Vitellogenin regulates hormonal dynamics in the worker caste of a eusocial insect

Karina R. Guidugli, Adriana M. Nascimento, Gro V. Amdam, Angel R. Barchuk, Stig Omholt, Zilá L.P. Simões, Klaus Hartfelder

1. Introduction

Understanding the proximate mechanisms that underlie division of labor in eusocial insect societies is a major issue in sociobiology. In particular, the honey bee has emerged as a key model for dissecting the regulatory architecture that gives rise to task partitioning in social groups. Honey bee workers undergo a characteristic progression in task performance. They care for brood during the first two weeks of adult life (“nursing”), and later start foraging for nectar and pollen. This behavioral switch is accompanied by an increase in the juvenile hormone (JH) titer and a decrease in the vitellogenin titers associated with the shift in worker function from the performance of nest-tasks to foraging for food, nectar, and pollen. Honey bee workers that have high vitellogenin and low JH titers are compatible with the physiological condition of nurse bees, whereas honey bees that have high JH and low vitellogenin titers are compatible with the physiological condition of forager bees. Based on available data, general life history considerations, and theory on the generic principles of stable states, a dynamic model of the forager transition was recently presented [2]. The mathematical model proposed that the shift from nest tasks to foraging is controlled by a two-repressor constellation that involves a positive regulatory feedback loop between JH and vitellogenin. Modulation of the parameter values of this feedback loop permitted simulations of co-variance patterns between the JH and vitellogenin titer that are observed under natural and experimental conditions. The simulation results supported the experimentally verified inhibitory effect of an elevated JH titer on the hemolymph vitellogenin level [4, 11], and thereby the hormone’s association with the shift in worker behavior from the performance of nest-tasks to foraging for

Season and colony age demography influence the JH titer and, consequently, the timing of onset of foraging behavior in individual honey bee workers [5, 6]. This plastic and adaptive individual level response to colony needs led to the perception that JH may function as a pacemaker of behavioral development in workers [7]. However, recent experiments on allatectomized bees [8] show that JH is not the only effector that drives the behavioral transition to foraging. Thus, a better understanding of the regulatory signature in the shift to foraging activity is required, also because insights into this signature may provide novel information on the evolutionary trajectory of pathways that control division of labor in social insects.

Regulation of female fertility in the social Hymenoptera is usually considered under the general paradigm of hormonal control of insect reproduction, where JH and ecdysteroids act as the main inducers of vitellogenin synthesis and uptake [3]. In adult honey bees, however, vitellogenin synthesis is not upregulated by these hormones [4], and it is only during the initiation of vitellogenin expression in the late pupal stages that JH acts as an inducer of vitellogenin production [9]. This unconventional association was partly explained when it was shown that vitellogenin has evolved functions beyond the restricted context of reproduction: putative vitellogenin receptors can be found in the royal jelly producing hypopharyngeal glands of workers, suggesting that vitellogenin is used to synthesize proteins that nurse bees feed to the larvae [10]. The conversion of a yolk protein to larval food proteins is compatible with the physiological condition of nurse bees, which have high vitellogenin and low JH titers [11].

© 2005 Federation of European Biochemical Societies. Published by Elsevier B.V. All rights reserved.

Keywords: Juvenile hormone; Vitellogenin; Ultraspiracle; Age polyethism; Honey bee

Abstract Functionally sterile honey bee workers synthesize the yolk protein vitellogenin while performing nest tasks. The subsequent shift to foraging is linked to a reduced vitellogenin and an increased juvenile hormone (JH) titer. JH is a principal controller of vitellogenin expression and behavioral development. Yet, we show here that silencing of vitellogenin expression causes a significant increase in JH titer and its putative receptor. Mathematically, the increase corresponds to a dynamic dose-response. This role of vitellogenin in the tuning of the endocrine system is uncommon and may elucidate how an ancestral pathway of fertility regulation has been remodeled into a novel circuit controlling social behavior.
pollen and nectar. This transition is a major behavioral change that is also accompanied by a reorganization of central nervous system structures involved in learning and memory formation [1,7].

Yet, in addition to supporting the inhibitory effect of JH on vitellogenin, the mathematical simulations further arrived at the non-intuitive prediction that forced repression of vitellogenin synthesis should trigger an increase in the JH titer, which subsequently causes the worker bee to become behaviorally and physiologically locked in the forager state. It is this combination of inhibitory interactions that constitutes the Double Repressor framework [2] for the regulation of a honey bee worker’s physiology during its adult life cycle. Thus, the “Double Repressor” hypothesis has implications well beyond those for the specific physiology of social insects due to its prediction of a regulatory association between JH and vitellogenin that is not commonly part of the reproductive architecture of female insects.

In the present study, we experimentally tested whether vitellogenin indeed exhibits this regulatory function in honey bee workers. We used RNA interference (RNAi) technology to silence vitellogenin expression and investigated the effect on the JH titer. In addition, we monitored the expression level of a putative JH receptor [12]. The study was performed on two genetically different bee sources kept under distinct social conditions to assess the robustness of a putative regulatory relationship involving vitellogenin and JH.

2. Materials and methods

2.1. Bees

Frames of sealed worker brood were retrieved from *Apis mellifera* colonies maintained at the Apiaries of the Norwegian University of Life Sciences, Aas, Norway (*Apis mellifera carnica*) and of the University of São Paulo at Ribeirão Preto, Brazil (Africanized hybrids). Bees emerging from brood cells were marked. The Norwegian source bees were introduced into one-story hives that contained an egg-laying queen. The Brazilian source bees were kept queenless in groups of 25–35 bees in an incubator. They received a pollen/sugar diet that supports vitellogenesis [13]. In both setups, the bees were retrieved after seven days for hemolymph and fat body sampling. Control foragers represent a random sample of bees that were collected from flowers near the apiary.

Seven days after the start of the experiment, the bees were retrieved and anesthetized on ice for hemolymph sampling [14] and dissection. For JH quantification, 1–3 μl of each hemolymph sample were drawn into 500 μl acetonitrile and stored at −20 °C. The remaining hemolymph was stored at −20 °C for protein electrophoresis. RNA was extracted from abdominal carcasses by a TRizol protocol.

2.2. Silencing of vitellogenin expression

Double-stranded honey bee vitellogenin RNA was prepared as previously described [15] following the protocol of the Promega Ribopure™ T7 system (Promega) and using as template the clone *ApaH5* that contains a partial sequence of the honey bee vitellogenin cDNA [16]. After phenol-chloroform extraction and heat treatment, the dsRNA was diluted with nuclease-free water to a final concentration of 5 μg/μl. Newly emerged workers were injected intra-abdominally with 1 μl dsRNA solution or 1 μl water (sham). Bees showing signs of hemolymph leakage after withdrawal of the needle were discarded.

2.3. SDS-PAGE and RT-PCR

Protein analysis was performed using samples of 1 μl hemolymph that were subjected to SDS-PAGE (7.5% gel) and stained with Coomassie Brilliant Blue. Gels were scanned and imported into tinamate (3.3.12a, Linux) to quantify staining intensity and size of the vitello-

2.4. Juvenile hormone titer analysis

The JH extraction procedure followed a protocol established for honey bee hemolymph [18]. Briefly, 1 ml NaCl (0.9%) and 1 ml hexane were added to the acetonitrile extract. After vigorous vortexing, the phases were separated by centrifugation (700 × g). The hexane phase was removed and the extraction was repeated twice. The pooled hexane phases were dried and the residue was redissolved in 50 μl toluene. Before starting the RIA, the solvent was removed by vacuum centrifugation.

The antisera used in this study was diluted to 1:1250 in phosphate buffer supplemented with bovine serum albumin (0.1%) and rabbit immunoglobulin G (0.1%). The assays were performed with [10-^3H(N)]-juvenile hormone III (spec. activity 19.4 Ci/mmol, NEN Life Science Products, Boston), diluted in the phosphate buffer to 6000–6500 cpm/50 μl. Juvenile hormone III (Fluka) was used as non-radioactive ligand. Standard curves were set up to cover a 50 pg to 10 ng range. The RIA procedure followed the protocol established by Goodman et al. [19]. JH titers of unknown samples are expressed as JH-III equivalents (pg/μl hemolymph).

3. Results and discussion

We found that vitellogenin gene knockdown by RNAi was associated with a significant increase in the JH titer of worker bees. This effect was observed consistently in both genetic backgrounds and social conditions, although the response was more apparent in the Africanized *A. mellifera* hybrids (Fig. 1A and B). In addition to the quantification of vitellogenin protein by densitometric analysis of the corresponding SDS-PAGE band in the full set of hemolymph samples (47 bees of Norwegian origin and 47 Africanized honey bees), we also monitored the vitellogenin transcript levels by RT-PCR in a subset of fatbody mRNA samples. These results confirmed the vitellogenin knockdown phenotype (Fig. 1C and D). The observed association between the downregulation of vitellogenin and upregulation of JH provides strong evidence for the previously hypothesized [2] regulatory feedback constellation between vitellogenin and JH. Our data also suggest that the relationship is quite robust to variation in the genetic origin and the social milieu of worker bees.

We further explored the dynamic linkage between vitellogenin and JH at the level of individual bees (Fig. 2A). This analysis performed on data from Africanized *A. mellifera* hybrids showed a significant negative correlation between the vitellogenin titer and the JH level in bees with the knockdown phenotype (r = −0.66, P = 0.02). The control groups, which consisted of untreated and sham-injected bees, displayed no such association (r = 0.04, P = 0.82), but a significant non-linear component was apparent when JH was modeled as a function of the hemolymph concentration of vitellogenin (r = 0.51, P = 0.01). This interpretation is supported by the non-linear threshold shift observed in the JH level of vitellogenin knockdown bees (Fig. 2A and B), which all exhibited JH titers above the 200 pg/μl upper bound of the control group.
The mathematical model of the Double Repressor framework [2] was built on the assumption that the JH titer shows a stringent inverse linear or non-linear dose–response relative to the vitellogenin level. In our study, this pattern was not played out as a continuous association. Rather, the strong inverse linear association was observed in the knockdown group, whereas the control showed the non-linear component. This result could be interpreted as being in conflict with the Double Repressor model. Yet, the explicit mapping of dose–response trajectories requires collection of individual-specific samples over time, and non-destructive repeated measurements of JH and vitellogenin cannot be obtained from insects of the size of bees. Also, worker bees may differ considerably in their relative response trajectories due to physiological and genetic heterogeneity [20], as outlined by a projection of predicted trajectories on our plotted individual data (Fig. 2B).

In conclusion, the JH concentrations measured for knockdown and control bees (Fig. 1) corresponded to relative levels typically observed in foragers and nurse bees, respectively [5,18]. Thus, the dynamics triggered by experimental vitellogenin downregulation are in accordance with the predicted signature of a double repressor association between JH and vitellogenin (see Fig. 2B and [2]), and may consequently represent the actual endocrine transition state that characterizes worker bees as they switch from nest-tasks to foraging activities [18,21].

Finally, to assess whether the increase in JH caused by vitellogenin downregulation triggered a normal response cascade, we analyzed the expression level of usp, which is a candidate JH receptor [22]. In honey bees, usp expression was recently shown to be rapidly upregulated by experimental JH-III application [12]. Thus, we selectively tested usp mRNA levels in workers with strongly contrasting relationships between vitellogenin and JH, i.e., individuals with high vitellogenin titer and low JH level in the control group, and individuals with low vitellogenin titer and high JH level in the experimental RNAi group. We found an approximately 45% increase in the level of usp expression in the knockdown bees (Fig. 3). This suggests that vitellogenin gene knockdown causes an operational increase in the JH titer, accompanied by the transcriptional modulation in a nuclear receptor implicated in the JH response of target tissues.

The proximate mechanism responsible for the inhibitory action of vitellogenin on JH was not determined in this study. However, it is likely that the effect is mediated through allatoregulatory peptides [23]. Of functional interest are also observations that the endocrine system of animals may show a highly sensitive response to changes in extracellular zinc levels [24]. Vitellogenin proved to be the major zinc carrier in honey bee hemolymph, and the plasma zinc concentration of a forager is, therefore, significantly lower than in a nurse bee [25]. Accordingly, a regulatory action of vitellogenin on JH would arise if JH synthesis in worker bees is conditional on the plasma zinc level.

Vitellogenin and JH are basic elements in the reproductive machinery of female insects. Yet, most studies on the relationship between the two components focus, unidirectionally, on hormonal regulation of vitellogenin induction and its shut...
down [3]. Only one study [26] has previously reported a reverse effect of vitellogenin on JH synthesis: explicitly, removal of the ovary in *Blattella germanica* resulted in an extreme accumulation of vitellogenin, subsequently inducing a drop in the JH titer. Vitellogenin clearly does not accumulate to such levels in worker bee hemolymph, yet we observed a similar inhibition of JH release at much lower vitellogenin levels than in *Blattella*.

These observations imply that an inhibitory effect of excessively high vitellogenin levels on JH production may constitute an ancient and possibly conserved mechanism in the reproductive physiology of female insects. This hypothesis is meaningful from the perspective that such a mechanism can protect against adverse effects of JH-stimulated vitellogenin production during periods when the female is unable to convert hemolymph proteins into eggs. Such situations are typically encountered in social insects where the subordinate caste of female workers foregoes individual reproduction to rear the mother’s or sister’s offspring.

In the honey bee, worker fertility is extremely low in the presence of an egg-laying queen. Yet, if the queen is lost, workers can activate follicle development through enhanced vitellogenesis and initiate egg laying [4]. The regulatory architecture of a worker bee’s physiology consequently encounters two quite distinct situations: one when the queen is present and the worker undergoes the characteristic transition between behavioral tasks; and one in the absence of the queen when egg-laying becomes a possibility and an important fitness component. Under the first condition of low fertility and alloparental function, the evolution of a worker caste seems to have been accompanied by a co-option of the reproductive hormone JH into a regulatory function in behavioral development [1,27]. In addition, our findings suggest that this evolutionary trajectory involved a regulatory shift to a lower response threshold in the inhibitory interactions between vitellogenin on the JH titer. This would imply that it is not solely JH that has been co-opted into a novel role, but rather
that an entire core control module of insect fertility (involving JH and vitellogenin) has been remodeled during the evolution of the worker caste in social insects.

Intriguingly, traces of this transition were recently detected in a study of the association between reproductive regulatory circuits and honey bee foraging behavior [28]. It was shown that the foraging behavior of workers, i.e., whether a bee preferentially collects nectar or pollen, is strongly influenced by an individual’s reproductive physiology. In the context of the present results, these considerations ultimately suggest that condition-sensitive synthesis and release of molecules derived from ancestral regulatory modules in reproductive pathways underlie at least part of the physiological and behavioral plasticity that is a hallmark of social insects.

Acknowledgments: We thank Walter Goodman for a generous supply of juvenile hormone antibody, and Robert Page, Xavier Bellés and Bert Hölldobler for critically reading and commenting on previous versions of this manuscript. Financial support by the Fundação de Amparo a Pesquisa do Estado de São Paulo (FAPESP, 1999/00719-6) and the Norwegian Research Council (Project no. 157851/432) is acknowledged.

References


