Description of a presumptive hepatopancreatic reovirus, and a putative gill parvovirus, in the freshwater crayfish *Cherax quadricarinatus*

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ABSTRACT: The redclaw freshwater crayfish *Cherax quadricarinatus* has a reputation for being hardy and resistant to handling stress. However, in recent years, possibly since 1996, *C. quadricarinatus* farmers in northern Queensland have noted a decrease in stress resistance in their stock. A presumptive reovirus in the hepatopancreas, and a putative parvovirus in the gills, were associated with chronic mortalities in *C. quadricarinatus* at one northern Queensland farm. Hypertrophic nuclei with margined chromatin were observed in gill epithelium in moribund crayfish which had recently been relocated to a laboratory from the holding tank facility on the farm. Affected nuclei appeared to be vacant or contained a faint granular basophilia in H&E stained sections. However, toluidine blue staining revealed a homogenously granular appearance of the nuclei. Transmission electron microscopy revealed approximately 20 nm diameter virus-like particles within the nucleus. Eosinophilic, Feulgen-negative, cytoplasmic inclusions were observed in distal hepatopancreatocytes in 1 moribund *C. quadricarinatus* collected from the same on-farm holding tank approximately 6 mo later. This crayfish did not display the gill lesions. Transmission electron microscopy showed that the inclusions contained icosahedral virus particles 35 to 40 nm in diameter. The histopathology and preliminary virus morphology of the presumptive hepatopancreatic reovirus, and the histopathology, ultrastructural pathology and preliminary virus morphology of the putative gill parovirus, are reported herein.

KEY WORDS: Freshwater crayfish*C. quadricarinatus* · Crustacea · Aquaculture · Reovirus · Parovirus · Disease · Pathology

INTRODUCTION

The redclaw freshwater crayfish *C. quadricarinatus* has been widely translocated from its natural habitat in the river systems of the Gulf of Carpentaria in Australia for aquaculture purposes. Aquaculture of *C. quadricarinatus* commenced in Australia in the mid 1980s and the industry is still in its infancy. *C. quadricarinatus* is farmed along the north-eastern coast of Australia, as well as in many other countries including Latin American and Carribean countries, China, Taiwan, South Africa, Zambia and the USA (Medley et al. 1994). Ecuador in particular has established large areas for *C. quadricarinatus* aquaculture, with 250 ha currently under cultivation (Romero 1998). *C. quadricarinatus* is typically farmed semi-intensively. In Australia, varying levels of pond management are employed, but in most cases monitoring of pond and stock conditions is less intensive than for prawn aquaculture.

*Cherax quadricarinatus* has frequently been promoted as 'disease-free'. However, health certification, histopathological surveys and studies on sick *C. quadricarinatus* have revealed a number of disease conditions and potential pathogens of the species, including viruses, rickettsia-like organisms and other

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bacteria, protozoans and fungi (Herbert 1987, Anderson & Prior 1992, Ketterer et al. 1992, Owens et al. 1992, Edgerton et al. 1994, 1995, 1996, Edgerton & Prior 1999). Though some of these pathogens have shown clear potential to have a deleterious effect at least at the individual level, none have been associated with widespread epizootic mortalities.

Histopathological surveys to determine the prevalence of pathogens in Cherax quadricarinatus in northern Queensland farms were conducted in 1992 and 1996 (Edgerton 1996, Edgerton & Owens 1999). A number of pathogens and pathological conditions were observed, and some had a very high prevalence in some farms. Notably, C. quadricarinatus bacilliform virus (CqBV) was common and its prevalence had increased markedly in the period between the 2 surveys. However, the lack of reports of significant mortalities on farms in which the prevalence of CqBV was very high (up to 70.5%) suggested that CqBV was not highly pathogenic. Nonetheless, the authors cautioned that CqBV may cause low-level mortalities and/or growth retardation, and may immunocompromise infected C. quadricarinatus.

Since 1994, disease issues have become increasingly important in Cherax quadricarinatus aquaculture in northern Queensland. Initial concern was focussed on the emergence of black spots on the cuticles of crayfish which affected marketability (Edgerton 1996, in press). However, an association between the black spots and poor production has not been proven. One northern Queensland farm had had continuing chronic mortalities since 1993 which have been associated with several pathogens. Initial studies by Edgerton et al. (1995) led these authors to conclude that a systemic rickettsia-like organism (RLO) was the most important pathogen in the mortality episode as it was considered to be the most likely cause of morbidity in approximately one-third of moribund C. quadricarinatus. Recent work by C. K. Tan (James Cook University, unpubl. research 1998, pers. comm.) showed that considerable numbers of moribund crayfish were still being detected on this farm, and that the RLO had an incidence of approximately 80% in sick and moribund crayfish.

In the early years of Cherax quadricarinatus aquaculture, farmed product was considered to be very hardy and was shipped to market for several days in crude conditions, relative to other aquatic species, with high survival rates. In mid-1997, it became apparent that many C. quadricarinatus farmers in northern Queensland had been concerned for some time about an apparent decrease in stress resistance in their stock which manifested as high mortalities of crayfish when in holding facilities and during transit, even when these periods were short. This paper describes a presumptive hepatopancreatic reovirus and a putative gill parovirus that were associated with chronic mortalities at one farm in northern Queensland.

MATERIALS AND METHODS

In October 1997, a group of approximately 30 subadult Cherax quadricarinatus were obtained from a farm in northern Queensland. Soon after relocation the C. quadricarinatus were used in a failed transmission trial with the gill-associated virus/lymphoid organ virus corona-like virus complex (referred to hereafter as GAV) which infects Penaeus monodon in Australia (Spann et al. 1995, 1997). Moribund crayfish detected prior to and during the experiment, and some apparently healthy crayfish at the termination of the experiment, were fixed by injection with Davidson’s fixative (Humason 1967). Gill tissue was removed from several crayfish and fixed in 2.5% glutaraldehyde for transmission electron microscopy.

Two moribund Cherax quadricarinatus were collected from the same on-farm holding tank in June 1998 and were fixed by injection with Davidson’s fixative.

Tissues for light microscopy were fixed for 48 h in Davidson’s fixative and were then trimmed into micr-ocassettes. Full sagittal sections of the cephalothorax were taken, as well as additional sections of antennal gland, heart and gill. Tissues were processed routinely for histology and haematoxylin and eosin-stained sections were prepared. Special stains were prepared according to Croft & Stevens (1992), and included Feulgen’s and Brown and Brenn’s Gram stain. Histological slides were examined using a Leitz Orthoplan microscope.

Gill tissues for transmission electron microscopy (TEM) were fixed for 1.5 h and then rinsed several times and stored at 4°C in 0.1 M cacodylate buffer (pH 7.4). Tissue from 1 of the control Cherax quadricarinatus were postfixed in 1% osmium tetroxide in 0.1 M cacodylate buffer for 2 h at room temperature, washed several times in 0.1 M cacodylate buffer, dehydrated through a series of ethanols and embedded in Epon resin. Hepatopancreatic tissues for TEM were removed from unstained 5 µm sections on glass slides. Tissues were deparaffinised in xylene for 15 min, rehydrated in a series of ethanol, fixed in 3% glutaraldehyde in 0.1 M cacodylate buffer for 15 min at room temperature, rinsed in 0.1 M cacodylate buffer, post-fixed in 1% osmium tetroxide in 0.1 M cacodylate buffer for 30 min, dehydrated in a series of ethanol and embedded in Epon resin. Ultrathin sections were cut using a Reichert-Jung Ultracut ultramicrotome, and were viewed using a Jeol 1010 transmission electron microscope.
RESULTS

Failed transmission trial in October 1997

One *Cherax quadricarinatus* was moribund, some others were lethargic, but the majority appeared healthy, on arrival at the laboratory. Several *C. quadricarinatus* died prior to the commencement of the experiment, as did many soon after being injected with extracts from GAV-infected *Penaeus monodon*. Several control *C. quadricarinatus* died prior to the termination of the trial at 24 d post inoculation.

Histopathology for *Cherax quadricarinatus* injected with the GAV extracts was consistent with the non-injected crayfish. Some *C. quadricarinatus* had light *Cherax Giardiavirus*-like virus (CGV) or light CqBV infections, and 1 had diffuse systemic nodules consistent with mild bacteraemia. The only consistent lesion in the *C. quadricarinatus* was multifocal hypertrophic nuclei in the gill epithelium. All moribund and 4 of 5 clinically normal *C. quadricarinatus* that were sampled displayed the lesions. A moribund *C. quadricarinatus* sampled prior to commencement of the transmission trial displayed the highest intensity (the greatest number of affected nuclei) and associated haemocytic infiltration. Foci of hypertrophic nuclei were clearly observed in cross-sectioned gill lamellae amongst normal lamellae, and occasionally the majority of gill epithelial nuclei in a longitudinal section of a lamella were affected (Figs. 1 & 2). The epithelium of heavily affected lamellae was more dense, as evidenced by darker staining with H&E and increased electron density, and had shrunken away from the cuticle.

Hypertrophic nuclei contained margined chromatin and had peripheral nucleoli (Fig. 3). The interior of affected nuclei in H&E sections appeared to be vacant or contained a faint granular basophilia. However, the interior of the nucleus appeared homogenously granular in 1 µm thick toluidine blue sections. Feulgen staining of affected nuclei was too light to be of value. Consistently sized, electron-dense, rounded virus-like particles occasionally showing angular profiles, approximately 20 nm in diameter, were scattered throughout the interior of the nucleus (Fig. 4). The cytoplasm of affected cells was more electron-dense than uninfected cells, and the mitochondria displayed irregular profiles and were swollen.

Cytopathology suggestive of a sequence of infection was observed. The earliest stage of infection was an in-
and unaffected epithelia in longitudinally sectioned tubules as affected epithelia displayed significantly less lipid vacuolation than normal epithelia and were flattened (Fig. 6). Affected tubules were surrounded by haemocytic infiltrates (Fig. 7). Inclusions were Feulgen-negative, often irregularly shaped presumably due to remaining lipid vacuoles, and often closely associated with the apparently normal nucleus (Figs. 8 & 9).

TEM revealed that the inclusions consisted of virogenic stroma containing regularly sized non-enveloped virions. The virions were 35 to 40 nm, had an angular appearance with hexagonal and pentagonal forms, and were spaced regularly in the virogenic stroma never in paracrystalline arrays (Fig. 3). Electron microscopy confirmed that the inclusions were intimately associated with the nucleus. However, the nucleus appeared unaltered.

**DISCUSSION**

The poor survival of *Cherax quadricarinatus* prior to the commencement of the transmission trial suggests

![Fig. 3. *Cherax quadricarinatus*. Hypertrophic nuclei in the gill epithelium. Affected nuclei (arrows) have marginated chromatin and displaced nucleoli, and contain granular material in the centre. Uranyl acetate and lead citrate. Scale bar = 2.35 μm](image)

![Fig. 4. *Cherax quadricarinatus*. Hypertrophic nuclei in the gill epithelium. Virus-like particles within the central portion of the nucleus (arrows). Note that some have angular profiles. Uranyl acetate and lead citrate. Scale bar = 66.7 nm](image)
that they were initially in poor health. Furthermore, later discussions with the farmer revealed that this group of *C. quadricarinatus* had been held in the on-farm holding tank for a considerable period prior to relocation to the laboratory, and that a significant number of deaths had occurred in that time. Moreover, poor survival in the holding tank, as well as in several tanks which housed broodstock for a controlled breeding programme, had been a constant problem for a considerable length of time. It is presumed that the *C. quadricarinatus* that died after being injected with extracts containing GAV did so due their suboptimal health state resulting in an inability to clear the foreign proteins.

The only consistent remarkable histopathological lesion in *Cherax quadricarinatus* in the transmission trial was the hypertrophic gill nuclei. Edgerton (1996) observed hypertrophic nuclei in gill epithelial cells in *C. quadricarinatus* from 4 farms during a histopathological survey in 1996 at a prevalence of up to 13.6%. The same lesions had not been detected previously by the author, either in a histopathological survey of the same farms conducted in 1992, or in a study of chronic mortality at one of the surveyed farms in northern Queensland in 1993 (Edgerton et al. 1995, Edgerton 1996, Edgerton & Owens 1999). The apparent 'emergence' of the lesion in *C. quadricarinatus* from farms in northern Queensland, the association of the lesions with mortalities in 1 group of *C. quadricarinatus*, in the absence of other consistent remarkable lesions, and the multifocal nature of the lesions suggest that it may have an infectious aetiology. Furthermore, consistently sized approximately 20 nm virus-like particles observed within the affected cells suggest that lesion is caused by a virus, possibly a parvovirus.

Similar hypertrophic nuclei occur in the gill epithelium of *Penaeus monodon* also from farms in northern Queensland. This industry has experienced significant widespread mortality episodes since 1994 which have been associated with a number of viruses, including GAV. However, few consistent lesions have been detected in moribund prawns. The name Midcrop Mortality Syndrome (MCMS) has been coined for the syndrome. The significance of the hypertrophic nuclei in gill epithelium in *P. monodon* in northern Queensland and their relationship to the similar lesions in *Cerax quadricarinatus* have yet to be determined. However, Owens et al. (1998) have associated a parvo-like virus similar or identical to spawner-isolated mortality virus (SMV) with MCMS. SMV has been associated with lesion similar to those reported here, in myocardium and striated muscle but not in gill epithelium (Fraser & Owens 1996, Owens 1997).

Sections from a moribund *Cherax quadricarinatus* fixed prior to the commencement of the trial and a moribund *C. quadricarinatus* injected with an extract containing GAV were both

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**Fig. 5. Cherax quadncarinatus. Hypertrophic nucleus (arrow) in an early stage of infection, containing a centrally developing viroplasm (V) in the gill epithelium. Uranyl acetate and lead citrate. Scale bar = 0.9 μm**

**Fig. 6. Cherax quadrcarinatus. Hepatopancreatic reo-like virus in the crayfish. Cytoplasmic inclusions in distal hepatopancreaticocytes. Note the strong demarcation (arrows) between infected (I) and normal (N) epithelium due to lipid reduction in the former. L = lumen of hepatopancreatic tubule. H&E. Scale bar = 62.5 μm**
gentia cytoplasmic inclusions in hepatopancreatocytes (Lightner 1996). Furthermore, the consistency of the penaeid reovirus virogenic stroma is similar to that observed in this study (cf. Fig. 28, Hukuhara & Bonami 1991). However, whilst the 50 to 70 nm penaeid reovirus is smaller than most reoviruses, which are typically 60 to 80 nm in diameter (Murphy et al. 1995), it is still considerably larger than the virus reported here. Reoviruses consist of an electron-dense core surrounded by a double-shelled capsid, with the diameter of the core being approximately 50 to 70% of that of the whole virion. The size of the virus reported herein is therefore similar to that of reovirus cores, and the virions visualised in these preparations may have been immature reovirus virions, without the double-shelled capsid, within the viroplasm.

The picornavirus Taura syndrome virus (TSV) accumulates in the cytoplasm of infected cells in penaeid prawns (Hasson et al. 1995). Other small cytoplasmic viruses which infect crustaceans include Chesapeake Bay virus (CBV; Johnson 1978) and Macrobrachium muscle virus (MMV; Tung et al. 1999, Arcier et al. 1999). TSV, CBV and MMV are all around 30 nm in diameter, which is considerably smaller than the virus reported here. Other similar-sized viruses which accu-

Fig. 7. Cherax quadricarinatus. Hepatopancreatic reo-like virus in the crayfish. Cross-section of infected hepatopancreatic tubule. Note the cytoplasmic inclusions within vacuoles (arrows) and the strong haemocytic infiltrate (H) surrounding the tubule. H&E. Scale bar = 28.6 µm

positive following in situ hybridisation with a gene probe for the parvo-like virus involved in MCMS (C. McElnea, James Cook University, unpubl. data). Positive signal was displayed in nuclei in the hepatopancreas, midgut and foregut epithelia, and gill epithelia, though the hypertrophic nuclei were not positive. Similarly, hypertrophic nuclei in gill epithelia in Penaeus monodon do not react positively following in situ hybridisation with the parvo-like virus gene probe (McElnea pers. comm.).

Similar hypertrophic nuclei with marginated chromatin, but without intranuclear inclusions obvious with light microscopy, are commonly observed in crustaceans infected by various other parvo-like viruses such as Cherax destructor systemic parvo-like virus (Edgerton et al. 1997) and infectious hypodermal and haematopoietic necrosis virus (J. R. Bonami, University of Montpellier 2, pers. comm.). In these cases, the lesions are presumed to represent early or chronically infected nuclei.

The hepatopancreatic virus reported herein has not been reported previously in Cherax quadricarinatus, nor has similar histopathology or cytopathology been reported previously in this species. The cytopathology is reminiscent of reovirus in penaeid prawns which consists of eosinophilic to ma-

Fig. 8. Cherax quadricarinatus. Hepatopancreatic reo-like virus in the crayfish. Higher magnification of infected hepatopancreatocytes. Note that the inclusions (arrows) are within cytoplasmic vacuoles and that some inclusions are closely associated with the nucleus. N = normal epithelium. H&E. Scale bar = 20 µm
Cherax quadricarinatus aquaculture commenced in Australia in the mid-1980s. After a rapid increase in production in the early 1990s, production has stagnated at around 60 t between 1994/95 and 1997/98 even though the number of producing farms has doubled in this time (Lobegeiger 1999). There is mounting evidence, both proven and circumstantial, to suggest that disease is now becoming a limiting factor in the development of C. quadricarinatus aquaculture. The emergence of disease issues should not be viewed as surprising, rather as expected, because this is consistent with the developmental sequence of other aquaculture and indeed agriculture industries.

Fig. 9. Cherax quadricarinatus. Hepatopancreatic reo-like virus in the crayfish. Electron micrograph of an infected hepatopancreatocyte. Note that the inclusion (arrow) is closely associated with the apparently normal nucleus (N). Uranyl acetate and lead citrate. Scale bar = 1.3 μm. (Tissues fixed in Davidson’s fixative)

mulate in the cytoplasm in arthropods include nodaviruses (~30 nm diameter) and tetraviruses (35 to 38 nm) (Garzon & Charpentier 1991, Reinganum 1991). Determination of the relatedness of this virus to other arthropod viruses will require further studies.

This paper presents the results of a minor and opportunistic study on Cherax quadricarinatus diseases. Like many studies resulting in the description of new pathogens in C. quadricarinatus, including many by the senior author, this study was too limited in scope and involved too few specimens to give any indication, but the most preliminary, as to the significance of the viruses described herein. Consequently, determination of the association of these viruses and other pathogens with the ongoing mortalities at this farm, and with the decrease in stress resistance of farmed C. quadricarinatus, requires further examination. Bacterial infections are common in freshwater crayfish holding facilities and are likely contributors to the mortalities seen at this farm, and in the wider C. quadricarinatus farming community, as regular cleaning of such facilities typically is not routine.

Cherax quadricarinatus still displays considerable potential for aquaculture as farms utilising best practice culture have been achieving 3 t ha⁻¹ yr⁻¹ across the total ponded area, with average production from growout ponds exceeding 4 t ha⁻¹ yr⁻¹ (Wingfield 1999). Research priorities need adjustment so that disease issues in C. quadricarinatus aquaculture can be overcome to allow sustainable development of the industry.

Fig. 10. Cherax quadricarinatus. Hepatopancreatic reo-like virus in the crayfish. Higher magnification of an inclusion showing electron-dense virions within viroplasm. Note that many of the virions have angular profiles (arrows). Uranyl acetate and lead citrate. Scale bar = 111 nm
Acknowledgements. Thanks to Drs Catriona McElnea and Leigh Owens, James Cook University, for conducting and analysing the in situ hybridisation of selected sections with their gene probe to the parvo-like virus from *Penaeus monodon* in Australia. Helpful discussions with Dr J. R. Bonami, University of Montpellier 2, Prof. D. V. Lightner, University of Arizona, and Dr Leigh Owens are greatly appreciated. Thanks also to 3 anonymous reviewers for their valuable comments.

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