NOTE

Does hyperthermia increase apoptosis in white spot syndrome virus (WSSV)-infected Litopenaeus vannamei?

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ABSTRACT: Apoptosis plays a critical role in development and maintenance of multicellular organisms. It has also been described as an anti-viral mechanism in both insects and vertebrates. In fact, to escape the immune system and to increase their spread, some viruses such as baculovirus produce anti-apoptotic molecules. Conversely, a recent report showing a positive correlation between the number of apoptotic cells and the severity of white spot syndrome virus (WSSV) infection in Penaeus monodon suggested that apoptosis might be the cause of death in viral-infected shrimp. Searching for the mechanisms involved in the beneficial effect of hyperthermia for WSSV-infected Litopenaeus vannamei (also called Penaeus vannamei) and considering that hyperthermia increases apoptosis in other experimental models, we investigated the presence of apoptosis by Tdt-mediated dUTP nick-end label (TUNEL), from 4 of their examined tissues or organs, compared to 62 (21%) for those at 32°C. Moreover, shrimp at 32°C also had a significantly higher overall mean apoptotic index (AI) than shrimp at 25°C (p < 0.05). Comparison of mean AI at 72, 96 and 120 h post-infection showed that individuals at 32°C presented a significantly higher values than those at 25°C. These results suggested that hyperthermia might facilitate apoptosis in WSSV-infected L. vannamei and might be one of the mechanisms responsible for increased survival of infected shrimp maintained at 32°C.

KEY WORDS: Apoptosis · White spot syndrome virus · Hyperthermia · Litopenaeus vannamei

In the last decade, many components of the apoptotic machinery have been characterized in vertebrates and invertebrates, elucidating part of the biochemical events that underlie the morphological alterations (Hengartner 2000). Apoptosis is mediated by the sequential activation of cysteine proteases or caspases that cleave other caspases and noncaspase substrates such as proteins of the DNA repair system and of the cytoskeleton (Nicholson 1999). This process generates structural and functional modifications related to cell death, including stepwise double-stranded DNA fragmentation that is considered the biochemical hallmark of apoptosis.

Apoptosis is essential in both the development and homeostasis of multicellular organisms and many of its components are remarkably conserved from worms to humans (Bergmann et al. 1998). Regarding the immune response, apoptosis has been described as an effector mechanism in invertebrates (Clem & Miller 1993, 1994), while in mammals it regulates the adaptive response (Koyama et al. 2000). Apoptosis has been proposed as the main anti-viral mechanism in invertebrates (Koyama et al. 2000). Studies on insect cell lines infected with mutant baculoviruses and adenoviruses suggested that the major significance of apoptosis is an abortion of viral multiplication by premature lysis of the infected cells (Pilder et al. 1984, Clem & Miller 1993). Other experimental models showed that several distinct viruses have developed mechanisms to block premature apoptosis of infected cells in order to prolong cell survival and to guarantee the production of viral progeny (Roulston et al. 1999).

Recently, we described a beneficial effect of hyperthermia upon the survival of WSSV-infected Litopenaeus vannamei (also called Penaeus vannamei) but the underlying mechanism is still unknown (Vidal et al. 2001). Taking into account that apoptosis is involved in the invertebrate antiviral response and that hyperthermia facilitates apoptosis in mammalian experimental environments, we investigated the presence of apoptosis by Tdt-mediated dUTP nick-end label (TUNEL) in WSSV-infected Litopenaeus vannamei (also called Penaeus vannamei) and considering that hyperthermia increases apoptosis in other experimental models, we investigated the presence of apoptosis by Tdt-mediated dUTP nick-end label (TUNEL). From 4 of their examined tissues or organs, compared to 62 (21%) for those at 32°C. Moreover, shrimp at 32°C also had a significantly higher overall mean apoptotic index (AI) than shrimp at 25°C (p < 0.05). Comparison of mean AI at 72, 96 and 120 h post-infection showed that individuals at 32°C presented a significantly higher values than those at 25°C. These results suggested that hyperthermia might facilitate apoptosis in WSSV-infected L. vannamei and might be one of the mechanisms responsible for increased survival of infected shrimp maintained at 32°C.

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Apoptosis is the genetically regulated mechanism by which individual cells orchestrate their own deletion in normal and diseased tissues (Kerr et al. 1972, Soini et al. 1998). Its initial description was based on typical morphological features such as shrunken organelles, condensed chromatin, formation of apoptotic bodies and absence of inflammation in the adjacent areas (Kerr et al. 1972, Wyllie et al. 1980).

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models (Hermisson et al. 2000, Yamamoto et al. 2000), we investigated the presence of apoptosis in WSSV-infected *Litopenaeus vannamei* maintained at ambient temperature and in heated tanks. In this study, apoptotic cells were detected by TdT-mediated-dUTP-Nick-End-Labeling (TUNEL), which measures nuclear DNA fragmentation.

**Materials and methods.**

**Experimental individuals and conditions:** *Litopenaeus vannamei* utilized in these experiments were bred in captivity at CENIACUA’s facilities located in Cartagena, Colombia (Atlantic Coast). Before the trials, they were acclimated for 20 d in the CENIACUA Bioassay Laboratory, Tumaco, Colombia (Pacific Coast). To rule out the presence of WSSV prior experimental infection, pleopods from 30 shrimp from the same batch (full siblings) were tested by nested PCR (Lo et al. 1996). Experimental infection was performed by injection of 0.1 ml (3 \( \times 10^7 \) viral copies determined by real time PCR) (Tang-infection was performed by injection of 0.1 ml (3 \( \times 10^7 \) viral copies determined by real time PCR) (Tang-infection was performed by injection of 0.1 ml (3 \( \times 10^7 \) viral copies determined by real time PCR) (Tang-infection was performed by injection of 0.1 ml (3 \( \times 10^7 \) viral copies determined by real time PCR) of our WSSV inoculum at the 4th abdominal segment, as previously reported (Vidal et al. 2001).

Shrimp of 1 to 2 g average weight were maintained in 40 l plastic tanks (10 individuals per tank) containing chlorinated and filtered seawater, 29 ppt salinity, under constant aeration and with 100% daily water exchange. The 2 water temperature regimens used in tanks were 24.9 ± 0.5°C (ambient temperature, hereinafter referred to as 25°C) and 31.6 ± 2°C (hereinafter referred to as 32°C) maintained with immersion heaters (Aquarium system VISI Thems Vtn 100, VISI Therm Tecnology). The shrimp were fed twice a day with a commercial shrimp diet (Purina 25% protein).

**Experimental design:** We studied 3 groups of 50 WSSV-free *Litopenaeus vannamei*. At Day 0, shrimp from Group 1 were injected with WSSV and kept (10 in each tank) at ambient temperature (25°C). Group 2 was also injected with WSSV but maintained at 32°C. Group 3 consisted of healthy uninjected individuals kept at 32°C.

Five shrimp each (1 from each tank of each group) were sacrificed and fixed at 4, 6, 18, 24, 48, 72, 96, 120, 144 and 168 h after infection. Due to the WSSV infection in the shrimp at 25°C, 8 individuals died before they could be sampled; therefore, this group was evaluated up to 120 h only. In this group, we also studied 2 extra moribund individuals fixed at 75 and 92 h post infection. All samples were preserved in Davidson’s fixative for 24 h and then transferred to 70% ethanol until processing as previously described (Bell & Lightner 1988). Routine histopathology (Bell & Lightner 1988) was performed on all samples to rule out concomitant diseases and to verify WSSV infection status. Infections were considered H&E positive when they showed typical WSSV histopathological features including hypertrophied nuclei with basophilic inclusions in the cuticular epithelium, connective tissue, antennal gland, lymphoid organ and/or hematopoietic tissue (Lightner 1996). In addition, selected samples from each time point were also analyzed using an *in situ* hybridization assay for WSSV (DiagXotics). Although we did not find mortalities within the individuals kept at 32°C, 7 shrimp from Group 2 and 1 shrimp from Group 3 were excluded from this study due to the presence of concomitant diseases such as hepatopancreatic parovirus and vibrosis. Thus, the total number of assayed shrimp was 42, 43 and 49 from Groups 1, 2 and 3, respectively.

**Detection of apoptotic cells:** We used the Dead-End™ Colorimetric Apoptosis Detection System (Promega) following the manufacturer’s protocol, except for increasing the hydrogen peroxide concentration from 0.3 to 6% to improve the blocking of endogenous peroxidases. The TUNEL assay is based on the incorporation of a biotinylated nucleotide at the 3’-OH end of fragmented DNA by the enzyme terminal deoxynucleotidyl transferase. These nucleotides are detected by the addition of horseradish-peroxidase-labeled streptavidin and the chromogen substrate diaminobenzidine that stain the apoptotic nuclei brown. After developing, slides were counterstained with Methyl Green (Vector Laboratories) for 1 min at 56°C, rinsed with deionized water and mounted in VectaMount™ (Vector).

Two researchers independently counted the number of positive cells in 10 high power magnification fields (400×) for each organ or tissue (de Jong et al. 2000). These included antennal gland, gills, hepatopancreas, lymphoid organ, heart, stomach, muscle, cuticular epithelium, connective tissue and pericardium for each shrimp. The apoptotic index was determined as the total number of apoptotic cells per shrimp (the sum of apoptotic cells in all high power fields analyzed for each individual. The mean apoptotic index represents the average of the apoptotic index from the 5 shrimp studied per time point. Statistical analysis was performed using ANOVA and Student’s *t*-test.

**Results and discussion.** The presence or absence of WSSV in tissues from experimental individuals was determined (Table 1) by routine histopathology (Lightner 1996) and by detection of WSSV DNA using *in situ* hybridization. Overall, a much higher percentage of WSSV infected shrimp showing typical WSSV nuclear inclusions was found at 25°C (i.e. 62% in Group 1) than at 32°C (i.e. 16% in Group 2). No WSSV inclusions were found in control Group 3 at 32°C. *In situ* hybridization revealed similar results with 54% positive specimens in Group 1 and 18% in Group 2 while Group 3 was negative. Many shrimp samples collected from challenge groups in the early stages of infection gave
These data suggest that hyperthermia might not be the only mechanism responsible for increasing the number of infected cells. We believe that apoptosis after 144 h, a finding that might reflect a decrease in shrimp held at 32°C showed a reduction in mean AI than the mean for shrimp kept at 25°C (Student’s t-test, p < 0.05). Furthermore, comparison of the mean of 8 AI means for each group (4 to 120 h) showed that the overall mean for shrimp kept at 32°C was significantly higher than the mean for shrimp kept at 25°C (Student’s t-test, p < 0.05). These data suggest that hyperthermia might facilitate the apoptotic process, which, in turn, could reduce viral replication and limit infection. Indeed, the shrimp held at 32°C showed a reduction in mean AI after 144 h, a finding that might reflect a decrease in the number of infected cells. We believe that apoptosis is not the only mechanism responsible for increasing survival of WSSV-infected Litopenaeus vannamei kept at 32°C, since all infected individuals analyzed had apoptotic cells. Moreover, apoptosis is not sufficient to eliminate the virus because reducing the water temperature for WSSV-infected shrimp that had been kept in heated tanks for 20 d resulted in mortality (data not shown).

After 48 h of infection, the mean AI of Group 1 started to increase, reaching the highest level at 120 h (Fig. 2). This increase in AI observed at the final stages of disease is consistent with the only report on apoptosis in WSSV-infected shrimp (Sahtout et al. 2001). These authors also found that intranuclear inclusion bodies were positive by fluorescent TUNEL in typical WSSV infected cells. In contrast, the apoptotic cells detected by us (colorimetric TUNEL) did not present hypertrophied nuclei, characteristic of WSSV-infected shrimp (Khanobdee et al. 2002). This discrepancy can be attributed, at least in part, to the high sensitivity of fluorescent dyes. However, the use of fluorescence does not permit the analysis of apoptotic cell morphology, increasing the risk of misinterpretation and false positive results due to necrosis, autolysis and tissue damage through processing (Labat-Moleur et al. 1998, Stadelmann & Lassmann 2000).

Based on the histological findings, i.e. the absence of inflammatory response, it has been suggested that apoptosis is involved in the pathogenesis of WSSV infection and that it might be the cause of death in infected shrimp (Flegel & Pasharawipas 1998). This hypothesis is supported by the results of Sahtout, that showed a positive correlation between the number of apoptotic cells and the severity of WSSV infection (Sahtout et al. 2001). Another study also shows a correlation between apoptosis and mortality in yellow head virus-infected Penaeus monodon and it is suggested that the high number of apoptotic cells may be the primary cause of death (Khanobdee et al. 2002). As reported (Sahtout et al. 2001), we found an increase of mean AI in WSSV-infected shrimp maintained at ambient temperature, but interestingly, this increment was lower than the one observed in individuals subjected to higher temperature conditions. Therefore, our data suggests that apoptosis is associated with survival of WSSV-infected shrimp. Our hypothesis is supported by the fact that apoptosis might be an important antiviral mechanism in insects (Clem & Miller 1993, 1994, Koyama et al. 2000) and that some viruses produce anti-apoptotic molecules such as p35 and IAP (Clem 2001, Roulston et al. 1999) in order to escape it. However, further investigations about the apoptotic routes in Litopenaeus vannamei and their interactions with WSSV molecules are needed to resolve this issue.

<table>
<thead>
<tr>
<th>Diagnostic test</th>
<th>Group 1 (25°C)</th>
<th>Group 2 (32°C)</th>
<th>Group 3 (32°C)</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>n positive/n total</td>
<td>n positive/n total</td>
<td>n positive/n total</td>
</tr>
<tr>
<td><strong>H&amp;E staining</strong></td>
<td>26/42</td>
<td>8/43</td>
<td>0/49</td>
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<tr>
<td>% positive</td>
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<td>0</td>
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<tr>
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<td>18 to 120 h</td>
<td>6 to 48 h</td>
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<tr>
<td><strong>In situ hybridization</strong></td>
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<td>5/27</td>
<td>0</td>
</tr>
<tr>
<td>% positive</td>
<td>54</td>
<td>18</td>
<td>0</td>
</tr>
<tr>
<td>Positive time points</td>
<td>24 to 120 h</td>
<td>45 to 48 h</td>
<td>0</td>
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Fig. 1. WSSV infecting *Litopenaeus vannamei*. Photomicrographs of TUNEL assays performed with tissue sections of WSSV-infected juveniles of *L. vannamei*. Apoptotic nuclei with fragmented and biotin-tagged DNA were visualized as dark brown due to binding of streptavidin labeled with horseradish-peroxidase that reacted with the chromogen diaminobenzidine. The slides were counterstained with methyl green which stained non apoptotic nuclei green. (A) Stomach epithelium (SE) of a shrimp maintained at ambient temperature (25°C) for 18 h post-infection. Black arrow points to an apoptotic nucleus (brown). (B) Stomach epithelium (SE) of a shrimp specimen kept at ambient temperature for 120 h post-infection. Black arrow points to the only apoptotic cell in this field. Red arrow points to an intranuclear inclusion (green), many of which were seen at this stage of infection. (C) Cuticular epithelium (CE) and connective tissue (CT) of a shrimp specimen maintained in heated tanks (32°C) for 96 h post-infection. Black arrow points to an apoptotic nucleus. In this field, only a few cells show intranuclear inclusions (red arrow). (D) Connective tissue (CT) and a vessel (VE) of a shrimp maintained in heated tanks for 120 h post-infection. The majority of the cells in this field are apoptotic (brown nuclei). HA: haemolymph
Looking at the distribution of apoptotic cells in tissues or organs from all individuals of each group, we found that cuticular epithelium (40%) was the preferential site at 25°C, while connective tissue (34%) was the preferential site at 32°C (Fig. 3). In both groups, cells undergoing apoptosis were detected within WSSV target tissues (Lightner 1996) and in hepatopancreatic interstitial cells, as described before (Sahtout 2001). These results indicate that WSSV is necessary to induce apoptosis, either indirectly by eliciting a host response (Clem & Miller 1993, 1994) or directly via a viral mediated mechanism (Roulston et al. 1999).

In summary, the significant increase in number of apoptotic cells in shrimp kept at 32°C suggests that hyperthermia facilitates apoptosis in WSSV-infected Litopenaeus vannamei. This would reduce viral replication, allowing the shrimp to control the disease and survive. This hypothesis is supported by previous reports showing that hyperthermia induces apoptosis by enhancing cytochrome c release in human glioma cells (Hermisson et al. 2000) and by redistribution of Bax in rat testes (Yamamoto et al. 2000). However, further research is needed to establish the role of these and other molecules in the shrimp apoptotic routes and the interactions between virus and host in this context.

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