RNA sequencing to characterize transcriptional changes of sexual maturation and mating in the female oriental fruit fly Bactrocera dorsalis

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Abstract

Background: Female reproductive potential plays a significant role in the survival and stability of species, and sexual maturation and mating processes are crucial. However, our knowledge of the reproductive genes involved in sexual maturation and mating has been largely limited to model organisms. The oriental fruit fly Bactrocera dorsalis is a highly invasive agricultural pest, known to cause major economic losses; thus, it is of great value to understand the transcriptional changes involved in sexual maturation and mating processes as well as the related genes. Here, we used a high-throughput sequencing method to identify multiple genes potentially involved in sexual maturation and mating in female B. dorsalis.

Results: We sequenced 39,999 unique genes with an average length of 883 bp. In total, 3264 differentially expressed genes (DEGs) were detected between mature virgin and immature Bactrocera dorsalis libraries, whereas only 83 DEGs were identified between flies that had mated or were mature virgins. These DEGs were functionally annotated using the GO and KEGG pathway annotation tools. Results showed that the main GO terms associated with the DEGs from the mature virgin vs. immature groups were primarily assigned to the metabolic and developmental processes, which we focused on, whereas those from the mated vs. mature virgin group largely belonged to the response to stimulus and immune system processes. Additionally, we identified multiple DEGs during sexual maturation that are involved in reproduction, and expression pattern analysis revealed that the majority DEGs detected were highly enriched in those linked to the ovaries or fat bodies. Several mating responsive genes differentially expressed after mating were also identified, and all antimicrobial peptides detected were highly enriched in fat body and significantly up-regulated approximately 2- to 10-fold at 24 h after mating.

Conclusion: This study supplied female reproductive genes involved in sexual maturation and the post-mating response in B. dorsalis, based on RNA-seq. Our data will facilitate molecular research related to reproduction and provide abundant target genes for effective control of this agricultural pest.

Keywords: RNAseq, Transcriptome, Tephritid, Maturity, Post-mating, Reproduction, Gene expression

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Background
Tephritid fruit flies, especially those belonging to the genus *Bactrocera*, are globally important due to their destructive impact on agriculture. Their high reproductive ability is an important factor that leads to enhanced survival rates and potential to multiply. Reproductive biology has been well studied in tephritid species, including *Bactrocera tryoni* (Froggatti), *Bactrocera cucurbitae* (Coquillett) and *Ceratitis capitata* (Wiedemann), but most reports have mainly focused on the processes at the physiological and behavioral levels [1–3]. *Bactrocera dorsalis* (Hendel) (the oriental fruit fly), which is one among the important *Bactrocera* species, is a highly invasive agricultural pest that is currently distributed across most Asian countries and in a number of Pacific Islands. Adults can lay eggs on various types of host plants, where the hatched larvae subsequently feed and cause crop loss [4]. Some factors that regulate the fecundity and mating behavior of *B. dorsalis* females have been previously studied, such as methyl eugenol-fed males and multiple mates, etc. [5]. Although the *B. dorsalis* transcriptome from different developmental stages of the whole body as well as specific tissue in males is available on NCBI, large-scale molecular analysis of reproduction in *B. dorsalis* females remains limited [6–10]. As an invasive agricultural pest with a wide host range and high fecundity, it is essential to investigate the reproductive biology of *B. dorsalis* females at the molecular level in order to formulate simple and effective strategies for agricultural pest control.

Recently, transcriptional changes in several genes that may be involved in female sexual maturity and mating were identified in *C. capitata* based on microarray data [11]; however, the cDNA library of *C. capitata* is limited to the head of female adult. Additionally, it is worth noting that the reproductive strategy of *B. dorsalis* is different from that of *C. capitata*. For instance, medfly reach sexual maturity rapidly within 2–3 days after emergence, whereas it takes approximately 2 weeks for the oriental fruit fly to complete the transformation after emergence to maturity [12, 13]. Therefore, our study is planned because both species have different mating strategy and that the methodology followed in previous study provided only limited resources.

Female reproductive potential significantly contributed to the multiply and stability of species, which mainly depends on the process of sexual maturation and mating [14]. In most holometabolous insects, adults emerge from the pupal stage as sexually immature individuals. After they ingest a food source, they become sexually mature within several days dependent on the species [14]. During the development of the oocyte, the ovary becomes mature and then when females reach sexual maturity they began to mate and oviposit [15]. Until now, studies on sexual maturity have mainly concentrated on the model species whose genomes have been sequenced except *C. capitata*, including *Drosophila melanogaster*, *Apis mellifera*, *Aedes aegypti* and *Blattella germanica* [16–20]. However, these reports on model species mainly focused on the protein vitellogenin and information on other proteins that play roles in reproduction is lacking. While genes involved in female adult reproductive maturation, including EcR, USP and InR, have been identified in *Tribolium castaneum* using microarrays, they are mainly the early genes in the signal transduction pathway, and little is known about the other genes, especially the late or effect genes [21].

Successful mating behavior and the following egg-laying is essential for sexual reproduction in insect, especially for its multiply. Female reproductive process has been revealed to be greatly affected by male reproductive gland secretion during copulation. Reports in *C. capitata* and *Bactrocera tryoni* showed the effect of mating on remating propensity, refractory period and fecundity of females [22–24]. In *D. melanogaster*, it was reported that mating regulated ovulation and spawning as well as other processes in mated females, such as, inducing innate immunity, acceleration of reproductive maturation and egg production [25–28]. The sex peptide receptor, which mediates the post-mating switch in females, has been reported in *D. melanogaster*, *Helicoverpa armigera* and *B. dorsalis* [4, 29–31]. However, the other factors involved in this process are still largely unknown. Genome-wide research into the post-mating response in females has been performed in model species such as *D. melanogaster*, the honeybee queen *Apis mellifera*, and *C. capitata*, which have revealed that the post-mating response differs among species as well as the time following mating [11, 25, 28, 32–34]. Regardless, there is no such information available for *B. dorsalis*, and the transcriptional changes and genes involved in this process is also still unknown.

In the present study, the transcriptome of the whole body of the female *B. dorsalis* were sequenced at three different physiological stages (immature, mature virgins and mated individuals) using the Illumina HiSeq 2500 system, and assembled based on our previously established transcriptome of *B. dorsalis* [6]. Furthermore, analysis of the DEGs between mature virgins and immature individuals was performed in order to identify the transcripts involved in the process of sexual maturation. An additional comparison between mated individuals and mature virgins was also carried out to identify the genes that respond to mating. Tissue specific expression profiles of the differentially expressed genes from these two comparisons were analyzed by qRT-PCR. Finally, the expression levels of the genes involved in the post-mating response at different time points were also obtained.
Methods
Insect culture
B. dorsalis were cultured in our lab under a 12 h light/dark cycle at 27 °C. Adult flies were reared on artificial diets composed of 25 % yeast extract and 75 % sugar [35].

Sample preparation
Newly emerged adults were collected and sexed within 12 h after emergence, and both sexes were subsequently maintained in separate cages. As male and female oriental fruit flies require 14 days to reach sexual maturity under standard rearing conditions, 1- to 2-day-old (1–2 days) individuals were considered immature while 14-day-old (14 days) individuals were considered sexually mature. Twenty-four hours after emergence, the immature female flies were chilled briefly and RNA was extracted from the whole bodies of 10 individuals using TRIzol regent (Invitrogen, Carlsbad, CA, USA). Similarly, mature virgin samples were collected 14 days after emergence and total RNA was isolated from the whole bodies of 10 mature virgin females.

To obtain mated flies, approximately 100 fourteen-day-old virgin flies of each sex were placed into a 30 cm² cage shortly before dark as B. dorsalis mating occurs during dusk [36]. When copulation was observed, mating pairs were isolated and placed in small vials about 14 cm in width and 23 cm in height with 11.5 cm diameter. Notably, only pairs that mated for at least 90 min were used in further experiments in order to avoid false mating with little or no sperm transfer. Copulation terminated naturally and the mated pairs were placed in separate cages according to sex. RNA was extracted from the whole bodies of 10 mated females the following day (24 h after mating) with the same procedure as was used for the virgin flies. RNA samples were treated to remove any contaminating DNA with the DNA-free kit (Ambion, Foster City, CA, USA).

Sequencing
Total RNA from each sample was isolated using TRIzol reagent (Invitrogen) following the manufacturer’s instructions. The concentration and integrity of total RNA were determined using the 2100 Bioanalyzer (Agilent, Santa Clara, CA, USA). Total RNA was then incubated with 10 U DNase I (Takara, Shiga, Japan) at 37 °C for 30 min, and purified using the Dynabeads® Oligo (dT)25 (Thermo Fisher, Waltham, MA, USA) following the manufacturer’s instructions. To ensure the accuracy of the results, we conducted two independent replicates for each sample, specifically immature (1, 2), mature virgin (1, 2) and mated (1, 2) individuals. For each case, the following validation described below was performed.

For cDNA library construction 100 ng purified mRNA was used using NEBNext® UltraTM RNA Library Prep Kit for Illumina sequencing (New England Biolabs, Ipswich, MA, USA). First-strand cDNA was synthesized using Protoscript II Reverse Transcriptase. After incubation, Second Strand Synthesis Enzyme Mix was added for the synthesis of the second strand of the cDNA. Double-stranded cDNA was then purified using AMPure XP Beads (Beckman Coulter, Brea, CA, USA), followed by an end repair step using End Prep Enzyme Mix. After adaptor ligation and further purification, the cDNA library was obtained. To verify the quality of the library, it was checked using three different methods, including Qubit (Thermo Fisher) quantitative analysis, 2 % agarose gel electrophoresis, and high-sensitivity DNA chip determination.

The cDNA libraries were then used for cluster generation with the TruSeq PE Cluster Kit (Illumina, San Diego, CA, USA) and sequenced on an Illumina HiSeq® 2500 instrument with paired-end sequencing.

Differential expression analysis
For the identification of genes that were possibly involved in reproduction, those that were differentially expressed at the various physiological stages of adults were analyzed. Two comparisons of the expression level of paired sets of the three samples were performed between mature virgin vs. immature and mated vs. mature virgin individuals. The number of reads for contigs from each sample was first converted to Reads per Kilobase per Million (RPKM), and then the MARS model (MA-plot-based method with Random Sampling model) in
the DEGseq package was used to calculate the expression abundance of each contig [40]. A false discovery rate (FDR) < 0.001 was considered to indicate significant expression abundance. Additionally, differentially expressed genes were further used for GO and KEGG analysis to identify the pathways that the DEGs were predicted to be involved in. Relativity analysis was performed to evaluate the stability and reliability of the replicates. GOEAST was used for GO term enrichment analysis [41]. The relative significant level of enrichment of the differently expressed gene to all genes was calculated with hypergeometric statistical test method, then the GO term whose FDR < 0.001 was considered to be significant. For pathway enrichment analysis, a similar method was used. The KEGG pathway was considered as a unit, and its relative significant level of enrichment of the differently expressed pathway to whole genome was calculated.

Quantitative real-time PCR (qRT-PCR) verification
Selected DEG data was validated by qRT-PCR using the SYBR Premix Ex Taq kit (Takara) according to the manufacturer’s instructions with a real-time thermal cycler (Bio-Rad, Hercules, CA). TRIzol reagent (Invitrogen) was used to extract total RNA from B. dorsalis at three different physiological stages, including the immature (newly emergent within 24 h), mature virgin (14-day-old before mating), and mated (14-day-old post-mating). At least 10 insects were collected for each sample. The first strand cDNA was obtained from 2 μg of total RNA using M-MLV Reverse Transcriptase (Takara) with the primer oligo-anchor R (5′-GACCACCGGTATCGATGTCGAC T16 (A/C/G)-3′). The primers used for qRT-PCR detection are listed in Additional file 1: Table S1. The relative gene expression data were analyzed using the 2^ΔΔCt method as described by Zheng et al. [42]. The results were analyzed using a one-way analysis of variance (ANOVA) statistical test. All quantitative PCR experiments were repeated in three biological replicates.

Tissue specific expression pattern analysis of DEGs involved in reproduction
Total RNA was extracted from different tissues of the mature female adults, including the head, thorax, ovary, fat body and midgut. The tissue samples were dissected from 30 individuals. After the cDNA was obtained, qRT-PCR was performed according to the methods described above. A total of eleven genes differentially expressed during maturation and six AMPs whose transcript abundance changed after mating were used in this study.

Change in transcriptional abundance of mating-responsive genes
To investigate the expression level of mating-responsive genes before and after mating, newly emergent individuals (1 day after emergence) of each sex were separated and the mated pairs were obtained as described above. RNA was extracted from the head or fat body of mated females 1, 12, 24 h after mating. At least 30 individuals were used for each sample and qRT-PCR experiments were repeated in three biological replicates.

Results and discussion
De novo sequence assembly
The cDNA samples from different stages were sequenced, and these raw data (Additional file 2: Table S2) were assembled into 39,999 unique genes (Additional file 3: Table S3). The mean contig size was 883 bp with lengths ranging from 201 to 27,791 bp (Additional file 3: Table S3 and Additional file 4: Figure S1). The contig size distribution revealed that more than half of the contigs (23,645; 59.11 %) were between 200 and 500 bp in length, whereas 34.88 % (13,951) were between 500 and 3000 bp in length (Additional file 4: Figure S1).

Gene ontology and clusters of orthologous group classification
Gene ontology (GO) analysis was performed for functional categorization of unigenes (Additional file 5: Figure S2). Within the biological processes, the main GO terms were grouped into cellular (3362; 19 %) and metabolic (2696; 15 %). As for the molecular function category, binding (3663; 50 %) constituted the largest group, followed by catalytic activity (2096; 29 %). The primary reason for this difference might be that the metabolic activities of insects during larval development is greater than in the adult stage. Many complicated physiological processes occurs during molting and metamorphosis, such as a molting cascade similar to the larval molt, histolysis of larval tissues, remodeling and formation of adult tissues, etc. [43, 44].

Assignments of clusters of orthologous groups (COG) were performed to further evaluate the completeness of the transcriptome and the effectiveness of our annotation process. The annotated sequences were grouped into 25 major functional classes (Additional file 6: Figure S3). Among them, the majority of the clusters were “Signal transduction mechanisms” (795; 13.01 %), “General function prediction only” (742; 12.14 %), whereas “Cell motility” (17; 0.28 %) and “Nuclear structure” (28; 0.46 %) represented the smallest groups (Additional file 6: Figure S3). Our transcriptome will aid further research focused on signal transduction in tephritid flies.
Transcriptional changes during female maturation

To identify transcripts that may play a role in female maturation, differentially expressed gene sequences between the mature and immature libraries were identified with a total of 3264 DEGs (FDR < 0.001, Fig. 1 & Additional file 7: File S1). Among them, 2567 transcripts were more abundant in the mature females, while 697 were more abundant in immature females. Our results are different from those revealed by the adult transcriptome of *C. capitata* obtained with a microarray, in which only 811 transcripts displayed significant changes during female maturation. Furthermore, in the transcriptome of *C. capitata*, the transcripts with greater abundance were found in the immature females (462) instead of the mature individuals (349) [11]. These differences may be attributed to the use of different sequencing methods and the different sexual maturation time utilized by these two species. For example, *B. dorsalis* is sexually mature after 14 days, while *C. capitata* can mate after only 4 days [12, 13]. Therefore, more genes are possibly implicated in the sexual maturation in *B. dorsalis*.

Analysis of enriched biological process GO terms with significant differences (p < 0.05) during female maturation was performed. We focused on the physiological processes that appear to play essential roles in maturation, including reproductive, signaling, developmental, metabolic, immune processes as well as response to stimulus (Fig. 2a). A table of the identified genes is provided in the supplementary data (Additional file 8: Table S4). Results revealed that differentially expressed genes in the metabolic and developmental processes are a much larger group than in other processes, which is similar to the results in *C. capitata* that indicated the most enriched terms in mature females were categorized in the metabolic process group [11]. These data indicate that a variety of metabolic activities and developmental affairs usually occur during oogenesis and ovary maturation. Additionally, enrichment analysis of the KEGG pathways for the differentially expressed genes during maturation was also performed. Results showed that the number of differentially expressed genes in the metabolism pathways are the largest groups, including Carbohydrate metabolism, Energy metabolism, Nucleotide metabolism, Lipid metabolism, Metabolism of cofactors and vitamins, Amino acid metabolism, etc. (Additional file 9: File S2). This is similar with the result of GO enrichment analysis.

Transcriptional changes of mating-responsive genes in females

To identify transcripts that may have been involved in female mating, differentially expressed genes between the mated and mature virgin females were identified. Out of 83 transcripts that displayed significant transcriptional changes between virgin mature and mated females 24 h post-mating, 65 (78 %) were more abundant in mated females, while only 18 (22 %) were more abundant in virgin mature females (FDR < 0.001, Fig. 1 & Additional file 7: File S1). This is similar with the result found in *C. capitata* and *D. melanogaster* [11, 27, 28]. Only 32 and 28 transcripts were altered in abundance 24 h after mating in *C. capitata* and *D. melanogaster*, respectively. However, in a study of *Drosophila* using DNA microarray analysis, 539 transcripts are differentially expressed in the lower reproductive tracts [25]. One reason for the variable changes in transcription could be that like previous studies by Lawniczak and Begun [28] in *Drosophila*, our study focused on broader changes of gene expression in the whole body, while a study by Mack et al. [25] in *Drosophila* concentrated on the lower reproductive tracts, where significant changes take place following mating. Furthermore, the time point for sample collection after mating in both our study and that of the medfly was 24 h [11], that in the Lawniczak and Begun [28] analysis was 1–3 h, whereas the Mack et al. [25] examined several time points (0, 3, 6 and 24 h post-mating). It was demonstrated in previous studies that post-mating transcriptional changes are highly variable at different time points, thus possibly explaining transcriptional differences among these studies [25].

Statistically significant differences (p < 0.05) among enriched biological process GO terms during mating included those involved in the response to stimulus, as well as immune system, developmental, cellular, biological regulation and metabolic processes. Of these transcripts, those that are related to the response to stimulus and immune system process are the largest group (Fig. 2b). A table of these genes is provided in the supplementary data (Additional file 10: Table S5).
Similar results were found in *A. mellifera* where the genes involved in stress response were significantly regulated following mating [45]. Additionally, an immune response was stimulated by mating in *Drosophila*, and it might arise from sexually antagonistic interaction between the sexes [46, 47]. Furthermore, enrichment analysis of KEGG pathways for the mating responsive genes was also performed (Additional file 9: File S2). Our results showed that the metabolism pathways were the main groups, including carbohydrate metabolism, transport and catabolism, glycan biosynthesis and metabolism, lipid metabolism, metabolism of cofactors and vitamins. The different results between KEGG enrichment and GO enrichment might be caused by the different database used in the analysis.

**Genes involved in oogenesis and ovary maturation**

From DEG analysis of the samples from immature (newly eclosed) vs. mature virgin (sexually mature before mating) individuals, we identified seven genes that reportedly participate in oogenesis and ovary maturation in insects (Additional file 8: Table S4). Interestingly, two genes that were previously found to play important roles in Wnt signaling pathways and an additional two involved in sex determination were also identified in this DEG library [48–51]. The subsequent qRT-PCR validated this result (Fig. 3). All of these genes were significantly highly expressed at the sexually mature and post-mating stage, which is consistent with our transcriptome data, indicating that they are involved in physiological processes of mature females as is the case in other insects [52–54] (Fig. 3). Additionally, we analyzed expression of these genes in several tissues to elucidate their roles and found that most are highly enriched in the ovary, including *hu-li tai shao* (*hts*), *oskar*, *mago nashi*, *yolkless* (*yl*), *disheveled*, *axin*, *sry* and *transformer-2* (Fig. 4). The expression level of *oskar* in the ovary is approximately 50 fold higher than that in the fat body and 300 fold in the midgut. Notably, three *vitellogenin* genes can be detected in the head, thorax, ovary, fat body, while they are more abundantly expressed in fat body (Fig. 4). These results are explained below by analyzing their roles determined for other insects.

*hts* was reported to modulate actin polymerization in neurons, sharing a conserved function with mammalian adducing in actin-capping activity [55]. One isoform of *hts*, Ovhts-RC, was shown to be distributed along a cortical anterior-posterior gradient during *Drosophila* late-oogenesis to modulate spatially restricted actin filament growth at the oocyte cortex while alterations in Ovhts-RC led to actin overgrowth in the oocyte [52]. A previous report indicated that *oskar* participated in regulating pole plasm assembly, and a long form of *oskar* played an important role in yolk endocytosis and F-actin projection at the posterior pole [56]. Recently, it was found that *oskar* functions upstream of D-EndoB in the polarized endocytic activity of the yolk uptake process in *Drosophila* oocytes [57]. It was also found that *mago nashi* interacts with Tsunagi/Y14 and other proteins, forming a multiprotein complex to play an essential role in localization of *oskar* mRNA within the oocyte in *Drosophila* [58]. Recent analyses demonstrated that *mago nashi* forms a complex with Tsunagi/Y14 and Ranshi to influence oocyte differentiation within the posterior pole of the presumptive oocyte in *Drosophila* [53]. *Vitellogenin*, also known as *yolk protein*, has been well studied in insects, especially in *Drosophila, Aedes aegypti, Anopheles gambiae* and *C. capitata* [54, 59–62]. In these species, it is expressed in the fat body, secreted into the hemolymph and subsequently sequestered by oocytes to facilitate the
transport of carbohydrates, lipids and other nutrients to the ovaries [63]. In the transcriptome we obtained, three vitellogenin genes were identified, including vitellogenin 1, vitellogenin 2 and vitellogenin 2 precursor, all of which are up-regulated in sexually mature females, indicating that they are essential for Bactrocera sexual maturation. The gene yolkless, commonly known as the vitellogenin receptor, belongs to the apolipoproteins receptor (LDLR) superfamily [64]. Drosophila yolkless is expressed very early during the development of the oocyte at both transcriptional and translational level, long before vitellogenesis begins. Previous studies found that yolkless mutants failed
to incorporate yolk proteins during oogenesis [65]. Our findings are validated in the qRT-PCR results, revealing that these genes were enriched in the ovary or fat body.

Furthermore, we found disheveled and axin (isoform A) in our transcriptome, both of which are core components of Wnt signaling pathways (Additional file 8: Table S4). They are also required for cell movement and FAK regulation during ovarian morphogenesis, which may explain why both genes are abundantly expressed in mature *B. dorsalis* females [48] (Fig. 3).

Notably, two genes involved in sex determination were also identified, including sex-determining region y protein (*sry*) and transformer-2 (*tra-2*). It has been reported that *sry* plays a role in mammalian sex determination, reduced or delayed *sry* expression impairs testis development. This indicates a potential role of *sry* dysregulation in human intersex disorders [49]. Additionally, *tra-2* was also reported to be an essential switch for the sex determination cascade in dipteran insect species [50, 51]. Knockdown of *tra-2* leads to the production of male-only progeny in *B. dorsalis* [66]. In the present study, we identified these two genes and found they were up-regulated in sexually mature females compared to their newly eclosed counterparts, which is the first
evidence that sex determination related genes may also
function during sex maturation.

**Genes involved in the post-mating response of females**
Upon mating, significant expression changes occurred in
genes involved in the immune or stress response (Fig. 2b).
The immune response stimulated by mating might arise
from sexually antagonistic interaction between the sexes
[27, 28, 46, 47]. Several up-regulated genes from the DEG
analysis showed significant increases in transcript abundance
in females, including two PRRs (peptidoglycan-recognition
protein SB1-like and peptidoglycan-recognition protein LB-like) and six antimicrobial peptides (AMPs)
(defensin, diptericin, phormicin-like, sapecin, cecropin-1 and
attacin-C-like) (Additional file 10: Table S5). To investigate
the expression pattern of these genes, the expression levels
of AMPs in different tissues were analyzed by qRT-PCR.
The results showed that all were highly enriched in the fat
body, which is usually an important organ involved in the
immune response in insects (Fig. 5). The expression pattern
of these genes indicates that they may participate in the im-
mune response after mating in *B. dorsalis*.

To further validate, the changes in expression at different
time points after mating were analyzed in both the
fat body and head. All genes except *attacin-C-like* and
*phormicin-like* were significantly up-regulated 24 h after
mating in the fat body approximately 2–10 fold (Fig. 6a).
This is partly supported by the previous studies in *D.
melanogaster*, which showed that the immune response
is activated in mated females by mating and associated
sex peptides transfer [28, 46, 47]. Upon mating, potent
signal molecules from the males enter the females and
the behavior and physiology of the mated females shifted
from reproductive receptivity to fecundity. The immune
response was induced in the females and it might arise
from sexually antagonistic interaction between the sexes
[45, 47]. However, this is different from the results of the
previous report on *C. capitata*, which revealed no sig-
nificant changes of transcript abundance were detected
in immune-related genes after mating, except for a large
reduction in *defensin* in the female abdomen [11]. This
data suggests that post mating changes differ among
species [33]. Additionally, the expression level of most
AMPs did not significantly change during the time
course in the female head 24 h post-mating, except for
*sapecin* and *cecropin-1*, both of which significantly in-
creased in transcript abundance, by 2.3 and 3.0 fold,
respectively (Fig. 6b). It is clear that the fat body is the
main tissue where the immune response takes place and
not the head.

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**Fig. 5** Expression pattern of the immune-related genes between mated females and the mature virgin individuals in different tissues, including the head, thorax, ovary, fat body and midgut. Relative transcript levels of six antimicrobial peptide are calculated by real-time PCR using the 16 s rRNA gene as the standard control. Different letters indicate significant differences (p < 0.05, Duncan’s test) among samples and three independent biological replicates were performed for each sample.
Fig. 6 The effect of mating on the expression level of immune-related genes in the fat body (a) and head (b) of B. dorsalis females. RNA was extracted from the head or fat body of mature virgin females as well as mated females at 1, 12, 24 h after mating. Relative transcript levels of six antimicrobial peptide are calculated by real-time PCR using the 16 s rRNA gene as the standard control. Various letters represent significant differences of the expression level of genes (p < 0.05, Duncan's test). Three biological replicates were performed.
Conclusions
This study significantly advances the characterization of the sexual maturation and mating process at the molecular level in the oriental fruit fly, a major invasive agricultural tephritid pest. Numerous transcripts displayed significant changes in abundance during maturation of the female oriental fruit fly and generally mirrored the physiological and behavioral alterations in female flies. Multiple reproductive genes were differentially expressed between mature virgin and immature individuals, notably including two genes involved in sex determination, possibly indicating they have important roles during sexual maturation. However, modest post-mating changes were also observed in females. The difference in transcriptional changes pertaining to sexual maturation and mating, compared to C. capitata, is perhaps a result of different reproductive strategies employed by these species.

Our data reveals the existence of additional potential molecular targets for B. dorsalis control, and may aid in future studies of reproductive mechanisms in fruit flies. Furthermore, our findings provide a comprehensive resource for exploring the complex molecular mechanisms driving the evolution of reproductive processes, and will likely facilitate the development of effective and eco-friendly pest control strategies.

Availability of supporting data
This Transcriptome Shotgun Assembly project has been deposited at DDBJ/EMBL/GenBank under the accession GEEAO0000000. Other data sets supporting the results of this article are included in the additional files.

Additional files

Additional file 1: Table S1. Primers used for qRT-PCR. (DOC 36 kb)
Additional file 2: Table S2. Summary of the Solexa sequencing statistics of B. dorsalis adult transcriptome. (DOCX 14 kb)
Additional file 3: Table S3. Summary of the B. dorsalis adult transcriptome. (DOCX 12 kb)
Additional file 4: Figure S1. Assembled contig length distribution of the transcriptome. The x-axis shows contig size and the y-axis indicates the number of contigs for each given size. (TIF 1138 kb)
Additional file 5: Figure S2. Gene ontology (GO) analyses of transcriptome sequences predicted to participate in (A) biological processes and (B) molecular functions. Data are presented as level 3 GO categorization for molecular function and level 2 GO categorization for biological process. Classified gene objects are showed as percentages (in brackets) of the total number according to the GO assignments. (ZIP 1448 kb)
Additional file 6: Figure S3. Histogram presentation of Clusters of orthologous groups (COG) classification. A total of 6111 predicted unigenes were classified in 25 COG categories. (TIF 1448 kb)
Additional file 7: File S1. Genes differentially expressed between mature virgin females and immature individuals, and between mated females and mature virgins. Reads, RPKM and the fold values for each comparison group (mature virgin vs. immature, and mated vs. mature virgin) is shown. (XLS 674 kb)

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
The study was conceived and designed by WWZ. DYL and FYW prepared the oriental fruit fly samples. FYW conducted the RNA extraction. WWZ performed the qRT-PCR. WWZ, JLW and HYZ analyzed the transcriptomic data. WWZ prepared the initial draft of the manuscript and HYZ modified the final version. All authors have read and approved the final version of the manuscript.

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