Social immunity in honeybees (Apis mellifera): transcriptome analysis of varroa-hygienic behaviour

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Abstract

Honeybees have evolved a social immunity consisting of the cooperation of individuals to decrease disease in the hive. We identified a set of genes involved in this social immunity by analysing the brain transcriptome of highly varroa-hygienic bees, who efficiently detect and remove brood infected with the Varroa destructor mite. The function of these candidate genes does not seem to support a higher olfactory sensitivity in hygienic bees, as previously hypothesized. However, comparing their genomic profile with those from other behaviours suggests a link with brood care and the highly varroa-hygienic Africanized honeybees. These results represent a first step toward the identification of genes involved in social immunity and thus provide first insights into the evolution of social immunity.

Keywords: honeybee, social immunity, hygienic behaviour, Varroa, gene expression.

Introduction

In insects, defence against pathogens relies mainly on an efficient innate immunity that is comprised of both cellular and humoral reactions (e.g. phagocytosis, melanization and secretion of antimicrobial peptides) (Hoffmann, 2003; Siva-Jothy et al., 2005). However, when living in groups, as do social insects, the presence of stored resources and the close living quarters increase both the attractiveness for pathogens and disease transmission (Schmid-Hempel, 1998). Therefore, higher capacities to respond to and defend against pathogens may be expected. However, a genome-wide analysis of immunity in the honeybee Apis mellifera showed that they possess only one-third of the number of immune response genes known for solitary insects (e.g. fruit fly, mosquito and moth) (Evans et al., 2006). As the reduction in the number of genes involved the different steps of the immune response, honeybees appear to have a reduced capacity to respond to and defend against pathogens.

Despite the wide range of pathogens to which social insects are exposed, they successfully resist disease, suggesting that other defence mechanisms may be involved. Indeed, in addition to individual defences, social insects have developed group-level strategies against parasites and pathogens. Such social immunity includes grooming, the use of antimicrobial materials for nest construction (e.g. resin) (Christe et al., 2003; Simone et al., 2009), social fever (Starks et al., 2000) and nest hygiene (see Cremer et al., 2007, for a review). Since their description, many studies have explored the behavioural mechanisms of these collective immune defences against pathogens (see the following reviews: Cremer et al., 2007; Cremer & Sixt, 2009; Wilson-Rich et al., 2009), but the molecular basis and pathways remain largely unknown.

The identification of genes that influence social immunity would not only improve our understanding of its mechanisms but also provide new insights into the evolution of collective defence in insect societies. In the honeybee genome, genes involved in social immunity might have replaced genes from individual immunity that have been
lost during evolution of sociality and might be key factors for defence against diseases.

In honeybees, a well-known behavioural trait to fight against pathogens is hygienic behaviour, which involves the identification and removal of dead or infected larvae. Using the honeybee genome, we attempted to identify genes involved in this well-characterized behaviour, a main component of social immunity. Hygienic behaviour is directed toward dead brood, but also those in the brood infected with bacteria or fungi (Boecking & Spivak, 1999), the greater wax moth, Galleria mellonella (Villegas & Villa, 2006; Corrêa-Marques & De Jong, 1998), the small hive beetle, Aethina tumida (Ellis et al., 2003; Neumann & Härtels, 2004) or the mite Varroa destructor, the parasite with the most pronounced effect on honeybee colonies. Indeed, development of varroa populations, which reproduce in brood cells, often leads to the death of a colony (Le Conte et al., 2010). As hygienic behaviour is genetically controlled (Rothenbuhler, 1964a,b), selective breeding for varroa resistance through increased varroa-hygienic behaviour offers a sustainable means for controlling mite parasitism amongst the other factors contributing to a stable parasite-host relationship (Rosenkranz et al., 2010). Accordingly, different varroa-hygienic bee lines have been successfully bred (Boecking & Spivak, 1999; Harbo & Harris, 1999, 2005a; Spivak & Reuter, 2001a; Büchler et al., 2010) with one of these characterized by a low percentage of reproducing varroa mites in the hive. Bees from these colonies display effective removal of varroa-infested pupae from capped brood cells, which limits the varroa infestation rate and reproduction (Harbo & Harris, 2005b, 2009). This genetic line, called varroa-sensitive hygiene (VSH), thus represents a good model for uncovering genes involved in social immunity. Harbo & Harris (2005a) suggested only a few genes to be involved in VSH behaviour; however, the molecular basis still remained to be deciphered. We therefore directly compared brain-specific gene expression profiles of bees selected for their high rate of hygienic behaviour (VSH+) to bees displaying a low rate of hygienic behaviour (VSH−). We used a honeybee oligonucleotide microarray, based on gene predictions and annotation from the honeybee genome sequencing project (Honeybee Genome Sequencing Consortium, 2006). Finally, in order to characterize the VSH trait further, we compared the brain gene expression profile of VSH bees to genomic profiles from other well-defined behavioural phenotypes.

Results

VSH behaviour

Four VSH+ and four VSH− colonies were selected from 24 colonies of a breeding programme to present high or low rates of varroa-sensitive hygienic expression. The percentage of varroa-infected brood removed by workers in each colony is shown in Table 1.

VSH genes

A total of 39 transcripts was found to be differentially expressed in the brains of VSH+ and VSH− bees at a false discovery rate < 0.05 (Table 2). Amongst them, 14 were significantly up-regulated and 25 down-regulated in VSH+. The magnitude of the differences in expression ranged from 1.61 to 2.75 for the up-regulated genes and from 1.47 to 2.69 for down-regulated genes. The significance of the variation in gene expression seemed robust because three exons of the Down syndrome cell adhesion molecule (Dscam) gene found to be differentially expressed were consistently down-regulated in VSH+. Microarrays were further validated by a quantitative real-time PCR (qRT-PCR) analysis (Fig. 1). Ratios of mean expression levels (VSH−/VSH+) from qRT-PCR analyses were similar to microarray ratios: Antdh: 1.47, lop1: 1.45 and Arrestin2: 1.35.

Overlap between VSH and other behavioural gene sets

To characterize the VSH genomic profile further, we compared the brain genomic profile of VSH bees to eight relevant gene sets that are associated with different behavioural phenotypes: foraging behaviour, performance of vibration signal (behavioural communication), bees stimulated by queen mandibular pheromone, brood pheromone or alarm pheromone and finally genes differentially expressed between Africanized and European honeybees (guard, soldier forager).

From three to 12 genes overlapped between the VSH and one of the behavioural gene sets (Table 3). We then determined whether the different overlaps were higher than the number of genes expected to overlap by chance alone. The VSH gene sets significantly overlapped with the gene set that is induced by the brood pheromone and

| Table 1. Microarray pair comparisons of four varroa-sensitive hygiene (VSH) colonies with low (VSH−) and high (VSH+) rates of hygienic behaviour |
|---|---|---|
| Pair comparisons | VSH+ (%) | VSH− (%) |
| 1 | 100 | 17 |
| 2 | 90 | 30 |
| 3 | 80 | 27 |
| 4 | 100 | 33 |

The percentage of varroa-infected brood removed by workers in each colony is indicated. The baseline population of mites in each colony was estimated in 200 worker-brood cells that were 0–3 days post-capping. Then, when the cells were 7–10 days post-capping, the number of uninfested cells was counted giving the brood removal rate (see Harbo & Harris, 2009, for more details).
the gene sets of vibrating bees and Africanized honeybees. As a result of the ‘low’ number of VSH genes, the number of overlapping genes was small and thus we could not perform statistical tests to determine the directional bias of the different overlaps. However, Table 2 indicates that there is a slight tendency for genes that were up-regulated in one of the gene sets to be down-regulated in the VSH+ sets and vice versa (Table 4).

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### Table 2. Probes differentially expressed in brains of varroa-sensitive hygiene (VSH) bees with low (VSH−) and high (VSH+) rates of hygienic behaviour

| Probe ID | Description | Drosophila orthologue | log2 ratio (VSH+/VSH−) | Gene ontology |
|----------|-------------|-----------------------|------------------------|---------------|
| AM01773  | NW_001253965.1 SET: UI_EST | / | 1.46 | |
| AM02265  | BB160015A20H04 | / | 1.17 | |
| AM01777  | NW_001253491.1 SET: UI_EST | / | 1.12 | |
| AM01915  | BB160006A1OF07 | / | 0.74 | |
| AM02211  | BB160014A10B11 | / | 0.71 | |
| AM04119  | GB111499 | CG31004 | 0.46 | Cell-matrix adhesion |
| AM05646  | GB13036 | / | 0.69 | |
| AM01622  | BB170002A10A06 | / | 0.58 | |
| AM00808  | BB773117 RIKEN full-length enriched honeybee cDNA library | / | 0.49 | |
| AM02131  | BB160011B20H01 | / | 0.63 | |
| AM03470  | GB10845 | PRL−1 | 0.54 | Protein tyrosine phosphatase activity; immediate early gene |
| AM01686  | BB170006B20H07 | / | 0.77 | |
| AM09325  | GB16747 | CG17323 | 0.81 | Glucuronosyltransferase activity; inter-male aggressive behaviour |
| AM01607  | NW_001253063.1 SET: UI_EST | / | 0.69 | |
| AM07547  | GB14956 | CG910 | −0.56 | Inositol oxygenase activity; oxidation reduction |
| AM12206  | GB19657 [long wavelength sensitiveopsin 1 (log1)] | Rhodopsin 6 | −0.79 | G-protein coupled photoreceptor activity; phototransduction |
| AM01085  | BB170007A20A04 | / | −0.60 | |
| AM02392  | NM_001040230.1 | / | −1.42 | |
| AM03333  | GB10708 | / | −0.99 | |
| AM03972  | GB12522 | Antdh | −0.68 | Carbonyl reductase nicotinamide adenine dinucleotide phosphate activity; oxidation reduction |
| AM05381  | GB12766 | Arrestin 2 | −0.74 | Adaptation of rhodopsin mediated signalling |
| AM00612  | DB739042 RIKEN full-length enriched honeybee cDNA library | / | −0.77 | |
| AM09032  | GB16453 | CG32645 | −0.70 | Transferase activity |
| AM00604  | DB738421 RIKEN full-length enriched honeybee cDNA library | / | −0.69 | |
| AM00955  | DB752711 RIKEN full-length enriched honeybee cDNA library | / | −0.61 | |
| AM06645  | GB30234 | Ets65A | −0.72 | Transcription factor activity |
| AM12005  | GB30242 [Odorant binding protein 3 (Obp3)] | / | −0.58 | Odorant binding; sensory perception of chemical stimulus |
| AM03226  | GB10599 | / | −0.80 | |
| AM06202  | GB13602 | CG10175 | −0.56 | Carboxylesterase activity; metabolic process |
| AM10277  | GB17704 | / | −0.50 | |
| AM02039  | BB160009B10D09 | / | −0.74 | |
| AM00103  | GB30209 [Down syndrome cell adhesion molecule (Dscam) exon 3] | Dscam | −0.77 | Axon guidance; mushroom body development |
| AM00166  | GB15141 (Dscam exon 10.9) | Dscam | −1.08 | Axon guidance; mushroom body development |
| AM04590  | GB11973 (Cyp4g11) | Cyp4g15 | −0.93 | Electron carrier activity; steroid biosynthetic process |
| AM01535  | BB170004B10F01 | / | −1.11 | |
| AM01750  | BB170027A10E09 | / | −1.43 | |
| AM01173  | BB170009B20C11 | / | −1.52 | |
| AM00167  | GB15141 (Dscam exon 10.10) | Dscam | −1.62 | Axon guidance; mushroom body development |
| AM01360  | BB170030B10C09 | / | −1.43 | |

Corresponding Drosophila orthologues, log2 ratio of expression values (VSH+/VSH−) and gene ontology based on fly orthologies (Flybase) are shown. Positive expression values indicate higher expression in VSH+ bees compared to VSH− bees. ‘/’ indicates the absence of clear Drosophila orthologues.
Discussion

As group-level defence against pathogens mostly involves collective behaviour, analysing genes involved in social immunity comes down to the identification of behavioural genes. By comparing VSH+ bees, characterized by a high performance level of hygienic behaviour, to VSH− bees, we expected to find some genes to be differentially expressed in VSH bees. The identification of 39 transcripts that are differentially expressed between VSH+ and VSH− bees confirmed this hypothesis.

Candidate genes for social immunity

VSH bees are characterized by their high ability to detect and remove varroa-parasitized brood (Harbo & Harris, 2005b). Harris (2007) suggested that VSH bees are either more sensitive to olfactory-based stimuli associated with parasitized brood members or have a lower response threshold to mite density (initiate hygienic behaviour at a lower mite density), which is not related to a higher olfactory sensitivity. This latter assumption comes from the fact that, at very low mite densities, there is no difference in hygienic behaviour between resistant (hygienic) Africanized bees and nonresistant European bees (Vandame et al., 2000). Hygienic behaviour would be initiated when a critical threshold of mite infestation rate is reached (Vandame et al., 2002); a threshold that is lower in resistant bees. The identification of genes that are differentially expressed between VSH and control bees might give some clues on the mechanisms of hygienic behaviour.

Amongst the genes up-regulated, PRL-1 encodes a protein tyrosine phosphatase. In Drosophila, its function is unknown but this gene belongs to the category of immediate-early genes (Diamond et al., 1994), which are genes that play an essential role in neural morphogenesis and functioning in mammals (Paul & Lombroso, 2003). This key regulatory component in signal transduction pathways might therefore be important to the development of hygienic behaviour. CG17323 has been found to be involved in diverse functions like circadian rhythm (Ceriani et al., 2002), aggression (Edwards et al., 2009) and response to ethanol exposure (Morozova et al., 2009). In addition, CG17323 and CG31004 (also up-regulated in VSH bees) are both affected by nutrient intake (Zinke et al., 2002). However, we do not have enough information on VSH behaviour to establish a link between this behaviour and these functions. The function of the cytochrome P450 Cyp4g11 is unknown. However, the Drosophila orthologue Cyp4g15 has been found to be predominantly expressed in the brain of Drosophila flies, where it might be involved in ecdysteroid metabolism rather than in detoxifying xenobiotics (Maibeche-Coisne et al., 2000). This suggests that in the bee brain Cyp4g11 might catalyse a reaction in some metabolic pathways that could be involved in hygienic behaviour. Another important gene is Dscam, an immunoglobulin superfamily member essential for wiring the brain. The molecular diversity of Dscam (38 016 alternative splicing forms in Drosophila) is essential for mediating axon guidance and neuronal wiring.
Table 3. Significance of overlap between the varroa-sensitive hygiene (VSH) and other behavioural gene sets

|                           | Expected no. | Observed no. | RF   | P-value |
|---------------------------|--------------|--------------|------|---------|
| Forager                   | 4.3          | 5            | 1.1  | 0.44    |
| Vibrating bee             | 3            | 10           | 3.3  | <0.001  |
| Queen mandibular pheromone| NA           | 8            | NA   | NA      |
| Brood pheromone           | 1.1          | 7            | 6.5  | <0.001  |
| Alarm pheromone           | 1.5          | 3            | 1.9  | 0.2     |
| AHB guard                 | 0.8          | 6            | 7.5  | <0.001  |
| AHB soldier               | 1.8          | 12           | 6.7  | <0.001  |
| AHB forager               | 0.19         | 5            | 26.5 | <0.001  |

Expected no., the number of genes expected to overlap between two gene sets by chance alone; RF, representation factor. As the gene set regulated by the queen mandibular pheromone was determined by using a different microarray platform (cDNA microarrays generated from brain expressed sequenced tags), its overlap with the VSH gene sets could not be calculated. Guard and forager describe specific tasks of bees: the first guard the hive at the nest entrance, the second are the first to react to a threat. AHB, Africanized honeybees; NA, not available.

Table 4. Overlap between the varroa-sensitive hygiene (VSH) and other behavioural gene sets

| Oligo ID   | Foraging bee | Vibrating bee | QMP  | BP  | Alarm pheromone | AHB guard | AHB soldier | AHB forager |
|------------|--------------|---------------|------|-----|-----------------|-----------|-------------|------------|
| AM01773    | Down         |               |      |     |                 |           |             |            |
| AM02265    |              |               |      |     |                 |           |             |            |
| AM01177    | Down         |               |      |     |                 |           |             |            |
| AM01915    | Down         |               |      |     |                 |           |             |            |
| AM02211    |              |               |      |     |                 |           |             |            |
| AM04119    |              |               |      |     |                 |           |             |            |
| AM05646    |              |               |      |     |                 |           |             |            |
| AM01622    |              |               |      |     |                 |           |             |            |
| AM0808     |              |               |      |     |                 |           |             |            |
| AM02131    |              |               |      |     |                 |           |             |            |
| AM03470    | Up           | Up            | Down |     |                 |           |             | Up         |
| AM01666    |              |               |      |     |                 |           |             |            |
| AM09325    |              |               |      |     |                 |           |             | Down       |
| AM01607    |              |               |      |     |                 |           |             | Down       |
| AM07547    | Up           | Up            | Up   |     |                 |           |             | Up         |
| AM12206    | Up           | Up            | Down |     |                 |           |             | Up         |
| AM01985    | Down         | Up            |      |     |                 |           |             |            |
| AM02392    | Up           | Up            | Down |     |                 |           |             | Down       |
| AM03333    | Up           | Up            | Up   |     |                 |           |             | Down       |
| AM03972    | Up           | Down          | Up   |     |                 |           |             | Down       |
| AM05381    | Up           | Down          | Up   |     |                 |           |             | Down       |
| AM06612    | Up           |               |      |     |                 |           |             |            |
| AM09032    |              |               |      |     |                 |           |             |            |
| AM06604    |              |               |      |     |                 |           |             |            |
| AM09955    | Up           |               |      |     |                 |           |             | Down       |
| AM06645    |              |               |      |     |                 |           |             |            |
| AM12005    |              |               |      |     |                 |           |             |            |
| AM03226    |              |               |      |     |                 |           |             |            |
| AM06202    |              |               |      |     |                 |           |             |            |
| AM10277    |              |               |      |     |                 |           |             |            |
| AM02039    |              |               |      |     |                 |           |             |            |
| AM00103    |              |               |      |     |                 |           |             |            |
| AM00166    | Up           |               |      |     |                 |           |             | Up         |
| AM04590    |              |               |      |     |                 |           |             |            |
| AM01535    |              |               |      |     |                 |           |             | Down       |
| AM01750    |              |               |      |     |                 |           |             | Up         |
| AM01173    |              |               |      |     |                 |           |             |            |
| AM00167    |              |               |      |     |                 |           |             | Up         |
| AM01360    |              |               |      |     |                 |           |             | Up         |

The upper and lower part of the table shows probes that are up- and down-regulated in VSH+ bees, respectively. ‘Up’ and ‘Down’ indicate whether probes are up- or down-regulated in the corresponding behavioural phenotype. AHB, Africanized honeybees; BP, brood pheromone; QMP, queen mandibular pheromone.

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specificity (Chen et al., 2006). The significant down-regulation of three Dscam exons suggests therefore a different wiring of neuronal networks in the brains of VSH bees.

Two other genes are both involved in visual signalling: long-wavelength sensitive opsin 1 and arrestin 2 (Dolph et al., 1993). In insects, arrestin 2 is also expressed in olfactory neurones and is believed to be important for a normal olfactory physiology (Merrill et al., 2002; Walker et al., 2008). Long-wave opsins have been described in the optic lobes (Lampel et al., 2005) but in honeybees this gene is solely expressed in the compound eyes (Velarde et al., 2005), which suggests that a small amount of retinal tissue was associated with the dissected brain tissue. As the brood is reared in the dark inside the hive, the down-regulation of the visual signalling cascades would indicate that VSH bees spend more time within the hive than nonhygienic bees of the same age.

The down-regulation of odorant binding protein 3 (obp3), a member of the Obp family first described as a carrier of odorant molecules in olfactory tissue, could support a role in the olfactory sensitivity of VSH bees. However, its expression in the brain and in other body parts, with the exception of the antennae (Foret & Maleszka, 2006), suggests that obp3 is involved in other physiological functions. A similar conclusion can be drawn with Antdh, which was first described in the antennae of Drosophila as being involved in odorant turnover (Wang et al., 1999). However, the down-regulation of Ets65A, a candidate gene likely to account for olfactory behaviour in the smell-impaired mutant lines 65A (Anholt & Mackay, 2001), suggests that VSH bees express a different level of odor-guided behaviour than other bees.

The lack or down-regulation of genes involved in olfaction does not seem to support the hypothesis that VSH bees are more sensitive to olfactory-based stimuli associated with parasitism of the brood. The alternative explanation focusing on the tolerance level to mite density would be more likely. However, to reject definitely the hypothesis of higher olfactory sensitivity, the analysis of peripheral tissues, like antennae, should be performed. Indeed, insect behaviour can be dramatically affected by changes in expression of genes that are antennal-specific (Wang et al., 2008).

Relationship between VSH and others behavioural phenotypes

The overlap analyses between different genomic profiles provided a better characterization of VSH behaviour. We did not find a significant overlap between VSH and forager brain gene expression profile. This could be a result of age differences between our VSH samples and old foragers; however, our findings that these two phenotypes are not linked is supported by Goode et al. (2006), who found that hygienic behaviour is independent of foraging ontogeny. Interestingly, the number of genes overlapping between VSH and vibrating bees was higher than expected by chance alone. This ‘modulatory communication signal’ induces a nonspecific increase in worker activity (Schneider & Lewis, 2004) and is produced by a restricted number of bees, primarily successful forager collecting food outside the hive (Schneider & Lewis, 2004). As VSH bees tend to display an opposite gene expression pattern to vibrating bees, this suggests that hygienic bees might spend more time in inside-hive activity. Another interesting result is the significant overlap between the genomic profiles of VSH bees and bees stimulated by brood pheromone (BP). This pheromone emitted by larvae stimulates brood care (feeding) (Le Conte et al., 2001) but also the capping of brood cells containing mature larvae (Le Conte et al., 1990). This raises the question of whether BP can affect hygienic behaviour. Interestingly, BP tends to inhibit genes that are up-regulated in VSH bees and inversely, which suggests that hygienic bees do not spend time on feeding larvae and/or have a higher propensity to not cap opened brood cells. Indeed, hygienic behaviour is performed on mite-infested pupae, especially young pupae (3–5 days post capping) (Harris, 2007). As the queen mandibular pheromone exhibits compelling similarities in effects on bee behavior compared to BP (Alaux et al., 2010), a similar conclusion could be drawn with this pheromone. Finally, our results indicate that Africanized honeybees (AHB) and VSH genomic profiles share a significant number of genes despite the age difference between bees from both data sets. This is particularly appealing given that AHB are tolerant to varroa compared to the European honeybee (EHB). One explanation highlights the fact that, similarly to VSH bees, AHB are more able to remove infested brood than EHB (Guzmán-Novoa et al., 1999; Vandame et al., 2000; but see Mondragon et al., 2005). The shared behavioural and molecular traits suggest that the mechanism underlying the tolerance to varroa is similar between AHB and VSH bees. However, amongst the overlapping genes, few are regulated in the same direction in both AHB and VSH bees. Genes that are common to both strains and follow the same pattern of regulation may represent key factors of the molecular resistance to varroa.

Comparisons to others studies

A previous study identified genes associated to another type of mite resistance displayed by some French colonies (Navajas et al., 2008). Only two genes were both differentially expressed in VSH and those Varroa surviving bees (VSB) that are naturally tolerant to Varroa infestation. BB160006A10F07 and Dscam exon 10.9, up-regulated and down-regulated, respectively, in VSH
were both down-regulated in VSB+ compared to VSB−. Although this low number was surprising, on the one hand the analysis of VSB individuals was performed on whole pupae and the present study was performed with bee brains, and on the other hand the mechanisms of varroa tolerance in the French strains has not yet been clearly established. It is possible that the mechanism differs from the hygienic behaviour performed by VSH bees. Nevertheless, as \textit{Dscam exon 10.9} was also down-regulated in VSH bees, this gene may be important for the resistance to varroa parasitism.

A different approach based on genetic mapping was used in order to identify the genetic component of hygienic behaviour. By performing a quantitative trait loci (QTLs) analysis, Lapidge \textit{et al.} (2002) first found that hygienic behaviour is influenced by many different loci but recently, in an attempt to provide marker-assisted selection for hygienic behaviour, Oxley \textit{et al.} (2010) identified three QTLs that influence the propensity of workers to perform hygienic tasks. QTLs are phenotypically defined genomic regions associated with variation in a phenotypic trait, which can be large and contain hundreds of candidate genes. However, QTL analysis does not indicate the expression pattern of these genes. Quantitative expression studies such as microarray analysis can be used to systematically reduce the list of candidate loci and reveal regulatory variation in genes and pathway signalling. Thus, combining QTL mapping with transcriptome analysis promises to identify positional candidate genes for a phenotype of interest whose expression varies amongst lines (Jansen & Nap, 2001; Wayne & McIntyre, 2002; Li & Burmeister, 2005). Unfortunately, none of the genes from these QTLs were found to be differentially expressed in VSH bees, indicating that further studies are needed to understand the genetic background of such behaviour. This lack of overlap could come from the different breeding lines of bees that were used in both studies. The QTL mapping was performed with the Minnesota (USA) Hygienic bee stock (Boecking & Spivak, 1999; Spivak & Reuter, 2001a) and the transcriptome analysis with VSH bees from Lousiana (USA) (Harbo & Harris, 1999). In addition, each line was obtained with different methods of selection: the VSH line was specifically selected for its resistance to varroa but the selection of the Minnesota line was not varroa specific and included a broad spectrum of pathogens, causing notably the American foulbrood (Spivak & Reuter, 2001b) and chalkbrood diseases (Spivak & Reuter, 1998). Thus, combining both techniques on the same bee lines may provide finer identification of candidate genes involved in hygienic behaviour.

**Conclusion**

It has been shown that social defence reduces the investment of individuals in their own immune function in ants and honeybees (Castella \textit{et al.}, 2008; Simone \textit{et al.}, 2009). This indicates that the evolution of sociality might have led to the loss of immune genes, as indicated by the analysis of the honeybee genome (Evans \textit{et al.}, 2006), to the expense of behavioural genes involved in group level defence. As behavioural genes are often pleiotropic (Greenspan, 2001; Sokolowski, 2001), those genes might be both involved in social immunity and other behavioural phenotypes (see Tables 3 and 4), which might reduce the physiological investment in the defence against pathogens compared to less pleiotropic immune genes. This study represents a first step towards understanding the genomic basis of social immunity. Future research will have to test the functions of those candidate genes in collective defences.

**Experimental procedures**

**Honeybee rearing and selective breeding**

The honeybee colonies used in this study presenting high or low rates of varroa-hygienic behaviour were the same as those previously used by Harbo & Harris (2009) for studying the responses to varroa by honeybees with different levels of varroa-sensitive hygiene. Briefly, we produced a group of 26 colonies of a European mix of \textit{A. mellifera} subspecies typically found in North America that presented different levels of varroa-sensitive hygiene. Fourteen colonies had queens produced from a line with 100% expression of VSH (high line, H), and 12 had queens produced from a line that did not express VSH (low line, L). Each of the 26 queens was backcrossed to a single drone produced by an HL queen (a daughter of both lines H and L). As the HL queen had half of the VSH alleles, she produced drones that ranged from having 0 to 100% of the VSH alleles. With this design, the 14 colonies in group H should have 50–100% of the alleles for VSH and the 12 in group L should have 0–50%. Therefore, we speculated that the lowest of the low group had none of the genes for VSH and the highest of the high group had all of the genes for VSH. Each queen was introduced into colonies that were evaluated for varroa hygienic ability as in Harbo & Harris (2009), after the test queens had been laying in their colonies for at least 6 weeks. In this way all worker bees that were between 0 and 3 weeks old were daughters of the test queen. The varroa hygienic ability corresponded to the percentage of mite-infested pupae that were removed by the colony. Thanks to this method, we were able to sample high and low varroa hygienic bees for gene expression analysis. Some bees were marked at emergence and returned to their colonies of origin. We then collected and flash froze in liquid nitrogen 14-day-old bees from four VSH+ colonies and four VSH− colonies (control) (Table 1). To avoid any bias toward a specific behaviour, bees found on the brood area were randomly collected. VSH+/VSH− colonies were directly compared with microarrays analysis using a dye swap method.

**Brain dissection and mRNA extraction**

Whole heads were partially freeze-dried at $-80\, ^\circ\mathrm{C}$ (0.0005 mmbar for 140 min) to facilitate brain dissection. Dissections were performed on dry ice to prevent brain thawing. For
each sample, 10 frozen bee brains were pooled and grounded on dry ice. We added 600 µl of RLT buffer containing 6 µl of β-mercaptoethanol to the powder to disrupt the tissue. RNA extraction was carried out as indicated in the Qiagen RNeasy kit for total RNA (Qiagen, Courtaboeuf, France). RNA isolated from 10 pooled brains was then used for microarrays analysis.

Microarrays and data analysis
For the preparation of the labelled Cy3- and Cy5- aRNA target, total RNA (1 µg) was amplified with an Amino Allyl MessageAmp II aRNA Amplification kit (Ambion, Courtaboeuf, France), according to the manufacturer’s instructions. Then, amino allyl RNA samples were dried in SpeedVac (Thermo Fisher Scientific, Courtaboeuf, France) and resuspended in 9 µl coupling buffer (0.1 M carbonate buffer pH 9). We added to each sample 11 µl of one of the N-hydroxysuccinimide ester dyes (Cy3 and Cy5) diluted in dimethylosulfoxide (CyDye Post-Labeling Reactive Dye, GE Healthcare, Montpellier, France). Samples were incubated at room temperature in the dark for 30 min with shaking and 4.5 µl 4 M hydroxylamine was added to each sample. Samples were incubated at room temperature for 15 min in the dark. Nuclease-free water was added to each sample to bring the volume to 100 µl. Then, we added 350 µl of aRNA Binding Buffer and 250 µl 100% ethanol to each aRNA sample. Samples were applied on the column and centrifuged 10 000 g for 1 min. We added 650 µl Wash Buffer and centrifuged for 1 min at 10 000 g. After discarding the flow-through, samples were centrifuged again for 1 min at 10 000 g. Samples were eluted twice in 50 µl nuclease-free water. Cy3 and Cy5 reactions were equally combined and fragmented according to the kit’s instructions.

Before hybridization, slides were passed quickly through steam and placed in a UV linker at 100 mJ. Before pre-hybridization, slides were plunged twice in 0.2% sodium dodecyl sulphate (SDS) and immediately shaken vigorously for 1 min. They were then washed twice in distilled water for 1 min. For the labelled aRNA, a 4 x hybridization buffer (GE Healthcare) was used. In a final concentration of 50% formamide, denatured at 95 °C for 5 min and applied to the microarrays in individual chambers of an automated slide processor (GE Healthcare). Hybridization was carried out at 37 °C for 16 h. Hybridized slides were washed at 52 °C successively with 0.1% probe saturation. Microarrays were immediately scanned at 10 µm resolutions in both Cy3 and Cy5 channels using a GenePix 4200AL scanner (Molecular Devices, St. Grégoire, France). The scanning was carried out with a variable photomultiplier tube (PMT) voltage to obtain maximal signal intensities (<0.1% probe saturation). ARRAYVISION software (GE Healthcare) was used for feature extraction. Spots with high local background or contamination fluorescence were flagged manually. A local background was calculated for each spot as the median values of the fluorescence intensities of four squares surrounding the spot. This background was subtracted from the foreground fluorescence intensity.

No background correction was performed. No spatial bias in the quality analysis was detected so a Loess normalization was performed for all microarrays to correct dye effect and technical bias. Tests of differential expression were conducted using the Siggenes package from Bioconductor and the Significance Analysis of Microarrays (SAM) proposed by (Tusher et al., 2001). SAM assigns a score to each gene based on the standard deviation of repeated gene expression measurements. Then, a false discovery rate is estimated by permutations of the repeated measurements to obtain a ranking of significantly expressed genes. The Bioarray Software Environment (BASE) (local installation: http://baseprod.igf.cnrs.fr/index.phtml) was used to visualize differential expression for each gene.

Verification by qRT-PCR
In order to validate the microarrays results, qRT-PCR was performed on each sample. The transcript abundance was measured for lop1, Arrestin2 and Andth with a Roche LightCycler 480 Real-Time PCR System (Roche Diagnostics, Meylan, France). Their expression levels were then normalized to a housekeeping gene (Bis511718) and to a control sample using 2(-Delta-Delta CT) values. Primer sequences (5’ to 3’) were lop1 forward: GTTCTCTCTCGGATGGGACTA, reverse: GGAGC GAAGTAAACCCCAAAAT; Arrestin2 forward: CCTTGAAGGAGG CGTAAATCGCTCA, reverse: TGGAGCTAACATTGTACCC; Andth forward: CAATTAGAGATTGGCCGTCC, reverse: TCCA GTATGAAAGGACCTC; Bis511718 forward: CTATCACTGTGT TGTTTCTCCTC, reverse: TCGTGGTGCCTCTTACGCTTGT.

Overlap between the VSH and other gene expression profiles
We compared the VSH genomic profile to different gene lists previously identified in other honeybee studies. These brain gene expression profiles are specific to foraging behaviour (Alaux et al., 2009b), bees specialized in vibration communication (strong arousal state) (Alaux et al., 2009a), Africanized honeybees (Alaux et al., 2009c) and finally bees stimulated by queen mandibular pheromone (Grozinger et al., 2003), brood pheromone (Alaux et al., 2009b) or alarm pheromone (Alaux et al., 2009c). We calculated a ‘representation factor’ (the number of observed overlapping genes divided by the expected number of overlapping genes) and used an exact hypergeometric probability test to determine whether the overlap between the VSH and another gene set was statistically significant. The expected number corresponds to the product of the number of genes in each list divided by the total number of genes analysed (Kim et al., 2001).

Web resource
Gene expression data meet Minimum Information About a Microarray Experiment (MIAME) standards and have been deposited at ArrayExpress (http://www.ebi.ac.uk/arrayexpress): E-TABM-1002.

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Various studies have been conducted to understand the mechanisms involved in the resistance of honey bees to Varroa mites. For instance, Neumann and Härtels (2004) conducted a study on the removal of small hive beetle (Aethina tumida) from honey bee colonies, highlighting the importance of this pest in honey bee health. 

In another study, Mondragon et al. (2005) investigated the multifaceted roles of honey bees in modulatory communication and the organization of labor in colonies. The authors emphasized the significance of honey bees as a model system for studying social organization and learning.

The interaction between honey bees and their natural enemies, such as Varroa mites, has been a focal point of research. Sokolowski (2001) conducted an in-depth genomic analysis of Drosophila melanogaster, revealing insights into the evolutionary ecology of insect immunity. This study underscores the importance of honey bees as a model system for understanding insect defense mechanisms.

Furthermore, research on the visual arrestins in Drosophila melanogaster has provided valuable insights into the mechanisms of visual perception. Carlson and Zwiebel (2002) explored the role of visual arrestins in these insects, shedding light on the evolutionary conservation of visual signaling pathways.

In summary, the studies mentioned above highlight the multifaceted roles of honey bees in various aspects of biology, including parasitism, behavior genetics, and immunity. These studies underscore the importance of honey bees as a model system for understanding complex ecological interactions and the underlying genetic and molecular mechanisms.