

Silkworm Immunity and Hormonal Regulation

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Abstract

Silkworm diseases are the most important disease that inflicts heavy loss to crops. Most losses in sericulture can be attributed directly to silkworm diseases.

In insects, it relies on both humeral and cellular responses that are mediated via certain recognizing receptors and activation of several pathways. Fat body and hemocytes are the origins for the production and secretion of antimicrobial agents and activators/regulators of cellular response, while cell mediated immunity in insects is performed by hemocytes. Humoral immunity acts as a front-line barrier against invading pathogens. The humoral response includes, the activation of the Phenol oxidase system, which triggers the synthesis of melanin and ii) the production of several immune effectors, such as lysozyme, reactive oxygen and nitrogen species, and antimicrobial peptides (AMPs).

Keywords: Humoral Immunity; Hemocytes; Haemolymph

Abbreviations: AMPs: Antimicrobial Peptides; PO: Prophenoloxidase; Ddc: Dopa Decarboxylase; FAAs: Free Amino Acids; JH: Juvenile Hormone; CA: Corpora Allata; NVP: Nucleopolyhedrovirus; ICP: Insecticidal Crystal Protein; BB: *Bacillus Bombysepticus*; BT: *Bacillus Thuringiensis*.

Introduction

Cellar Immunity

Insect haemolymph contains several types of mesodermal cells that are suspended in plasma, attached to various organs or attached to the inner surface of integument. Circulating haemocyte, carry out an immune response, or cellular defense, via phagocytosis, nodulation, encapsulation and melanization.

These processes appear to be discrete immune responses in terms of gene expression and outcome.

However, these certain immune responses share a number of common elements that function in concert to clear pathogens from the hemolymph.

Phagocytosis is induced when phagocyte surface receptors, are activated by target cells. It must be noted that the hemocyte response to various bacteria differs. For example, in *A. aegypti* hemocytes respond to *Escherichia coli* with phagocytosis, whereas to *Micrococcus luteus* with melanization [1-5].

Gad and Alzahofi [6] explained that hyperphagocytic hemocytes are involved in nodule formation and phagocytosis to clear large numbers of bacteria at the early stages of infection, while plasmatocytes and granulocytes are involved in phagocytosis of bacterial cells remained in circulation. Also, demonstrated that the immune response against bacterium is the same in pathogenic and non pathogenic bacteria.

Nodulation

Nodulation refers to multicellular hemocytic aggregates that entrap a large number of bacteria. Melanized or non-melanized nodules are formed in response to a number of invaders. Nodule formation appears to be related with eicosanoids in many insect species [7] or prophenoloxidase (PO) and dopa decarboxylase (Ddc) in medfly hemocytes [8].

Encapsulation

Encapsulation refers to the binding of hemocytes to larger targets, such as parasites, protozoa, and nematodes. Encapsulation can be observed when parasitoid wasps lay their eggs in the hemocel of *Drosophila larvae*. Hemocytes after binding to their target they form a multilayer capsule around the invader, which is ultimately accompanied by melanization. Within the capsule the invader is killed, by the local production of cytotoxic free radicals ROS and RNS, or by asphyxiation [5,9].

Humoral Immunity

Humoral immunity acts as a front-line barrier against invading pathogens, and the majority of the components are widely conserved among species. Regulation of innate immunity is important for overcoming infections and preventing self-damaging sepsis.

Defense peptides are key factors in humoral immunity against bacteria and fungi in invertebrates. Antimicrobial peptides play a crucial role in fighting against invading pathogens. They are synthesized in response to microbial infection or septic body injury mainly in insect fat body and in certain blood cells, and then rapidly released into the haemolymph where they act synergistically against microorganisms [10]. Anti-microbial peptide homologous to *B. mori*, cecropin D and a proline-rich peptide of unique amino acid sequence were purified by Morishima, et al. [11].

To understand the role of such network of amino acids in the defense mechanism against bacterial infection, it is important to know about many antibacterial peptides which were isolated from different species of insects. Such peptides can be classified into five major groups: Cecropins, insect defensins, attacin-like protein, proline rich peptides and lysozyme.

In *B. mori*, cecropins and lysozyme have been reported as antibacterial peptides by Abraham, et al. [12] According to Hara and Yamakawa, et al. [13] insect defenses are highly effective against Gram-positive bacteria, including pathogenic bacteria such as

Staphylococcus aureus, whereas they do not exhibit strong activity against gram negative bacteria. Also, they isolated antibacterial peptide from *B. mori* larvae which was immunized with gram-positive bacteria *Staphylococcus aureus* called moricin that consists of 42 amino acids.

Some amino acid sequences of the moricin peptide are as follows: Lys-Ala-Ile-Asp-Gly-Val-Arg-Leu-Phe-Pro-Thr-His-Ser. Moricin is believed to have antibacterial activity against both gram negative and some pathogenic positive bacteria such as *Bacillus cereus*. Furthermore, [13] isolated three antibacterial peptides from *B. mori* larvae which were immunized with gram-negative bacteria *E. coli*. These peptides were 32 amino acids long and characteristically rich in proline residues. Some of amino acid of the novel peptides was Pro, Leu, Tyr, Arg, Asp, Lys, Ile, Gly, Phe, Met, Val and Thr. These peptides are effective against gram-negative bacteria such as: *E. coli* and *Acinetobacter* sp.

Gad et al. [14] evaluated the effect of *Escherichia coli* and *Bacillus thuringiensis* infection on amino acids in the fifth larval instar of *B. mori*, 16 free amino acids (FAAs) were observed, these amino acids were: Aspartic acid, Threonine, Serine, Glutamic acid, Proline, Glycine, Alanine, Valine, Methionine, Isoleucine, leucine, Tyrosine, Phenylalanine, Histidine, Lysine and arginine in the control and treatment. There was a marked increase in some FAAs, eg: Threonine, Serine, Proline, Glycine, Valine, Methionine, leucine, Phenylalanine, Histidine, Lysine and arginine after 24 and 48hrs. These FAAs formed the major sequences in moricin and other novel antibacterial peptides [13]. On the other hand, the concentrations of aspartic, glutamic, glycine, alanine, tyrosine and lysine in the haemolymph of immunized larvae were relatively low. These amino acids are six of the known eight major FAAs forming a fibroin molecule of the silk filament [15].

These results can be explained that the larval metabolism was busy exhibiting immunological responses against bacterial infection producing anti-bacterial peptides in addition to cellular immunity while silk protein was greatly decreased.

The Hormonal Regulation of Silkworm Immunity

Insect humoral immunity can be affected by juvenile hormone (JH) and 20-hydroxyecdysone (20E), but how humoral immunity is developmentally regulated by these two hormones in insects and has not yet been elucidated. Fat body produces humoral response molecules and hence is considered as the major organ involved in innate immunity [16]. Tian, et al. [17] suggested that JH plays a

positive role in the regulation of innate immunity in the larval fat body. Also, Riddiford, et al. [18] suggested that JH has a significant role in the control of immune humoral function. A number of studies in *Drosophila* imply that 20E induces AMP mRNA expression and acts as an immune-activator Silverman, et al. [19] while JH acts as an immune-suppressor by antagonizing 20E signaling [20]. Also, Flatt, et al. [21] found that juvenile hormone regulate the immunity of *Drosophila* by inhibiting phenoloxidase (Po) synthesis and prevents cuticular melanization.

Gad, Abdel-Megeed, et al. [22] demonstrated that injection of *B. mori* fifth larval instar with *B. thuringiensis* caused a sharp decrease in the corpora allata (CA) surface area up to 96 h post-infection to reach the minimum value of about 54.5% less than the control which is followed by a marked increase in the CA surface area at 120 h about 12.18% over the control. This result may due to the effect of bacteria on CA activity during a few days of infection and the increase of the CA activity after that may be due to the release of antibacterial peptides as an immunity response against bacteria which helped the gland to reprogrammed itself in its cycle during the last larval instar. The corpora allata (CA) volume was used as an indicator of the juvenile hormone (JH) level [23].

Mutagenic Effects of Pathogenic Bacteria on Silkworm

It is important to explain the side effects and the mode of action by which the injection of the bacteria can affect the physiology and genetics of *B. mori*.

A molecular marker of *B. mori* DNA is the most important method for determining the affected region on DNA since they reveal DNA polymorphisms among genetically related individuals [24]. A similar strategy has been used to identify the nucleopolyhedrovirus (NPV), another important silkworm genotype virus [25]. *B. thuringiensis* is the most widely used microbial pesticides. The biochemical basis of the pesticide is an insecticidal crystal protein (ICP), which is produced by the bacterium as a 133-kDa protoxin that requires proteolytic cleavage in the insect gut for activation. The mutagenic effect of the bacterial injection produced several proteins that enable it to kill insects through the alteration of the physiological processes [26].

Gad, Abdel-Megeed, et al. [22] observed that injection of *B. mori* fifth larval instar with *B. thuringiensis* and *E. coli* caused genotoxicity in *B. mori* DNA band region, which composed of several structural domains that are

disrupted by the toxin secreted by both *B. thuringiensis* and *E. coli* against *B. mori*.

Huang, et al. [27] suggested that injection of *B. mori* with *Bacillus bombysepticus* (Bb) caused a lot of basal metabolic pathways which were significantly modulated. Furthermore, genes of juvenile hormone synthesis and related metabolism showed up regulation, suggesting that juvenile hormone participate in host modulation during the infection. Moreover, host cellular and systemic immune responses are also induced. Similar to *B. thuringiensis* (Bt), Bb can also induce a silkworm poisoning-related response.

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