000
Carbon metabolism	0.0000
Apoptosis	0.0000
Phagosome	0.0000
Methane metabolism	0.0001
Lysosome	0.0001
Pentose and glucuronate interconversions	0.0001
Glutathione metabolism	0.0002
Fructose and mannose metabolism	0.0010
Metabolism of xenobiotics by cytochrome P450	0.0014
Cysteine and methionine metabolism	0.0019
Glyoxylate and dicarboxylate metabolism	0.0020
Arachidonic acid metabolism	0.0020
Drug metabolism – cytochrome P450	0.0045
Glycine, serine and threonine metabolism	0.0059
Glycolysis/gluconeogenesis	0.0084
Galactose metabolism	0.0086
PPAR signalling pathway	0.0116
Autophagy – animal	0.0165
Pentose phosphate pathway	0.0165

Pfams associated with ribosomal proteins were removed from the table. No enriched terms with FDR <0.05 were identified in the upregulated unigenes.

Development and reproduction

The expression levels for unigenes coding for enzymes linked to development and reproduction were extensively impacted on by the acquisition of WSMV, including many with potential roles in hormone biosynthesis, binding and metabolism. Arthropod development, moulting and sexual maturity are tightly regulated by several hormonal molecules, including ecdysteroids [64]. In this study, the downregulation of unigenes associated with steroid biosynthesis pathway, such as hydroxyl- δ -5-steroid dehydrogenase and cholesteryl ester hydrolase (one upregulated and one downregulated), could be linked to reduced synthesis of ecdysteroids, suggesting that infection with WSMV may alter the ecdysis and reproductive maturity of WCMs. Signalling is the next most important step downstream to hormonal biogenesis. The normal activity of ecdysteroids is mediated by juvenile hormone (JH), which is involved in maintaining the larval stage and also has roles in regulating metamorphosis and reproduction [65, 66]. JHBP is the first and the most important signal transmitter of JH [67] and also protects JH against haemocoelic esterase-mediated hydrolysis [68]. More than 99.8 % of JH exists as JH–JHBP complexes in the haemolymph [69]. Thus, the downregulation of eight unigenes encoding JHBP may result in JH depletion, which may expedite development [70], resulting in precocious sexual maturity and promoting population

expansion, which was previously observed in WSMV-exposed mites [71, 72].

Also related to growth and development, a majority of unigenes encoding cuticular proteins, which provide structural stability to the epicuticle and cuticle of the chitinous exoskeleton [73], were downregulated in viruliferous mites, although a smaller number of these unigenes were upregulated. The large degree of differential expression in unigenes encoding these proteins suggests that cuticle formation and chitin metabolism were likely affected in viruliferous mites. Supporting this hypothesis, a unigene encoding GH 18 chitinase was downregulated in viruliferous mites. Additionally, lysosomal cysteine proteases called cathepsins can be involved in hydrolyzing cuticle proteins and are often linked with arthropod developmental events, including moulting and eclosion [74]. In this study, 19 unigenes encoding cathepsin inhibitors, which can suppress or delay cathepsin enzyme activity during earlier life stages [75], were exclusively downregulated in viruliferous mites, and this downregulation may expedite WCM development under the influence of WSMV.

Behaviour and signalling

Behavioural manipulation of the vector is another mechanism by which viruses can facilitate transmission [24]. In WCM, unigenes encoding numerous components of signalling pathways and components of the nervous system showed differential expression in viruliferous WCMs. For example, acetylcholinesterase (AChE) is one of the highly conserved carboxylesterases involved in terminating nerve impulses by rapidly hydrolyzing acetylcholine (ACh) in neural synapses [76]. The upregulation of a unigene encoding AChE suggests that neurotransmission may be altered in WCM under the influence of WSMV. Related to neuronal activity, a unigene encoding a ryanodine receptor was upregulated. These receptors can play pivotal roles in intracellular calcium flux between the cytosol and endoplasmic reticulum of myocytes and neurons [77] and may similarly alter neural activity in WCMs. Other unigenes encoding proteins related to cell signalling were downregulated, including unigenes encoding components of the MAPK, sphingolipid, TOL and IMD, and Rap1 signalling pathways. While the differential expression of unigenes encoding proteins associated with these pathways could have a variety of effects on the physiology of WCMs, it is possible that these expression changes could alter the behaviour of viruliferous mites, as many of these genes have been linked to increased locomotion, spermatophore acquisition and altered oviposition behaviours in other studies [78, 79].

Nutrition

Many unigenes related to digestion, nutrient uptake and storage were downregulated in viruliferous WCMs. For example, many unigenes associated with the degradation of fatty acids were downregulated in viruliferous mites. Lipids are known to play critical roles in energy homeostasis, membrane integrity and signalling [80], and the

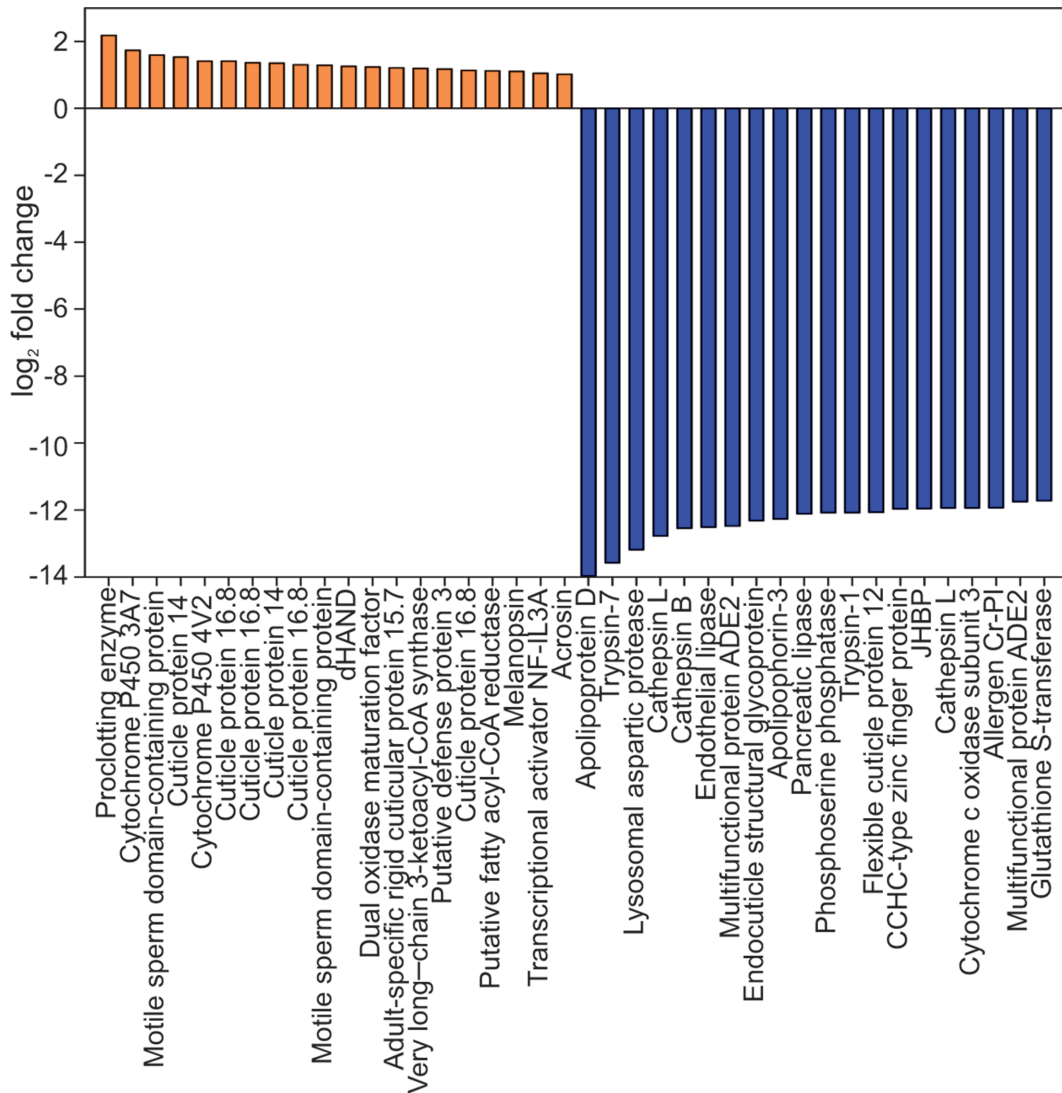


Fig. 6. Top 20 up- and downregulated unigenes in viruliferous wheat curl mites. The bars depict the log₂ fold changes in upregulated (orange) and downregulated (blue) unigenes.

downregulation of unigenes linked to lipid and fatty acid degradation may support increased rate of growth and potentially the increased demand for signal-transducing lipoproteins due to the upregulation of several signalling pathways in viruliferous mites, as discussed in the previous section. Unigenes encoding proteins that bind and sequester minerals and other nutrients were also affected in viruliferous WCMs. For example, several unigenes encoding ferritin (involved in iron uptake, transport and storage [81]), calsequestrin domains (involved in the binding and release of calcium ions [82]) and lipocalin proteins (involved in transportation of lipids, vitamins and building up energy reserves in fat bodies [83]) were all downregulated in viruliferous mites. Likewise, a unigene encoding a perilipin protein, which safeguards lipid droplets in *Drosophila* fat bodies [84], was downregulated in WSMV viruliferous mites,

suggesting that virus acquisition could change fat storage and/or more broadly influence energy metabolism in WCMs. The broad downregulation of unigenes associated with nutrition, glycolysis, gluconeogenesis and the citric acid cycle suggests that energy production could be significantly altered in WSMV viruliferous WCMs. Plausibly, resources were redirected for other processes, such as development, and were not available for anabolism or storage.

Ubiquitination

The acquisition of WSMV was associated with lower expression levels of unigenes encoding enzymes and proteins related to apoptosis and ubiquitination. Ubiquitins are regulatory proteins that are covalently attached post-translationally to damaged, overexpressed and misfolded proteins targeted for degradation via proteasomes [85, 86]. As an

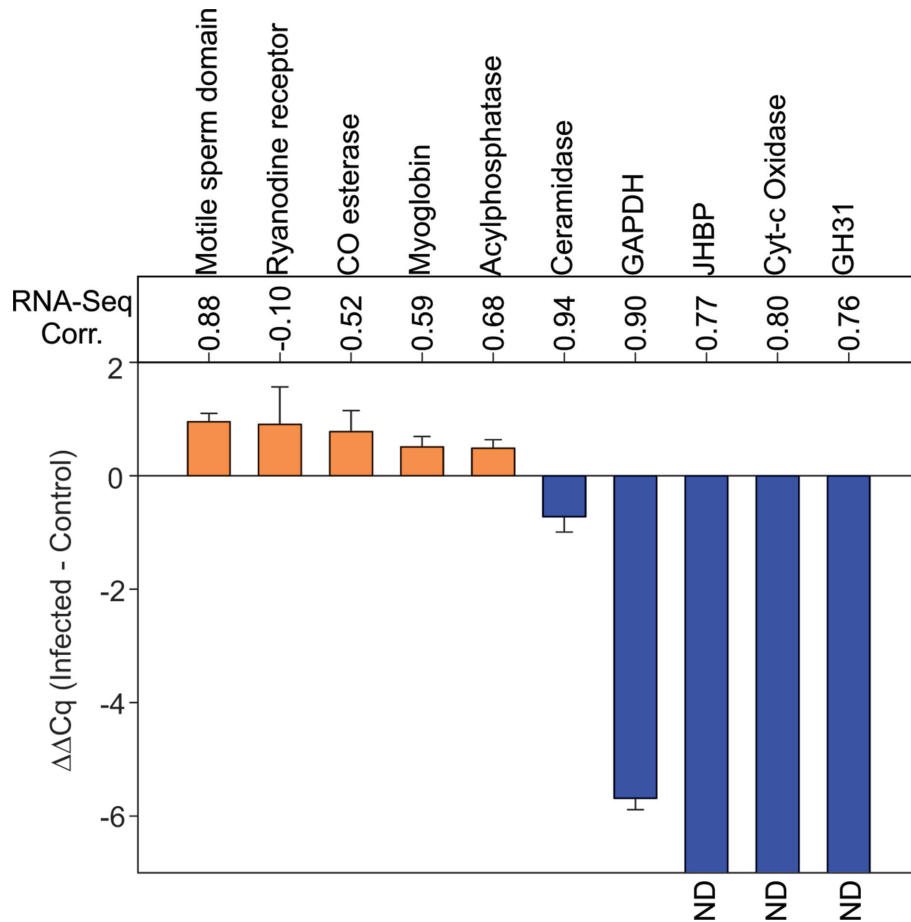


Fig. 7. Reverse-transcription real-time PCR (RT-qPCR) validation of selected wheat curl mite unigenes. Five each of up- and downregulated unigenes observed in the viruliferous mites detected in the RNA-seq analysis were validated using RT-qPCR. Technical duplicates of the same three biological replicates of viruliferous (infected) and control mites that were used in the RNA-seq analysis were used for RT-qPCR validation. The $\Delta\Delta C_t$ method was used with actin and G6PDH as internal reference genes. ND indicates that expression was not detected in viruliferous mite samples. The bars represent standard error. Pearson's correlations between the RT-qPCR and RNA-seq data were also calculated for each unigene.

additional factor, unigenes encoding translocons and secY translocases, and membrane proteins, which typically maintain ER–cytosol traffic [87], were downregulated in viruliferous mites. Overall, many other unigenes encoding enzymes related to protein biosynthesis were downregulated in viruliferous WCMs, consistent with lowered rates of ubiquitination and protein degradation by 26S proteasomes. In other virus–vector systems, ubiquitination, proteolysis and protein biosynthesis are common targets of viral manipulation that can result in enhanced viral nucleoprotein accumulation and titre [88]. Supporting this hypothesis, other factors that could be linked to viral replication and virus–vector associations were observed in viruliferous WCMs. For example, a unigene encoding serpin (serine protease inhibitor) was highly downregulated in viruliferous WCMs, which may promote activity of the WSMV P1 serine protease, required for posttranslational polyprotein processing during viral infection [89, 90].

Immunity and stress response

Overall, the majority of differentially expressed unigenes that encode proteins related to stress response and immunity were downregulated in viruliferous WCMs. For example, several unigenes encoding apolipoprotein-III (apoLp-III) were downregulated. In addition to its role in lipid transport, apoLp-III can act as a pattern recognition protein that can bind to pathogen-associated molecular patterns (PAMPs) to induce innate immunity in other systems [91]. Additionally, insect cells often overexpress apoLp-III during programmed cell death [92]. Four unigenes encoding Reeler domain proteins were also differentially expressed in viruliferous WCMs (three downregulated and one upregulated) as compared to aviruliferous WCMs. Reeler domain proteins have also been linked to defence responses and can possess nonspecific affinity towards haemocytes and PAMPs, resulting in nodulation, which is one of the most important arthropod immune responses [93, 94]. Other unigenes, such

as one encoding cupins and one encoding an LRR, were downregulated, indicative of modulated defence response in viruliferous mites. In addition, the majority of the proteolytic enzymes currently identifiable in the WCM transcriptome were extensively downregulated in viruliferous WCM. WSMV viruliferous mites also showed greater downregulation of unigenes encoding CYP450s, although five copies showed upregulation. In addition to roles in stress response and detoxification, CYP450s have been shown to play roles in membrane synthesis, cuticular biosynthesis and hormone metabolism [95, 96]. Although the downregulation of CYP450s in viruliferous mites was consistent with the overall downregulation of stress-related transcripts observed after viral acquisition, it is possible that some of these CYP450 unigenes were associated with other metabolic processes, such as cuticular and/or membrane synthesis. Overall, the downregulation of unigenes encoding canonical proteins and enzymes linked to immunity and stress responses suggests that viral acquisition potentially modulates innate immune responses in WCMs.

Oxidative stress

A single unigene encoding dual oxidase maturation factor (DUOX) was upregulated in WSMV viruliferous WCMs. DUOX is an NADPH-oxidative transmembrane enzyme that is involved in reactive oxygen species (ROS) generation and has previously been shown to be involved in antimicrobial defence mechanisms [97]. In tandem, the coordinated downregulation of unigenes that encode ROS-mitigating enzymes in viruliferous WCMs, such as superoxide dismutase, peroxiredoxin, and glutathione peroxidases, uricase [98], thioredoxin-like domain-containing proteins, alkyl hydroperoxidases and catalases, is suggestive of greater oxidative stress after viral acquisition in this system. In addition, 11 unigenes encoding GSTs were extensively downregulated in viruliferous mites, which also can serve as inhibitors of oxidative stress. This finding is in contrast to the findings of previous studies on *Bombyx mori* [99], where infections with *B. mori* nuclear polyhedrosis virus (*BmNPV*) and *B. mori* densovirus (*BmDENV*) were associated with an upregulation of GSTs. Why the responses of GSTs differ in the WCM-WSMV and *B. mori*-virus systems is not known; however, both *BmNPV* and *BmDENV* are pathogens to their hosts, whereas WSMV is not known to cause disease in WCMs. Additional factors, such as differences in the ages and nutritional statuses of the hosts used in the studies, or exposure to pesticides or other abiotic stresses, can also influence the expression levels of GSTs and could also account for the differences observed between the two studies.

Conclusions

This study represents the first transcriptome-level analysis of the WCM and contributes significantly to our understanding of the influence of WSMV on WCM biology. Transcriptomic analyses to study virus–vector interactome are essential to understand the cascade of events starting from WSMV acquisition through transmission.

Comparative analyses of WCM transcriptomes from WSMV viruliferous and aviruliferous mites suggested that transcriptional changes in viruliferous mites may inhibit WCM immune response, and growth and development. The downregulation of unigenes coding for proteins and enzymes linked to stress response, detoxification and immunity could reduce innate immune responses in viruliferous mites and prolong viral association with WCMs. Furthermore, the downregulation of unigenes coding for enzymes involved in hormone biosynthesis and JHBPs may expedite WCM development to enhance population expansion, as has been previously observed [71, 72]. In addition to studying differential expression under WSMV influence, this study also provides a catalogue of annotated WCM unigenes that can facilitate functional genomics studies and may help to devise novel strategies of vector control as a component of WSMV disease management.

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Conflicts of interest

The authors declare that there are no conflicts of interest.

Ethical statement

This article does not contain any studies involving human participants or animals.

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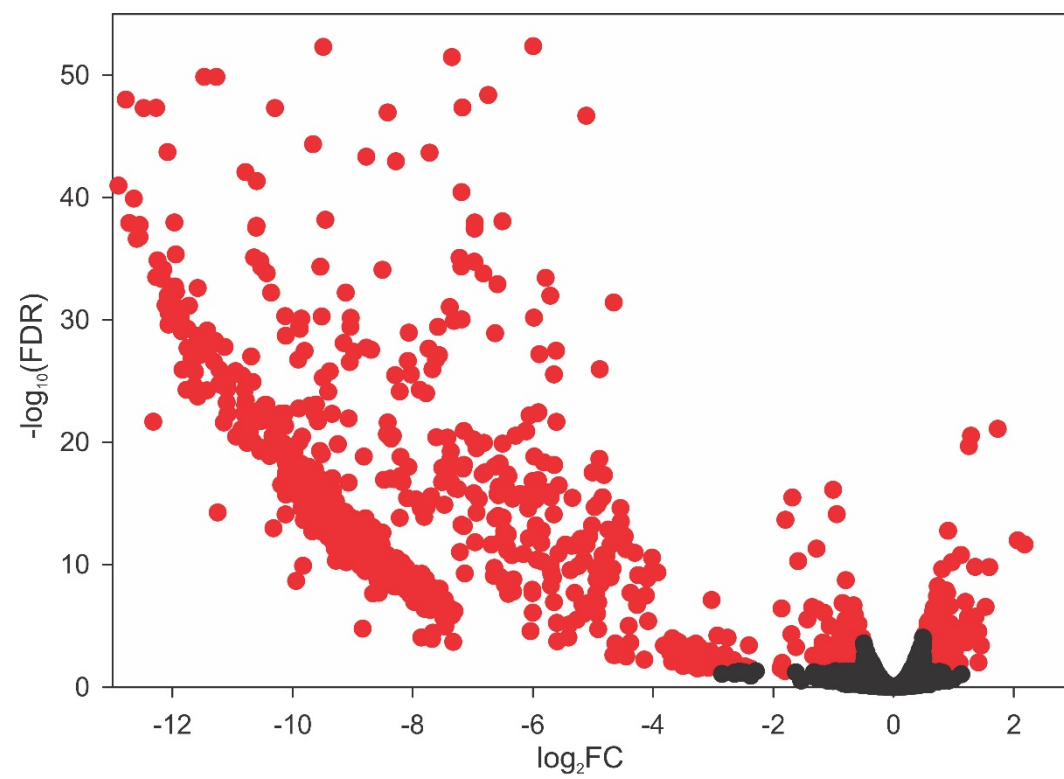


Fig. S1. Volcano plot of expressed unigenes in *Wheat streak mosaic virus* (WSMV) viruliferous and aviruliferous wheat curl mites (WCM). Differentially expressed unigenes (1,020; red) were identified with an FDR cutoff of ≤ 0.05 and a $\log_2FC \geq 0.5$. Non-differentially expressed unigenes (6,271) are shown in black.

Table S1: Primers used for qRT-PCR of representative candidates to validate RNA-Seq analysis of wheat curl mite transcriptome in response to WSMV

Unigene ID	Gene	Primer (5' to 3') forward / reverse
TR6223 c6_g1	Actin	GCCACAACAATCACAACCC/ CGCGATTCCGGATCAGCGG
TR2714 c0_g1	Glucose-6-phosphate dehydrogenase (G6p	ATGTATGTATTTGTACATTTGG/ CCTGTTCTTTGTGTAAGG
TR6258 c1_g2	Glyceraldehyde 3-phosphate dehydrogen	TCCAAGATCGGAATTAACGG/ GGTAGTGAAGACACCACTGG
TR6295 c13_g1	Ceramidase	GGCATTGCCGACATAACAGG/ GAGCATGTAAGTAACC
TR1462 c0_g1	Cyt-C Oxidase	CCCTACTGTAAGAACAGACC/ CTTGAATACGAATACAGAGCC
TR4699 c0_g2	Glycosyl hydrolase family 31 gene (GH31)	AGTCGCCGGCCAGATTTGC/ TGTCGTCGATCTCGAGCTGG
TR6354 c5_g1	Juvenile Hormone Binding Protein (JHBP)	GCGGCACACAGAAACAAACAG/ GATGTTAAGGTCCGAGAACTGC
TR5487 c1_g1	Acylphosphatase	AGCGCCAAATCCAGGAGCC/ TTATCTTATTCTAAAGTCGC
TR6320 c2_g1	Carboxylesterase	ACATCAGCATCAGCATCATC/ TGAGAAAGGTGTATAGGTGC
TR6383 c4_g1	Cral-trio	CAACGATTGCAAAACGGGCTC/ CCGTTTCGGTTGCAACCGGC
TR6108 c11_g2	Globin-1	ATGGCTTTGAGCGAGGCCGACG/ TTGGTGACCACGAACAAGG
TR4132 c0_g1	Ryanodie receptor	CGGTGGAGGGGCCACTGGTCC/ GCGAGCTTGACCGTTCCAAAGG

Table S2: GO Enrichment Results

GOs Enriched in Downregulated DEGs					GOs enriched in Upregulated DEGs						
category	ontology	numDEInCa	numInCat	FDR	term	category	ontology	numDEInCa	numInCat	FDR	term
GO:0006754	BP	18	24	0.0209443	ATP biosynthetic process	GO:0042302	MF	10	47	1.94E-06	structural constituent of cuticle
GO:0015991	BP	10	19	0.053052	ATP hydrolysis coupled proton transport	GO:0005506	MF	9	61	0.0016772	iron ion binding
GO:0046034	BP	22	31	0.0054352	ATP metabolic process	GO:0020037	MF	8	60	0.00963	heme binding
GO:0015986	BP	18	24	0.0209443	ATP synthesis coupled proton transport	GO:0046906	MF	8	60	0.00963	tetrapyrrole binding
GO:0008150	BP	535	3016	0.0005353	biological_process	GO:0006629	BP	11	124	0.0124731	lipid metabolic process
GO:0009058	BP	202	524	2.03E-09	biosynthetic process						
GO:1901137	BP	27	66	0.0497099	carbohydrate derivative biosynthetic process						
GO:1901135	BP	46	146	0.0234577	carbohydrate derivative metabolic process						
GO:0005975	BP	40	131	0.0059463	carbohydrate metabolic process						
GO:0098655	BP	28	47	0.0019922	cation transmembrane transport						
GO:0044249	BP	191	463	2.36E-09	cellular biosynthetic process						
GO:0034645	BP	158	322	8.86E-10	cellular macromolecule biosynthetic process						
GO:0044237	BP	268	1171	0.00082	cellular metabolic process						
GO:0009987	BP	309	1494	0.001426	cellular process						
GO:0044267	BP	159	529	0.0027708	cellular protein metabolic process						
GO:0015074	BP	2	3	0.0452803	DNA integration						
GO:0015988	BP	10	19	0.053052	energy coupled proton transmembrane transport, against electrochemical gradient						
GO:0015985	BP	18	24	0.0209443	energy coupled proton transport, down electrochemical gradient						
GO:1901659	BP	25	44	0.0252596	glycosyl compound biosynthetic process						
GO:1901657	BP	37	74	0.0056436	glycosyl compound metabolic process						
GO:1902600	BP	28	43	0.0009853	hydrogen ion transmembrane transport						
GO:0006818	BP	30	45	0.0006111	hydrogen transport						
GO:0098662	BP	28	47	0.0019922	inorganic cation transmembrane transport						
GO:0098660	BP	28	47	0.0019922	inorganic ion transmembrane transport						
GO:0034220	BP	28	60	0.0200742	ion transmembrane transport						
GO:0009059	BP	158	347	3.76E-08	macromolecule biosynthetic process						
GO:0043170	BP	267	1173	0.0031632	macromolecule metabolic process						
GO:0008152	BP	469	1962	4.30E-14	metabolic process						
GO:0032787	BP	18	35	0.0452803	monocarboxylic acid metabolic process						
GO:0015672	BP	30	61	0.0093801	monovalent inorganic cation transport						
GO:0055086	BP	43	113	0.0068775	nucleobase-containing small molecule metabolic process						
GO:0009163	BP	25	44	0.0252596	nucleoside biosynthetic process						
GO:0009116	BP	37	74	0.0056436	nucleoside metabolic process						
GO:0009124	BP	20	33	0.0300997	nucleoside monophosphate biosynthetic process						
GO:0009123	BP	24	40	0.0066678	nucleoside monophosphate metabolic process						
GO:0006753	BP	38	99	0.0134533	nucleoside phosphate metabolic process						
GO:0009142	BP	21	28	0.0079491	nucleoside triphosphate biosynthetic process						
GO:0009141	BP	31	49	0.0014631	nucleoside triphosphate metabolic process						
GO:0009117	BP	38	98	0.0127394	nucleotide metabolic process						
GO:1901576	BP	197	479	4.26E-10	organic substance biosynthetic process						
GO:0071704	BP	374	1550	8.86E-10	organic substance metabolic process						
GO:1901566	BP	30	92	0.0523246	organonitrogen compound biosynthetic process						
GO:0019637	BP	39	127	0.0445914	organophosphate metabolic process						
GO:0055114	BP	74	288	0.0190219	oxidation-reduction process						
GO:0044238	BP	352	1473	1.31E-08	primary metabolic process						
GO:0019538	BP	241	853	1.27E-05	protein metabolic process						
GO:0015992	BP	30	45	0.0006111	proton transport						
GO:0042451	BP	25	40	0.0101366	purine nucleoside biosynthetic process						
GO:0042278	BP	36	62	0.0005921	purine nucleoside metabolic process						
GO:0009127	BP	20	29	0.0093801	purine nucleoside monophosphate biosynthetic process						
GO:0009126	BP	24	36	0.0019922	purine nucleoside monophosphate metabolic process						
GO:0009145	BP	21	28	0.0079491	purine nucleoside triphosphate biosynthetic process						
GO:0009144	BP	31	48	0.0008959	purine nucleoside triphosphate metabolic process						
GO:0006164	BP	24	45	0.0242488	purine nucleotide biosynthetic process						
GO:0006163	BP	35	67	0.0010383	purine nucleotide metabolic process						
GO:0046129	BP	25	40	0.0101366	purine ribonucleoside biosynthetic process						
GO:0046128	BP	36	62	0.0005921	purine ribonucleoside metabolic process						
GO:0009168	BP	20	29	0.0093801	purine ribonucleoside monophosphate biosynthetic process						
GO:0009167	BP	24	36	0.0019922	purine ribonucleoside monophosphate metabolic process						
GO:0009206	BP	21	28	0.0079491	purine ribonucleoside triphosphate biosynthetic process						
GO:0009205	BP	31	48	0.0008959	purine ribonucleoside triphosphate metabolic process						
GO:0009152	BP	23	42	0.0358981	purine ribonucleotide biosynthetic process						
GO:0009150	BP	34	64	0.0015429	purine ribonucleotide metabolic process						
GO:0072522	BP	28	52	0.0053624	purine-containing compound biosynthetic process						
GO:0072521	BP	39	74	0.0002143	purine-containing compound meta						Could pick one of these categories related to purine metabolism and mention that several categories related to this process were enriched.
GO:0042455	BP	25	44	0.0252596	ribonucleoside biosynthetic process						
GO:0009119	BP	36	66	0.0012269	ribonucleoside metabolic process						
GO:0009156	BP	20	33	0.0300997	ribonucleoside monophosphate biosynthetic process						
GO:0009161	BP	24	40	0.0066678	ribonucleoside monophosphate metabolic process						
GO:0009201	BP	21	28	0.0079491	ribonucleoside triphosphate biosynthetic process						
GO:0009199	BP	31	48	0.0008959	ribonucleoside triphosphate metabolic process						
GO:0009260	BP	23	46	0.0741131	ribonucleotide biosynthetic process						
GO:0009259	BP	34	68	0.0031552	ribonucleotide metabolic process						
GO:0046390	BP	23	46	0.0741131	ribose phosphate biosynthetic process						
GO:0019693	BP	37	78	0.0019922	ribose phosphate metabolic process						
GO:0044711	BP	39	150	0.0810334	single-organism biosynthetic process						
GO:0044712	BP	21	57	0.0977539	single-organism catabolic process						
GO:0044710	BP	143	655	0.00082	single-organism metabolic process						
GO:0044699	BP	188	1061	0.0084868	single-organism process						
GO:0044281	BP	71	237	0.0034864	small molecule metabolic process						
GO:0006412	BP	139	216	1.29E-12	translation						
GO:0006414	BP	18	26	0.0148561	translational elongation						
GO:0044444	CC	193	405	1.64E-10	cytoplasmic part						
GO:0005622	CC	102	260	0.013371	intracellular						
GO:0043232	CC	146	253	5.29E-11	intracellular non-membrane-bounded organelle						
GO:0043229	CC	160	479	0.0019922	intracellular organelle						
GO:0044424	CC	222	835	0.0103311	intracellular part						
GO:0030529	CC	146	240	1.43E-11	intracellular ribonucleoprotein complex						
GO:0032991	CC	210	534	1.86E-08	macromolecular complex						
GO:0043228	CC	146	253	5.29E-11	non-membrane-bounded organelle						
GO:0043226	CC	160	479	0.0019922	organelle						
GO:0033178	CC	10	18	0.0445914	proton-transporting two-sector ATPase complex, catalytic domain						
GO:0005840	CC	146	226	3.49E-13	ribosome						
GO:0015078	MF	36	57	0.0005921	hydrogen ion transmembrane transporter activity						
GO:0016787	MF	158	854	0.0510499	hydrolase activity						
GO:0022890	MF	36	103	0.062887	inorganic cation transmembrane transporter activity						
GO:0015077	MF	36	75	0.0074209	monovalent inorganic cation transmembrane transporter activity						
GO:0016491	MF	92	352	0.0179365	oxidoreductase activity						
GO:0016616	MF	19	56	0.0209443	oxidoreductase activity, acting on the CH-OH group of donors, NAD or NADP as acceptor						
GO:0070011	MF	80	304	0.0815075	peptidase activity, acting on L-amino acid peptides						
GO:0042302	MF	27	47	0.0514632	structural constituent of cuticle						

GO:0003735	MF	146	225	3.12E-13	structural constituent of ribosome
GO:0005198	MF	173	287	7.66E-14	structural molecule activity
GO:0003746	MF	11	16	0.0491896	translation elongation factor activity

Table S3: Pfam Enrichment Results

	Pfam Enriched in Downregulated DEGs					Pfam Enriched in Upregulated DEGs							
	PFAM_Name	PFAM_desc	total	downregula	pval	fd	PFAM_Name	PFAM_desc	total	Upregulate	pval	fd	
PF00379	Chitin_bind_4	Insect cuticle protein	47	22	6.51E-09	3.93E-06	PF00379	Chitin_bind_4	Insect cuticle protein	47	10	5.77E-08	3.49E-05
PF08246	Inhibitor_I29	Cathepsin propeptide inhibit	43	19	2.23E-07	6.73E-05	PF00067	p450	Cytochrome P450	35	5	0.000975	0.29456
PF06585	JHBP	Haemolymph juvenile hormo	9	8	4.46E-07	8.99E-05	PF00487	FA_desaturase	Fatty acid desaturase	11	3	0.001571	0.316357
PF00061	Lipocalin	Lipocalin / cytosolic fatty-aci	12	9	1.06E-06	0.000128	PF00246	Peptidase_M14	Zinc carboxypeptidase	14	3	0.0033	0.408534
PF00248	Aldo_ket_red	Aldo/keto reductase family	12	9	1.06E-06	0.000128	PF07993	NAD_binding_4	Male sterility protein	15	3	0.004058	0.408534
PF02798	GST_N	Glutathione S-transferase, N	16	10	3.33E-06	0.00022	PF03098	An_peroxidase	Animal haem peroxidase	15	3	0.004058	0.408534
PF13417	GST_N_3	Glutathione S-transferase, N	16	10	3.33E-06	0.00022	PF01370	Epimerase	NAD dependent epimerase/	17	3	0.00587	0.49406
PF00467	KOW	KOW motif	16	10	3.33E-06	0.00022	PF01607	CBM_14	Chitin binding Peritrophin-A	34	4	0.006544	0.49406
PF00828	Ribosomal_L18e	Ribosomal protein L18e/L15	6	6	3.65E-06	0.00022	PF00011	HSP20	Hsp20/alpha crystallin famil	7	2	0.009631	0.580119
PF08212	Lipocalin_2	Lipocalin-like domain	6	6	3.65E-06	0.00022	PF00151	Lipase	Lipase	21	3	0.010758	0.580119
PF01112	Peptidase_C1	Papain family cysteine prote	67	22	1.02E-05	0.000559	PF00650	CRAL_TRIO	CRAL/TRIO domain	21	3	0.010758	0.580119
PF03723	Hemocyanin_C	Hemocyanin, ig-like domain	9	7	1.29E-05	0.00065	PF07716	bZIP_2	Basic region leucine zipper	8	2	0.012655	0.580119
PF03722	Hemocyanin_N	Hemocyanin, all-alpha doma	5	5	2.95E-05	0.00137	PF00571	CBS	CBS domain	8	2	0.012655	0.580119
PF00428	Ribosomal_60s	60s Acidic ribosomal protein	10	7	3.84E-05	0.001658	PF00089	Trypsin	Trypsin	88	6	0.013446	0.580119
PF00240	ubiquitin	Ubiquitin family	24	11	5.54E-05	0.002232	PF02719	Polysacc_synt_2	Polysaccharide biosynthesis	9	2	0.016034	0.605299
PF00043	GST_C	Glutathione S-transferase, C	17	9	6.55E-05	0.002472	PF00170	bZIP_1	bZIP transcription factor	9	2	0.016034	0.605299
PF00736	EF1_GNE	EF-1 guanine nucleotide excl	8	6	8.17E-05	0.002803	PF00640	PID	Phosphotyrosine interaction	10	2	0.019753	0.701798
PF14560	Ubiquitin_2	Ubiquitin-like domain	14	8	8.35E-05	0.002803	PF03015	Sterile	Male sterility protein	12	2	0.028139	0.894533
PF03392	OS-D	Insect pheromone-binding fi	6	5	0.000159	0.004589	PF04083	Abhydro_lipase	Partial alpha/beta-hydrolyse	12	2	0.028139	0.894533
PF00164	Ribosomal_S12_S2	Ribosomal protein S12/S23	6	5	0.000159	0.004589	PF00010	HLH	Helix-loop-helix DNA-binding	31	3	0.030939	0.934366
PF11976	Rad60-SLD	Ubiquitin-2 like Rad60 SUMC	15	8	0.00016	0.004589	PF01073	3beta_HSD	3-beta hydroxysteroid dehy	13	2	0.032776	0.942708
PF05193	Peptidase_M16	Peptidase M16 inactive dom	12	7	0.000202	0.005549	PF00100	Zona_pellucida	Zona pellucida-like domain	17	2	0.053939	1
PF00026	Asp	Eukaryotic aspartyl protease	16	8	0.000285	0.007471	PF01151	ELO	GNS1/SUR4 family	23	2	0.092087	1
PF00372	Hemocyanin_M	Hemocyanin, copper contain	7	5	0.000498	0.011572							
PF00177	Ribosomal_S7	Ribosomal protein S7p/S5e	7	5	0.000498	0.011572							
PF01849	NAC	NAC domain	7	5	0.000498	0.011572							
PF00578	AhpC-TSA	AhpC/TSA family	14	7	0.000698	0.015625							
PF03953	Tubulin_C	Tubulin C-terminal domain	11	6	0.000965	0.017497							
PF00213	OSCP	ATP synthase delta (OSCP) si	5	4	0.001072	0.017497							
PF02772	S-AdoMet_synt_I	S-adenosylmethionine synth	5	4	0.001072	0.017497							
PF13409	GST_N_2	Glutathione S-transferase, N	5	4	0.001072	0.017497							
PF00573	Ribosomal_L4	Ribosomal protein L4/L1 far	5	4	0.001072	0.017497							
PF01090	Ribosomal_S19e	Ribosomal protein S19e	5	4	0.001072	0.017497							
PF01157	Ribosomal_L21e	Ribosomal protein L21e	5	4	0.001072	0.017497							
PF01158	Ribosomal_L36e	Ribosomal protein L36e	5	4	0.001072	0.017497							
PF01159	Ribosomal_L6e	Ribosomal protein L6e	5	4	0.001072	0.017497							
PF00956	NAP	Nucleosome assembly prote	5	4	0.001072	0.017497							
PF00128	Alpha-amylase	Alpha amylase, catalytic don	8	5	0.001192	0.018002							
PF00210	Ferritin	Ferritin-like domain	8	5	0.001192	0.018002							
PF10587	EF-1_beta_acid	Eukaryotic elongation factor	8	5	0.001192	0.018002							
PF01248	Ribosomal_L7Ae	Ribosomal protein L7Ae/L30	9	5	0.002409	0.035482							
PF00012	HSP70	Hsp70 protein	17	7	0.002824	0.038901							
PF13881	Rad60-SLD_2	Ubiquitin-2 like Rad60 SUMC	13	6	0.002875	0.038901							
PF00675	Peptidase_M16	Insulinase (Peptidase family	13	6	0.002875	0.038901							
PF00464	SHMT	Serine hydroxymethyltransf	6	4	0.002898	0.038901							
PF01576	Myosin_tail_1	Myosin tail	10	5	0.004327	0.056812							
PF00431	CUB	CUB domain	14	6	0.004508	0.057933							
PF00725	3HCDH	3-hydroxyacyl-CoA dehydrog	7	4	0.006098	0.070835							
PF02874	ATP-synt_ab_N	ATP synthase alpha/beta fan	7	4	0.006098	0.070835							
PF00056	Ldh_1_N	lactate/malate dehydrogena	7	4	0.006098	0.070835							
PF00255	GSHPx	Glutathione peroxidase	7	4	0.006098	0.070835							
PF02866	Ldh_1_C	lactate/malate dehydrogena	7	4	0.006098	0.070835							
PF00089	Trypsin	Trypsin	88	19	0.010594	0.118692							
PF00160	Pro_isomerase	Cyclophilin type peptidyl-prc	12	5	0.010984	0.118692							
PF00306	ATP-synt_ab_C	ATP synthase alpha/beta chi	8	4	0.011005	0.118692							
PF01479	S4	S4 domain	8	4	0.011005	0.118692							
PF00085	Thioredoxin	Thioredoxin	27	8	0.014072	0.134372							
PF00137	ATP-synt_C	ATP synthase subunit C	5	3	0.015795	0.134372							
PF01294	Ribosomal_L13e	Ribosomal protein L13e	5	3	0.015795	0.134372							
PF00252	Ribosomal_L16	Ribosomal protein L16p/L10	5	3	0.015795	0.134372							
PF00318	Ribosomal_S2	Ribosomal protein S2	5	3	0.015795	0.134372							
PF00327	Ribosomal_L30	Ribosomal protein L30p/L7e	5	3	0.015795	0.134372							
PF00380	Ribosomal_S9	Ribosomal protein S9/S16	5	3	0.015795	0.134372							
PF00416	Ribosomal_S13	Ribosomal protein S13/S18	5	3	0.015795	0.134372							
PF00838	TCTP	Translationally controlled tu	5	3	0.015795	0.134372							
PF01246	Ribosomal_L24e	Ribosomal protein L24e	5	3	0.015795	0.134372							
PF01283	Ribosomal_S26e	Ribosomal protein S26e	5	3	0.015795	0.134372							
PF01929	Ribosomal_L14e	Ribosomal protein L14	5	3	0.015795	0.134372							
PF03719	Ribosomal_S5_C	Ribosomal protein S5, C-term	5	3	0.015795	0.134372							
PF07650	KH_2	KH domain	5	3	0.015795	0.134372							
PF14543	TAXI_N	Xylanase inhibitor N-termina	5	3	0.015795	0.134372							
PF01442	Apolipoprotein	Apolipoprotein A1/A4/E don	9	4	0.017881	0.150005							
PF00091	Tubulin	Tubulin/FtsZ family, GTPase	14	5	0.022466	0.185884							
PF00456	Transketolase_N	Transketolase, thiamine dipt	6	3	0.028693	0.219371							
PF02221	E1_DerP2_DerF2	ML domain	6	3	0.028693	0.219371							
PF00333	Ribosomal_S5	Ribosomal protein S5, N-terr	6	3	0.028693	0.219371							
PF00338	Ribosomal_S10	Ribosomal protein S10p/S20	6	3	0.028693	0.219371							
PF00687	Ribosomal_L1	Ribosomal protein L1p/L10e	6	3	0.028693	0.219371							
PF01201	Ribosomal_S8e	Ribosomal protein S8e	6	3	0.028693	0.219371							
PF00153	Mito_carr	Mitochondrial carrier protei	32	8	0.038158	0.288091							
PF00006	ATP-synt_ab	ATP synthase alpha/beta fan	7	3	0.045638	0.32816							
PF01459	Porin_3	Eukaryotic porin	7	3	0.045638	0.32816							
PF13917	zf-CCHC_3	Zinc knuckle	7	3	0.045638	0.32816							
PF13019	Telomere_Sde2	Telomere stability and silenc	7	3	0.045638	0.32816							
PF03143	GTP_EFTU_D3	Elongation factor Tu C-termi	12	4	0.051847	0.368422							
PF00183	HSP90	Hsp90 protein	8	3	0.066418	0.461111							
PF08534	Redoxin	Redoxin	8	3	0.066418	0.461111							
PF00180	Iso_dh	Isoctrate/isopropylmalate d	9	3	0.090687	0.60192							
PF12763	EF-hand_4	Cytoskeletal-regulatory com	9	3	0.090687	0.60192							
PF14497	GST_C_3	Glutathione S-transferase, C	9	3	0.090687	0.60192							
PF03446	NAD_binding_2	NAD binding domain of 6-ph	9	3	0.090687	0.60192							
PF03144	GTP_EFTU_D2	Elongation factor Tu domain	20	5	0.093159	0.611611							

Table S4: Differentially Expressed Unigenes at FDR < 0.05 and log2FC > 0.5

Gene Model	log2FC (FDR)	transcript_id	sprot	Top %	Best BlastX results					Best BlastP results					Additional Annotations			
					e-val	gene_name	species	prot_id	sprot	Top %	e-val	gene_name	species	Pfam		gene_ontology	KEGG	
TR6271_c0_01	-14.30	1.90E-83 TR6271_c0_01	AP0D_HUI	35.56	1.00E-15	APOD	Apolipoprotein A	Human	TR6271_c0_01	11m.9392	AP0D_HUI	35.17	8.00E-17	APOD	Apolipoprotein A	Human	PF00661.1	
TR6274_c0_01	-13.98	7.74E-95 TR6274_c0_01	AP0D_HUI	35.56	1.00E-15	APOD	Apolipoprotein A	Human	TR6274_c0_01	11m.3780	AP0D_HUI	35.17	8.00E-17	APOD	Apolipoprotein A	Human	PF00661.1	
TR6289_c0_01	-13.58	7.52E-68 TR6289_c0_01	TRYP_AED	38.11	2.00E-50	TRYP7	Agmatase	Human	TR6289_c0_01	11m.15989	TRYP_AED	38.11	4.00E-51	TRYP7	Agmatase	Human	PF00089.2	Neuroactive ligand-receptor interaction
TR6295_c0_01	-13.19	4.00E-59 TR6295_c0_01	ASPP_AED	60.12	1.00E-14	AEL00061E	Lysosomal Aedes aeg	Human	TR6295_c0_01	11m.14316	ASPP_AED	59.48	3.00E-16	AEL00061E	Lysosomal Aedes aeg	Human	PF00026.1	Apoptosis/Autophagy - animal
TR6296_c0_01	-12.90	1.00E-41 TR6296_c0_01	RS18_SPOF	91.85	2.00E-71	RpS18	40S ribosomal S18	Drosophila	TR6296_c0_01	11m.8358	RS18_SPOF	92.76	8.00E-102	RpS18	40S ribosomal S18	Drosophila	PF00416.1	GO:0003732mol
TR6319_c5_01	-12.78	9.86E-49 TR6319_c5_01	CATL_SARF	49.07	2.00E-29	CATL	Cathepsin L	Human	TR6319_c5_01	11m.18098	CATL_SARF	48.09	4.00E-29	CATL	Cathepsin L	Human	PF00026.1	Apoptosis/Autophagy - animal
TR6321_c5_01	-12.73	1.11E-38 TR6321_c5_01	RSP_APMI	87.37	2.00E-105	RpS5	40S ribosomal S5	Drosophila	TR6321_c5_01	11m.15457	RSP_APMI	81.73	1.00E-123	RpS5	40S ribosomal S5	Drosophila	PF00201.1	Ribosome
TR6337_c0_01	-12.64	1.26E-40 TR6337_c0_01	RL13_SPOF	79.93	4.00E-98	RpL13	60S ribosomal L13	Drosophila	TR6337_c0_01	11m.17667	RL13_SPOF	79.93	5.00E-112	RpL13	60S ribosomal L13	Drosophila	PF01294.1	GO:0003735mol
TR6339_c0_01	-12.60	2.39E-37 TR6339_c0_01	RL13_RAT*	75.21	1.00E-53	RpL13	60S ribosomal L13	Rattus	TR6339_c0_01	11m.1158	RL13_RAT*	80	1.00E-60	RpL13	60S ribosomal L13	Rattus	PF01294.1	GO:0003735mol
TR6340_c0_01	-12.55	1.73E-37 TR6340_c0_01	RL27_RAT*	67.65	7.00E-60	RpL27	60S ribosomal L27	Rattus	TR6340_c0_01	11m.20300	RL27_RAT*	67.65	3.00E-59	RpL27	60S ribosomal L27	Rattus	PF04672.1	GO:0003735mol
TR6343_c1_01	-12.55	1.75E-38 TR6343_c1_01	CATB_BOV	60.06	8.00E-125	CTSB	Cathepsin B	Human	TR6343_c1_01	11m.8408	CATB_PIG*	57.4	2.00E-133	CTSB	Cathepsin B	Human	PF01112.1	GO:0004197mol
TR6349_c0_01	-12.52	1.45E-80 TR6349_c0_01	LIPE_HUM	4.00	5.00E-39	LIPG	Endothelial lipase	Human	TR6349_c0_01	11m.2894	LIPE_MOUSE	36.79	2.00E-40	Lipg	Endothelial lipase	Human	PF00151.1	
TR6351_c0_01	-12.48	4.80E-48 TR6351_c0_01	PURE_DRO	71.88	6.00E-77	RpL4	60S ribosomal L4	Drosophila	TR6351_c0_01	11m.9297	PURE_DRO	62.43	7.00E-70	RpL4	60S ribosomal L4	Drosophila	PF00573.1	GO:0006189mol
TR6353_c0_01	-12.32	2.09E-22 TR6353_c0_01	CIUD2_SCH	49.5	2.00E-26	Endoectodisc	Schistocerca	Human	TR6353_c0_01	11m.6222	CIUD2_SCH	49.5	3.00E-25	Endoectodisc	Schistocerca	Human	PF00379.1	GO:0042302mol
TR6354_c0_01	-12.27	3.26E-34 TR6354_c0_01	RS13_SPOF	90.73	4.00E-93	RpS13	40S ribosomal S13	Drosophila	TR6354_c0_01	11m.7129	RS13_SPOF	90.73	2.00E-95	RpS13	40S ribosomal S13	Drosophila	PF00312.1	GO:0003735mol
TR6357_c0_01	-12.27	4.70E-48 TR6357_c0_01	ASPL1_GALU	32.07	7.00E-12	Asolipoph	Galleria	Human	TR6357_c0_01	11m.6550	ASPL1_GALU	32.07	6.00E-09	Asolipoph	Galleria	Human	PF01442.1	GO:0002889mol
TR6361_c0_01	-12.25	3.35E-35 TR6361_c0_01	RL12_MOUSE	78.53	3.00E-90	RpL12	60S ribosomal S12	Human	TR6361_c0_01	11m.20469	RL12_MOUSE	78.53	7.00E-93	RpL12	60S ribosomal S12	Human	PF00298.1	GO:0003735mol
TR6363_c0_01	-12.18	4.53E-34 TR6363_c0_01	RL35A_POF	60.91	2.00E-47	RpL35A	60S ribosomal P35A	Drosophila	TR6363_c0_01	11m.3145	RL35A_POF	60.91	8.00E-44	RpL35A	60S ribosomal P35A	Drosophila	PF01247.1	GO:0003735mol
TR6365_c0_01	-12.15	7.83E-35 TR6365_c0_01	RL4_DROV	61.26	7.00E-77	RpL4	60S ribosomal L4	Drosophila	TR6365_c0_01	11m.9297	RL4_DROV	62.43	7.00E-70	RpL4	60S ribosomal L4	Drosophila	PF00573.1	GO:0003735mol
TR6366_c0_01	-12.12	6.34E-32 TR6366_c0_01	LIPE_MOUSE	34.12	2.00E-35	LIPE	Lipase	Human	TR6366_c0_01	11m.1719	LIPE_MOUSE	33.59	8.00E-38	LIPE	Lipase	Human	PF0151.1	
TR6367_c0_01	-12.08	1.93E-42 TR6367_c0_01	SERB_BOV	58.06	5.00E-51	PSPH	Phosphoserine	Human	TR6367_c0_01	11m.17230	SERB_BOV	58.06	9.00E-55	PSPH	Phosphoserine	Human	PF00702.2	GO:0016791mol
TR6368_c0_01	-12.08	1.06E-32 TR6368_c0_01	CTRL_HUM	37.45	3.00E-36	CTRL	Chromatin	Human	TR6368_c0_01	11m.7307	CTRL_HUM	34.01	1.00E-39	CTRL	Chromatin	Human	PF00881.1	GO:004252mol
TR6374_c0_01	-12.07	2.43E-30 TR6374_c0_01	RL15_CHIT	81.15	8.00E-107	RpL15	60S ribosomal L15	Drosophila	TR6374_c0_01	11m.6947	RL15_CHIT	81.15	4.00E-114	RpL15	60S ribosomal L15	Drosophila	PF00827.1	GO:0003735mol
TR6375_c0_01	-12.07	2.47E-31 TR6375_c0_01	LIJ2_HYA	50.6	4.00E-20	CP12	Flexible cy	Human	TR6375_c0_01	11m.9039	LIJ2_HYA	49.41	3.00E-19	CP12	Flexible cy	Human	PF00379.1	GO:0042302mol
TR6380_c28_01	-12.07	1.90E-30 TR6380_c28_01	RL35_RAT*	75.61	2.00E-55	RpL35	60S ribosomal L35	Rattus	TR6380_c28_01	11m.9292	RL35_RAT*	75.61	1.00E-55	RpL35	60S ribosomal L35	Rattus	PF00881.1	GO:0003735mol
TR6383_c1_01	-11.97	8.90E-30 TR6383_c1_01	Y3800_DRO	50.3	1.00E-48	CG3800	CyC-type	Drosophila	TR6383_c1_01	11m.7381	Y3800_DRO	50.89	4.00E-44	CG3800	CyC-type	Drosophila	PF00098.1	GO:0003676mol
TR6384_c0_01	-11.96	1.18E-31 TR6384_c0_01	TR6354_MOUSE					Human	TR6354_MOUSE	11m.18173	TR6354_MOUSE						PF06585.6	
TR6389_c0_01	-11.96	1.86E-33 TR6389_c0_01	RL23A_RAT	67.95	6.00E-65	RpL23A	60S ribosomal R23A	Rattus	TR6389_c0_01	11m.1082	RL23A_RAT	73.81	3.00E-58	RpL23A	60S ribosomal R23A	Rattus	PF00276.1	GO:0003735mol
TR6391_c0_01	-11.94	4.55E-36 TR6391_c0_01	CATL_SARF	40.74	2.00E-19	Cathepsin	Sarcophaga	Human	TR6391_c0_01	11m.16994	CATL_SARF	40.74	1.00E-18	Cathepsin	Sarcophaga	Human	PF08246.7	
TR6396_c8_01	-11.94	5.01E-33 TR6396_c8_01	COX3_COA	58.39	1.00E-46	COH	Cytochrome	Human	TR6396_c8_01	11m.19477	COX3_COA	60.34	3.00E-41	MT-COX3	Cytochrome	Human	PF00510.1	GO:0015002mol
TR6398_c5_01	-11.93	4.82E-107 TR6398_c5_01	CPII_PERA	40.17	9.00E-131	CP	Allegren	Human	TR6398_c5_01	11m.12477	CPII_PERA	40.17	9.00E-131	CP	Allegren	Human	PF00372.1	
TR6411_c0_01	-11.87	3.10E-34 TR6411_c0_01	RL13A_CH	78.52	3.00E-94	RpL13A	60S ribosomal L13A	Drosophila	TR6411_c0_01	11m.12669	RL13A_CH	77.48	4.00E-83	RpL13A	60S ribosomal L13A	Drosophila	PF00573.1	GO:0003735mol
TR6415_c0_01	-11.86	5.29E-31 TR6415_c0_01	RS6_MANS	84.12	2.00E-128	RpS6	40S ribosomal MANS	Drosophila	TR6415_c0_01	11m.6308	RS6_MANS	84.86	4.00E-145	RpS6	40S ribosomal MANS	Drosophila	PF01092.1	GO:0003735mol
TR6419_c0_01	-11.85	8.58E-30 TR6419_c0_01	RL9_SPOF	80.42	1.00E-104	RpL9	60S ribosomal SPOF	Drosophila	TR6419_c0_01	11m.13197	RL9_SPOF	80.42	6.00E-110	RpL9	60S ribosomal SPOF	Drosophila	PF00347.1	GO:0003735mol
TR6420_c0_01	-11.83	1.17E-26 TR6420_c0_01	RL36_DROV	73.04	9.00E-50	RpL36	60S ribosomal DROV	Drosophila	TR6420_c0_01	11m.2290	RL36_DROV	73.04	2.00E-49	RpL36	60S ribosomal DROV	Drosophila	PF01158.1	GO:0003735mol
TR6421_c0_01	-11.83	1.18E-31 TR6421_c0_01	RL6_PIG**	48.12	2.00E-69	RpL6	60S ribosomal S6	Drosophila	TR6421_c0_01	11m.372	RL6_PIG**	48.95	4.00E-66	RpL6	60S ribosomal S6	Drosophila	PF01159.1	GO:0003735mol
TR6423_c0_01	-11.77	5.26E-25 TR6423_c0_01	RL17A_DRO	90.22	6.00E-47	RpL17A	60S ribosomal L17A	Drosophila	TR6423_c0_01	11m.1915	RL17A_DRO	90.22	3.00E-46	RpL17A	60S ribosomal L17A	Drosophila	PF01159.1	GO:0003735mol
TR6424_c0_01	-11.75	2.05E-28 TR6424_c0_01	PURE_DRO	71.23	6.00E-77	RpL4	60S ribosomal L4	Drosophila	TR6424_c0_01	11m.9297	PURE_DRO	71.23	6.00E-77	RpL4	60S ribosomal L4	Drosophila	PF00573.1	GO:0003735mol
TR6425_c0_01	-11.72	7.11E-32 TR6425_c0_01	GST1_BLA	55.5	4.00E-68	Glutathion	Battellata	Human	TR6425_c0_01	11m.3070	GST1_BLA	55.5	3.00E-68	Glutathion	Battellata	Human	PF00043.2	GO:0005515mol
TR6476_c1_01	-11.68	1.41E-27 TR6476_c1_01	IF4A2_RAT	76.41	5.00E-57	EIF4A2	Eukaryotic	Human	TR6476_c1_01	11m.8068	IF4A2_RAT	76.14	4.00E-16	EIF4A2	Eukaryotic	Human	PF00270.2	GO:0003676mol
TR6479_c0_01	-11.64	2.65E-25 TR6479_c0_01	RL4O_DRO	96.88	5.00E-87	RpL4O	Ubiquitin-D	Drosophila	TR6479_c0_01	11m.4818	RL4O_DRO	96.88	1.00E-87	RpL4O	Ubiquitin-D	Drosophila	PF00240.1	GO:0005515mol
TR6481_c0_01	-11.63	1.76E-26 TR6481_c0_01	RL3A_AEDV	73.95	2.00E-55	RpL3A	60S ribosomal AEDV	Drosophila	TR6481_c0_01	11m.1769	RL3A_AEDV	73.95	3.00E-54	RpL3A	60S ribosomal AEDV	Drosophila	PF01199.1	GO:0003735mol
TR6482_c0_01	-11.59	5.00E-31 TR6482_c0_01	RS16_DROV	44.07	2.00E-31	RpS16	40S ribosomal S16	Drosophila	TR6482_c0_01	11m.7670	RS16_DROV	44.07	2.00E-31	RpS16	40S ribosomal S16	Drosophila	PF00573.1	GO:0003735mol
TR6486_c0_01	-11.58	1.81E-24 TR6486_c0_01	RS16_DRO	86.99	1.00E-90	RpS16	40S ribosomal S16	Drosophila	TR6486_c0_01	11m.543	RS16_DRO	86.99	5.00E-91	RpS16	40S ribosomal S16	Drosophila	PF00380.1	GO:0003735mol
TR6532_c0_01	-11.58	2.12E-29 TR6532_c0_01	RL32_APIV	85.82	2.00E-80	RpL32	60S ribosomal S16	Drosophila	TR6532_c0_01	11m.5858	RL32_APIV	85.82	6.00E-82	RpL32	60S ribosomal S16	Drosophila	PF01655.1	GO:0003735mol
TR6531_c6_01	-11.57	8.11E-29 TR6531_c6_01	RS19A_DRI	66.41	1.00E-51	RpS19A	40S ribosomal S19A	Drosophila	TR6531_c6_01	11m.11042	RS19A_DRI	61.04	3.00E-62	RpS19A	40S ribosomal S19A	Drosophila	PF01090.1	GO:0003735mol
TR6529_c12_01	-11.55	9.75E-28 TR6529_c12_01	ATPG_DRO	73.17	3.00E-138	ATPSynan	ATP synthase	Drosophila	TR6529_c12_01	11m.15203	ATPG_DRO	73.17	1.00E-149	ATPSynan	ATP synthase	Drosophila	PF00231.1	GO:0046933mol
TR6566_c0_01	-11.54	5.07E-29 TR6566_c0_01	RL4_DROV	74.11	1.00E-98	RpL4	60S ribosomal DROV	Drosophila	TR6566_c0_01	11m.9296	RL4_DROV	74.11	7.00E-101	RpL4	60S ribosomal DROV	Drosophila	PF00573.1	GO:0003735mol
TR6572_c4_01	-11.53	4.40E-34 TR6572_c4_01	RL2_HUM	94.92	4.00E-77	RpL2A	60S ribosomal AEDV	Human	TR6572_c4_01	11m.9291	RL2_HUM	95.28	6.00E-84	RpL2A	60S ribosomal AEDV	Human	PF00928.1	GO:0003735mol</

T86096c5_g1	-10.05	3.46E-19	TR6096c5_g1	MDCC_CHI	74.24	1.00E-98	MDH1 MD1	MD Malate dehydrogenase PFO0056.1 G:0016491 mol	MDHC_HU	73.74	1.00E-102	MDH1 MD Malate dehydrogenase PFO0056.1 G:0016491 mol	
T88990c1_g1	-10.04	5.35E-18	T88989c1_g1									Carbon metabolism 'Citrate cycle (TCA cycle)'	
T86226c7_g1	-10.03	6.85E-19	T86226c7_g1									Cystyl	
T86442c16_g1	-10.02	4.24E-17	T86442c16_g1									Carbon metabolism 'Citrate cycle (TCA cycle)'	
T82372c01_g1	-10.01	1.74E-27	T82372c01_g1	ATPS_DRC	66.36	3.00E-40	ATPSynO ATP synthase Drosophila PFO0213.1 G:00046933 mol					Oxidative phosphorylation	
T81665c10_g1	-10.07	6.20E-20	T81665c10_g1	IDHC_MIC	75.93	2.00E-147	IDH1 IDP2	Isocitrate C-microtrans PFO0180.1 G:00166166 mol				2-Oxocarboxylic acid metabolism	
T88897c01_g1	-10.07	8.81E-18	T88897c01_g1	ATPS_DRC	45.45	5.00E-24	ATPSynC6 ATP synthase Drosophila PFO0511.6 G:00150787 mol					Oxidative phosphorylation	
T85488c1_g1	-10.04	2.13E-09	T85488c1_g1										
T84271c01_g1	-10.01	2.53E-16	T84271c01_g1	TPIS_CULT	73.28	2.00E-129	TPi	Triosephosphate Culex tarsalis PFO0121.1 G:00048077 mol				Biosynthesis of amino acids 'Carbon metabolism'	
T86261c2_g1	-10.01	5.43E-17	T86261c2_g1	LPHI_RAB1	91.67	3.00E-34	LPHI	Lipase meso-cyctolipase PFO0155.1 G:00048083 mol				Protein processing in endoplasmic reticulum	
T83937c01_g1	-10.01	8.26E-19	T83937c01_g1	R515A_DR	62.61	3.00E-74	R515Aa	60S ribosomal protein L515a PFO0010.1 G:00037355 mol				Ribosome	
T83231c1_g1	-10.01	4.09E-18	T83231c1_g1	NPL14_MU	52.04	3.00E-104	Nap14	Nucleosomus mus mus PFO0556.1 G:00060334 mol					
T85888c01_g1	-10.01	1.75E-23	T85888c01_g1	CU2D_SCH	45.28	1.00E-23	Endocuticle Schistocerca gregaria PFO0379.1 G:00042302 mol						
T86087c1_g1	-10.01	4.30E-30	T86087c1_g1	YELL_DROE	41.33	2.00E-100	Protein Yell Drosophila PFO0202.1 G:00042302 mol						
T8789c01_g1	-10.01	9.38E-21	T8789c01_g1	NSA2_HUA	73.71	7.00E-118	NSA2	Nucleosomus mus mus PFO0120.1 G:00042302 mol					
T86365c01_g1	-10.01	5.40E-17	T86365c01_g1	CU2D_SCH	45.28	1.00E-23	Endocuticle Schistocerca gregaria PFO0379.1 G:00042302 mol						
T81828c01_g1	-10.01	9.87E-15	T81828c01_g1	GLYC_RAB1	55.93	3.00E-64	SHMT1	Serine hydroxymethyltransferase PFO0454.1 G:00040372 mol					Biosynthesis of amino acids 'Carbon metabolism'
T8754c1_g1	-10.01	8.85E-09	T8754c1_g1	VDAC_DRC	57.8	7.00E-118	SHMT1	Serine hydroxymethyltransferase PFO0454.1 G:00040372 mol					Biosynthesis of amino acids 'Carbon metabolism'
T85346c01_g1	-10.01	2.98E-15	T85346c01_g1	HEXA_BA	36.96	5.00E-156	Hexamerin Baboer de PFO0364.1 G:00150787 mol						
T82322c1_g1	-10.01	3.58E-21	T82322c1_g1	DNA1A_PO	59.74	3.00E-119	DNA1A	Dna1 homologue PFO0382.1 G:00031072 mol					Protein processing in endoplasmic reticulum
T88306c01_g1	-10.01	1.22E-10	T88306c01_g1										
T85128c1_g1	-10.01	3.75E-17	T85128c1_g1	CY1_BOVH	65.06	2.00E-96	CY1	Cytochrome b-taurin PFO02167.1 G:00055056 mol					Oxidative phosphorylation
T86169c1_g1	-10.01	2.35E-14	T86169c1_g1	CTRB1_LY	48.2	3.00E-41	Chymotrypsin-like trypsin PFO0042.1 G:00042527 mol						
T82030c01_g1	-10.01	6.50E-17	T82030c01_g1	UCRL_MOL	69.23	2.00E-96	UCRFs3	Cytochrome oxidase subunit 3 PFO0355.2 G:00081211 mol					Oxidative phosphorylation
T85837c01_g1	-10.01	3.42E-28	T85837c01_g1	RL28_SPOF	66.67	4.00E-55	Rpl28	60S ribosomal protein S28 PFO1778.1 G:00037355 mol					Ribosome
T83339c2_g1	-10.01	9.27E-18	T83339c2_g1	FRL_XENU	27.11	3.00E-14	Ferritin-like protein Xenusus PFO0210.1 G:00081999 mol						
T81255c01_g1	-10.01	1.66E-14	T81255c01_g1										
T8757c01_g1	-10.01	9.80E-19	T8757c01_g1	SERS_DRO	37.39	6.00E-14	Sam5 M2	Serine PFO0089.2 G:00044252 mol					
T85386c01_g1	-10.01	7.97E-18	T85386c01_g1	CBPE_ASTJ	44.2	2.00E-50	Carboxypeptidase Astacus PFO0246.1 G:00040818 mol						
T85125c01_g1	-10.01	8.75E-17	T85125c01_g1	SODC_CER	73.86	1.00E-75	Superoxide dismutase Ceratitis PFO0089.2 G:00044252 mol						Longevity regulating pathway - multiple species 'Pe'
T84550c01_g1	-10.01	2.73E-16	T84550c01_g1	FIBC_LUMG	37.85	4.00E-21	Fibrinolytic Lumbricus PFO0089.2 G:00044252 mol						
T85874c01_g1	-10.01	1.03E-23	T85874c01_g1	PPIA_DRO1	84.15	2.00E-98	Cyp1	Cyp1 Peptidyl-prolyl isomerase PFO0160.1 G:00037355 mol					Calcium signaling pathway
T85346c01_g1	-10.01	1.14E-16	T85346c01_g1	CRPI_PERA	37.08	1.00E-143	Allergen Cr Periplaneta PFO0372.1 G:00044252 mol						
T8436c01_g1	-10.01	9.00E-18	T8436c01_g1	COX42_TH	41.35	3.00E-22	Cytochrome Thaumaspis PFO0293.6 G:00044129 mol						Oxidative phosphorylation
T85830c01_g1	-10.01	1.92E-13	T85830c01_g1	LPHI_RAB1	91.67	3.00E-34	LPHI	Lipase meso-cyctolipase PFO0155.1 G:00048083 mol					Protein processing in endoplasmic reticulum
T81910c01_g1	-10.01	6.55E-15	T81910c01_g1	YCYT_DRO	38.23	2.00E-24	Cy1	Cyp1 Peptidyl-prolyl isomerase PFO0160.1 G:00037355 mol					Arachidonic acid metabolism
T87625c10_g1	-10.01	4.43E-06	T87625c10_g1	R3L_DROM	84	0	Rpl34	60S ribosomal protein L34 PFO0297.1 G:00037355 mol					Ribosome
T84653c01_g1	-10.01	6.65E-18	T84653c01_g1	ACD5B_MU	64.47	0	Acad5b	Short/branched mus mus PFO0441.1 G:00166277 mol					Fatty acid degradation 'Fatty acid metabolism'
T87020c01_g1	-10.01	6.65E-18	T87020c01_g1	ATPS_POI	50	2.00E-20	ATPS	ATP synthase Pongosa PFO0418.1 G:00150787 mol					Oxidative phosphorylation
T85894c01_g1	-10.01	6.65E-18	T85894c01_g1	TKT_PONA	69.31	3.00E-88	TKT	Transketolase Pongosa PFO0456.1 G:00080661 mol					Biosynthesis of amino acids 'Carbon metabolism'
T8436c01_g1	-10.01	3.57E-16	T8436c01_g1	COX42_TH	41.35	3.00E-22	Cytochrome Thaumaspis PFO0293.6 G:00044129 mol						Oxidative phosphorylation
T8696c01_g1	-10.01	2.00E-13	T8696c01_g1	MDHC_HU	73.74	1.00E-102	MDH1	MD Malate dehydrogenase PFO0056.1 G:0016491 mol					Carbon metabolism 'Citrate cycle (TCA cycle)'
T86350c01_g1	-10.01	6.63E-13	T86350c01_g1	RS2E_ANO	77.39	2.00E-57	Rps26	80S ribosomal protein S26 Agas Anopheles PFO0283.1 G:00037355 mol					Ribosome
T86350c01_g1	-10.01	6.63E-13	T86350c01_g1	ATPA_DRO	68	2.00E-34	ATPS	ATP synthase Pongosa PFO0418.1 G:00150787 mol					Oxidative phosphorylation
T85890c01_g1	-10.01	6.63E-13	T85890c01_g1	SUI1_ANO	97.27	6.00E-73	AGAP0064	Protein translocase Anopheles PFO0123.1 G:00037355 mol					RNA transport
T81045c01_g1	-10.01	6.73E-17	T81045c01_g1	GSLT_BLA	35.78	3.00E-43	Glutathione S-transferase Blattella PFO0443.1 G:00037355 mol						Arachidonic acid metabolism
T84647c01_g1	-10.01	8.25E-14	T84647c01_g1	AC011_TR	55.92	1.00E-136	D1105 PGY	Acyl-CoA Drosophila PFO0447.1 G:00044252 mol					Biosynthesis of unsaturated fatty acids 'Fatty acid'
T82896c01_g1	-10.01	1.05E-14	T82896c01_g1	RL24_PLU2	69.23	2.00E-92	Rpl24	60S ribosomal protein L24 PFO0297.1 G:00037355 mol					Ribosome
T85800c1_g1	-10.01	9.59E-19	T85800c1_g1	URIC_DRO	50.16	4.00E-97	Uro UO	Uro UO G.A. Uracilase Drosophila PFO0184.1 G:00044252 mol					Purine metabolism
T81733c01_g1	-10.01	5.93E-17	T81733c01_g1	RAN_DRO	95.53	3.00E-129	Ran	Ran GTP-binding Drosophila PFO0025.1 G:00055255 mol					Ribosome biogenesis in eukaryotes 'RNA transport'
T8642c15_g1	-10.01	1.95E-22	T8642c15_g1	CU2D_SCH	56.19	3.00E-34	Endocuticle Schistocerca gregaria PFO0379.1 G:00042302 mol						
T87379c01_g1	-10.01	5.58E-10	T87379c01_g1										
T86186c01_g1	-10.01	1.20E-14	T86186c01_g1	ADT1_ANC	77.7	1.00E-78	AGAP0067	ADP-ATP C Anopheles PFO0153.2 G:00044252 mol					Calcium signaling pathway
T86317c02_g1	-10.01	4.92E-13	T86317c02_g1	MR2_DRO	62.98	2.00E-79	Sam5 M2	Serine Pongosa PFO0089.2 G:00044252 mol					Biosynthesis of amino acids 'Cysteine and methion'
T85902c01_g1	-10.01	5.12E-25	T85902c01_g1	RL22_DRO	74.94	3.00E-28	Rpl22	60S ribosomal protein L22 Drosophila PFO1776.1 G:00037355 mol					Ribosome
T85902c01_g1	-10.01	5.12E-25	T85902c01_g1	AQD_AEDA	35.98	3.00E-29	AEL00351	Aquaporin Aedes aegypti PFO0230.1 G:00052515 mol					
T82372c01_g1	-10.01	5.95E-15	T82372c01_g1	ATPO_DRC	50.5	4.00E-24	ATPSynO	ATP synthase Drosophila PFO0213.1 G:00046933 mol					Oxidative phosphorylation
T89530c01_g1	-10.01	4.02E-14	T89530c01_g1	DPGN_DIP	58.33	1.00E-05	Ferrinone Diplolepis PFO0010.1 G:00055155 mol						
T81316c01_g1	-10.01	5.62E-17	T81316c01_g1	ATPS_DRC	47.52	2.00E-23	ATPSynC6	ATP synthase Drosophila PFO1316.1 G:00151115 mol					Oxidative phosphorylation
T86754c01_g1	-10.01	5.20E-31	T86754c01_g1	RS25_SPOF	84	3.00E-37	Rps25	40S ribosomal protein S25 Drosophila PFO0297.1 G:00037355 mol					Ribosome
T88107c01_g1	-10.01	3.81E-17	T88107c01_g1	CXGA1_BA	38.89	6.00E-12	CXGA1	CC Cytochrome b-taurin Drosophila PFO0297.1 G:00044252 mol					Oxidative phosphorylation
T87859c01_g1	-10.01	9.68E-20	T87859c01_g1	MRP_BA	67.63	1.00E-95	PMPBC	Mitochondrial Pongosa PFO0193.1 G:00044252 mol					
T82200c01_g1	-10.01	5.17E-15	T82200c01_g1	FIBC_LUMG	37.85	4.00E-21	Fibrinolytic Lumbricus PFO0089.2 G:00044252 mol						
T84550c01_g1	-10.01	5.36E-26	T84550c01_g1	TR0P_HUA	35.07	3.00E-06	TRO K1A1	Trophin Homosapiens PFO0089.2 G:00044252 mol					
T85987c01_g1	-10.01	4.99E-53	T85987c01_g1	RLA2_DRO	74.24	1.00E-24	Rpl2	80S ribosomal protein L2 Drosophila PFO0587.1 G:00037355 mol					Ribosome
T83350c01_g1	-10.01	4.80E-13	T83350c01_g1	ADT1_ANC	77.7	1.00E-78	AGAP0067	ADP-ATP C Anopheles PFO0153.2 G:00044252 mol					Calcium signaling pathway
T84857c01_g1	-10.01	4.35E-16	T84857c01_g1	APOD_HU	27.49	2.00E-11	APOD	Apolipoprotein Homosapiens PFO0601.1 G:00055155 mol					Drug metabolism - cytochrome P450
T84479c01_g1	-10.01	4.85E-14	T84479c01_g1	CTRB2_LY	37.43	3.00E-24	Chymotrypsin-like trypsin Drosophila PFO0089.2 G:00044252 mol						
T84261c01_g1	-10.01	9.47E-16	T84261c01_g1	METK_DRC	81.01	2.00E-87	Sam5 M2	Serine Pongosa PFO0089.2 G:00044252 mol					Biosynthesis of amino acids 'Cysteine and methion'
T8884c01_g1	-10.01	9.47E-16	T8884c01_g1	ACH2_BO	38.61	4.00E-01	Antichymotrypsin Bombyx PFO0089.2 G:00044252 mol						
T85262c1_g1	-10.01	9.47E-16	T85262c1_g1	PURH_CHIC	69.88	4.00E-169	ATIC	Purine bifunctional Gausax PFO0178.1 G:00039377 mol					One carbon pool by folate Purine metabolism
T85890c1_g1	-10.01	9.47E-16	T85890c1_g1	RS2A_PERA	84.15	2.00E-98	Cyp1	Cyp1 Peptidyl-prolyl isomerase PFO0160.1 G:00037355 mol					Ribosome
T84894c1_g1	-10.01	6.44E-16	T84894c1_g1	YMPF_CAE	39.93	4.00E-33	K12H47	Putative Caenorhabditis PFO0326.1 G:00028365 mol					
T87351c01_g1	-10.01	9.54E-17	T87351c01_g1	DYF_MAN	37.02	4.00E-14	Putative de Caenorhabditis PFO0214.1 G:00037355 mol						
T88351c01_g1	-10.01	4.52E-16	T88351c01_g1	CRUST_PA	50								

TR4920 c2_g1	-7.96	8.34E-09	TR4920 c2_g1.1	4EBP2_HU	59.46	5.00E-26	E4f4e2p	Eukaryotic Homo sapi	TR4920 c2_g1.1 1m.5137	4EBP2_MO	65.15	2.00E-26	E4f4e2p	Eukaryotic Mus mus	PF05456.6 G0:0008190 mol	Longevity regulating pathway - multiple species RN
TR5099 c4_g1	-7.96	8.47E-08	TR5099 c4_g1.1	AMPN_PL	31.82	2.00E-10	APN1	Amniote Pterodroma vyl	TR5099 c4_g1.1 1m.5434	AMPN_DR	31.82	1.00E-10	APN1	Amniote Pterodroma vyl	PF11838.3	
TR3356 c1_g1	-7.96	5.88E-10	TR3356 c1_g1.1	ANXN9_DR	62.8	3.00E-56	ANXN9	Amniote Anaxinix Be Drospila	TR3356 c1_g1.1 1m.3164	ANXN9_DR	62.8	1.00E-61	ANXN9	Amniote Anaxinix Be Drospila	PF01139.1 G0:0005509 mol	
TR8485 c1_g1	-7.95	2.53E-08	TR8485 c1_g1.1	PRX6_RK	52.46	6.00E-36	PRX6	Perovexio Gallus gallu	TR8485 c1_g1.1 1m.4275	PRX6_RK	52.46	1.00E-35	PRX6	Alipi Perovexio Rattus non	PF10417.4 G0:0005192 mol	
TR5894 c1_g1	-7.95	4.65E-04	TR5894 c1_g1.1	TKTL2_HU	45.16	3.00E-43	TKTL2	Transketol Homo sapi	TR5894 c1_g1.1 1m.7239	TKTL2_HU	45.16	2.00E-43	TKTL2	Transketol Drospila	PF0356.1 G0:0003355 mol	Calcium signaling pathway
TR4040 c1_g1	-7.95	2.65E-16	TR4040 c1_g1.1	TPIS_CULT	72.87	2.00E-128	TPIS	Trospheos Culex tarsa	TR4040 c1_g1.1 1m.3870	TPIS_CULT	72.87	6.00E-134	TPIS	Trospheos Culex tarsa	PF02121.1 G0:0004807 mol	Biosynthesis of amino acids/Carbon metabolism/P
TR5756 c5_g1	-7.95	1.00E-07	TR5756 c5_g1.1					TR5756 c5_g1.1 1m.6212								
TR2891 c0_g2	-7.93	8.52E-10	TR2891 c0_g2.1	TAKT_DRO	21.6	3.00E-07	CG1185	Protein tak Drospila	TR2891 c0_g2.1 1m.2698	TAKT_DRO	22.22	3.00E-07	CG1185	Protein tak Drospila	PF06858.6	
TR6791 c0_g2	-7.91	6.99E-09	TR6791 c0_g2.1	NDUB7_CB	43.48	2.00E-19	NDUB7	NADH dehyd Drospila	TR6791 c0_g2.1 1m.6303	NDUB7_CB	43.48	9.00E-18	D2030.4	NADH dehyd Caenorhab	PF05678.6 G0:0003954 mol	Oxidative phosphorylation
TR2382 c0_g2	-7.91	5.16E-25	TR2382 c0_g2.1	CATM_MAC	52.31	7.00E-119	C758	Catepsin B Drospila	TR2382 c0_g2.1 1m.2614	CATM_MAC	52.31	1.00E-114	C758	Catepsin B Drospila	PF01121.1 G0:0004107 mol	Apoptosis/Autophagy - animal/Lysosome
TR5874 c2_g1	-7.89	3.96E-05	TR5874 c2_g1.1	PKA_DRO	91.67	5.00E-43	CyclI	Protein kinase Drospila	TR5874 c2_g1.1 1m.3829	PKA_DRO	80.23	7.00E-45	CyclI	Protein kinase Drospila	PF0356.1 G0:0003355 mol	Calcium signaling pathway
TR5894 c1_g2	-7.88	9.76E-09	TR5894 c1_g2.1	TKTL2_HU	65.81	4.00E-59	TKTL2	Transketol Homo sapi	TR5894 c1_g2.1 1m.7388	TKTL2_HU	65.81	4.00E-66	TKTL2	Transketol Homo sapi	PF02779.1 G0:0003824 mol	Biosynthesis of amino acids/Carbon metabolism/P
TR5956 c6_g2	-7.87	1.96E-08	TR5956 c6_g2.1					TR5956 c6_g2.1 1m.7749								
TR4715 c0_g1	-7.87	3.51E-09	TR4715 c0_g1.1					TR4715 c0_g1.1 1m.4755								
TR5621 c0_g1	-7.87	2.69E-15	TR5621 c0_g1.1	CUD2_SCH	45.69	1.00E-25		Endocytic Schistocen	TR5621 c0_g1.1 1m.6507	CUD2_SCH	45.69	4.00E-26		Endocytic Schistocen	PF03079.1 G0:0042302 mol	
TR6232 c2_g2	-7.86	1.01E-09	TR6232 c2_g2.1	MYS4_DRO	77.62	8.00E-53	Mhc CG175	Myosin he Drospila	TR6232 c2_g2.1 1m.17091	MYS4_DRO	77.08	2.00E-46	Mhc CG175	Myosin he Drospila	PF00612.2 G0:0005515 mol	
TR4821 c1_g1	-7.86	8.96E-05	TR4821 c1_g1.1	PKS_ABAT	55.97	4.00E-50	At1g75040	Pathogene Arabidopsi	TR4821 c1_g1.1 1m.4854	PKS_ABAT	55.97	2.00E-46	At1g75040	Pathogene Arabidopsi	PF03414.1	
TR4735 c1_g1	-7.85	5.02E-10	TR4735 c1_g1.1	GLYM_HU	59.23	3.00E-46	SHMT2	Serine hyd Homo sapi	TR4735 c1_g1.1 1m.4787	GLYM_HU	59.23	4.00E-40	SHMT2	Serine hyd Homo sapi	PF04064.2 G0:00040372 mol	Biosynthesis of amino acids/Carbon metabolism/GP
TR5250 c0_g1	-7.82	1.27E-07	TR5250 c0_g1.1	SEI2A_MC	90.11	1.00E-112	Sec61a2	Protein tra Mus mus	TR5250 c0_g1.1 1m.5731	SEI2A_MC	90.11	3.00E-116	Sec61a2	Protein tra Mus mus	PF03044.2 G0:0015031 mol	Phagosome/Protein export/Protein processing in e
TR3813 c0_g1	-7.81	1.27E-14	TR3813 c0_g1.1	COSX5_BO	36.44	1.00E-18	Cox5b	Cytochrom Mus mus	TR3813 c0_g1.1 1m.3625	COSX5_MC	36.44	7.00E-20	Cox5b	Cytochrom Mus mus	PF02115.1 G0:0004419 mol	Oxidative phosphorylation
TR5925 c0_g2	-7.81	4.09E-07	TR5925 c0_g2.1	CUD1_SCH	42.73	3.00E-14		Endocytic Schistocen	TR5925 c0_g2.1 1m.7493	CUD1_SCH	42.73	7.00E-20		Endocytic Schistocen	PF02239.1 G0:0004632 mol	
TR4020 c0_g1	-7.79	3.26E-08	TR4020 c0_g1.1	CATL_DRO	69.03	3.00E-54	Cst1 f[2]CS2	Cathepsin Drospila	TR4020 c0_g1.1 1m.3863	CATL_DRO	69.05	6.00E-55	Cst1 f[2]CS2	Cathepsin Drospila	PF00428.2 G0:0002834 mol	Apoptosis/Autophagy - animal/Lysosome/Phagosome
TR3647 c1_g1	-7.79	1.55E-09	TR3647 c1_g1.1	PKS_ABAT	55.97	4.00E-50	At1g75040	Pathogene Arabidopsi	TR3647 c1_g1.1 1m.4854	PKS_ABAT	55.97	4.00E-50	At1g75040	Pathogene Arabidopsi	PF03414.1	
TR5804 c1_g1	-7.78	8.48E-05	TR5804 c1_g1.1	RL44_OCH	92.41	4.00E-48	Rpl44	60S ribosom Ochlerlat	TR5804 c1_g1.1 1m.7880	RL44_OCH	92.41	1.00E-40	Rpl44	60S ribosom Ochlerlat	PF00935.3 G0:0003735 mol	Oxidative phosphorylation
TR6071 c2_g1	-7.77	2.00E-15	TR6071 c2_g1.1	PTGR1_BO	53.23	2.00E-69	PTGR1	Protein tro Drospila	TR6071 c2_g1.1 1m.8096	PTGR1_BO	53.61	1.00E-62	PTGR1	Protein tro Drospila	PF01007.2 G0:00040270 mol	
TR6087 c2_g1	-7.75	3.55E-08	TR6087 c2_g1.1	MRP3_AP				TR6087 c2_g1.1 1m.9855	MRP3_AP	33.33	2.00E-06	MRP3	Major roya Apis mellif			
TR5479 c0_g1	-7.75	2.20E-07	TR5479 c0_g1.1	PP03_DRO	41.45	1.00E-31	PP03	Phenoloxi Drospila	TR5479 c0_g1.1 1m.6154	PP03_DRO	41.18	5.00E-31	PP03	Phenoloxi Drospila	PF03023.5	
TR4712 c0_g1	-7.74	2.21E-28	TR4712 c0_g1.1	RL10A_SPC	88.94	2.00E-124	Rpl10A	60S ribosom Drospila	TR4712 c0_g1.1 1m.4749	RL10A_SPC	88.94	7.00E-129	Rpl10A	60S ribosom Drospila	PF00687.1	Ribosome/Ribosome
TR6390 c4_g1	-7.72	2.20E-44	TR6390 c4_g1.1	RL4B_PAK	79.55	3.00E-152	Rpl4b	60S ribosom Drospila	TR6390 c4_g1.1 1m.18873	RL4B_PAK	79.71	7.00E-156	Rpl4b	60S ribosom Drospila	PF00428.2 G0:00042254 mol	Ribosome
TR3651 c0_g1	-7.71	7.00E-07	TR3651 c0_g1.1	IPYR_DRO	68.09	5.00E-77	Ruff1-38	CyC inorganic Drospila	TR3651 c0_g1.1 1m.3827	IPYR_DRO	68.09	1.00E-76	Ruff1-38	CyC inorganic Drospila	PF03023.5	Oxidative phosphorylation
TR6071 c2_g1	-7.70	2.64E-16	TR6071 c2_g1.1	ATC1_ANO	87.23	1.00E-153	Ca-P60A	Ca Calcium-tr Drospila	TR6071 c2_g1.1 1m.16645	ATC1_ANO	85.61	2.00E-168	Ca-P60A	Ca Calcium-tr Anopheles	PF00689.1	Calcium signaling pathway
TR5137 c0_g1	-7.69	7.54E-08	TR5137 c0_g1.1	MKNK1_XE	56.95	2.00E-51	mknk1	MAP kinase Xenopus tr	TR5137 c0_g1.1 1m.5479	MKNK1_XE	61.07	1.00E-52	mknk1	MAP kinase Xenopus tr	PF00669.2 G0:00040472 mol	MAPK signaling pathway
TR8826 c0_g1	-7.68	1.19E-04	TR8826 c0_g1.1	AL1L_LEPD	47.58	4.00E-34		Mite gro Lepidoptera	TR8826 c0_g1.1 1m.20887	AL1L_LEPD	47.58	5.00E-37		Mite gro Lepidoptera	PF02211.1	Lysosome
TR886 c0_g1	-7.67	1.06E-26	TR886 c0_g1.1	CPK61_BA	38.98	2.00E-121	CPYK61	Cytochrom Blattella ge	TR886 c0_g1.1 1m.746	CPK61_BA	39.17	1.00E-126	CPYK61	Cytochrom Blattella ge	PF00067.1 G0:0005056 mol	
TR1817 c0_g1	-7.67	3.83E-05	TR1817 c0_g1.1	LSO2_DRO	28.96	7.00E-21	CG92	Lipid dro Drospila	TR1817 c0_g1.1 1m.173	LSO2_DRO	29.48	9.00E-20	CG92	Lipid dro Drospila	PF03036.1	PPAR signaling pathway
TR5959 c0_g1	-7.67	1.82E-08	TR5959 c0_g1.1	NDU4A_PK	84.35	6.00E-10	NDUFA4	NADH dehyd Pongo pyp	TR5959 c0_g1.1 1m.3827	NDU4A_PK	84.35	1.00E-10	NDUFA4	NADH dehyd Pongo pyp	PF03421.1	Oxidative phosphorylation
TR9346 c0_g1	-7.65	6.27E-07	TR9346 c0_g1.1	CRYAA_CB	42.73	2.00E-17	CRYAA	Alpha-cryz Choleopus	TR9346 c0_g1.1 1m.21171	CRYAA_ER	42.48	2.00E-18	CRYAA	Alpha-cryz Eriacurus	PF00011.1	Longevity regulating pathway - multiple species/Prc
TR6054 c6_g1	-7.62	1.64E-08	TR6054 c6_g1.1	ECHM_BO	57.02	6.00E-39	ECH51	Elong-CoA Bos taurus	TR6054 c6_g1.1 1m.9042	ECHM_BO	55.28	2.00E-40	ECH51	Elong-CoA Bos taurus	PF03078.1 G0:0003824 mol	beta-Alanine metabolism/Biutanoate metabolism/C
TR5786 c1_g1	-7.62	1.74E-27	TR5786 c1_g1.1	EF1G_ART	72.75	0		Elongation Artemia sa	TR5786 c1_g1.1 1m.7011	EF1G_ART	73.71	0		Elongation Artemia sa	PF00043.2 G0:0005515 mol	
TR5962 c0_g1	-7.62	2.48E-07	TR5962 c0_g1.1	APLP_LOCP	43.58	2.00E-59		Apolipoplo Drospila	TR5962 c0_g1.1 1m.8257	APLP_LOCP	43.58	5.00E-60		Apolipoplo Drospila	PF00072.4 G0:0004332 mol	Biosynthesis of amino acids/Carbon metabolism/P
TR6008 c2_g1	-7.61	9.53E-09	TR6008 c2_g1.1	ALIP_DROM	70.3	3.00E-36	Alid G6059	Fructose-b Drospila	TR6008 c2_g1.1 1m.7859	ALIP_DROM	65.35	3.00E-36	Alid G6059	Fructose-b Drospila	PF00274.4 G0:0004332 mol	Biosynthesis of amino acids/Carbon metabolism/P
TR4515 c0_g1	-7.61	3.93E-12	TR4515 c0_g1.1	ENO_DRO	87.22	0	ENO	Eno CG176 Drospila	TR4515 c0_g1.1 1m.4505	ENO_DRO	87.22	0	ENO	Eno CG176 Drospila	PF00112.1 G0:00020278 mol	Oxidative phosphorylation
TR5856 c1_g1	-7.59	3.00E-08	TR5856 c1_g1.1	CBPZ_SIMA	50	2.00E-20		Zinc carbo Simulium u	TR5856 c1_g1.1 1m.7276	CBPZ_SIMA	48.98	4.00E-20		Zinc carbo Simulium u	PF02046.2 G0:0004181 mol	
TR6126 c0_g1	-7.58	6.57E-57	TR6126 c0_g1.1	PABP1_HU	65.57	0	pabpc1	Polyadenyl Homo sapi	TR6126 c0_g1.1 1m.10853	PABP1_HU	66.51	0	pabpc1	Polyadenyl Homo sapi	PF00076.1 G0:0003676 mol	mRNA surveillance pathway/mRNA surveillance pat
TR5875 c0_g1	-7.58	3.79E-28	TR5875 c0_g1.1	RS9_DROM	96.13	4.00E-119	RpS9	60S ribosom Drospila	TR5875 c0_g1.1 1m.7330	RS9_DROM	95.15	5.00E-121	RpS9	60S ribosom Drospila	PF00163.1 G0:0001983 mol	Ribosome
TR5715 c0_g1	-7.57	7.69E-28	TR5715 c0_g1.1	RS27A_DRI	78.21	9.00E-54	RpS27A	Ubiquitin-D Drospila	TR5715 c0_g1.1 1m.6794	RS27A_DRI	78.21	3.00E-71	RpS27A	Ubiquitin-D Drospila	PF02040.1 G0:0005515 mol	Ribosome
TR5937 c0_g1	-7.55	9.00E-09	TR5937 c0_g1.1	PNPH_HU	64.75	1.00E-50	PMP	Purine nuc Homo sapi	TR5937 c0_g1.1 1m.7858	PNPH_HU	64.75	3.00E-50	PMP	Purine nuc Homo sapi	PF01048.2 G0:0003824 mol	Nicotinate and nicotinamide metabolism/Purine m
TR4740 c0_g1	-7.55	1.00E-10	TR4740 c0_g1.1	TRX1_HU	58.35	1.00E-10	TRX1	Translocati Xenopus tr	TR4740 c0_g1.1 1m.3827	TRX1_HU	60.87	1.00E-11	TRX1	Translocati Xenopus tr	PF00932.1 G0:0003912 mol	Oxidative phosphorylation
TR2145 c0_g1	-7.55	1.24E-18	TR2145 c0_g1.1	TRAM1_BO	49.24	1.00E-19	TRAM1	Translocati Xenopus tr	TR2145 c0_g1.1 1m.1785	TR11_XEN	45.43	7.00E-11	TRAM1	Translocati Xenopus tr	PF03798.1 G0:0016021 mol	Cell Protein processing in endoplasmic reticulum
TR4305 c0_g1	-7.50	1.97E-06	TR4305 c0_g1.1	CY1_BOVIN	66.92	6.00E-38	Cyc1	Cytochrom Mus mus	TR4305 c0_g1.1 1m.4187	CY1_BOVIN	66.14	4.00E-57	CYC1	Cytochrom Bos taurus	PF00112.1 G0:0005056 mol	Oxidative phosphorylation
TR3962 c0_g1	-7.49	9.28E-07	TR3962 c0_g1.1	GRP75_PO	76.98	6.00E-65	HSPA9	Stress-70 Pongo abe	TR3962 c0_g1.1 1m.3784	GRP75_PO	69.94	6.00E-69	HSPA9	Stress-70 Pongo abe	PF00217.1	RNA degradation
TR8215 c0_g1	-7.47	3.28E-15	TR8215 c0_g1.1	AT1F2_CAE	46.43	9.00E-47	mal-2 B054	ATPase inh Caenorhab	TR8215 c0_g1.1 1m.20522	AT1F2_CAE	49.23	6.00E-16	mal-2 B054	ATPase inh Caenorhab	PF04588.7 G0:0004857 mol	
TR5962 c0_g2	-7.47	1.06E-08	TR5962 c0_g2.1	APLP_LOCP	42.75	3.00E-58		Apolipoplo Drospila	TR5962 c0_g2.1 1m.7859	APLP_LOCP	42.75	1.00E-58		Apolipoplo Drospila	PF00072.4 G0:0004332 mol	Biosynthesis of amino acids/Carbon metabolism/P
TR4182 c0_g1</																

TR6008[0]_g1	-6.31	2.83E-16	TR6008[0]_g1	ALF_DROM	84.59	6.00E-172	Alf CG6051 Fructose-6-Drospothila PF00274.1	GO:0004332mol Biosynthesis of amino acids	Biosynthesis of amino acids	
TR9391[0]_g1	-6.29	2.90E-21	TR9391[0]_g1	RS17_SPOF	87.79	3.00E-77	Rp517	40S ribosom Spodoptera	PF00833.1	GO:0003735mol Ribosome
TR9545[0]_g1	-6.22	1.69E-16	TR9545[0]_g1	RS2_DROM	91.67	6.00E-79	Rp52 soc C	40S ribosom Drospothila	TR9545[0]_g1	GO:0003735mol Ribosome
TR673[0]_g1	-6.11	1.36E-11	TR673[0]_g1	RS3O_RAT	83.05	1.00E-19	Fau	40S ribosom Rattus	TR673[0]_g1	GO:0003735mol Ribosome
TR6201[0]_g1	-6.12	1.36E-17	TR6201[0]_g1	MSY1_DR	75.16	1.00E-17	Pm CG593 Paramyosin Drospothila	TR6201[0]_g1	GO:0003735mol Ribosome	
TR8420[0]_g1	-6.11	1.30E-21	TR8420[0]_g1	RL7_DROM	68.16	2.00E-10	RpL7 CG48 60S ribosom Drospothila	TR8420[0]_g1	GO:0003735mol Ribosome	
TR8469[0]_g1	-6.08	2.54E-15	TR8469[0]_g1	FRI_AEAE	47.09	6.00E-46	FERH AEL Ferritin sub Aedes aeg	TR8469[0]_g1	GO:0003735mol Ribosome	
TR8849[0]_g1	-6.07	6.79E-13	TR8849[0]_g1	LOLA2_DRO	43.22	4.00E-56	loia CG120 Longitudin Drospothila	TR8849[0]_g1	GO:0003735mol Ribosome	
TR5673[0]_g1	-6.06	6.04E-13	TR5673[0]_g1	RLA1_DRO	80.95	3.00E-29	RpL2 60S acid: Drospothila	TR5673[0]_g1	GO:0003735mol Ribosome	
TR6044[0]_g1	-6.04	1.34E-16	TR6044[0]_g1	RS5_HUMJ	93.26	7.00E-126	RP55	40S ribosom Homo	TR6044[0]_g1	GO:0003735mol Ribosome
TR8963[0]_g1	-6.04	2.72E-05	TR8963[0]_g1	AMP1_ME	48.78	3.00E-07	Antimicrob Acrosin	TR8963[0]_g1	GO:0003735mol Ribosome	
TR5654[0]_g1	-6.01	1.13E-08	TR5654[0]_g1	ARF1_LOCO	99.02	6.00E-66	ARF1 ADP-ibosy Locusta	TR5654[0]_g1	GO:0003735mol Ribosome	
TR7683[0]_g1	-6.01	2.13E-08	TR7683[0]_g1	TBA1_DRO	100	1.00E-105	alphaTub8 Tubulin aeg Drospothila	TR7683[0]_g1	GO:0003735mol Ribosome	
TR5762[0]_g1	-6.01	1.50E-17	TR5762[0]_g1	RL19_DRO	84.29	2.00E-95	RpL19 M12 60S ribosom Drospothila	TR5762[0]_g1	GO:0003735mol Ribosome	
TR4363[0]_g1	-6.00	8.52E-07	TR4363[0]_g1	TXTP_CAEE	72.39	8.00E-82	K11H3.3 Putative Tr Caenorhab	TR4363[0]_g1	GO:0003735mol Ribosome	
TR1356[0]_g1	-5.98	4.47E-13	TR1356[0]_g1	EFZ_DROM	92.3	0	EFZ E2D o Elongation Drospothila	TR1356[0]_g1	GO:0003924mol	
TR1895[0]_g1	-5.98	2.00E-01	TR1895[0]_g1	MLR_DRO	48	2.00E-40	Mysin reg Bombyx m	TR1895[0]_g1	GO:0005099mol	
TR6186[0]_g1	-5.98	1.27E-15	TR6186[0]_g1	RL3A_CA	61.16	3.00E-45	rpL33 F10E 60S ribosom Caenorhab	TR6186[0]_g1	GO:0003735mol Ribosome	
TR6181[0]_g1	-5.98	1.97E-16	TR6181[0]_g1	RL13A_CH	71.57	2.00E-103	RpL13A 60S ribosom Choristone	TR6181[0]_g1	GO:0003735mol Ribosome	
TR5332[0]_g1	-5.96	4.89E-16	TR5332[0]_g1	RL32_APIH	85.61	7.00E-79	RpL32 rp45 60S ribosom Ascaris	TR5332[0]_g1	GO:0003735mol Ribosome	
TR6173[0]_g1	-5.96	6.09E-14	TR6173[0]_g1	CUD2_SCH	47.12	9.00E-28	Endocutic Schistocen	TR6173[0]_g1	GO:0004230mol	
TR4967[0]_g1	-5.94	4.05E-11	TR4967[0]_g1	OB10_DRO	38.64	4.00E-116	OB10 O5-D C Putative o Drospothila	TR4967[0]_g1	GO:0003924mol	
TR5958[0]_g1	-5.93	4.51E-23	TR5958[0]_g1	TCTP_DROM	78.31	3.00E-92	Tcp Translator Bombyx m	TR5958[0]_g1	GO:0003823mol	
TR6355[0]_g1	-5.92	1.24E-17	TR6355[0]_g1	ATPA_PIG	83.56	9.00E-73	ATPSA3 At ATP synth S crofa	TR6355[0]_g1	GO:0006820mol Oxidative phosphorylation	
TR6325[0]_g1	-5.91	3.56E-23	TR6325[0]_g1	MYSR_DRC	84.8	0	Mhc CG175 Mysosin Drospothila	TR6325[0]_g1	GO:0003735mol Ribosome	
TR2896[0]_g1	-5.90	1.53E-13	TR2896[0]_g1	RL24_SPOF	84.55	8.00E-54	RpL24 CG9 60S ribosom Spodoptera	TR2896[0]_g1	GO:0003735mol Ribosome	
TR6128[0]_g1	-5.89	6.45E-28	TR6128[0]_g1	CUD2_SCH	43.64	1.00E-17	Endocutic Schistocen	TR6128[0]_g1	GO:0004230mol	
TR5803[0]_g1	-5.88	1.75E-11	TR5803[0]_g1	CAIM_LOCO	100	8.00E-101	0 Calmodulm Locusta	TR5803[0]_g1	GO:0005509mol Calcium signaling pathway	
TR6044[0]_g1	-5.85	5.61E-11	TR6044[0]_g1	RS5_HUMJ	93.26	7.00E-126	RP55	40S ribosom Homo	TR6044[0]_g1	GO:0003735mol Ribosome
TR5476[0]_g1	-5.85	6.03E-11	TR5476[0]_g1	PDI_DROM	59.93	2.00E-115	Pdi CG698 Protein Dis Drospothila	TR5476[0]_g1	GO:0004554mol Protein processing in endoplasmic reticulum	
TR5697[0]_g1	-5.82	4.36E-19	TR5697[0]_g1	RS2O_NEM	85.47	3.00E-63	rp20 40S ribosom Xenopus	TR5697[0]_g1	GO:0003735mol Ribosome	
TR6232[0]_g1	-5.79	3.74E-34	TR6232[0]_g1	PTPM_BLAG	93.99	2.00E-136	0 Troponmy: Blatella g	TR6232[0]_g1	GO:0003735mol Ribosome	
TR6171[0]_g1	-5.73	1.24E-16	TR6171[0]_g1	IFSA_SPOF	91.25	4.00E-104	efF-5A eF5 Eukaryotic Spodoptera	TR6171[0]_g1	GO:0003735mol Ribosome	
TR6355[0]_g1	-5.71	1.08E-12	TR6355[0]_g1	ATPA_DRO	84.46	0	blw ATP5Y ATP synth Drospothila	TR6355[0]_g1	GO:0005992mol Oxidative phosphorylation	
TR1244[0]_g1	-5.71	3.14E-16	TR1244[0]_g1	RL3E_RAT	71.54	3.00E-162	RP35 60S ribosom Rattus	TR1244[0]_g1	GO:0003735mol Ribosome	
TR6235[0]_g1	-5.71	1.10E-10	TR6235[0]_g1	CUJA2_TE	41.25	2.00E-19	0 Larval cutic Tenobryx m	TR6235[0]_g1	GO:0003735mol Ribosome	
TR6629[0]_g1	-5.70	5.27E-09	TR6629[0]_g1	PRDX5_HU	57.49	4.00E-28	PRDX5 AC6 Peroxiredoxin Homo sapi	TR6629[0]_g1	GO:0006191mol Peroxisome	
TR4576[0]_g1	-5.70	3.06E-10	TR4576[0]_g1	PDI_DROM	65.07	2.00E-57	Pdi CG698 Protein Dis Drospothila	TR4576[0]_g1	GO:0004554mol Protein processing in endoplasmic reticulum	
TR5476[0]_g1	-5.70	3.06E-10	TR5476[0]_g1	VATL_DRO	91.33	3.00E-89	Vha16-1 V1-V type pro Drospothila	TR5476[0]_g1	GO:0005784mol Protein processing in endoplasmic reticulum	
TR5430[0]_g1	-5.67	1.12E-09	TR5430[0]_g1	EF1B2_BO	84.85	4.00E-53	0 Elongation Bombyx m	TR5430[0]_g1	GO:0003735mol Ribosome	
TR5889[0]_g1	-5.66	7.88E-15	TR5889[0]_g1	CUJA2_TE	55.56	6.00E-16	0 Pupal cutic Tenobryx m	TR5889[0]_g1	GO:0003735mol Ribosome	
TR5889[0]_g1	-5.66	7.88E-15	TR5889[0]_g1	CUJA2_TE	55.56	6.00E-16	0 Pupal cutic Tenobryx m	TR5889[0]_g1	GO:0003735mol Ribosome	
TR6275[0]_g1	-5.65	2.90E-26	TR6275[0]_g1	RL3_DROM	84.25	0	RpL3 CG48 60S ribosom Drospothila	TR6275[0]_g1	GO:0003735mol Ribosome	
TR1002[0]_g1	-5.65	7.32E-19	TR1002[0]_g1	RS4_CARGI	90.77	4.00E-174	RP54 40S ribosom Carabus g	TR1002[0]_g1	GO:0003735mol Ribosome	
TR6248[0]_g1	-5.62	3.29E-28	TR6248[0]_g1	RS15_ELAC	77.33	7.00E-83	RP515 40S ribosom Eleais oleif	TR6248[0]_g1	GO:0003735mol Ribosome	
TR700[0]_g1	-5.61	2.20E-22	TR700[0]_g1	RL4_DROM	70.73	3.00E-174	RpL4 RpL1 60S ribosom Drospothila	TR700[0]_g1	GO:0003735mol Ribosome	
TR4448[0]_g1	-5.60	5.64E-06	TR4448[0]_g1	RS11_RAT	66.43	9.00E-65	RP11 40S ribosom Rattus	TR4448[0]_g1	GO:0003735mol Ribosome	
TR1212[0]_g1	-5.58	1.24E-11	TR1212[0]_g1	PROBAC_DRO	92.47	1.00E-108	Probabact Drospothila	TR1212[0]_g1	GO:0004550mol Purine metabolism	
TR6251[0]_g1	-5.58	1.25E-11	TR6251[0]_g1	GRP_DRO	88.03	8.00E-90	Rack1 CG7 Guanine n Drospothila	TR6251[0]_g1	GO:0005515mol	
TR5976[0]_g1	-5.57	3.29E-17	TR5976[0]_g1	IF4A1_RAB	89.17	1.00E-56	EIF4A1 DD: Eukaryotic Oryctolagu	TR5976[0]_g1	GO:0003735mol Ribosome	
TR5430[0]_g1	-5.46	2.33E-12	TR5430[0]_g1	EF1B2_BO	67.39	4.00E-94	0 Elongation Bombyx m	TR5430[0]_g1	GO:0003735mol Ribosome	
TR1001[0]_g1	-5.40	8.97E-05	TR1001[0]_g1	LECB_CAEE	54.1	9.00E-56	lec-8 R07B: Probable g Caenorhab	TR1001[0]_g1	GO:0003246mol	
TR9043[0]_g1	-5.40	8.00E-06	TR9043[0]_g1	GRX2_HU	51.76	1.00E-24	GLRX2 GRX Glutaredoxin Homo sapi	TR9043[0]_g1	GO:0003055mol	
TR5625[0]_g1	-5.39	2.00E-15	TR5625[0]_g1	PE8B_DRO	68.00	2.00E-88	Rack1 CG7 Guanine n Drospothila	TR5625[0]_g1	GO:0005515mol	
TR6305[0]_g1	-5.38	4.65E-12	TR6305[0]_g1	PE8B_DRO	47.75	3.00E-32	Ebp1B1 Ebp1: Eukaryotic Drospothila	TR6305[0]_g1	GO:0003932mol	
TR5765[0]_g1	-5.34	3.51E-16	TR5765[0]_g1	TNTP_PER	80.59	1.00E-166	TNTP Troponin T Periplanet	TR5765[0]_g1	GO:0005861mol cell	
TR6353[0]_g1	-5.31	8.46E-56	TR6353[0]_g1	EF1A2_DRO	95.27	0	EF1alpha10 Elongation Drospothila	TR6353[0]_g1	GO:0003924mol Protein transport	
TR5476[0]_g1	-5.31	2.00E-115	TR5476[0]_g1	PIA1_DROM	59.93	2.00E-115	Pdi CG698 Protein Dis Drospothila	TR5476[0]_g1	GO:0004554mol Protein processing in endoplasmic reticulum	
TR6087[0]_g1	-5.28	3.27E-16	TR6087[0]_g1	VELL_DROE	41.57	7.00E-93	Protein vcl Manduca s	TR6087[0]_g1	GO:0003924mol	
TR5532[0]_g1	-5.27	2.28E-10	TR5532[0]_g1	YATB_HELI	98.17	4.00E-156	Y-type pro Manduca s	TR5532[0]_g1	GO:0005224mol mTOR signaling pathway	
TR6235[0]_g1	-5.27	1.13E-07	TR6235[0]_g1	CUD2_SCH	47.12	9.00E-28	Endocutic Schistocen	TR6235[0]_g1	GO:0004230mol	
TR5602[0]_g1	-5.24	1.28E-10	TR5602[0]_g1	ACOL11_SPI	38.69	5.00E-54	0 Acyl-CoA D Spodoptera	TR5602[0]_g1	GO:0006629mol Biosynthesis of unsaturated fatty acids	
TR6032[0]_g1	-5.21	8.03E-13	TR6032[0]_g1	RL8_SPOFF	88.6	4.00E-70	RpL8 60S ribosom Rattus	TR6032[0]_g1	GO:0003735mol Ribosome	
TR5555[0]_g1	-5.18	1.62E-07	TR5555[0]_g1	A1A1A_DRO	55.34	1.00E-78	Akr1a1 Akr Alodrospothila	TR5555[0]_g1	GO:0003735mol Ribosome	
TR5803[0]_g1	-5.16	4.37E-11	TR5803[0]_g1	CAIM_LOCO	100	8.00E-101	0 Calmodulm Locusta	TR5803[0]_g1	GO:0005509mol Calcium signaling pathway	
TR6160[0]_g1	-5.15	1.17E-16	TR6160[0]_g1	EF1A2_OSC	88.92	0	ef-4 Elongation Ochusius t	TR6160[0]_g1	GO:0003924mol Protein transport	
TR6474[0]_g1	-5.11	2.00E-04	TR6474[0]_g1	ACT3_ART	90.50E-07	2.00E-5	cyn-5 Cyp5: Peptidyl-tr Caenorhab	TR6474[0]_g1	GO:0003735mol Ribosome	
TR6438[0]_g1	-5.11	1.07E-07	TR6438[0]_g1	ACT_MAN	98.14	0	0 Actin, mms Drospothila	TR6438[0]_g1	GO:0003735mol Ribosome	
TR6322[0]_g1	-5.11	1.25E-07	TR6322[0]_g1	MYSR_DRC	76.43	1.00E-69	Mhc CG175 Mysosin Drospothila	TR6322[0]_g1	GO:0003735mol Ribosome	
TR5879[0]_g1	-5.11	2.57E-17	TR5879[0]_g1	RL27A_BO	69.64	9.00E-51	RpL27a 60S ribosom Rattus	TR5879[0]_g1	GO:0003735mol Ribosome	
TR6107[0]_g1	-5.07	4.79E-13	TR6107[0]_g1	RL13_SPOF	73.11	1.00E-96	RpL13 60S ribosom Spodoptera	TR6107[0]_g1	GO:0003735mol Ribosome	
TR6265[0]_g1	-5.07	1.99E-13	TR6265[0]_g1	YG31B_YE	30.27	1.00E-100	Y318-G-VG Transposon Saccharom	TR6265[0]_g1	GO:0005074mol	
TR5747[0]_g1	-5.07	1.99E-13	TR5747[0]_g1	YG31B_YE	30.27	1.00E-100	Y318-G-VG Transposon Saccharom	TR5747[0]_g1	GO:0005074mol	
TR5869[0]_g1	-5.01	2.95E-18	TR5869[0]_g1	RL18_TMB	80.85	5.00E-109	RpL18 60S ribosom Timarcha t	TR5869[0]_g1	GO:0003735mol Ribosome	
TR5976[0]_g1	-4.97	1.99E-15	TR5976[0]_g1	IF4A2_RAT	78.84	0	Ef4a2 Eukaryotic Rattus	TR5976[0]_g1	GO:0003735mol Ribosome	
TR5682[0]_g1	-4.95	1.75E-10	TR5682[0]_g1	GPI_LOCO	36.05	1.00E-41	Gpi Lettzymine Glossina a	TR5682[0]_g1	GO:0004252mol	
TR5888[0]_g1	-4.95	2.22E-08	TR5888[0]_g1	CUD2_SCH	42.65	2.00E-10	0 Endocutic Schistocen	TR5888[0]_g1	GO:0004230mol	
TR6164[0]_g1	-4.94	1.70E-09	TR6164[0]_g1	HSP70_MD	82.83	1.00E-49	0 Heat shock Manduca s	TR6164[0]_g1	GO:0003924mol	
TR6131[0]_g1	-4.93	3.28E-03	TR6131[0]_g1	CYPS_CAEE	78.48	1.00E-66	cyn-5 Cyp5: Peptidyl-tr Caenorhab	TR6131[0]_g1	GO:0003735mol Ribosome	
TR4595[0]_g1	-4.92	1.21E-15	TR4595[0]_g1	RL13_SPOF	76.61	6.00E-61	RP13 60S ribosom Spodoptera	TR4595[0]_g1	GO:0003735mol Ribosome	
TR6322[0]_g1	-4.92	1.21E-07	TR6322[0]_g1	MYSR_DRC	80.86	1.0				

TR1026 c0_1	-3.31	5.47E-04	rp5-23 F28i 405 riboso	Caenorhab TR1026 c0_1 1m.847	RS23_CAEE	91.61	6.00E-91	rp5-23 F28i 405 riboso	Caenorhab PF00164.2 1 GO:0003735 mol	Ribosome	
TR4299 c0_2	-3.29	2.05E-03	AGAP0067 ADP.ATP c	Anopheles TR4299 c0_2 1m.4183	ADT_DROA			ADP.ATP c	Drosophila PF01533.2	Calcium signaling pathway	
TR8242 c0_1	-3.29	2.84E-04	Rp515A	Rp515A RAE	RS15A_RA	87.5	2.00E-71	Rp515A	Rp515A RAE	0003735 mol	Ribosome
TR1790 c0_1	-3.28	2.89E-03	rp1-26 F28i 605 riboso	Caenorhab TR1790 c0_1 1m.1539	RL32_CAEE	88.03	4.00E-87	rp1-26 F28i 605 riboso	Caenorhab PF00467.2	Ribosome	
TR7275 c0_1	-3.27	2.70E-02	rp1-27a rp1-605 riboso	Ochelus TR7275 c0_1 1m.19089	RL27A_OSC	94.48	1.00E-96	rp1-27a rp1-605 riboso	Ochelus PF00828.1	Ribosome	
TR4299 c0_1	-3.27	5.18E-03	AGAP0067 ADP.ATP c	Anopheles TR4299 c0_1 1m.4182	ADT_DROA			ADP.ATP c	Drosophila PF01533.2	Calcium signaling pathway	
TR1585 c0_1	-3.26	3.07E-02	rack-1 K04I Guanine ri	Caenorhab TR1585 c0_1 1m.1344	GRBP_CAEE	83.39	6.00E-165	rack-1 K04I Guanine ri	Caenorhab PF04002.0	GO:0005515 mol	
TR6851 c0_1	-3.24	3.68E-03	rp5-23 F28i 405 riboso	Caenorhab TR6851 c0_1 1m.19755	RS33_ORY1	76.27	4.00E-133	Fau	0005113 mol	Ribosome	
TR6610 c0_1	-3.24	2.05E-02	rp5-5 T05E 405 riboso	Caenorhab TR6610 c0_1 1m.19641	RS5_CAEE	83.18	1.00E-120	rp5-5 T05E 405 riboso	Caenorhab PF01278.1	Ribosome	
TR8134 c0_1	-3.24	5.51E-04	rp5-1 F54C3 605 riboso	Caenorhab TR8134 c0_1 1m.20465	RS5_CAEE	76.38	1.00E-142	rp5-1 F54C3 605 riboso	Caenorhab PF00828.1	GO:0003735 mol	Ribosome
TR4834 c0_1	-3.22	2.70E-02	asp-6 F21F Aspartic pr	Caenorhab TR4834 c0_1 1m.4979	ASPE_CAEE	43.05	5.00E-24	asp-6 F21F Aspartic pr	Caenorhab PF00026.1 1 GO:0004190 mol	Apoptosis/Autophagy - animal Lysosome/Sphingolipid	
TR8241 c0_1	-3.18	2.51E-03	rp1-26 F28i 605 riboso	Caenorhab TR8241 c0_1 1m.20627	RL26_CAEE	84.46	3.00E-117		000281.1 1 GO:0003735 mol	Ribosome	
TR251 c0_1	-3.15	2.22E-02	rp1-24.1 D1 605 riboso	Caenorhab TR251 c0_1 1m.23	RL24_CAEE	73.98	9.00E-36	rp1-24.1 D1 605 riboso	Caenorhab PF01246.1	Ribosome	
TR3513 c1_1	-3.13	1.02E-02	rp5-23 F28i 405 riboso	Caenorhab TR3513 c1_1 1m.3303	RS3_CAEE	84.21	3.00E-147	rp3-C23G 405 riboso	Caenorhab PF01089.1 1 GO:0003735 mol	Ribosome	
TR5585 c2_1	-3.12	5.46E-03	rp5-23 F28i 405 riboso	Caenorhab TR5585 c2_1 1m.6424	RL4O_CAEE	81.79	4.00E-167	rp5-23 F28i 405 riboso	Caenorhab PF00428.1 1 GO:0042554 biol	Ribosome	
TR8671 c0_1	-3.12	4.98E-03	rp2-2 C49H 405 riboso	Caenorhab TR8671 c0_1 1m.20789	RS2_CAEE	89.95	5.00E-142	rp2-2 C49H 405 riboso	Caenorhab PF00333.1 1 GO:0003735 mol	Ribosome	
TR8119 c0_1	-3.11	5.29E-03	rp1-18 C8G 605 riboso	Caenorhab TR8119 c0_1 1m.20951	RL18_CAEE	79.14	1.00E-95	rp1-18 C8G 605 riboso	Caenorhab PF00828.1	Ribosome	
TR8195 c0_1	-3.11	2.84E-03	rp1-7A Y24I 605 riboso	Caenorhab TR8195 c0_1 1m.20509	RL7A_CAEE	76.89	4.00E-127	rp1-7A Y24I 605 riboso	Caenorhab PF01248.2	Ribosome	
TR6336 c16_1	-3.10	6.99E-03	rp1-26 F28i 605 riboso	Caenorhab TR6336 c16_1 1m.17606	RS10_SPOF	57.14	5.00E-52	Rp510	000515 mol	Ribosome	
TR1621 c0_1	-3.08	2.70E-02	rp1-20 E04F 605 riboso	Caenorhab TR1621 c0_1 1m.159	RL16A_CAEE	85.56	7.00E-112	rp1-20 E04F 605 riboso	Caenorhab PF01775.1 1 GO:0003735 mol	Ribosome	
TR8240 c0_1	-3.08	3.98E-03	rp1-28 R11I 605 riboso	Caenorhab TR8240 c0_1 1m.20626	RL28_CAEE	54.89	1.00E-34	rp1-28 R11I 605 riboso	Caenorhab PF01778.1	Ribosome	
TR8203 c0_1	-3.08	2.19E-03	rp1-28 R11I 605 riboso	Caenorhab TR8203 c0_1 1m.20368	RL4_CAEE	79.26	0	rp1-B0041 605 riboso	Caenorhab PF00573.1 1 GO:0003735 mol	Ribosome	
TR3034 c0_1	-3.07	1.16E-03	rp1-28 R11I 605 riboso	Caenorhab TR3034 c0_1 1m.206	RL4_CAEE	98.1	2.00E-68	rp1-28 R11I 605 riboso	Caenorhab PF00654.1 1 GO:0003735 mol	Ribosome	
TR9428 c0_1	-3.05	6.39E-03	Rp114	Rp114 PIG*	RL14_PIG*	50.75	3.00E-41	Rp114	000515 mol	Ribosome	
TR1362 c0_1	-3.03	9.30E-03	rp1-36 F37C 605 riboso	Caenorhab TR1362 c0_1 1m.1151	RL36_CAEE	76.7	3.00E-45	rp1-36 F37C 605 riboso	Caenorhab PF01158.1 1 GO:0003735 mol	Ribosome	
TR5712 c0_1	-3.03	7.79E-08	Rp523	Rp523 SPF	RS23_SPOF	95.1	7.00E-96	Rp523	000614.2 1 GO:0003735 mol	Ribosome	
TR9334 c0_1	-3.01	7.26E-03	rp1-27 C53I 605 riboso	Caenorhab TR9334 c0_1 1m.21162	RL27_CAEE	84.33	6.00E-80	rp1-27 C53I 605 riboso	Caenorhab PF00467.2 1 GO:0003735 mol	Ribosome	
TR9186 c0_1	-3.01	3.22E-03	rp1-28 R11I 605 riboso	Caenorhab TR9186 c0_1 1m.21062	RS8_CAEE	81.73	7.00E-124	rp8-F42C 405 riboso	Caenorhab PF01201.1	Ribosome	
TR8170 c0_1	-2.99	6.88E-03	rp1-26 F28i 605 riboso	Caenorhab TR8170 c0_1 1m.19754	RS14_CAEE	91.49	1.00E-91	rp1-26 F28i 605 riboso	Caenorhab PF00411.1 1 GO:0003735 mol	Ribosome	
TR6458 c11_1	-2.93	5.63E-04	ACT5C act1 Actin-5C	(F Anopheles TR6458 c11_1 1m.19342	ACT5C_AN	100	6.00E-88	ACT5C act1 Actin-5C	(F Anopheles PF00223.1	Adhesion junction/Apoptosis/Focal adhesion/Hippo	
TR9203 c0_1	-2.91	1.96E-02	rp1-31 W09 605 riboso	Caenorhab TR9203 c0_1 1m.21072	RL31_CAEE	84.3	2.00E-62	rp1-31 W09 605 riboso	Caenorhab PF01198.1 1 GO:0003735 mol	Ribosome	
TR1790 c0_2	-2.90	2.30E-02	rp1-26 F28i 605 riboso	Caenorhab TR1790 c0_2 1m.1540	RL26_CAEE	87.05	9.00E-84	rp1-26 F28i 605 riboso	Caenorhab PF00467.2	Ribosome	
TR1047 c0_1	-2.89	5.28E-03	rp1-10a rp1-605 riboso	Caenorhab TR1047 c0_1 1m.867	RL10A_CAEE	83.64	3.00E-109	rp1-10a rp1-605 riboso	Caenorhab PF00687.1	Ribosome	
TR7848 c0_1	-2.86	1.74E-02	rp5-23 F28i 405 riboso	Caenorhab TR7848 c0_1 1m.20290	RS26_CAEE	93.2	7.00E-65	rp5-23 F28i 405 riboso	Caenorhab PF01283.1 1 GO:0003735 mol	Ribosome	
TR2514 c0_1	-2.86	1.07E-02	rp1-17 Y48I 605 riboso	Caenorhab TR2514 c0_1 1m.1395	RL17_CAEE	82.00	2.00E-110	rp1-17 Y48I 605 riboso	Caenorhab PF00237.1 1 GO:0003735 mol	Ribosome	
TR6172 c15_1	-2.85	4.00E-03	rp1-20 E04F 605 riboso	Caenorhab TR6172 c15_1 1m.12478	RS20_CAEE	81.19	6.00E-55	rp1-20 E04F 605 riboso	Caenorhab PF00338.1	Ribosome	
TR5244 c0_2	-2.84	1.90E-02	rp5-25 K02 405 riboso	Caenorhab TR5244 c0_2 1m.5708	RS25_CAEE	81.67	3.00E-46	rp5-25 K02 405 riboso	Caenorhab PF03297.1	Ribosome	
TR1728 c0_1	-2.83	5.07E-03	Rp132	Rp132 SPOF	RS12_SPOF	57.81	6.00E-48	Rp132	000515 mol	Ribosome	
TR5274 c1_1	-2.81	2.01E-03	rp1-25 F54I 405 riboso	Caenorhab TR5274 c1_1 1m.5767	RL25_CAEE	78.68	1.00E-70	rp1-25 F54I 405 riboso	Caenorhab PF01248.2	Ribosome	
TR7371 c0_1	-2.80	1.08E-02	rp5-23 F28i 405 riboso	Caenorhab TR7371 c0_1 1m.20023	RS18_DAN	80.92	1.00E-89	rp5-23 F28i 405 riboso	Caenorhab PF00416.1 1 GO:0003735 mol	Ribosome	
TR1587 c0_1	-2.80	1.67E-02	rp1-28 R11I 605 riboso	Caenorhab TR1587 c0_1 1m.1345	CA1L_SARF	59.06	2.00E-120	rp1-28 R11I 605 riboso	Caenorhab PF01121.1 1 GO:0002334 mol	Apoptosis/Autophagy - animal Lysosome/Phagosome	
TR3351 c0_1	-2.79	4.17E-02	rp5-23 F28i 405 riboso	Caenorhab TR3351 c0_1 1m.3371	EF2_CAEE	97.96	7.00E-234	ef2-2 F23H E2F3	Caenorhab PF00318.1 1 GO:0005525 mol	Ribosome	
TR6055 c4_1	-2.76	3.05E-02	twy5-2c1 tRNA wbyD	Danio reric TR6055 c4_1 1m.9295	TYWS_DAN	41.05	4.00E-10	twy5-2c1 tRNA wbyD	Danio reric PF13621.1		
TR781 c0_1	-2.73	2.27E-02	rp1-25 F52D1 14-3-3-like	Caenorhab TR781 c0_1 1m.73	L43D2_CAEE	95.69	8.00E-162	rp1-25 F52D1 14-3-3-like	Caenorhab PF00244.1	Cell cycle/Hippo signaling pathway - fly MAPK signaling	
TR8931 c0_1	-2.73	3.47E-02	rp1-27 C53I 605 riboso	Caenorhab TR8931 c0_1 1m.21150	RS17_CAEE	72.8	6.00E-63	rp1-27 C53I 605 riboso	Caenorhab PF00833.1 1 GO:0003735 mol	Ribosome	
TR8995 c0_1	-2.68	2.98E-02	rp1-15 F36I 405 riboso	Caenorhab TR8995 c0_1 1m.20943	RS15_CAEE	84.56	2.00E-86	rp1-15 F36I 405 riboso	Caenorhab PF00203.1 1 GO:0003735 mol	Ribosome	
TR8864 c0_1	-2.68	3.07E-02	rp1-22 C27I 605 riboso	Caenorhab TR8864 c0_1 1m.744	RL22_CAEE	76.47	7.00E-64	rp1-22 C27I 605 riboso	Caenorhab PF01776.1 1 GO:0003735 mol	Ribosome	
TR8986 c0_1	-2.67	1.89E-02	rp1-26 F28i 605 riboso	Caenorhab TR8986 c0_1 1m.20935	RS16_CAEE	61.94	1.00E-161	rp1-26 F28i 605 riboso	Caenorhab PF00338.1	Ribosome	
TR8524 c0_1	-2.65	1.88E-02	rp1-6 R15I 605 riboso	Caenorhab TR8524 c0_1 1m.19576	RL6_CAEE	66.82	1.00E-94	rp1-6 R15I 605 riboso	Caenorhab PF01159.1 1 GO:0003735 mol	Ribosome	
TR81808 c0_1	-2.64	2.34E-02	rp1-23 B03I 605 riboso	Caenorhab TR81808 c0_1 1m.1549	RL23_CAEE	98.37	6.00E-81	rp1-23 B03I 605 riboso	Caenorhab PF00238.1 1 GO:0003735 mol	Ribosome	
TR8470 c0_1	-2.63	4.18E-02	rp1-20 E04F 605 riboso	Caenorhab TR8470 c0_1 1m.20642	RS24_TAKF	70.99	8.00E-59	rp1-20 E04F 605 riboso	Caenorhab PF01282.1 1 GO:0003735 mol	Ribosome	
TR7853 c0_1	-2.60	4.96E-02	icd-1 C56C Transcrip	Caenorhab TR7853 c0_1 1m.20293	BT29_CAEE	83.85	7.00E-100	icd-1 C56C Transcrip	Caenorhab PF01849.1	Apoptosis - fly	
TR1355 c0_1	-2.50	3.03E-02	rp1-28 R11I 605 riboso	Caenorhab TR1355 c0_1 1m.1143	MUR2_CAEE	90.51	6.00E-101	rp1-28 R11I 605 riboso	Caenorhab PF13202.1 1 GO:0005509 mol	Focal adhesion/Regulation of actin cytoskeleton/T	
TR6395 c9_1	-2.47	2.00E-02	ubq-2 Ubiquitin-c	Caenorhab TR6395 c9_1 1m.16588	UBQ2_BO	41.52	8.00E-2	ubq-2 Ubiquitin-c	Caenorhab PF00055.1 1 GO:0003735 mol	Ribosome	
TR8127 c0_1	-2.47	2.99E-02	rp1-28 R11I 605 riboso	Caenorhab TR8127 c0_1 1m.20459	FK81A_XET	72.38	3.00E-50	FK81A_XET	Caenorhab PF00254.2 1 GO:0004567 biol	Ribosome	
TR8201 c0_1	-2.42	2.74E-02	rp1-10 F10E 605 riboso	Caenorhab TR8201 c0_1 1m.17182	RL10O_CAEE	86.45	1.00E-138	rp1-10 F10E 605 riboso	Caenorhab PF00252.1 1 GO:0003735 mol	Ribosome	
TR6306 c3_1	-2.41	3.98E-04	Polyubiquitin Strongylo	Caenorhab TR6306 c3_1 1m.16554	UBQP2_STF	97.22	7.00E-69	Polyubiquitin Strongylo	Caenorhab PF00240.1 1 GO:0005515 mol	PPAR signaling pathway	
TR8868 c0_1	-2.38	3.10E-02	rp1-21 C14I 605 riboso	Caenorhab TR8868 c0_1 1m.20906	RL21_CAEE	76.88	2.00E-84	rp1-21 C14I 605 riboso	Caenorhab PF01157.1 1 GO:0003735 mol	Ribosome	
TR6245 c6_1	-1.88	2.95E-02	rp1-25 F54I 405 riboso	Caenorhab TR6245 c6_1 1m.14801	CNTNS_HU	27.27	3.00E-118	CNTNS	Caenorhab PF00047.1	Apoptosis - fly	
TR6444 c26_1	-1.86	3.78E-07	Heat shock B05 taup	Caenorhab TR6444 c26_1 1m.17960	HSF1_BOV	45.02	5.00E-54	HSF1	Caenorhab PF00047.1	GO:0003700 mol	
TR2168 c0_1	-1.85	4.07E-02	rp1-28 R11I 605 riboso	Caenorhab TR2168 c0_1 1m.1959	CA1L_SARF	59.06	2.00E-120	rp1-28 R11I 605 riboso	Caenorhab PF01121.1 1 GO:0002334 mol	Apoptosis/Autophagy - animal Lysosome/Phagosome	
TR6156 c3_1	-1.80	4.97E-02	Wdfy3	Wdfy3 KIA	WDYF3_XE	51.06	4.00E-27	Wdfy3	Caenorhab PF00216.1 1 GO:0051260 mol	Apoptosis/Autophagy - animal Lysosome/Sphingolipid	
TR6163 c19_1	-1.80	2.13E-14	CaTspn1 L1thobates	Caenorhab TR6163 c19_1 1m.12202	CATE_LUTC	26.27	1.00E-24	CaTspn1 L1thobates	Caenorhab PF00026.1 1 GO:0004490 mol	Apoptosis/Autophagy - animal Lysosome/Sphingolipid	
TR6338 c9_1	-1.70	4.85E-05	TR6338 c9_1 1m.17697							GO:0006810 biol	
TR6207 c11_2	-1.68	3.24E-16	TR6207 c11_2 1m.13438							PF01433.1 1 GO:0008237 mol	
TR1016 c1_1	-1.62	5.43E-04	TR1016 c1_1 1m.874							PF00433.1 1 GO:0008237 mol	
TR6363 c1_1	-1.59	5.21E-11	Adult-sc Scarecrow of	TR6363 c1_1 1m.3681	CUS5_ARA	48.57	2.00E-13	Adult-sc Scarecrow of	TR6363 c1_1 1m.3681	PF00239.2 1 GO:0002392 mol	
TR6278 c5_1	-1.54	2.98E-06	CaTspn1 Sarphagus	TR6278 c5_1 1m.1854	CA1L_SARF	38.75	2.00E-63	CaTspn1 Sarphagus	TR6278 c5_1 1m.1854	Apoptosis/Autophagy - animal Lysosome/Phagosome	
TR8591 c4_1	-1.35	2.99E-07	TR8591 c4_1 1m.8180							PF0431.1	
TR6287 c35_1	-1.34	2.77E-03	Cp1 f(2)5C3 Catepsin	Drosophila TR6287 c35_1 1m.15982							

TR6107c20_g1	-0.70	1.78E-02	Tena-C64; Teneurin-A Drosophila	TR6107c20_g1_1	TENA_DRO	32.69	5.00E-41	Tena-C64; Teneurin-A Drosophila	PF07974.8	
TR6116c3_g1	-0.69	5.98E-03	TR6116c3_g1_1		GAWKY_DI	38.33	1.00E-05	gw GW182 Protein Gm Drosophila	PF12938.2	
TR6242c25_g1	-0.68	2.95E-02	slc17a8 ygf Vesicular G Danio rerio	TR6242c25_g1_1	VGLU3_DA	22.8	3.00E-15	slc17a8 ygf Vesicular G Danio rerio	PF04063.1; G0:0055085tbl	
TR6286c1_g1	-0.67	4.12E-03	SHANK3 KH SH3 and m Homo sapi	TR6286c1_g1_1	SHANK3_HU	39.71	2.00E-13	SHANK3 KH SH3 and m Homo sapi	PF00023.2; G0:0055157tbl	
TR6236c3_g1	-0.67	8.11E-03	AEL00964 Ectopic P 4 Aedes aeg	TR6236c3_g1_1	ASPG_SPO	52.19	1.00E-07	N(4)(Beta) Spodopoter TR5210 c3_g1_1	PF01112.1; G0:0016737tbl	
TR6287c8_g1	-0.67	2.37E-02	TR6287c8_g1_1		TR6287c8_g1_1	m.15966				
TR6345c0_g1	-0.67	2.16E-07	TR6345c0_g1_1		TR6345c0_g1_1	m.17965				
TR6295c10_g1	-0.66	8.61E-04	fam46c3 zgc Protein FA Danio rerio	TR6295c10_g1_1	FA46A_HU	48.18	4.00E-27	FAM46A C Protein FA Homo sapi	PF07984.7	
TR6233c7_g1	-0.66	1.56E-03	TR6233c7_g1_1		TR6233c7_g1_1	m.14224				
TR6120c1_g1	-0.65	8.11E-03	TR6120c1_g1_1		TR6120c1_g1_1	m.19648				
TR6702c5_g1	-0.65	2.53E-02	TR6702c5_g1_1		TR6702c5_g1_1	m.9432				
TR6254c29_g1	-0.64	1.94E-02	Tbtk2 Bbv I TAU-ubiquitin Homo sapi	TR6254c29_g1_1	TTRK2_HU	63.49	1.00E-125	Tbtk2 Bbv I TAU-ubiquitin Homo sapi	PF00069.2; G0:0004672tbl	
TR6418c12_g1	-0.64	1.26E-02	TR6418c12_g1_1		TR6418c12_g1_1	m.19101				
TR6345c0_g2	-0.64	1.46E-06	TR6345c0_g2_1		TR6345c0_g2_1	m.17966				
TR6299c0_g1	-0.64	6.70E-04	cfad DDB, Counting f Dicytosteg	TR6299c0_g1_1	CFAD_DIC1	37.53	2.00E-74	cfad DDB, Counting f Dicytosteg	PF01121.1; G0:0008234tbl	
TR6237c8_g1	-0.63	1.72E-02	TR6237c8_g1_1		TR6237c8_g1_1	m.14773				
TR6118c0_g1	-0.63	4.12E-03	TR6118c0_g1_1		TR6118c0_g1_1	m.10640				
TR6286c1_g2	-0.62	1.36E-02	TR6286c1_g2_1		TR6286c1_g2_1	m.15891				
TR6225c0_g1	-0.62	1.52E-02	TR6225c0_g1_1		TR6225c0_g1_1	m.14026				
TR6222c10_g1	-0.60	1.38E-02	TR6222c10_g1_1		TR6222c10_g1_1	m.13944				
TR2734c0_g1	-0.60	1.13E-02	TR2734c0_g1_1		TR2734c0_g1_1	m.2433				
TR5974c30_g1	-0.60	3.04E-03	TR5974c30_g1_1		TR5974c30_g1_1	m.8053				
TR6293c0_g1	-0.60	1.02E-02	TR6293c0_g1_1		TR6293c0_g1_1	m.16060				
TR5492c1_g1	-0.60	1.16E-03	TR5492c1_g1_1		TR5492c1_g1_1	m.6192				
TR8665c0_g1	-0.60	3.51E-04	TR8665c0_g1_1		TR8665c0_g1_1	m.20784				
TR6189c29_g1	-0.59	4.66E-03	TR6189c29_g1_1		TR6189c29_g1_1	m.12890				
TR6107c29_g1	-0.59	4.66E-03	TR6107c29_g1_1		TR6107c29_g1_1	m.10350				
TR1259c0_g1	-0.59	1.29E-02	TR1259c0_g1_1		TR1259c0_g1_1	m.1065				
TR1060c1_g1	-0.59	4.43E-04	TR1060c1_g1_1		TR1060c1_g1_1	m.879				
TR6582c0_g1	-0.58	7.10E-06	TR6582c0_g1_1		TR6582c0_g1_1	m.19627				
TR6216c1_g2	-0.58	3.60E-04	TR6216c1_g2_1		TR6216c1_g2_1	m.13639				
TR6069c1_g2	-0.57	1.06E-02	TR6069c1_g2_1		TR6069c1_g2_1	m.9353				
TR5950c20_g1	-0.57	3.45E-04	TR5950c20_g1_1		TR5950c20_g1_1	m.1391				
TR6240c38_g1	-0.57	4.23E-02	TR6240c38_g1_1		TR6240c38_g1_1	m.14524				
TR6282c9_g1	-0.57	4.84E-02	TR6282c9_g1_1		TR6282c9_g1_1	m.15796				
TR6197c15_g1	-0.56	2.25E-04	TR6197c15_g1_1		TR6197c15_g1_1	m.13125				
TR6183c6_g1	-0.56	6.61E-03	TR6183c6_g1_1		TR6183c6_g1_1	m.12697				
TR473c0_g2	-0.56	1.21E-02	TR473c0_g2_1		TR473c0_g2_1	m.431				
TR6259c13_g1	-0.56	5.44E-02	TR6259c13_g1_1		TR6259c13_g1_1	m.16129				
TR6384c7_g2	-0.55	6.37E-03	TR6384c7_g2_1		TR6384c7_g2_1	m.18661				
TR6116c3_g2	-0.55	4.76E-03	TR6116c3_g2_1		TR6116c3_g2_1	m.10569				
TR254c0_g1	-0.54	3.93E-02	TR254c0_g1_1		TR254c0_g1_1	m.219				
TR6425c11_g1	-0.53	1.02E-02	TR6425c11_g1_1		TR6425c11_g1_1	m.19161				
TR2687c0_g1	-0.53	2.90E-02	TR2687c0_g1_1		TR2687c0_g1_1	m.2401				
TR6111c22_g1	-0.53	1.09E-02	TR6111c22_g1_1		TR6111c22_g1_1	m.15721				
TR6111c22_g2	-0.53	1.09E-02	TR6111c22_g2_1		TR6111c22_g2_1	m.15721				
TR4148c0_g1	-0.53	3.08E-02	TR4148c0_g1_1		TR4148c0_g1_1	m.4025				
TR6255c23_g1	-0.53	2.16E-03	TR6255c23_g1_1		TR6255c23_g1_1	m.3519				
TR265c0_g1	-0.52	9.66E-05	TR265c0_g1_1		TR265c0_g1_1	m.2319				
TR6019c2_g1	-0.52	5.72E-03	TR6019c2_g1_1		TR6019c2_g1_1	m.8350				
TR6035c5_g1	-0.52	1.07E-02	TR6035c5_g1_1		TR6035c5_g1_1	m.17433				
TR6249c9_g1	-0.52	3.70E-05	TR6249c9_g1_1		TR6249c9_g1_1	m.14902				
TR6294c19_g1	-0.52	1.72E-02	TR6294c19_g1_1		TR6294c19_g1_1	m.13051				
TR6197c21_g1	-0.51	4.75E-04	TR6197c21_g1_1		TR6197c21_g1_1	m.16198				
TR6324c34_g1	-0.51	2.38E-02	TR6324c34_g1_1		TR6324c34_g1_1	m.17209				
TR6058c7_g1	-0.51	4.78E-02	TR6058c7_g1_1		TR6058c7_g1_1	m.9136				
TR6220c18_g1	-0.50	2.15E-02	TR6220c18_g1_1		TR6220c18_g1_1	m.13863				
TR3745c1_g1	-0.50	7.19E-03	TR3745c1_g1_1		TR3745c1_g1_1	m.3573				
TR5956c13_g1	-0.50	1.75E-02	TR5956c13_g1_1		TR5956c13_g1_1	m.1890				
TR6344c2_g1	-0.50	7.97E-02	TR6344c2_g1_1		TR6344c2_g1_1	m.17930				
TR1942c0_g1	-0.50	1.58E-02	TR1942c0_g1_1		TR1942c0_g1_1	m.1656				
TR6800c8_g1	-0.50	2.85E-07	TR6800c8_g1_1		TR6800c8_g1_1	m.9674				
TR111c0_g1	-0.46	4.05E-04	TR111c0_g1_1		TR111c0_g1_1	m.1053				
TR3694c0_g1	-0.44	1.01E-02	TR3694c0_g1_1		TR3694c0_g1_1	m.3513				
TR6437c11_g1	-0.44	7.00E-11	TR6437c11_g1_1		TR6437c11_g1_1	m.13215				
TR6408c0_g2	-0.43	1.79E-06	TR6408c0_g2_1		TR6408c0_g2_1	m.19043				
TR6309c3_g1	-0.43	1.52E-10	TR6309c3_g1_1		TR6309c3_g1_1	m.16709				
TR6261c19_g1	-0.43	9.34E-05	TR6261c19_g1_1		TR6261c19_g1_1	m.15268				
TR6408c0_g1	-0.43	3.63E-06	TR6408c0_g1_1		TR6408c0_g1_1	m.19042				
TR6383c4_g1	-0.43	1.29E-29	TR6383c4_g1_1		TR6383c4_g1_1	m.18637				
TR6389c0_g1	-0.43	3.00E-11	TR6389c0_g1_1		TR6389c0_g1_1	m.20606				
TR1187c0_g1	-0.42	1.98E-06	TR1187c0_g1_1		TR1187c0_g1_1	m.1000				
TR6138c15_g1	-0.42	2.33E-04	TR6138c15_g1_1		TR6138c15_g1_1	m.11330				
TR6110c14_g2	-0.42	1.21E-07	TR6110c14_g2_1		TR6110c14_g2_1	m.10415				
TR6241c5_g1	-0.42	1.18E-08	TR6241c5_g1_1		TR6241c5_g1_1	m.14546				
TR2567c0_g2	-0.41	1.80E-04	TR2567c0_g2_1		TR2567c0_g2_1	m.2945				
TR3128c0_g1	-0.41	1.12E-06	TR3128c0_g1_1		TR3128c0_g1_1	m.2945				
TR5949c5_g1	-0.41	5.17E-05	TR5949c5_g1_1		TR5949c5_g1_1	m.7669				
TR111c1_g1	-0.40	1.08E-02	TR111c1_g1_1		TR111c1_g1_1	m.107				
TR6299c30_g1	-0.40	3.00E-02	TR6299c30_g1_1		TR6299c30_g1_1	m.16319				
TR6057c18_g1	-0.40	1.72E-04	TR6057c18_g1_1		TR6057c18_g1_1	m.9102				
TR6080c0_g1	-0.40	1.03E-08	TR6080c0_g1_1		TR6080c0_g1_1	m.9652				
TR2117c0_g2	-0.40	1.35E-03	TR2117c0_g2_1		TR2117c0_g2_1	m.1853				
TR6080c0_g2	-0.40	1.03E-08	TR6080c0_g2_1		TR6080c0_g2_1	m.9653				
TR6383c1_g2	-0.40	1.02E-78	TR6383c1_g2_1		TR6383c1_g2_1	m.18632				
TR6383c1_g1	-0.40	1.09E-03	TR6383c1_g1_1		TR6383c1_g1_1	m.18631				
TR6141c41_g1	-0.38	2.68E-02	TR6141c41_g1_1		TR6141c41_g1_1	m.11527				
TR5970c25_g1	-0.37	6.42E-11	TR5970c25_g1_1		TR5970c25_g1_1	m.7990				
TR5949c6_g1	-0.37	1.72E-02	TR5949c6_g1_1		TR5949c6_g1_1	m.7612				
TR6351c0_g2	-0.37	6.42E-02	TR6351c0_g2_1		TR6351c0_g2_1	m.18236				
TR6295c24_g1	-0.34	6.53E-06	TR6295c24_g1_1		TR6295c24_g1_1	m.16141				
TR9233c0_g1	-0.34	2.44E-07	TR9233c0_g1_1		TR9233c0_g1_1	m.21095				
TR6097c22_g1	-0.34	8.45E-04	TR6097c22_g1_1		TR6097c22_g1_1	m.10182				
TR5664c1_g1	-0.32	9.00E-06	TR5664c1_g1_1		TR5664c1_g1_1	m.6647				
TR6336c31_g1	-0.32	2.32E-03	TR6336c31_g1_1		TR6336c31_g1_1	m.17622				
TR6320c2_g1	-0.32	4.76E-02	TR6320c2_g1_1		TR6320c2_g1_1	m.1326				
TR6108c11_g2	-0.29	2.04E-08	TR6108c11_g2_1		TR6108c11_g2_1	m.10377				
TR2592c0_g1	-0.29	4.40E-02	TR2592c0_g1_1		TR2592c0_g1_1	m.2291				
TR6136c8_g1	-0.28	2.20E-02	TR6136c8_g1_1		TR6136c8_g1_1	m.11206				
TR6108c11_g3	-0.28	8.89E-08	TR6108c11_g3_1		TR6108c11_g3_1	m.10379				
TR6195c28_g1	-0.27	1.25E-08	TR6195c28_g1_1		TR6195c28_g1_1	m.13093				
TR6155c11_g1	-0.27	3.70E-05	TR6155c11_g1_1		TR6155c11_g1_1	m.12029				

TR6205 c10_g1	0.66	3.72E-03	TR6205 c10_g1_1		TR6205 c10_g1_1 m.13343			PF0100.11		
TR6243 c9_g1	0.65	2.74E-02	TR6243 c9_g1_1	DERP3 DE	35.07	5.00E-38	DERP3	Mite allerg Dermatop	PF00089.2:G0:0004252/mol	
TR264 c0_g1	0.65	1.79E-03	TR264 c0_g1_1	HSP11_CAI	40.43	1.00E-10	hsp-16.1 h	Heat shock Caenorhab	PF00011.11	
TR6075 c11_g1	0.64	2.38E-05	TR6075 c11_g1_1	TENS3_MC	53.69	5.00E-34	Tns3 Tens1 Tensin-3	T Mus musc	PF00017.2:G0:0005515/mol	
TR3688 c0_g1	0.64	1.55E-02	TR3688 c0_g1_1	TTIN_HUA	25.17	2.00E-22	sls ttin CG: Titin [D-Tit	Drosophila	PF00047.21	
TR6131 c3_g1	0.64	1.31E-02	TR6131 c3_g2_1	TTIN_DRO	26.3	1.00E-22	sls ttin CG: Titin [D-Tit	Drosophila	PF00047.21	
TR6344 c2_g1	0.64	2.79E-03	TR6344 c2_g1_1	CP9C1_DR1	29.03	4.00E-22	Cyp9c1 CG: Cytochrom	Drosophila	PF00067.1:G0:0005506/mol	
TR6050 c12_g1	0.64	7.01E-04	TR6050 c12_g1_1				TR6050 c12_g1_1 m.8951	PF01607.1:G0:0008061/mol		
TR6190 c0_g1	0.63	1.67E-02	TR6190 c0_g1_1	CLCN2_CA'	59.21	3.00E-20	CLCN2	Chloride ch Cavia porc	TR6190 c0_g1_1 m.12894	
TR6190 c0_g2	0.61	2.57E-02	TR6190 c0_g2_1	CLCN2_CA'	59.21	3.00E-20	CLCN2	Chloride ch Cavia porc	TR6190 c0_g2_1 m.12895	
TR6285 c3_g1	0.61	5.97E-03	TR6285 c3_g1_1				TR6285 c3_g1_1 m.15864			
TR6315 c3_g1	0.61	4.06E-04	TR6315 c3_g1_1	LN10_CAE	58.79	2.00E-129	lin-10 C09F: Protein lin-	Caenorhab	TR6315 c3_g1_1 m.16856	
TR6344 c4_g2	0.61	1.55E-04	TR6344 c4_g2_1				TR6344 c4_g2_1 m.17934			
TR6206 c27_g1	0.61	2.10E-03	TR6206 c27_g1_1	TGFR1_BO	41.56	7.00E-15	TGFR1	TGF-beta r Bos taur	TR6206 c27_g1_1 m.13407	
TR6131 c3_g2	0.60	3.41E-03	TR6131 c3_g2_1	LN10_CAE	58.03	2.00E-130	lin-10 C09F: Protein lin-	Caenorhab	TR6131 c3_g2_1 m.16857	
TR6277 c4_g1	0.60	1.18E-04	TR6277 c4_g1_1				TR6277 c4_g1_1 m.15730			
TR423 c1_g1	0.60	4.17E-02	TR423 c1_g1_1	CBPB_ASTJ	39.87	2.00E-64	0 Carboxype	Astacus ac	TR423 c1_g1_1 m.4211	
TR6147 c7_g1	0.60	1.15E-03	TR6147 c7_g1_1	NRFE_CAE	20.81	4.00E-13	nrf-6 C08B: Nose resist	Caenorhab	TR6147 c7_g1_1 m.11769	
TR6416 c17_g1	0.60	4.62E-05	TR6416 c17_g1_1	F214A_XEP	35	7.00E-32	fam214a s1 Protein FA	Danio rer	TR6416 c17_g1_1 m.13077	
TR6180 c40_g1	0.60	4.63E-06	TR6180 c40_g1_1	UBQL1_RA	81.82	4.00E-12	Ubqln1 Da- Ubiquilin-1	Rattus nor	TR6180 c40_g1_1 m.12263	
TR1217 c0_g1	0.60	1.74E-05	TR1217 c0_g1_1	CATS_CAN	33.66	5.00E-31			TR1217 c0_g1_1 m.1028	
TR6131 c4_g1	0.60	7.98E-05	TR6131 c4_g1_1				TR6131 c4_g1_1 m.11037			
TR6155 c1_g2	0.59	3.10E-03	TR6155 c1_g2_1	LIPE_AMOU	27.3	2.00E-19	LIggs	Endothelia Mus musc	TR6155 c1_g2_1 m.12200	
TR6145 c7_g2	0.59	8.56E-05	TR6145 c7_g2_1	MFD5A_DA	24.57	5.00E-16	mfsd5a mf Major facil	Danio rer	TR6145 c7_g2_1 m.11667	
TR2592 c0_g1	0.59	2.95E-02	TR2592 c0_g2_1	ENTK_HUA	29.37	1.00E-18	TPMRSS15	Enteropog	TR2592 c0_g2_1 m.2292	
TR6222 c5_g1	0.59	3.87E-06	TR6222 c5_g1_1	SCD5_BOV	54.36	3.00E-103	SCD5	Stearoyl-Cr Bos taur	TR6222 c5_g1_1 m.13929	
TR6301 c6_g1	0.58	1.68E-04	TR6301 c6_g1_1				TR6301 c6_g1_1 m.16390			
TR6235 c7_g2	0.58	8.73E-05	TR6235 c7_g2_1	HDACA_CH	47.28	4.00E-126	HDACA	Histone de Gallus gall	TR6235 c7_g2_1 m.14308	
TR6385 c8_g1	0.58	1.89E-04	TR6385 c8_g1_1	LIPI_HUMU	29.15	2.00E-12	LIPI	Lipase mer Homo sapi	TR6385 c8_g1_1 m.18702	
TR6301 c28_g1	0.58	2.63E-04	TR6301 c28_g1_1	SOX14_DA	87.18	7.00E-41	sox14 zgc:1	Transcript	Danio rer	TR6301 c28_g1_1 m.16424
TR6139 c13_g1	0.57	3.65E-05	TR6139 c13_g1_1	UMD_DICC	35.63	4.00E-07	limD limD1 LIM domal	Dicystostel	TR6139 c13_g1_1 m.11380	
TR6344 c4_g1	0.57	8.33E-05	TR6344 c4_g1_1				TR6344 c4_g1_1 m.17932			
TR6385 c4_g1	0.57	1.33E-05	TR6385 c4_g1_1	NCAH_DRC	85.48	4.00E-107	Nca CG764 Neurocalci	Drosophila	TR6385 c4_g1_1 m.18692	
TR6416 c13_g1	0.57	6.03E-04	TR6416 c13_g1_1	SW5_DR0F	58.11	2.00E-156	sws CG221	Neuropath	TR6416 c13_g1_1 m.13073	
TR5990 c1_g2	0.57	4.65E-04	TR5990 c1_g2_1				TR5990 c1_g2_1 m.8144			
TR6234 c0_g1	0.57	5.66E-06	TR6234 c0_g1_1	RPTOR_MC	37.74	5.00E-43	Rptor Rapt	Regulatory Mus musc	TR6234 c0_g1_1 m.17175	
TR9193 c0_g1	0.56	6.65E-03	TR9193 c0_g1_1				TR9193 c0_g1_1 m.21066			
TR6138 c17_g1	0.56	3.33E-05	TR6138 c17_g1_1	MIR_BOM	51.97	1.00E-46	0 Myosin reg Bombyx m		TR6138 c17_g1_1 m.11331	
TR6139 c12_g1	0.56	7.15E-04	TR6139 c12_g1_1	KCC1A_HU	71.79	4.00E-10	CAMK1	Calcium/ca Homo sapi	TR6139 c12_g1_1 m.17920	
TR5487 c1_g1	0.56	3.67E-03	TR5487 c1_g1_1	ACYP2_HU	40	3.00E-19	ACYP2	ACY Acylphosphol	TR5487 c1_g1_1 m.6180	
TR6248 c11_g1	0.56	1.61E-04	TR6248 c11_g1_1	RDHEZ_HU	44.73	1.00E-77	SDR16C5 R	Epidermal Homo sapi	TR6248 c11_g1_1 m.14882	
TR6512 c4_g1	0.56	3.62E-04	TR6512 c4_g1_1	ASTE1_POI	24.56	1.00E-14	Aste1	Protein ast Mus musc	TR6512 c4_g1_1 m.12970	
TR5418 c0_g1	0.55	8.35E-05	TR5418 c0_g2_1	TM163_BO	33.92	1.00E-22	TMEM163	Transmem Homo sapi	TR5418 c0_g2_1 m.6009	
TR5418 c0_g2	0.55	8.29E-05	TR5418 c0_g1_1	TM163_HU	33.92	1.00E-22	TMEM163	Transmem Homo sapi	TR5418 c0_g1_1 m.6008	
TR6343 c9_g2	0.55	8.75E-04	TR6343 c9_g2_1	CMK1_CAE	68.61	8.00E-152	cmk-1 CBG Calcium/ca	Caenorhab	TR6343 c9_g2_1 m.17906	
TR92 c0_g1	0.55	7.93E-03	TR92 c0_g1_1				TR92 c0_g1_1 m.81			
TR6298 c3_g1	0.55	3.71E-02	TR6298 c3_g1_1	ELF1_DR0F	49.45	7.00E-19	grh eF1 CG	Protein gra Drosophila	TR6298 c3_g1_1 m.16224	
TR6071 c4_g2	0.55	6.02E-05	TR6071 c4_g2_1	CAH14_MC	31.29	9.00E-35	Ca14 Car14	Carbonic a Mus musc	TR6071 c4_g2_1 m.9385	
TR6286 c9_g2	0.55	6.64E-03	TR6286 c9_g1_1	GSC1L_MO	38.14	5.00E-15	Gtsrcl1 Kia GLTSCR1 h	Mus musc	TR6286 c9_g1_1 m.15909	
TR6286 c9_g1	0.55	1.82E-04	TR6286 c9_g1_1	GSC1L_MO	38.14	7.00E-15	Gtsrcl1 Kia GLTSCR1 h	Mus musc	TR6286 c9_g1_1 m.15906	
TR6145 c7_g3	0.54	2.08E-04	TR6145 c7_g3_1	CFRPA_DA	24.57	6.00E-16	mfsd5a mf Major facil	Danio rer	TR6145 c7_g3_1 m.11668	
TR6080 c3_g1	0.54	1.82E-02	TR6080 c3_g1_1	CMTA1_HL	45.61	1.00E-08	CAMTA1 K	Calmodulin Homo sapi	TR6080 c3_g1_1 m.9663	
TR6324 c28_g1	0.54	8.45E-04	TR6324 c28_g1_1	DORS_DRC	53.82	4.00E-108	dj CG6667	Embryonic Drosophila	TR6324 c28_g1_1 m.17205	
TR5904 c0_g2	0.53	1.47E-02	TR5904 c0_g2_1	CALCR_RAI	32.59	1.00E-09	CALCR	Calcitonin r Cytocloga	TR5904 c0_g2_1 m.7421	
TR6150 c5_g1	0.53	1.27E-04	TR6150 c5_g1_1	ZASP_DR0	32.89	5.00E-22	Zasp52 Zas PDZ and LI	Drosophila	TR6150 c5_g1_1 m.11891	
TR6242 c2_g1	0.53	7.72E-03	TR6242 c2_g1_1	SMC6_MO	41.58	3.00E-16	Smc6 Kiaa4 Structural i	Mus musc	TR6242 c2_g1_1 m.14590	
TR6292 c6_g2	0.53	3.04E-03	TR6292 c6_g2_1				TR6292 c6_g2_1 m.16017			
TR6238 c15_g1	0.53	2.49E-03	TR6238 c15_g1_1	S16C6_BOI	38.6	2.00E-53	SDR16C6	Short-chain Bos taur	TR6238 c15_g1_1 m.14407	
TR3322 c1_g1	0.53	6.02E-05	TR3322 c1_g1_1	AQP9_MO	31.73	3.00E-29	Aqp9	Aquaporin Mus musc	TR3322 c1_g1_1 m.3129	
TR5969 c7_g1	0.53	1.56E-04	TR5969 c7_g1_1	HSPB1_CH	47.67	2.00E-16	HSPB1	Heat shock Gallus gall	TR5969 c7_g1_1 m.7925	
TR6222 c11_g2	0.53	7.32E-04	TR6222 c11_g2_1	FAESA_DA	45.81	3.00E-97			TR6222 c11_g2_1 m.13949	
TR6378 c4_g1	0.52	1.97E-02	TR6378 c4_g1_1	LPN_HUM	23.91	9.00E-15	LPN	LIPN	TR6378 c4_g1_1 m.18523	
TR4871 c0_g1	0.52	1.82E-03	TR4871 c0_g1_1	PAHX_HUA	25.42	3.00E-17	PHYH PAH	Phytanoyl- Homo sapi	TR4871 c0_g1_1 m.50531	
TR5969 c6_g1	0.52	4.04E-02	TR5969 c6_g1_1	NFIL_PIGW	78.95	6.00E-84	NFI	Nuclear fac Sus scrofa	TR5969 c6_g1_1 m.7923	
TR5954 c2_g2	0.52	8.57E-05	TR5954 c2_g2_1	CBPM_MO	48.91	2.00E-107	Cpm	Carboxype Mus musc	TR5954 c2_g2_1 m.7718	
TR6244 c2_g2	0.52	4.64E-04	TR6244 c2_g2_1	ABCG8_DR	30.08	3.00E-51	abcg20 DD ABC	transp Dicotyostel	TR6244 c2_g2_1 m.14711	
TR6131 c0_g1	0.52	1.61E-04	TR6131 c0_g1_1				TR6131 c0_g1_1 m.12564			
TR6085 c18_g1	0.52	5.17E-05	TR6085 c18_g1_1				TR6085 c18_g1_1 m.9822			
TR6297 c23_g1	0.51	4.51E-04	TR6297 c23_g1_1	MTG1_KTI	39.34	3.00E-57	mtg1	Mitochond itakurap s	TR6297 c23_g1_1 m.16199	
TR6228 c7_g1	0.51	2.17E-03	TR6228 c7_g1_1	PERC_ANO	36.75	4.00E-97	pxt AGAPO	Chorion pe Anopheles	TR6228 c7_g1_1 m.14082	
TR1101 c0_g1	0.51	1.14E-02	TR1101 c0_g1_1	ALKMO_RA	45.48	1.00E-123	Agmo Tme Alkylglycer	Rattus nor	TR1101 c0_g1_1 m.918	
TR6068 c0_g2	0.51	1.83E-04	TR6068 c0_g2_1	LIPIH_DANI	24.76	7.00E-14	liph zgc:91	Lipase mer Danio rer	TR6068 c0_g2_1 m.9317	
TR5954 c2_g1	0.51	1.07E-04	TR5954 c2_g1_1	CBPM_MO	48.91	2.00E-107	Cpm	Carboxype Mus musc	TR5954 c2_g1_1 m.7717	
TR6163 c12_g1	0.50	1.02E-02	TR6163 c12_g1_1	MLXPX_RA	41.79	3.00E-42	Mlxlp1 Wbs Carbolydr	Rattus nor	TR6163 c12_g1_1 m.12192	
TR6139 c4_g3	0.50	9.06E-03	TR6139 c4_g3_1				TR6139 c4_g3_1 m.11358			
TR5455 c1_g1	0.50	6.34E-04	TR5455 c1_g1_1	CP3AC_CA	26.89	3.00E-34			TR5455 c1_g1_1 m.6086	
TR6137 c15_g1	0.50	1.22E-04	TR6137 c15_g1_1	PERC_ANO	34.68	3.00E-92	pxt AGAPO	Chorion pe Anopheles	TR6137 c15_g1_1 m.11271	
TR6286 c8_g2	0.50	1.65E-02	TR6286 c8_g2_1	SCR11_HU	62.16	2.00E-60	SCR11	Transcripti Mus musc	TR6286 c8_g2_1 m.15904	
TR6191 c7_g1	0.50	3.72E-03	TR6191 c7_g1_1	MACF1_RA	31.98	1.00E-136	Macf1 Acf1	Microtubu Rattus nor	TR6191 c7_g1_1 m.12926	