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Testing may help increase IHHNV tolerance in Pacific white shrimp

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Clinical syndromes are species-specific



Shrimp families were maintained in individual net cages during the study.

The infectious hypodermal and hematopoietic necrosis virus (IHHNV) was first discovered in Pacific blue shrimp, *Penaeus stylirostris*, in 1981, when it caused epizootics associated with mass mortality. IHHNV is the smallest known penaeid shrimp virus. Containing single-stranded DNA with an estimated size of 4.1 kb, it is one of the most prevalent and widespread shrimp viruses. IHHNV can be transmitted both vertically and horizontally.

Once a population is infected with IHHNV, it is very difficult to eradicate the virus. Both the World Organization for Animal Health and United States Marine Shrimp Farming Program list IHHNV as a major pathogenic agent excluded from specific pathogen-free status.

Various shrimp species can be infected by IHHNV in all developmental stages. However, the clinical syndromes are species-specific. Unlike the effects shown in Pacific blue shrimp, infection of Pacific white shrimp, *P. vannamei*, with IHHNV does not result in mass mortality. Instead, animals show runt deformity syndrome (RDS) and large size variation at harvest. For *P. stylirostris*, an IHHNV refractory line was developed and successfully applied in commercial production during the late 1990s.

IHHNV study

To assess the possibility of selecting IHHNV-free shrimp from an infected population and the feasibility of developing an IHHNV refractory line for *P. vannamei*, the dominant shrimp species currently cultured worldwide, the authors conducted a large-scale study to investigate family effects on the prevalence of IHHNV in *P. vannamei* broodstock within the context of a family-based genetic selection program based on non-SPF populations.

The specific objectives were to estimate the IHHNV prevalence in the existing population, investigate the genetic variability of IHHNV susceptibility among families and evaluate the quantitative effects of IHHNV prevalence on shrimp production performance traits. Technical support for the study was

provided by Linda Nunan, Solangel Navarro, Brenda Noble and Dr. Donald Lightner of the Aquaculture Pathology Laboratory at the University of Arizona in the United States.

Experimental design

The study was conducted in an enclosed facility in Mexico where IHHNV infection was known to be present for over two years, using second-generation shrimp from a breeding program. For performance testing, 174 families were divided into three groups with 58 families per group and evaluated in raceways. Shrimp from each group were evenly distributed in four raceways, with about 40

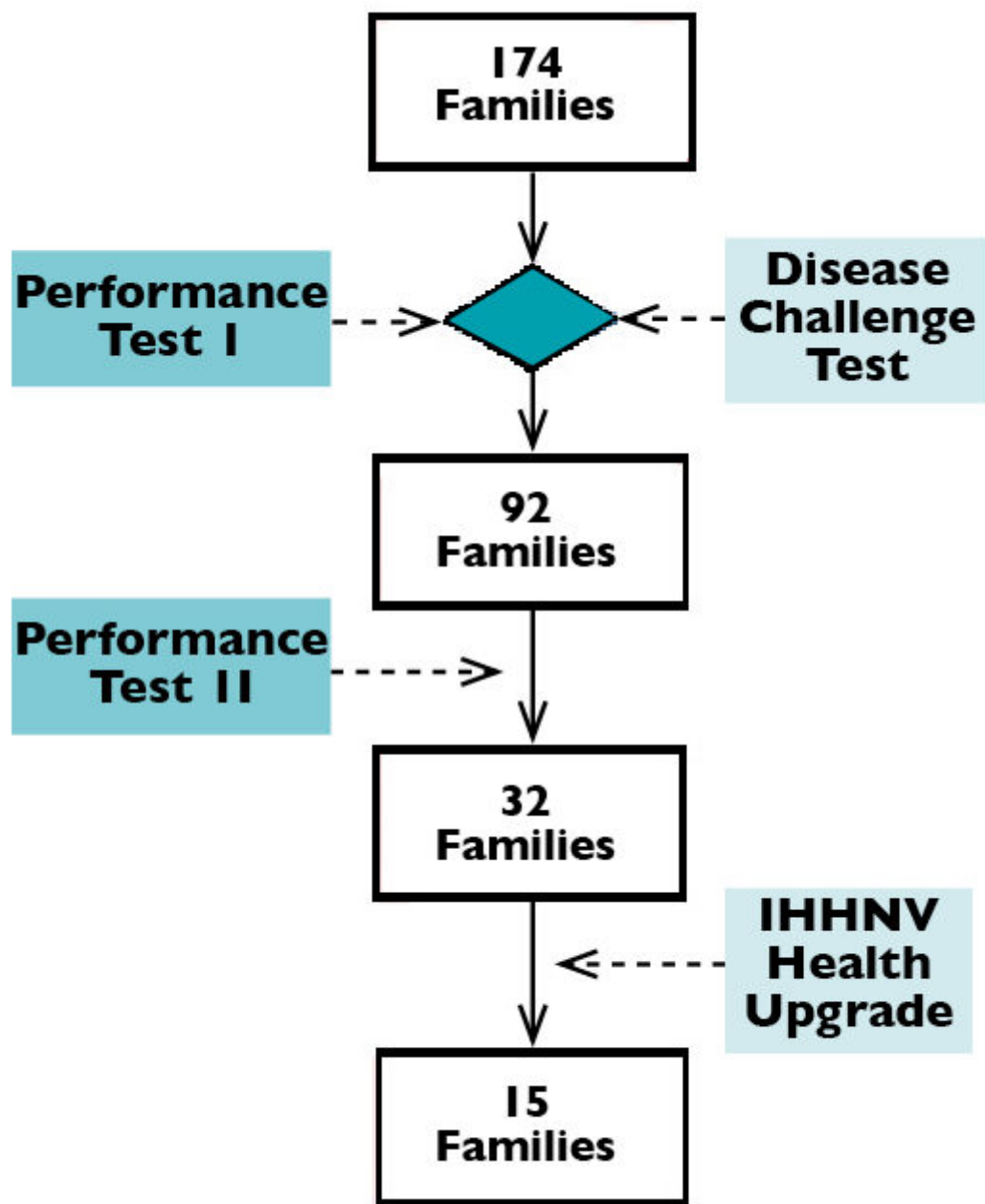


Fig. 1: Genetic selection scheme.

shrimp/family/raceway.

The genetic selection scheme implemented is outlined in Fig. 1. Family selection was based on the results of two-phase performance testing with target harvest weights of 15 and 28 to 30 grams, respectively, as well as the results of disease challenge tests at the juvenile stage with Taura syndrome

virus, white spot syndrome virus and infectious myonecrosis virus that were conducted in the Aquaculture Pathology Laboratory.

The IHNV study was carried out for the 32 selected families after the second performance test phase. All selected survivors were individually weighed, eye tagged and pooled together by family.

About 30 shrimp per family were selected for IHNV testing based on their sizes and visual health. Selected shrimp were held in flow-through raceways at a density of 15 per square meter to minimize the chance of further contamination.

Hemolymph was sampled from individual shrimp and submitted for dot blot hybridization testing on site. Shrimp found negative for IHNV were then held separately in individual tanks. One week later, additional hemolymph samples were sent to a laboratory for polymerase chain reaction (PCR) tests.

Results

Of the total 1,029 shrimp screened for IHNV by dot blot, 585 individuals showed negative. Of the 233 shrimp further tested by PCR, only 47 showed negative. An estimated 11.6 percent of the selected animals were negative by the first PCR.

Statistical analysis showed no significant effect related to testing group, raceway and shrimp sex, but family had a significant effect on the IHNV prevalence, as measured by percentages of the shrimp found negative by dot blot and PCR. Large variations of IHNV prevalence were found among families (Fig. 2), with dot blots showing 3 to 100 percent and PCR levels ranging 0 to 100 percent. Although prevalence by PCR was only 11.6 percent overall, three families had more than 60 percent of their individuals negative to IHNV, which suggested a fair chance to select IHNV-free animals from the population.

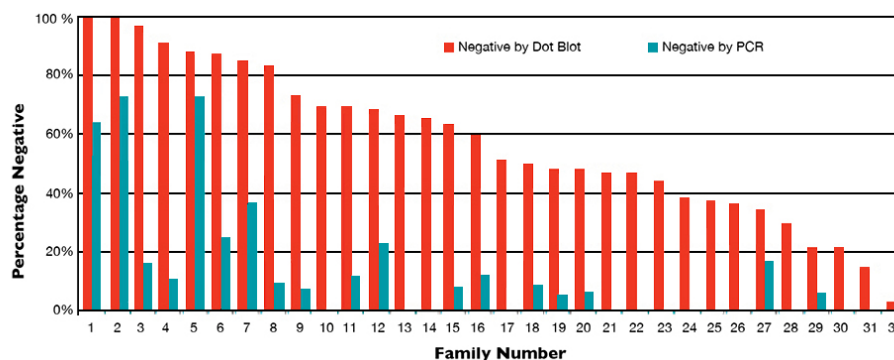


Fig. 2: Variations of IHNV prevalence by dot blot and PCR among shrimp families.

Performance test results showed positive correlation between growth rates from the first and second phases, and slightly negative correlation between growth and survival rate. Although no correlation between the growth and tolerance of Taura syndrome virus was evident, there was strong positive correlation between growout survival and TSV tolerance.

The least square mean of harvest weight for shrimp found negative either by dot blot or PCR was significantly higher than that for positive shrimp and those not selected for testing (Fig. 3). Lack of IHNV was positively correlated with harvest weight and negatively correlated with the coefficient of

variation, but had no apparent effect on survival and TSV resistance.

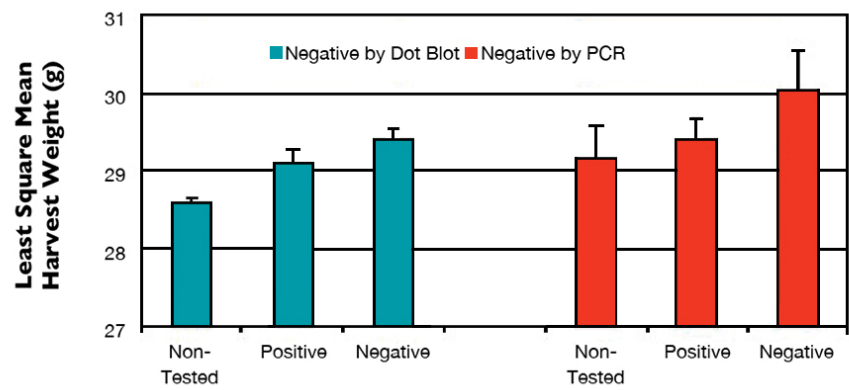


Fig. 3: Effects of severity of IHNV infection on shrimp harvest weight.

These results indicated that the severity of RDS and shrimp size variations were affected by IHNV prevalence. At the individual level, shrimp without IHNV infection or less viral load were larger than the severely infected shrimp. At family level, severe IHNV infection resulted in lower growth and higher coefficient of variation.

It is worth noting that the IHNV study was conducted using selected individuals from the best performance families, so the average 14 percent within-family coefficient of variation was smaller than the coefficient of 30 percent or more for a population with typical RDS. The PCR- percent (11.6 percent) didn't reflect the whole population. Nevertheless, the high percentage of IHNV-negative individuals in some families offered an opportunity to screen out IHNV-free individuals from an infected population.

Perspectives

Practicality remains one of the major challenges to minimize potential contamination at each stage and to utilize clean animals, which typically are handled many times during screening for successful reproduction. The process requires systematic planning to incorporate many important aspects, as well as detailed step-by-step implementation.

IHNV tolerance and resistance are two distinct genetic traits. Large between-family variation in IHNV susceptibility, as shown in this study, may provide a genetic basis to increase IHNV tolerance and to explore the potential of developing a line refractory to IHNV infection via genetic selection.

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