



# Comparative genomics and transcriptomics of host–pathogen interactions in insects: evolutionary insights and future directions

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## **Comparative genomics and transcriptomics of host-pathogen interactions in insects: evolutionary insights and future directions**

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### **Abstract**

Classical evolutionary studies of protein-coding genes have established that genes in the canonical immune system are often among the most rapidly evolving within and between species. As more genomes and transcriptomes across insects are sequenced, it is becoming clear that duplications and losses of immune genes are also a likely consequence of host-pathogen interactions. Furthermore, particular species respond to diverse pathogenic challenges with a wide range of challenge-specific responses that are still poorly understood. Transcriptional studies, using RNA-seq to characterize the infection-regulated transcriptome of diverse insects, are crucial for additional progress in understanding the ecology and evolution of the full complexity of the host response.

### **Highlights**

- Expansions and losses of immune genes are an important component of evolutionary change across species
- Transcriptional response to infection involves diverse processes beyond canonical immune pathways

- Careful RNA-seq studies across multiple species are needed to understand how inducible responses evolve

## Introduction

Genes involved in immune defense have long been recognized as hotspots for rapid evolution in many organisms including insects [1,2]. In *Drosophila*, early population genetic [3,4] and comparative genomic [5,6] studies demonstrated that key components of the innate immune system experienced substantially more adaptive protein evolution than typical genes in the genome. More recently, a combination of functional and comparative analysis showed a probable role for balancing selection in the maintenance of sequence diversity in antimicrobial peptides in *Drosophila* [7,8]. While *Drosophila* has received the most research attention, evidence for positive selection in insect immune genes is common in other groups as well [9–11]. These studies have established rapid, adaptively driven sequence evolution of immune proteins as a fundamental tenet of insect immunity [12,13].

Over the past decade, increasing evidence has accumulated that adaptive changes in protein sequence are far from the only important evolutionary dynamics occurring in insect immune systems. Comparisons of gene content in the immune system between different insect orders [14–16], distantly related dipterans [17] and more closely related drosophilids [6] all suggest substantial gains (often via gene duplication) and losses of immune system genes, especially outside signaling pathways. While determining the functional impact of these changes is not easy, transcriptomics provides a way forward, especially in non-model taxa where genetic tools are not readily accessible [18].

Transcriptional studies have been particularly important for understanding the full complexity of the insect immune response. The canonical innate immune system of insects consists of a set of receptor molecules that detect infection (usually via pathogen-associated or danger-associated molecular patterns) and trigger (generally) conserved signaling cascades (in particular Toll, imd, JAK/STAT, and JNK pathways) that ultimately serve to control the

transcription of a variety of downstream effectors [19]. Early microarray studies in *Drosophila melanogaster* [20–23] were crucial in establishing this picture of insect immunity, although as discussed below recent evidence suggests this may be substantially incomplete. Most early transcriptome studies were limited to model systems, due in large part to the requirement that microarrays need to be designed based on known transcript sequences. With the advent of RNA-sequencing, this limitation was removed, and in the past several years transcriptional studies of non-model insects have become extremely common [18] (Supplemental Table 1).

In this review, I discuss several insights into insect immunity that have been facilitated by transcriptional and genomic studies, focusing on their evolutionary implications. First, I will review recent work on the evolutionary dynamics of immune systems beyond sequence evolution, focusing on gene duplication, gene loss, rewiring of signaling pathways, and changes in the transcriptional response to infection. While this area of study is still in its infancy, increasingly affordable sequencing is poised to make these studies widespread. Second, I review the role of transcriptome studies in revising our understanding of the complexity of the transcriptional response to infection, both within and between species. Infection results in differential regulation of a wide array of pathways outside canonical immune genes, a process that RNA-seq studies have made abundantly clear, although whether these represent novel mechanisms of tolerance or resistance, or physiological consequences of infection with little or no benefit to the host, remains an open question in most cases. Taken together, these lines of research suggest that we are still far from understanding the entire process of how immune systems evolve and function.

### **Beyond protein evolution: evolutionary dynamics of gene content in insect immune systems**

While most studies of rapid evolution in insect immune genes have focused on identifying adaptive changes in protein sequence, changes in copy number and/or gene content (e.g., via gene duplication, gene loss, or *de novo* gene origination) are often of selective importance [24,25] and can be associated with functionally important traits in insects [26,27]. Early comparative genomic studies in holometabolous insects pointed to deep conservation of

canonical signaling components (the Toll, Imd, JAK/STAT, and JNK signal transduction pathways), while also suggesting that upstream recognition and downstream effector genes may be more likely to change copy number or experience species-specific expansion [6,14,17,28]. In recent years, dramatic increases in the number of sequenced insect genomes, as well as increasing feasibility of direct measures of infection-regulated genes using RNA-seq, have added substantial new details to this picture.

### *Depauperate immune systems in insects?*

Insects have a diverse array of life histories and ecologies, including some that have been proposed to be associated with reduced investment in individual immune defense. Most prominently, social insects engage in a variety of behaviors to reduce the spread of infection in colonies, such as removal of infected individuals from colonies (“social immunity”, [29]). Other insects, such as aphids, have microbial symbionts that may provide defense against pathogens [30,31]. If these forms of immunity reduce the need for individual defense pathways, it is reasonable to hypothesize that, as a consequence, genes required for immune defense may be lost or pseudogenized. Indeed, the first genome-wide studies in honeybees [14] and pea aphids [16] showed evidence for reduced number of homologs of canonical immune genes, based on comparisons to genes annotated in *Drosophila* (Figure 1). The honeybee genome, while containing homologs of all major signaling pathways identified in Diptera, encodes far fewer copies of several families of recognition and effector genes [14]. The pea aphid genome is missing homologs for many components of the Imd signaling pathway, in addition to a dramatic reduction in effector gene family content and complete loss of peptidoglycan recognition proteins (PGRPs) and other recognition proteins [16].

Hymenoptera now have among the most sequenced genomes of any insect order [32], and additional genomes of both social and solitary species have revealed that a reduced complement of homologs of dipteran immune genes does not appear to be associated with sociality [15,33,34] (Figure 1). Although many fewer termite genomes are available, similar lack of differentiation in immune gene content is apparent between termites on different extremes

of sociality [35]; as more Blattodea genomes become available [36] these comparisons will have increasing power.

Comparisons that focus primarily on homologs of canonical immune genes first identified in dipterans may not represent the full picture of hymenopteran immunity. Transcriptome studies using RNA-seq to characterize genes regulated by infection showed that a substantial number of infection-regulated genes in the parasitic wasp *Nasonia vitripennis* [37], and in multiple ant species [38–40], do not have clearly identifiable homologs in dipterans. This suggests that the observation of reduced numbers of homologs of dipteran immune genes in bees and other hymenopterans may, in part, be an artifact of homology-based annotation and the evolutionary distance between Diptera and Hymenoptera. The *de novo* evolution of Hymenoptera-restricted immune genes, the recruitment of existing genes to have an immune role in Hymenoptera, or rapid sequence evolution hindering our ability to recognize homolog could all produce a reduced annotation in distant species. Any of these models are plausible, although extreme caution is warranted in interpreting lack of detectable homologs as *prima facie* evidence for *de novo* gene origination [41,42]. Recruitment of existing genes to a novel immune role has been shown to occur in tetrapods: recent comprehensive RNA-seq study of the interferon response across ten tetrapods (nine mammals plus chicken) revealed, in addition to a core set of conserved interferon-responsive genes, evidence for the evolution of new transcriptional responsiveness across the phylogeny [43].

In the pea aphid, genomic evidence suggesting the absence of the Imd signaling pathways and a number of recognition proteins, including a complete absence of PGRPs [16] is complemented by some functional genomic data. Early transcriptome studies using suppression subtractive hybridization and EST screens failed to identify induced antimicrobial peptides in pea aphids [44]; proteomic screens of hemolymph after artificial infection with *E. coli* also did not detect strong evidence for inducible antimicrobial peptides [16]. Furthermore, pea aphids do not appear to pay a physiological cost from exposure to natural bacterial pathogens, supporting a role for a limited immune response to bacteria [45].

Nonetheless, the sequencing of genomes and transcriptomes of a number of additional hemimetabolous insects and more diverged arthropods has revealed that absence of Imd

pathway components, and reductions or absence of PGRPs, are not uncommon (Figure 1). Indeed, few arthropods outside holometabolous insects appear to have a completely intact Imd pathway, and many are missing all transmembrane PGRPs or all PGRPs entirely [46–49]. Although the existing data does not definitively resolve whether the Imd pathway arose and acquired an immune function in the ancestor of holometabolous insects [46], or represents an older pathway independently lost in many lineages [47], it appears clear that large-scale reorganization in the identity of major immune pathways is plausible across long evolutionary timescales. Ultimately, the combination of more complete genomic representation outside Holometabola and direct measures of infection-regulated transcription using RNA-seq in diverse insects will be required to fully understand the degree to which particular species truly have reduced repertoires of immune genes in their genomes.

#### *Expansion of innate immune genes*

In many cases, genomic and transcriptomic studies in insects have revealed evidence for expanded repertoires of immune genes, often through gene duplication and typically involving genes encoding recognition or effector proteins. The idea that host-pathogen arms races can drive copy number changes in critical genes of the immune system has a long history [50,51]. In insects, comparative genomic comparisons showed that effector genes in particular were prone to diversification across dipterans [6,17]. These observations support biochemical evidence for lineage-restricted antimicrobial peptides, including diapausin [52] and gambicin [53], although simple absence of homology can also be explained by rapid sequence evolution, especially in short proteins.

A particularly striking example of immune gene expansion is in the Harlequin ladybird *Harmonia axyridis*, a species native to central and eastern Asia that has become highly invasive in many regions, to the point of displacing native ladybird competitors. A *de novo* transcriptome assembly of *H. axyridis* after infection with several immune elicitors identified more than 50 putative antimicrobial peptides, a very large number for any insect [54]. Comparative transcriptomics with close relatives further showed that many of these AMPs are specifically

expanded in *H. axyridis* [55]. Functional studies have shown that *H. axyridis* has a more potent antimicrobial defense response than the native *Coccinella septempunctata* [56], and it is resistant to a parasitic microsporidian that is lethal to native competitors [55,57]. While transgenic manipulations to test the role of the expanded immune repertoire directly have not been possible in ladybirds, comparative evidence is strongly suggestive that the observed immune gene expansions are functionally related to improved defense against pathogens and parasites, potentially facilitating the invasive potential of *H. axyridis* [55,57] (but see [58]).

Gene duplications and immune gene family expansions have also proposed to arise as a consequence of insect ecology, particularly in response to high pathogen burdens. Increased copy number can both increase the speed of humoral immune response and facilitate functional diversification, albeit potentially increasing autoimmune or other costs of resistance. A transcriptome study using EST sequencing in drone fly maggots, which live in highly septic, contaminated water, showed a particularly high diversity of AMP transcripts [59]. This led the authors to hypothesize that environments particularly rich in bacteria may lead to selection for increased diversity of immune components. More recently, both genomic and RNA-seq studies from *Musca domestica* -- an insect that also inhabits particularly septic environments both as larvae and adults -- revealed unusually large expansions of certain immune gene families, including the thioester-containing proteins and the cecropin AMP family [60,61] (Figure 1). Further work characterizing immune gene family evolution in more species with diverse ecologies and associated bacterial communities will continue to refine this hypothesis.

### **Complexity of the transcriptional response to infection**

The advent of RNA-sequencing technology more than a decade ago has had a major impact on the study of insect immunity, with over 75 studies looking for differential gene expression after an infectious challenge published to date (Supplemental Table 1). In general, this wide range of studies has focused on annotating immune pathways in species (often of agriculture importance) without high quality genomes (e.g., [62,63]), elucidation of host-pathogen interactions of ecological interest (e.g., [38,64–66]), and characterization of



pathogen-vector interactions of medical relevance (e.g., [67,68]). There has been relatively less focus on explicitly comparative studies, either comparing multiple pathogens in the same species, or comparative studies across species, albeit with a few exceptions (e.g., [38,55,69–74]).

The degree to which the transcriptional response to infection varies across different pathogenic challenges in the same species is a critical question for understanding host-pathogen interactions in an ecological and evolutionary context. Two key recent studies – in *D. melanogaster* [70] and in *A. mellifera* [75] – have tackled this question in detail. While a number of microarray and RNA-seq studies have been done in *D. melanogaster* (e.g., [20,23]), previous work has typically focused on one or a few model bacteria, often non-pathogenic. To overcome this limitation, Troha and colleagues [70] infected *D. melanogaster* with ten different bacterial species across a range of pathogenicity, including several bacterial strains that have been recovered from wild fruit flies, and used RNA-seq to measure the transcriptional response to infection. They showed that, while a core of canonical immune genes are induced in most conditions, a substantial number of consistently regulated genes were involved in cellular and metabolic homeostasis, some of which may have functions related to immune tolerance [76], while others may reflect physiological consequences of infection. Furthermore, the majority of genes regulated in any particular infection were not part of the core, universally regulated response, implying a high degree of host-pathogen specificity in the transcriptional response to infection, largely outside the well understood immune pathways.

In honeybees, Doublet and colleagues [75] re-analyzed published microarray and RNA-seq studies examining the transcriptional response to infection with a variety of different pathogens. This work identified many similar patterns, albeit with reduced power and precision due to the need to rely on a heterogeneous collection of previously published studies varying in quality, as opposed to generating a consistent new dataset as in [70]. Despite these limitations this work also revealed both a core set of commonly regulated genes, including upregulation of canonical immune pathways and downregulation of metabolic genes, and a set of pathogen restricted genes enriched for functions like apoptosis.

Similar results have been seen in previous studies, including comparisons of different viral infections in *Aedes* mosquitos [72,73,77], comparisons of diverse pathogenic challenges in *Culex* mosquitoes [78], and comparisons between different fungal pathogens in ants [38]. Taken together, these studies imply that the canonical immune response is only a small part of the full suite of transcriptional changes associated with infection, and furthermore that a substantial fraction of the insect response to infection is highly specific to particular host-pathogen (or host-microbe) interactions. Importantly, it remains an open question the extent to which the broad transcriptional response to infection outside canonical immune pathways represents novel mechanisms of tolerance or resistance. Infection is likely to perturb a number of cellular and physiological processes, and much of the wider transcriptional response may reflect instead the physiological burden of disease. Ultimately, additional functional work (e.g., gene knockouts as in [70]) will be required to conclusively establish the role of genes transcriptionally regulated by infection.

While transcriptomic studies are increasingly using ecologically relevant pathogens, less attention has focused on comparative studies across species. Theoretical and computational advances in explicitly phylogenetic approaches for modeling gene expression data [79–81] are maturing and offer important benefits over pairwise tests, which can be misleading [82]. While incorporating ecologically relevant infectious challenges into comparative work will lead to better understanding of the full degree of evolutionary divergence in infection-regulated gene expression, these studies will also pose an experimental design challenge to keep costs and sample sizes manageable. Previous work has often used few or no biological replicates, potentially to increase experimental design complexity while minimizing costs, but this severely reduces both power and reproducibility [83,84]. Lack of adequate reference genomes also poses a challenge to comparative work, as methods for *de novo* transcriptome assembly often produce highly fragmented and duplicated reference transcriptomes that perform poorly for differential expression analysis in the absence of significant post-processing [85]. Fortunately, initiatives such as the i5K project [86] and declining sequencing costs are rapidly increasing the number of assembled insect genomes available, and new approaches such as Tag-seq [87], in

which only a short tag for each gene is sequenced, are making large-scale sequencing experiments including adequate replication increasingly feasible.

## Conclusions

Both genomic and transcriptomic studies have contributed substantially to our understanding of how insect immune systems evolve. Over the past decade, these studies have shown that, while adaptive evolution of protein sequence in canonical immune genes are clearly important, changes in gene content at short and long time scales, as well as transcriptional responses outside canonical pathways, are also significant components of the eco-evolutionary dynamics of host-pathogen interactions. Insects, with a rapidly expanding number of sequenced genomes and a vast arrange of host ecologies and life histories, will continue to provide new insights into host-pathogen evolutionary dynamics.

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## Figure Legends

### **Figure 1. Summary of changes in gene content across insect immune systems.**

Cladogram of major insect genera with characterized immune systems (topology following [88]). Status of major signaling pathways is indicated in the far right column: dark purple = complete, with paler shades proportional to the number of missing genes. Clades missing the Imd gene are marked with a red "X". Counts of key effector and recognition gene families are shown for a subset of species. Blocks are colored orange/red if counts are higher than typical and blue if lower than typical; gray indicates not analyzed. Data are summarized from [15,16,46,47,61]. All silhouettes from phylopic.org.

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