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This document was created in response to a Freedom of Information request made to CSIRO.

FOI Number: FOI2011/69

Date: 31 July 2012

Request: Documents relating to the outbreak of a herpes-like virus in Victoria abalone, now known as Abalone Viral Ganglioneuritis ("AVG") which commenced during December 2005 (the "Victorian Outbreak")

Documents: Part 3 – Documents 51-76

For more information, please refer to CSIRO's FOI disclosure log at [www.csiro.au/FOILog](http://www.csiro.au/FOILog)

## **DOCUMENT 51**

**EXEMPT IN FULL – s 47B(a)**

## **DOCUMENT 52**

**EXEMPT IN FULL – s 47B(a)**

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**From:** Crane, Mark (LI, Geelong)  
**Sent:** Wednesday, 7 June 2006 10:28 AM  
**To:** Elliott, Nick (CMAR, Hobart)  
**Subject:** RE: Abaloen virus / any progress

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Hi Nick,

We had negotiated that if we could get \$30,000.00 from any source and put it through Victoria State then FRDC could match it to make a total of \$60,000.00 for this financial year. The industry has not been able to raise any funds to date but I believe that it is still under discussion - but the financial year is now almost over. Certainly FRDC preproposal(s) have been prepared and whether they are processed out-of-cycle is also being considered - but nothing definite.

Mark

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
**CSIRO Livestock Industries**  
**Private Bag 24**  
**Geelong Vic 3220**

**International Phone: +61 3 52 275118**  
**International Fax: +61 3 52 275555**  
**email: [mark.crane@csiro.au](mailto:mark.crane@csiro.au)**

-----Original Message-----

**From:** Elliott, Nick (CMAR, Hobart)  
**Sent:** Wednesday, 7 June 2006 09:24  
**To:** Crane, Mark (LI, Geelong)  
**Subject:** Abaloen virus / any progress

Hi Mark

I have had no response from the FFF office on possible support. I will check with them.

Have you and others had any luck with FRDC?  
Nick

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**From:** Crane, Mark (LI, Geelong)  
**Sent:** Thursday, 8 June 2006 3:31 PM  
**To:** 'Mehdi.Doroudi@dpi.vic.gov.au'  
**Cc:** malcolm.lancaster@dpi.vic.gov.au; Corbeil, Serge (LI, Geelong)  
**Subject:** RE: URGENT - Comments on the pre-proposal  
**Attachments:** FRDC Abalone herpesvirus pre-proposal 060608.doc

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Mehdi,

I have revised the preproposal as discussed. Included are all the costs for AAHL. I have included salaries for technical support (salaries and operating) for Attwood. You will need to put in values for your components which will increase the total.

The breakdown is as follows:

AAHL Salaries: \$193 661  
AAHL Operating: \$41 339  
AAHL Travel: \$2 000  
AAHL In-kind: \$151 842

Attwood Salaries: \$42 000  
Attwood Operating: \$10 000  
Attwood Travel: \$2 000  
Attwood in-kind: \$21 000

Total to date: \$463 843

FRDC's contribution is \$291 000. If we get any cash contributions then this will be reduced.

Hope this makes sense.

Cheers

Mark

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
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**Geelong Vic 3220**

**International Phone: +61 3 52 275118**  
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-----Original Message-----

**From:** Mehdi.Doroudi@dpi.vic.gov.au [mailto:Mehdi.Doroudi@dpi.vic.gov.au]  
**Sent:** Thursday, 8 June 2006 14:03  
**To:** Crane, Mark (LI, Geelong)

**Cc:** malcolm.lancaster@dpi.vic.gov.au; Corbeil, Serge (LI, Geelong)  
**Subject:** RE: URGENT - Comments on the pre-proposal

Mark,

Agreed as discussed.

Regards  
Mehdi Doroudi - DVM, PhD  
Research Director  
Marine & Freshwater Systems - PIRVic  
Department of Primary Industries

Ph: +613 5258 0272 Fax: +613 5258 0270  
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Email: mehdi.doroudi@dpi.vic.gov.au  
2A Bellarine Hwy, Queenscliff VIC 3225 Australia  
PO Box 114, Queenscliff VIC 3225 Australia

Mark.Crane@csiro.au

08/06/2006 01:36 PM

To: Mehdi.Doroudi@dpi.vic.gov.au  
cc: malcolm.lancaster@dpi.vic.gov.au, Serge.Corbeil@csiro.au  
Subject: RE: URGENT - Comments on the pre-proposal

How about this Mehdi:

A 2-year project to achieve:

1. Development of management strategies and control options
2. Sequence genes and develop conventional PCR (1.0 FTE for 24 months at AAHL)
3. Use sequence to develop ISH (1.0 FTE for 12 months at VicDPI)

The real-time PCR would be dropped from this project to reduce the size of the project (but we may develop it in-house - anticipating future surveillance needs)

If you agree, I can revise our part of the preproposal and the research budget.

Mark

**MARK CRANE Ph.D.**  
**Project Leader**  
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**Australian Animal Health Laboratory**  
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**email: mark.crane@csiro.au**

-----Original Message-----

**From:** Mehdi.Doroudi@dpi.vic.gov.au [mailto:Mehdi.Doroudi@dpi.vic.gov.au]

**Sent:** Thursday, 8 June 2006 13:18

**To:** Crane, Mark (LI, Geelong)

**Cc:** malcolm.lancaster@dpi.vic.gov.au; Corbeil, Serge (LI, Geelong)

**Subject:** RE: URGENT - Comments on the pre-proposal

Mark,

Thanks for the message.

I understand the importance of reliable diagnostic tests (specifically as confirmatory tests). However, when it comes to PCR, it is more important to have access to a rapid test (than reliable) or tests that can detect virus while abalone is not clinically affected. This will definitely help in the development of management strategies, however, control measures could still be developed in the absence of a PCR test.

To be able to go forward from here, we need to make a decision how we want to pursue this pre-proposal. The possible options are:

1) To submit two separate pre-proposals and wait for the FRABs and FRDC's comments. This means that you will need to submit the current pre-proposal as it stands to VICFRAB by tomorrow and I will take care of the management concept.

2) To incorporate my comments into the current pre-proposal and add additional person to deliver on diagnostic work within 12 months or to extend the diagnostic side of the project for another 12 months.

Please indicate which approach would you prefer "option 1 or 2".

Regards

Mehdi Doroudi - DVM, PhD  
Research Director  
Marine & Freshwater Systems - PIRVic  
Department of Primary Industries

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Mark.Crane@csiro.au

08/06/2006 11:26 AM

To: Mehdi.Doroudi@dpi.vic.gov.au  
cc: Serge.Corbeil@csiro.au, malcolm.lancaster@dpi.vic.gov.au  
Subject: RE: URGENT - Comments on the pre-proposal

Mehdi,

Apologies for the lack of effective communication - I made the assumption that I was communicating with Victoria via Malcolm but in retrospect we should have cc'd you on everything.

The only problem I have with this proposal is that it will take more than 12 months to develop a rapid and reliable detection and identification procedure (PCR-based) - minimum 2 years would be required. Information we have to-date indicates that this is a new virus - the ostreid herpesvirus PCR primers do not recognise this abalone herpesvirus and therefore we would have to start from ground zero. If everything went smoothly it might be quicker (18 months) but rarely do things go smoothly in research. One option would be to put more

than one person on the project here at AAHL.

Unfortunately without the diagnostic tools everything else is more difficult. I do not know how you can put in fully effective management systems without rapid and reliable diagnostics.

Mark

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
**CSIRO Livestock Industries**  
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**Geelong Vic 3220**

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**email: mark.crane@csiro.au**

-----Original Message-----

**From:** Mehdi.Doroudi@dpi.vic.gov.au [mailto:Mehdi.Doroudi@dpi.vic.gov.au]  
**Sent:** Thursday, 8 June 2006 10:33  
**To:** Crane, Mark (LI, Geelong); Malcolm.Lancaster@dpi.vic.gov.au  
**Subject:** URGENT - Comments on the pre-proposal

Mark / Malcolm,

Your preproposal is about a long-term achievement (4 years of study) and is just focused on diagnostic techniques. I understand that diagnosis is your area of expertise and you may not have the answers to other questions, however, both industry and Vic DPI are after short-term resolutions to this outbreak. Today our stakeholders are after management solutions. There would always be a chance to develop long-term programs on this virus (PhD, POSTDOC etc.) the same as QX, Perkinsus, Haplosporidium etc. From my perspective, the current focus should be on what could be achieved in a short period of time (max 12 months).

I was under the impression that you guys will discuss with me any further development of FRDC proposals on this matter ( I think that we agreed on that previously) but obviously it did not happen until I approached you a couple of days ago. It is important to communicate effectively as we achieve more if we share our resources and expertise.

I have asked VicFRAB to extend the deadline for this proposal until lunchtime this Friday. There may be a chance to have an early start if the final proposal is approved by FRDC. Cash contribution could be provided through both Vic government and industry. I need you to go through my comments as soon as possible and let me know if we have a consensus on what I have proposed below or that you would prefer to pursue your preproposal as it is now.

**Specific Comments on Pre-Proposal:**

**Title:**

Developing Management Strategies for Herpes-Like Virus Infection in Abalone

**Leading Agency:**

Vic DPI (PIRVic)

**Collaborative agencies:**

AAHL, CSIRO and Industry (both aquaculture and wild sector).

**Principle Investigators:**

Dr Mehdi Doroudi

Dr Serge Corbeil (Mark, please advise if we could have more than one PI)

**Proposed Co-Investigators:**

Dr Mark Crane

Dr Malcolm Lancaster

**Duration of the project:**

12 months

**Budget:**

To be reviewed if we agree on the proposed changes.

**Need:**

First paragraph, last sentence could be changed to "If appropriate management strategies including rapid diagnostic techniques are not developed this emerging virus may have the potential to adversely impact Australian abalone industry".

Prior to the last paragraph add following as a new paragraph:

In order to protect Victoria's valuable abalone industry, there is a need to develop specific management strategies which incorporates disease monitoring, detection, response and control measures both in farm and wild stocks. These include the implementation of routine health management procedures and a system for the exclusion of virus. Health management strategies for both farmed and wild abalone sectors and processing plants could be developed to improve productivity of farmed sector and to protect the health status of wild population of abalone. The application of effective detection and exclusion or control methods incorporates into a workable biosecurity plan to minimise the impacts of this virus.

The last paragraph of 'Need' should be simplified.

**Objectives:**

To improve physical, chemical and biological measures of biosecurity within abalone farming systems to prevent the introduction and spread of virus

To develop a "rapid and reliable diagnostic and detection method"

To develop a "code of practice" for commercial divers to avoid further spread of virus in wild population

To develop a "practical biosecurity program" for abalone processing plants

Mark, to save each other's time, please let me know if you are happy with my suggestions made so far then I will have more to contribute into Industry and Management Consultation, Direct Benefits, Design and Methodology etc.

The deadline is tomorrow. Your prompt response will be appreciated.

Regards

Mehdi Doroudi - DVM, PhD  
Research Director  
Marine & Freshwater Systems - PIRVic  
Department of Primary Industries

Ph: +613 5258 0272 Fax: +613 5258 0270  
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PO Box 114, Queenscliff VIC 3225 Australia

# AQUATIC ANIMAL HEALTH SUBPROGRAM

## 2007-08 Preliminary Research Proposal



### Project Title

Aquatic Animal Health Subprogram: Development of management strategies for herpes-like virus infection of abalone

### FRDC Strategic challenge identification

Natural Resources Sustainability

### Co-Principal Investigators Contact Details

Title: Dr Mehdi Doroudi Organisation: Vic DPI (PIRVic) Mailing Address: PO Box 114, Queenscliff VIC 3225  
Phone No: 03 5258 0272 Fax No: 03 5258 0270 Email: serge.corbeil@csiro.au

Title: Dr Serge Corbeil Organisation: CSIRO Mailing Address: Private Bag 24, Geelong, Vic, 3220  
Phone No: 03 5227 5254 Fax No: 03 5227 5555 Email: serge.corbeil@csiro.au

### Commencement and completion date

Commencement date: 01-02-2007

Completion date: 31-01-2009

### Preliminary Budget

FRDC Contribution	2006-07	2007-08	2008-09	2009-10	TOTAL \$\$s
Salaries and on costs (AAHL)	46 149	95 990	51 522		193 661
Salaries and on costs (DPI Vic)		20 400	21 600		42 000
Travel		2 000	2 000		4 000
Operating	9 851	25 010	16 478		41 339
Capital (equipment)	0	0	0		
<b>FRDC TOTAL</b>	<b>56 000</b>	<b>143 400</b>	<b>91 600</b>		<b>291 000</b>
Research Organisation contribution (AAHL)	35 893	75 857	40 093		151 842
Research Organisation contribution (DPI Vic)		10 000	11 000		21 000
Total of Industry & Other Funding	x	x	x		x
<b>GRAND TOTAL in \$\$s</b>					<b>463 842</b>

### Need

In December 2005/January 2006, a disease outbreak caused high mortality rates in abalone from two farms in Victoria. A third Victorian farm also experienced disease but to a lesser extent. The abalone species affected by the outbreak are *Haliotis laevigata*, *H. rubra*, and *H. laevigata* x *H. rubra* hybrid. Histopathology performed on moribund animals indicated a ganglioneuritis – infiltration of haemocytes in multiple ganglia and nerves (cerebral ganglion affected, also other ganglia and nerves). Examination by electron microscopy revealed the presence of a herpes-like virus in the pleuropedal ganglion. Preliminary transmission studies, carried out within AAHL's high biosecurity facility, indicated that this emerging virus is highly pathogenic and can be transmitted to healthy abalone through the water column. In addition, the viral suspension remains infectious after undergoing a dilution up to 1 in 100. So far, the virus has not grown in tissue culture. Attempts by the farm managers to eliminate the disease from the abalone farms have failed partly due to the lack of detection methods specific to the aetiological agent which would allow early diagnosis of infected animals. If appropriate management strategies including rapid diagnostic techniques are not developed this emerging virus may have the potential to have a significant adverse impact on the Australian abalone industry.

In order to protect Victoria's valuable abalone industry, there is a need to develop specific management strategies which incorporate disease monitoring, detection, response and control measures both in farm and wild stocks. These include the

implementation of routine health management procedures and a system for the exclusion of virus. Health management strategies for both farmed and wild abalone sectors and processing plants could be developed to improve productivity of farmed sector and to protect the health status of wild population of abalone. The application of effective detection and exclusion or control methods incorporates into a workable biosecurity plan to minimise the impacts of this virus.

Access to diagnostic tests that are rapid, reliable and sensitive is of fundamental importance for effective control/management of disease outbreaks. In addition to surveillance tools such as PCR, better procedures/reagents for disease diagnosis are required. Presence of histological lesions provides a presumptive diagnosis. The development of in situ hybridisation probe(s) for the localisation of abalone herpesvirus within histological lesions will provide a means for definitive diagnosis to be made. The development of molecular tools and reagents will allow researchers and industry to rapidly and specifically detect and locate the virus in abalone tissues therefore providing a vital means for diagnosis and facilitating a better understanding of the epidemiology of this disease, leading to more efficient management of disease outbreaks.

### Objectives

1. To improve physical, chemical and biological measures of biosecurity within abalone farming systems to prevent the introduction and spread of virus.
2. To develop a "code of practice" for commercial divers to avoid further spread of virus in wild populations.
3. To develop a "practical biosecurity program" for abalone processing plants.
4. To identify nucleic acid sequences of the emerging abalone herpesvirus (via PCR-based gene amplification and sequencing) necessary for the development of diagnostic tools (e.g. conventional PCR, TaqMan PCR, ISH probe).
5. To develop and validate PCR assays for the detection of the abalone virus.
6. To develop an in situ hybridisation assay specific for the abalone herpesvirus
7. To document an Australian and New Zealand Standard Diagnostic Procedure and submit for external review.

### Industry and Management consultation

The abalone Growers Association of Victoria strongly supports the proposal.

VicFRAB

Abalone Subprogram

AAH Subprogram

NSW FRAB

QFIRAC

SAFRAB

WAFRAB

Tas FRAB

### Direct benefits and beneficiaries

1. All sectors of the abalone wild-capture and aquaculture industries (farmers, divers, processors) will directly benefit from increased biosecurity developed through the implementation of this project.
2. The diagnostic tests will be available to detect and identify the emerging virus in abalone (*Haliotis* spp.), and other potential host mollusc species.
3. Export certification services will be available to industries that wish to develop export markets as well as translocate farmed stock between regions without transmitting disease. In addition, should pathogenic agents be detected during health surveys, industries and State officers will be able to make informed decisions with regards to brood stock translocation, stock destruction etc. Specifically, mollusc aquaculture industries and State agencies will be able to develop health surveillance programs in collaboration with AFDL. In addition, Australia will be better prepared to negotiate with international trading partners on issues concerned with the importation of disease free molluscs from Australian sources.

### Estimated Flow of Benefits

Fisheries and aquaculture managed by States/Territories	
NSW	4.5%
Qld	1%
SA	15%
Tas	25%
Vic	50%
WA	4.5%
AFMA managed fisheries	0%
Other beneficiaries	0%
Total for all fisheries	100%

## Project Design and Methodology

- Training workshops for government and industry representatives will be held to develop practical biosecurity measures that are agreed to by both government and industry.
- The subtractive DNA hybridisation method will be used to clone viral gene fragments from infected abalone tissues. Cloned genes (fragments) will be sequenced and blasted against gene data banks in order to find existing homologous viral genes.
- Primers will be synthesized to perform a walking PCR strategy in order to get appropriate gene length for the development of PCR assays.
- Herpesvirus nucleic acid sequences suitable for use as a diagnostic probe will be developed for in situ hybridisation.
- Infection trials of healthy abalone will be performed within the AAHL biosecure facility to determine which abalone tissues are the most appropriate for sampling during an active surveillance program of, for example, wild broodstock.
- Procedures for the detection and identification of the abalone virus will be incorporated into an Australian and New Zealand Standard Diagnostic Procedure (ANZSDP) and submitted to SCAHLS for review and publication.

## Research Capability and Experience

Mehdi Doroudi

Serge Corbeil: BSc MSc PhD. Eleven years experience in aquatic animal disease research and diagnosis (viral, protozoan and bacterial diseases). Nine years experience in molecular diagnosis (conventional PCR, real-time PCR, gene sequencing) and immunodiagnosis of aquatic animal diseases. Five years experience in vaccine R&D for mammalian and fish diseases.

### Previous FRDC Projects

- Corbeil, S. and Crane, M. St. J. Aquatic Animal Health Subprogram: development of diagnostic procedures for the detection and identification of *Piscirickettsia salmonis*. Fisheries Research and Development Corporation Project Number 2001/624.
- Corbeil, S. and Crane, M. St. J. Aquatic Animal Health Subprogram: Development of molecular diagnostic expertise for the mollusc pathogen *Bonamia* sp. Fisheries Research and Development Corporation Project Number 2003/622.

### Relevant Publications

- Corbeil, S., McColl, K. A. and Crane, M. St. J. (2003) Development of a TaqMan quantitative PCR assay for the identification of *Piscirickettsia salmonis*. *Bull. Eur. Ass. Fish Pathol.* 23: 95-101.
- Corbeil, S., Hyatt, A. D. and Crane, M. St. J. (2005) Characterisation of an emerging rickettsia-like organism in Tasmanian farmed Atlantic salmon *Salmo salar*. *Dis. Aquat. Org.* 64: 37-44.
- Corbeil, S., Arzul, I., Robert, M., Berthe, F. C. J., Besnard-Cochennec, N. and Crane, M. St. J. (2006) Molecular characterisation of an Australian isolate of *Bonamia* isolate from *Ostrea angasi*. *Dis. Aquat. Org.* In press.
- Corbeil, S., Arzul, I., Diggles, B., Heasman, M., Chollet, B., Berthe, F. C. J. and Crane, M. St. J. (2006) Development of a TaqMan PCR assay for the detection of *Bonamia* species. *Dis. Aquat. Org.* In press.

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**Please forward the Preliminary Research Proposal in a MS-Word format via e-mail to:**

**Ms Joanne Slater**

**Coordinator, Aquatic Animal Health Subprogram**

**E-MAIL: [joanne.slater@csiro.au](mailto:joanne.slater@csiro.au)**

**No later than cob on Friday 23 June 2006.**

## **DOCUMENT 55**

**EXEMPT IN FULL – s 47B(a)**

**From:** Mehdi.Doroudi@dpi.vic.gov.au  
**Sent:** Tuesday, 13 June 2006 12:47 PM  
**To:** Crane, Mark (LI, Geelong)  
**Cc:** malcolm.lancaster@dpi.vic.gov.au; Corbeil, Serge (LI, Geelong)  
**Subject:** RE: URGENT - Comments on the pre-proposal  
**Attachments:** FRDC Abalone herpesvirus pre-proposal 060613.doc

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Thanks Mark. We will further discuss it with the industry today. Lets hope there will be a significant level of cash contribution from both industry and DPI. I will keep you guys posted with the outcomes. Regards

Mehdi Doroudi - DVM, PhD  
Research Director  
Marine & Freshwater Systems - PIRVic  
Department of Primary Industries

Ph: +613 5258 0272 Fax: +613 5258 0270  
Mobile: +61400 845 406

Email: mehdi.doroudi@dpi.vic.gov.au  
2A Bellarine Hwy, Queenscliff VIC 3225 Australia  
PO Box 114, Queenscliff VIC 3225 Australia

Mark.Crane@csiro.au

13/06/2006 12:35 PM

To: Mehdi.Doroudi@dpi.vic.gov.au  
cc: malcolm.lancaster@dpi.vic.gov.au, Serge.Corbeil@csiro.au  
Subject: RE: URGENT - Comments on the pre-proposal

Mehdi,

I have accepted most of your changes - corrected some minor grammatical/typographical errors - and reduced so me blurb so that it was less than 3 pages. I have also added some extra salary for 1.0 FTE at Attwood for one year, based on Malcolm's email.

Hope this is the fimal version :-)

Cheers

Mark

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
**CSIRO Livestock Industries**  
**Private Bag 24**  
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**email: [mark.crane@csiro.au](mailto:mark.crane@csiro.au)**

# AQUATIC ANIMAL HEALTH SUBPROGRAM

## 2007-08 Preliminary Research Proposal



### Project Title

Aquatic Animal Health Subprogram: Development of management strategies for herpes-like virus infection of abalone

### FRDC Strategic challenge identification

Natural Resources Sustainability

### Co-Principal Investigators Contact Details

Title: Dr Mehdi Doroudi    Organisation: Vic DPI (PIRVic)    Mailing Address: PO Box 114, Queenscliff VIC 3225  
 Phone No: 03 5258 0272    Fax No: 03 5258 0270    Email: mehdi.doroudi@dpi.vic.gov.au

Title: Dr Serge Corbeil    Organisation: CSIRO LI    Mailing Address: Private Bag 24, Geelong, Vic, 3220  
 Phone No: 03 5227 5254    Fax No: 03 5227 5555    Email: serge.corbeil@csiro.au

### Commencement and completion date

Commencement date: 01-02-2007

Completion date: 31-01-2009

### Preliminary Budget

FRDC Contribution	2006-07	2007-08	2008-09	2009-10	TOTAL \$\$s
Salaries and on costs (PIRVic – MAFS)	30 721	30 721			61 443
Salaries and on costs (AAHL)	46 149	95 990	51 522		193 661
Salaries and on costs (PIRVic – Animal Health)		40 400	41 600		82 000
Travel	10 000	12 000	2 000		24 000
Operating	24 851	40 000	16 478		71 339
Capital (equipment)	0	0	0		
<b>FRDC TOTAL</b>	<b>111 721</b>	<b>219 121</b>	<b>111 600</b>		<b>432 443</b>
Research Organisation Contribution (PIRVic – MAFS)	11 828	11 828			23 656
Research Organisation contribution (AAHL)	35 893	75 857	40 093		151 842
Research Organisation contribution (PIRVic – Animal Health)		10 000	11 000		21 000
Total of Industry & Other Funding	x	x	x		x
<b>GRAND TOTAL in \$\$s</b>					<b>628941</b>

### Need

Recent disease outbreaks in Victorian abalone farms associated with high mortality rates have been shown to be caused by a previously unknown herpes-like virus. The abalone species affected by the outbreak are *Haliotis laevisgata*, *H. rubra*, and *H. laevisgata* x *H. rubra* hybrid. The disease has now been detected in wild abalone populations in the vicinity of one of the affected farms.

Histopathology performed on moribund animals indicated a ganglioneuritis – infiltration of haemocytes in multiple ganglia and nerves (cerebral ganglion affected, also other ganglia and nerves). Examination by electron microscopy revealed the presence of a herpes-like virus in the pleuropedal ganglion. Preliminary transmission studies, carried out within AAHL's high biosecurity facility, indicated that this emerging virus is highly pathogenic and can be transmitted to healthy abalone through the water column. In addition, the viral suspension remains infectious after undergoing a dilution up to 1 in 100. So far, the virus has not grown in tissue culture.

Attempts by the farm managers to eliminate the disease from the abalone farms have failed partly due to the lack of detection methods specific to the aetiological agent which would allow early diagnosis of infected animals. If appropriate management strategies including rapid diagnostic techniques are not developed this emerging virus may have the potential

to have a significant adverse impact on the Australian abalone industry.

In order to protect Australia's valuable abalone industry, there is a need to develop specific management strategies which incorporate disease monitoring, detection, response and control measures both in farm and wild stocks. These include the implementation of routine health management procedures and a system for the exclusion of virus. Health management strategies for both farmed and wild abalone sectors and processing plants could be developed to improve productivity of farmed sector and to protect the health status of wild populations of abalone. The application of effective detection and exclusion or control methods incorporates into a workable biosecurity plan to minimise the impacts of this virus.

Access to diagnostic tests that are rapid, reliable and sensitive is of fundamental importance for effective control/management of disease outbreaks. In addition to surveillance tools such as PCR, better procedures/reagents for disease diagnosis are required. Presence of histological lesions provides a presumptive diagnosis. The development of in situ hybridisation probe(s) for the localisation of abalone herpesvirus within histological lesions will provide a means for definitive diagnosis to be made. The development of molecular tools and reagents will allow researchers and industry to rapidly and specifically detect and locate the virus in abalone tissues therefore providing a vital means for diagnosis and facilitating a better understanding of the epidemiology of this disease, leading to more efficient management of disease outbreaks.

### Objectives

1. To improve physical, chemical and biological measures of biosecurity for abalone farms to prevent the introduction and spread of virus.
2. To develop a "code of practice" for commercial divers to avoid the introduction and further spread of virus in wild populations of abalone.
3. To develop a "practical biosecurity program" for abalone processing plants.
4. To identify nucleic acid sequences of the emerging abalone herpesvirus (via PCR-based gene amplification and sequencing) necessary for the development of diagnostic tools (e.g. conventional PCR, TaqMan PCR, ISH probe).
5. To develop and validate PCR assays for the detection of the abalone virus.
6. To develop an in situ hybridisation assay specific for the abalone herpesvirus
7. To document an Australian and New Zealand Standard Diagnostic Procedure and submit for external review.

### Industry and Management consultation

The following stakeholders support the development of management strategies for herpes-like virus infection of abalone including the need to develop rapid diagnostic techniques:

The Victorian abalone Growers Association, VicFRAB, National Aquaculture Council, Seafood Industry Victoria, Abalone and AAH Sub-programs of FRDC and Victorian Department of Primary Industries including both Fisheries and Biosecurity Victoria.

### Direct benefits and beneficiaries

1. All sectors of the abalone wild-capture and aquaculture industries (farmers, divers, processors) will directly benefit from increased biosecurity developed through the implementation of this project.
2. The diagnostic tests will be available to detect and identify the emerging virus in abalone (*Haliotis* spp.), and other potential host mollusc species.
3. Export certification services will be available to industries that wish to develop export markets as well as translocate farmed stock between regions without transmitting disease. Australia will be better prepared to negotiate with international trading partners on issues concerned with the importation of disease free molluscs from Australian sources.
4. Mollusc aquaculture industries and State agencies will be able to develop health surveillance programs in collaboration with AAHL Fish Diseases Laboratory. Should virus be detected during health surveys, industries and government agencies will be able to make informed decisions with regards to translocation of live abalone, eradication or control programs.

### Estimated Flow of Benefits

Fisheries and aquaculture managed by States/Territories	
NSW	4.5%
Qld	1%
SA	15%
Tas	25%
Vic	50%
WA	4.5%
AFMA managed fisheries	0%
Other beneficiaries	0%
Total for all fisheries	100%

## Project Design and Methodology

- All available and relevant information on the current and best management practices in biosecurity for intensive mollusc culture will be reviewed by conducting a literature search and contacting experts in specific fields.
- All information will be used to conduct expertise based risk assessments for farming systems, commercial fishery and processing plants.
- All available information and the outcome of the risk assessments will be used to prepare draft management strategies.
- Training workshops (or disease simulation exercise) for government and industry representatives will be held to effectively communicate the practical management strategies that will be developed through the above process.
- The subtractive DNA hybridisation method will be used to clone viral gene fragments from infected abalone tissues. Cloned genes (fragments) will be sequenced and blasted against gene data banks in order to find existing homologous viral genes.
- Primers will be synthesized to perform a walking PCR strategy in order to get appropriate gene length for the development of PCR assays.
- Herpesvirus nucleic acid sequences suitable for use as a diagnostic probe will be developed for in situ hybridisation.
- Infection trials of healthy abalone will be performed within the AAHL biosecure facility to determine which abalone tissues are the most appropriate for sampling during an active surveillance program of, for example, wild broodstock.
- Procedures for the detection and identification of the abalone virus will be incorporated into an Australian and New Zealand Standard Diagnostic Procedure (ANZSDP) and submitted to SCAHLS for review and publication.

## Research Capability and Experience

**Mehdi Doroudi** DVM, PhD. Mehdi has over 17 years experience in aquatic sciences with an emphasis on the health management of fish and shellfish both in government (research and policy) and commercial operations (aquaculture industry). Mehdi has developed and completed a number of research programs and policies in relation to the management of aquatic animal health including disease prevention, control and eradication of exotic or notifiable diseases, diagnostic investigation, monitoring and surveillance. He is the first author of 31 peer-reviewed, industry and government publications on aquaculture and aquatic animal health.

**Serge Corbeil** BSc MSc PhD. Serge has eleven years experience in aquatic animal disease research and diagnosis (viral, protozoan and bacterial diseases), nine years experience in molecular diagnosis (conventional PCR, real-time PCR, gene sequencing) and immunodiagnosis of aquatic animal diseases and five years experience in vaccine R&D for mammalian and fish diseases.

## Previous FRDC Projects

- Corbeil, S. and Crane, M. St. J. Aquatic Animal Health Subprogram: development of diagnostic procedures for the detection and identification of *Piscirickettsia salmonis*. Fisheries Research and Development Corporation Project Number 2001/624.
- Corbeil, S. and Crane, M. St. J. Aquatic Animal Health Subprogram: Development of molecular diagnostic expertise for the mollusc pathogen *Bonamia* sp. Fisheries Research and Development Corporation Project Number 2003/622.

## Relevant Publications

- Corbeil, S., McColl, K. A. and Crane, M. St. J. (2003) Development of a TaqMan quantitative PCR assay for the identification of *Piscirickettsia salmonis*. *Bull. Eur. Ass. Fish Pathol.* 23: 95-101.
- Corbeil, S., Hyatt, A. D. and Crane, M. St. J. (2005) Characterisation of an emerging rickettsia-like organism in Tasmanian farmed Atlantic salmon *Salmo salar*. *Dis. Aquat. Org.* 64: 37-44.
- Corbeil, S., Arzul, I., Robert, M., Berthe, F. C. J., Besnard-Cochennec, N. and Crane, M. St. J. (2006) Molecular characterisation of an Australian isolate of *Bonamia* isolate from *Ostrea angasi*. *Dis. Aquat. Org.* In press.
- Corbeil, S., Arzul, I., Diggles, B., Heasman, M., Chollet, B., Berthe, F. C. J. and Crane, M. St. J. (2006) Development of a TaqMan PCR assay for the detection of *Bonamia* species. *Dis. Aquat. Org.* In press.

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**From:** Crane, Mark (LI, Geelong)  
**Sent:** Wednesday, 14 June 2006 11:31 AM  
**To:** Ann Fleming  
**Cc:** Mehdi Doroudi (mehdi.doroudi@dpi.vic.gov.au); Malcolm Lancaster (malcolm.lancaster@dpi.vic.gov.au); Corbeil, Serge (LI, Geelong)  
**Subject:** FRDC Revised Preproposal: Abalone herpes-like virus

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Ann,

As discussed by telephone, Victoria and AAHL have revised the preproposal to include development of specific management strategies which incorporate disease monitoring, detection, response and control measures in both farm and wild stocks.

In order to keep costs as low as possible we have limited the development of diagnostic tests to the most essential (AAHL may develop further tests in-house for other specific purposes). Nevertheless, the project remains relatively big with a significant budget. Industry cash contributions will be required to reduce the cost to FRDC and to ensure that this project proposal is funded. Mehdi (representing the lead agency) will be following up on these financial issues.

Serge will be logging the preproposal on FISHNET some time today and I will forward the preproposal to other relevant FRABs.

Regards

Mark



FRDC Abalone  
herpesvirus pre-p..

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
**CSIRO Livestock Industries**  
**Private Bag 24**  
**Geelong Vic 3220**

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**email: [mark.crane@csiro.au](mailto:mark.crane@csiro.au)**

# AQUATIC ANIMAL HEALTH SUBPROGRAM

## 2007-08 Preliminary Research Proposal



### Project Title

Aquatic Animal Health Subprogram: Development of management strategies for herpes-like virus infection of abalone

### FRDC Strategic challenge identification

Natural Resources Sustainability

### Co-Principal Investigators Contact Details

Title: Dr Mehdi Doroudi Organisation: Vic DPI (PIRVic) Mailing Address: PO Box 114, Queenscliff VIC 3225  
 Phone No: 03 5258 0272 Fax No: 03 5258 0270 Email: mehdi.doroudi@dpi.vic.gov.au

Title: Dr Serge Corbeil Organisation: CSIRO LI Mailing Address: Private Bag 24, Geelong, Vic, 3220  
 Phone No: 03 5227 5254 Fax No: 03 5227 5555 Email: serge.corbeil@csiro.au

### Commencement and completion date

Commencement date: 01-02-2007

Completion date: 31-01-2009

### Preliminary Budget

FRDC Contribution	2006-07	2007-08	2008-09	2009-10	TOTAL \$\$s
Salaries and on costs (PIRVic – MAFS)	30 721	30 721			61 443
Salaries and on costs (AAHL)	46 149	95 990	51 522		193 661
Salaries and on costs (PIRVic – Animal Health)		40 400	41 600		82 000
Travel	10 000	12 000	2 000		24 000
Operating	24 851	40 000	16 478		71 339
Capital (equipment)	0	0	0		
<b>FRDC TOTAL</b>	<b>111 721</b>	<b>219 121</b>	<b>111 600</b>		<b>432 443</b>
Research Organisation Contribution (PIRVic – MAFS)	11 828	11 828			23 656
Research Organisation contribution (AAHL)	35 893	75 857	40 093		151 842
Research Organisation contribution (PIRVic – Animal Health)		10 000	11 000		21 000
Total of Industry & Other Funding	x	x	x		x
<b>GRAND TOTAL in \$\$s</b>					<b>628941</b>

### Need

Recent disease outbreaks in Victorian abalone farms associated with high mortality rates have been shown to be caused by a previously unknown herpes-like virus. The abalone species affected by the outbreak are *Haliotis laevigata*, *H. rubra*, and *H. laevigata* x *H. rubra* hybrid. The disease has now been detected in wild abalone populations in the vicinity of one of the affected farms.

Histopathology performed on moribund animals indicated a ganglioneuritis – infiltration of haemocytes in multiple ganglia and nerves (cerebral ganglion affected, also other ganglia and nerves). Examination by electron microscopy revealed the presence of a herpes-like virus in the pleuropedal ganglion. Preliminary transmission studies, carried out within AAHL's high biosecurity facility, indicated that this emerging virus is highly pathogenic and can be transmitted to healthy abalone through the water column. In addition, the viral suspension remains infectious after undergoing a dilution up to 1 in 100. So far, the virus has not grown in tissue culture.

Attempts by the farm managers to eliminate the disease from the abalone farms have failed partly due to the lack of detection methods specific to the aetiological agent which would allow early diagnosis of infected animals. If appropriate management strategies including rapid diagnostic techniques are not developed this emerging virus may have the potential

to have a significant adverse impact on the Australian abalone industry.

In order to protect Australia's valuable abalone industry, there is a need to develop specific management strategies which incorporate disease monitoring, detection, response and control measures in both farm and wild stocks. These include the implementation of routine health management procedures and a system for the exclusion of virus. Health management strategies for both farmed and wild abalone sectors and processing plants could be developed to improve productivity of farmed sector and to protect the health status of wild populations of abalone. The application of effective detection and exclusion or control methods incorporates into a workable biosecurity plan to minimise the impacts of this virus.

Access to diagnostic tests that are rapid, reliable and sensitive is of fundamental importance for effective control/management of disease outbreaks. In addition to surveillance tools such as PCR, better procedures/reagents for disease diagnosis are required. Presence of histological lesions provides a presumptive diagnosis. The development of in situ hybridisation probe(s) for the localisation of abalone herpesvirus within histological lesions will provide a means for definitive diagnosis to be made. The development of molecular tools and reagents will allow researchers and industry to rapidly and specifically detect and locate the virus in abalone tissues therefore providing a vital means for diagnosis and facilitating a better understanding of the epidemiology of this disease, leading to more efficient management of disease outbreaks.

### Objectives

1. To improve physical, chemical and biological measures of biosecurity for abalone farms to prevent the introduction and spread of virus.
2. To develop a "code of practice" for commercial divers to avoid the introduction and further spread of virus in wild populations of abalone.
3. To develop a "practical biosecurity program" for abalone processing plants.
4. To identify nucleic acid sequences of the emerging abalone herpesvirus (via PCR-based gene amplification and sequencing) necessary for the development of diagnostic tools (e.g. conventional PCR, TaqMan PCR, ISH probe).
5. To develop and validate PCR assays for the detection of the abalone virus.
6. To develop an in situ hybridisation assay specific for the abalone herpesvirus
7. To document an Australian and New Zealand Standard Diagnostic Procedure and submit for external review.

### Industry and Management consultation

The following stakeholders support the development of management strategies for herpes-like virus infection of abalone including the need to develop rapid diagnostic techniques:

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  - Corbeil, S., Arzul, I., Robert, M., Berthe, F. C. J., Besnard-Cochennec, N. and Crane, M. St. J. (2006) Molecular characterisation of an Australian isolate of *Bonamia* isolate from *Ostrea angasi*. *Dis. Aquat. Org.* In press.
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**From:** Elliott, Nick (CMAR, Hobart)  
**Sent:** Tuesday, 11 July 2006 11:37 AM  
**To:** Crane, Mark (LI, Geelong)  
**Subject:** Abalone virus

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Hi Mark

I have just returned from a meeting in France and while there I visited the IFREMER laboratories at La Tromblade to see their oyster research. During various discussions the issue of viruses came up and some discussion was held on abalone herpes-like virus. My contact there Pierre Boudry mentioned one of his colleagues (not there at the time) had worked on this issue both in oysters and in abalone in Taiwan. I cannot find my notes right now with his name (may have been Tristan Renault) but I did ask Pierre to pass on your contact details to him – so he may already have contacted you. If not, and if you are not already aware of that group I can follow things up with Pierre.

In your plans with the Vic DPI and industry, has thought been given to the development of a suitable biosecure challenge facility that could be utilised for the breeding program work?

How are the plans going for funding?

Cheers  
Nick

---

**From:** Williams, Nette (LI, Geelong)  
**Sent:** Tuesday, 11 July 2006 3:01 PM  
**To:** Crane, Mark (LI, Geelong)  
**Subject:** RE:

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Only research they were the creatures that we got directly from Portland to infect the naive abs with in the LAF and I thought it was worth a go as they were the freshest samples we would ever get..

-----Original Message-----

**From:** Crane, Mark (LI, Geelong)  
**Sent:** Tuesday, 11 July 2006 2:42 PM  
**To:** Williams, Nette (LI, Geelong)  
**Subject:** RE:

Nette,

I do not have any info on 06-0442. Is this research only rather than a diagnostic sample?

Mark

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
**CSIRO Livestock Industries**  
**Private Bag 24**  
**Geelong Vic 3220**

**International Phone: +61 3 52 275118**  
**International Fax: +61 3 52 275555**  
**email: [mark.crane@csiro.au](mailto:mark.crane@csiro.au)**

-----Original Message-----

**From:** Williams, Nette (LI, Geelong)  
**Sent:** Tuesday, 11 July 2006 14:35  
**To:** Crane, Mark (LI, Geelong)  
**Subject:**

06 0442

Abalone from "known infected abalone from Portland" that I tried to isolate virus form.

Do I need to do a report or is all this covered previously?

Nette

---

**From:** Crane, Mark (LI, Geelong)  
**Sent:** Tuesday, 25 July 2006 3:09 PM  
**To:** Malcolm Lancaster (malcolm.lancaster@dpi.vic.gov.au)  
**Subject:** Abalone herpesvirus preproposal

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Hi Malcolm,

The FRDC Aquatic Animal Health Subprogram rec'd the DPI preproposal. I noted that there is no ISH probe development in it - does this mean that DPI VIC are no longer interested in a probe? As Subprogram Leader I am going to have to give advice to the preproposal Pis, following assessment of the preproposals and Serge is going to be asking for some direction.

Mark

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
**CSIRO Livestock Industries**  
**Private Bag 24**  
**Geelong Vic 3220**

**International Phone: +61 3 52 275118**  
**International Fax: +61 3 52 275555**  
**email: [mark.crane@csiro.au](mailto:mark.crane@csiro.au)**

---

**From:** Crane, Mark (LI, Geelong)  
**Sent:** Wednesday, 9 August 2006 4:51 PM  
**To:** 'Mehdi.Doroudi@dpi.vic.gov.au'  
**Subject:** RE: Abalone Up-date

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Mehdi,

We can always tailor our proposals to a specific budget. Getting a PCR is easy. Getting a PCR that does what you want and demonstrating that is the case is not so easy. I am not sure how Attwood can do that especially since they seem to be under the misapprehension that aquatic herpes-like viruses are like mammalian herpesviruses.

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
**CSIRO Livestock Industries**  
**Private Bag 24**  
**Geelong Vic 3220**

**International Phone: +61 3 52 275118**  
**International Fax: +61 3 52 275555**  
**email: [mark.crane@csiro.au](mailto:mark.crane@csiro.au)**

-----Original Message-----

**From:** Mehdi.Doroudi@dpi.vic.gov.au [<mailto:Mehdi.Doroudi@dpi.vic.gov.au>]  
**Sent:** Wednesday, 9 August 2006 16:46  
**To:** Crane, Mark (LI, Geelong)  
**Subject:** Re: Abalone Up-date

Mark,

I agree with the points that you have made in relation to the evaluation of PCR techniques etc. However, we need to remember that the main reason for industry to ask for a new proposal (from Attwood) was the costs associated with the development of the techniques.

I will discuss your email with the abalone reference group on next Monday but I am sure that they will say the same thing that AAHL proposal is too expensive.

You may want to discuss the possibility of collaborative work between AAHL and Attwood with either Malcolm or Catherine Ainsworth prior to Adelaide meeting. I will probably be in Adelaide.

Regards

Mehdi Doroudi - DVM, PhD  
Research Director  
Marine & Freshwater Systems - PIRVic  
Department of Primary Industries

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Mobile: +61400 845 406

Email: [mehdi.doroudi@dpi.vic.gov.au](mailto:mehdi.doroudi@dpi.vic.gov.au)  
2A Bellarine Hwy, Queenscliff VIC 3225 Australia  
PO Box 114, Queenscliff VIC 3225 Australia

[Mark.Crane@csiro.au](mailto:Mark.Crane@csiro.au)

09/08/2006 03:58 PM

To: [mehdi.doroudi@dpi.vic.gov.au](mailto:mehdi.doroudi@dpi.vic.gov.au)  
cc:  
Subject: Abalone Up-date

Hi Mehdi,

I just thought I would up-date you on the latest concerning FRDC and abalone herpesvirus diagnosis. Because there were two potentially competing applications (Malcolm's and Serge's) the FRDC Aquatic Animal Health Subprogram committee deferred evaluation. It is my impression that this issue will be discussed at the Abalone Workshop in Adelaide and, maybe, some progress can be made there. I explained to the committee as well as I could what had been happening re: DPI VIC/AAHL etc. AAHS recommended revised application(s) to be submitted after the Adelaide Conference.

AAHL is still keen to work on the virus - evaluation of any PCR technique will require experimental infections as a source of infected material. Infectivity trials will need to be undertaken to titrate the virus and come up with some semi-quantitative infective dose. Without this the specificity and sensitivity of the diagnostic methods cannot be assessed.

Crispian (FRDC) recommended coordination through Peter Appleford's group.

What is your opinion on how we can progress this issue sensibly? Malcolm has left it up to me to present the papers at the Adelaide Conference - he has a conflict in his diary and cannot make it. I am not sure how other interested States/industry groups/FRDC will view this. As you know, I am happy/keen to collaborate but there needs to be open communication up front rather than what happened previously.

Mark

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
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**email: [mark.crane@csiro.au](mailto:mark.crane@csiro.au)**

**From:** Young, John (LI, Geelong)  
**Sent:** Friday, 11 August 2006 2:21 PM  
**To:** Corbeil, Serge (LI, Geelong)  
**Subject:** RE: san

<b>Tracking:</b>	<b>Recipient</b>	<b>Recall</b>
	Corbeil, Serge (LI, Geelong)	Failed: 11/08/2006 2:36 PM

Hi Serge,

I have found that there were two lots of abalone in the February trial. SAN 06-00442 were the diseased specimens from Portland and 06-00441 were healthy specimens from Indented Head. They were both used in the transmission trial. As I understand it all of the specimens under SAN 06-00441 were used in the February trial. Of the specimens under SAN 06-00442 Mark has noted on the file that 10x were to be used live in the co-habitation trial, 10x were to be sacrificed and have their ganglia dissected out for use in that trial, 20x were to be frozen intact, 2x were to be sent to EM for examination, 2x were to go to Histology for examination and 6x were to be sacrificed and inoculated onto cell cultures in an attempt to grow the herpes virus.

Therefore the SAN that these samples come from depends on whether the abalone are from Indented Head (06-00441) or from Portland (06-00442). A new number would be reasonable to catalogue specimens created for each separate experiment. That is too late for the experiments already conducted, but it looks like SAN 06-00441 would cover any samples taken from the Indented Head specimens both before and after they were included in the co-habitation trial. I expect the specimens from SAN 06-00442 were just used to deliver the virus and not sampled at any stage. If so the whole experiment could easily be catalogued under SAN 06-00441.

Does that answer your question?

John

-----Original Message-----

**From:** Corbeil, Serge (LI, Geelong)  
**Sent:** Friday, 11 August 2006 12:20  
**To:** Young, John (LI, Geelong)  
**Subject:** RE: san

Hi John,

The samples were from the February cohab transmissibility exp. Therefore I will tell Diane that the san number is 06-00442, can you confirm this?

Serge

-----Original Message-----

**From:** Young, John (LI, Geelong)  
**Sent:** Friday, 11 August 2006 11:13  
**To:** Corbeil, Serge (LI, Geelong)  
**Subject:** RE: san

Hi Serge,

I sure can, but would like more information. I entered into LIMS a job 06-00442 in February. Though I have nothing recorded on what that was for I guess it was the original cohabitation transmissibility and infection work. I also entered jobs 60-1154 to 60-1156 in April to test the transmissibility of the abalone herpes virus through contaminated water. No samples have been recorded in LIMS for these experiments. If these samples are from those experiments I can add them as extra samples for analysis by the histo group. If they are new experiments we should give the whole experiment a new number then add these samples to that.

Regards,

John

-----Original Message-----

**From:** Corbeil, Serge (LI, Geelong)

**Sent:** Friday, 11 August 2006 10:49  
**To:** Young, John (LI, Geelong)  
**Subject:** san

Hi John,

I have 3 pieces of abalone, originating from the recent infection experiments carried out in the LAF, that I have fixed and brought to the histo lab for embedding and sectioning. Can you provide me with a san number for these even though they were not diagnostic samples sent to us?

Thanks,

Serge

---

**From:** Corbeil, Serge (LI, Geelong)  
**Sent:** Wednesday, 16 August 2006 10:13 AM  
**To:** Crane, Mark (LI, Geelong)  
**Subject:** research costing herpes virus DEST 16-08-06.xls  
**Attachments:** research costing herpes virus DEST 16-08-06.xls

**Follow Up Flag:** Follow up  
**Flag Status:** Completed





## PRICING DECISION STATEMENT

CONTRACT:  
 CUSTOMER:  
 PROJECT:

0

Type of Activity <input type="radio"/> Consulting or Technical Service <input type="radio"/> Research - Contract <input checked="" type="radio"/> Research - Collaborative - subject to CN <input type="radio"/> Research - Collaborative - NOT subject to CN
---

<b>COSTS:</b>	Direct Costs (less Costs of Sub-contracts)	26,322	
	Indirect Costs	8,023	
	Cost of Sub-Contracted Activities	-	
	<b>Full-cost Price</b>	<b>34,345</b>	
	Competitive Neutrality	5,220	
	Full-cost Price plus competitive neutrality	39,565	
<b>PRICE:</b>	<b>Cash Contribution</b>	<b>15,400</b>	59% DIRECT COSTS ; 45% TOTAL COSTS
	<b>CSIRO - Livestock Industries In-kind Contributions</b>	<b>18,945</b>	
	Value of Other Considerations (1)		
	<b>Justification (if applicable) (2):</b>		

Prepared by: \_\_\_\_\_ Recommended by: www  
 Project Leader Commercial Manager

Approved by: \_\_\_\_\_ Date: \_\_\_\_\_  
 Delegate

**NOTES:**

- (1) Other forms of Consideration include long term value of licence fees and royalties
- (2) Justifications for prices lower than Full-cost Price include National Interest (specify it) and commercial considerations



**From:** Crane, Mark (LI, Geelong)  
**Sent:** Friday, 18 August 2006 2:46 PM  
**To:** Elliott, Nick (CMAR, Hobart)  
**Cc:** Walker, Peter (LI, Geelong); Corbeil, Serge (LI, Geelong); McColl, Ken (LI, Geelong)  
**Subject:** Further developments re: Abalone herpesvirus

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Nick,

Harry Peters (EO, Western Victorian Abalone Divers Association) and I managed to catch up yesterday. Apparently, with funding from the industry, the association is trying to get three mollusc disease experts (Carolyn Friedman, Mike Hine and Tristan Renault) from overseas to come to Australia for a workshop on the abalone herpesvirus. I informed Harry that we had already been in contact with Tristan and were attempting to set up an initial collaboration and I also offered AAHL to host the workshop. Harry is trying to arrange for the o/s experts to visit the Victorian farms and then follow up the farm visits with laboratory sessions (histopathology, electron microscopy). He is hoping to do this for the week of 16-23 September. It is likely that at least one day would be spent at Attwood but he understood that it would be useful to have the visitors come to AAHL.

He also stated that the industry are more confident in CSIRO delivering the PCR than Attwood (but CSIRO was too expensive) - he seemed to think that there were internal conflicts within DPI VIC - for example, he knows that an initial report on the epidemiology of the disease was prepared in January but was not released until much later - around 6 months later - and suggested that the industry would have responded differently had they known the contents of the report. I also indicated that we could submit a "PCR-only" proposal which would be cheaper than the all-encompassing project. I suggested that he needed to let FRDC know what industry wanted and which organisation should be contracted to undertake the work.

In addition, I had a phone call today from Kevin Ellard (DPIW, Tasmania) - he needed some clarification on some of our results for a report he was writing for the CVO. He said that the department had met with industry earlier this week to discuss the department's management of the situation. Interestingly, the industry asked the department who should undertake the development of a diagnostic PCR and AAHL was recommended.

Kevin and I discussed other basic questions that needed to be answered and he was in favour of a much broader project rather than a PCR-only project. Some basic questions needing to be answered are, of course, stability of the virus in the environment, chemical inactivation, comparison with the Taiwanese virus etc. He seemed to think that we could get direct funding from the industry. Clearly, this issue is, now, seen as an industry-wide problem and not just a Victorian problem - perhaps, contrary to Victoria's belief, Victoria do not need to be the lead agency. Moreover, outside of DPI Victoria, stakeholders seem to be saying that the Victorian response to the virus has, in fact, added to the problem rather than help resolve it.

Perhaps the meeting in Adelaide will be instrumental in clarifying the research needs and the potential sources of funding even more than I originally thought. Perhaps, since the wild-capture industry is much bigger than the aquaculture industry and they are deeply concerned about this virus, they will be more influential and may contribute more to the discussion as well as to the funding debate.

Will Serge receive an evaluation of the preproposal from TASFRAB, at some stage?

Cheers

Mark

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
**CSIRO Livestock Industries**  
**Private Bag 24**  
**Geelong Vic 3220**

**International Phone: +61 3 52 275118**  
**International Fax: +61 3 52 275555**  
**email: [mark.crane@csiro.au](mailto:mark.crane@csiro.au)**

---

**From:** Crane, Mark (LI, Geelong)  
**Sent:** Wednesday, 9 August 2006 3:59 PM  
**To:** Mehdi Doroudi (mehdi.doroudi@dpi.vic.gov.au)  
**Subject:** Abalone Up-date

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Hi Mehdi,

I just thought I would up-date you on the latest concerning FRDC and abalone herpesvirus diagnosis. Because there were two potentially competing applications (Malcolm's and Serge's) the FRDC Aquatic Animal Health Subprogram committee deferred evaluation. It is my impression that this issue will be discussed at the Abalone Workshop in Adelaide and, maybe, some progress can be made there. I explained to the committee as well as I could what had been happening re: DPI VIC/AAHL etc. AAHS recommended revised application(s) to be submitted after the Adelaide Conference.

AAHL is still keen to work on the virus - evaluation of any PCR technique will require experimental infections as a source of infected material. Infectivity trials will need to be undertaken to titrate the virus and come up with some semi-quantitative infective dose. Without this the specificity and sensitivity of the diagnostic methods cannot be assessed.

Crispian (FRDC) recommended coordination through Peter Appleford's group.

What is your opinion on how we can progress this issue sensibly? Malcolm has left it up to me to present the papers at the Adelaide Conference - he has a conflict in his diary and cannot make it. I am not sure how other interested States/industry groups/FRDC will view this. As you know, I am happy/keen to collaborate but there needs to be open communication up front rather than what happened previously.

Mark

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
**CSIRO Livestock Industries**  
**Private Bag 24**  
**Geelong Vic 3220**

**International Phone: +61 3 52 275118**  
**International Fax: +61 3 52 275555**  
**email: [mark.crane@csiro.au](mailto:mark.crane@csiro.au)**

---

**From:** emma.young@dpi.vic.gov.au on behalf of Mehdi.Doroudi@dpi.vic.gov.au  
**Sent:** Friday, 25 August 2006 1:40 PM  
**To:** Crane, Mark (LI, Geelong)  
**Subject:** Visiting Abalone Disease Expert  
**Attachments:** Itinerary - Dr Anna Mouton - Sept 2006.doc

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Mark,

Please find attached a copy of Anna Mouton's Itinerary while she is in Australia. The basis of her visit to Australia is for her to provide DPI with an independent review of the epidemiology study undertaken to date.

On Wednesday, 20 September 2006 both Anna and myself intend to visit AAHL from 12:00-1:00pm. Can you please confirm this time is suitable for yourself.

Regards,

Mehdi Doroudi - DVM, PhD  
Research Director  
Marine & Freshwater Systems - PIRVic  
Department of Primary Industries

Ph: +613 5258 0272 Fax: +613 5258 0270  
Mobile: +61400 845 406

Email: [mehdi.doroudi@dpi.vic.gov.au](mailto:mehdi.doroudi@dpi.vic.gov.au)  
2A Bellarine Hwy, Queenscliff VIC 3225 Australia  
PO Box 114, Queenscliff VIC 3225 Australia

(See attached file: Itinerary - Dr Anna Mouton - Sept 2006.doc)

# Dr Anna Mouton

## ITINERARY

Department of Primary Industries

### SATURDAY, 16 SEPTEMBER 2006

TBA Dr Mouton flight Capetown, South Africa to Melbourne, Australia

### SUNDAY, 17 SEPTEMBER 2006

TBA Dr Mouton arrives at Tullamarine, Melbourne Airport  
Mehdi Doroudi and Hugh Millar to collect Dr Mouton and drive her  
to accommodation.  
Accommodation: Airport Motel, Attwood

### MONDAY, 18 SEPTEMBER 2006

9-10:30am Introductory meeting with Mehdi Doroudi, Peter Appleford, Hugh  
Millar, (Sally Ridge, Andrew Cameron), Attwood

10:30-12:30pm Mehdi Doroudi & Anna Mouton – Drive to Portland to visit CS Farm

5:00pm Drive to Warrnambool. Stay in Warrnambool Monday night

### TUESDAY, 19 SEPTEMBER 2006

8-11:00am Visit to SOM and affected abalone reefs

11-12:00pm Catch up with DPI Staff at Warrnambool (Graeme Hanel, Martin  
Roche)

2:00pm Drive back to Melbourne from Warrnambool  
Accommodation: Airport Motel, Attwood

### WEDNESDAY, 20 SEPTEMBER 2006

9-12:00pm Meeting with WADA team in Geelong

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2-4:00pm Internal DPI workshop –suggested participants, in addition to  
introductory meeting, invite Malcolm Lancaster, Mike Jeffers, Dallas  
D'Silva, Anthony Forster, Harry Gorfine

4-5:00pm Visit to Attwood

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**THURSDAY, 21 SEPTEMBER 2006**

9-10:00am                Discussions with Paul Hardy-Smith

10 - 3:00pm             Technical Abalone Disease Expert Forum – invite National  
colleagues

**FRIDAY, 22 SEPTEMBER 2006**

9-11:00am                Meet with Abalone Reference Group

11-5:00pm                Dr Mouton to work on consultancy report

**SATURDAY, 23 SEPTEMBER 2006**

TBA                        Return Flight to South Africa

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**From:** Crispian Ashby [Crispian@frdc.com.au]  
**Sent:** Sunday, 3 September 2006 5:03 PM  
**To:** Crane, Mark (LI, Geelong)  
**Subject:** FW: Visiting Abalone Disease Expert  
**Attachments:** Anna Mouton - Resume.doc; Itinerary - Dr Anna Mouton - Sept 2006.doc

Mark

As outlined below Anna Mouton is going to be in Australia. We were wondering if she may be appropriate for the epidemiology for the Perkinsus project, PI Geoff Liggins.

Your thoughts?

Regards

Crispian

-----Original Message-----

**From:** [emma.young@dpi.vic.gov.au](mailto:emma.young@dpi.vic.gov.au) [<mailto:emma.young@dpi.vic.gov.au>] On Behalf Of [Mehdi.Doroudi@dpi.vic.gov.au](mailto:Mehdi.Doroudi@dpi.vic.gov.au)  
**Sent:** Friday, 25 August 2006 1:23 PM  
**To:** Crispian Ashby  
**Cc:** Patrick Hone; [Peter.Appleford@dpi.vic.gov.au](mailto:Peter.Appleford@dpi.vic.gov.au); [Hugh.Millar@dpi.vic.gov.au](mailto:Hugh.Millar@dpi.vic.gov.au)  
**Subject:** Visiting Abalone Disease Expert

Crispian,

As requested please find attached a copy of Anna Mouton's brief CV and her Itinerary while she is in Australia.

The basis of this invitation is for her to provide DPI with an independent review of the epidemiology study undertaken to date.

We are going to form a technical scientific forum on Thursday, 21 September in order to invite relevant abalone disease experts from other jurisdictions to share our knowledge and seek their input in the further research and development required to overcome this disease. This exercise will have significant input into the current short-term FRDC project (development of management strategies for herpes virus infection in abalone).

As you know the wild abalone sector is planning to invite overseas experts at the same time. If you need further information on the nature of this invitation and the terms of reference for them you would need to speak to Harry Peeters (ph: 0417 119 577). Harry, as I understand, has been organising this visit. I encouraged Harry Peeters to liaise with us in terms of the maximum achievement from bringing all these experts together, however I personally believe this exercise is unnecessary.

Regards,

Mehdi Doroudi - DVM, PhD  
Research Director  
Marine & Freshwater Systems - PIRVic  
Department of Primary Industries

Ph: +613 5258 0272 Fax: +613 5258 0270  
Mobile: +61400 845 406

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2A Bellarine Hwy, Queenscliff VIC 3225 Australia PO Box 114, Queenscliff  
VIC 3225 Australia

(See attached file: Anna Mouton - Resume.doc)(See attached file:  
Itinerary  
- Dr Anna Mouton - Sept 2006.doc)

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The Fisheries Research and Development Corporation plans, invests in and manages fisheries research and development throughout Australia. It is a federal statutory authority jointly funded by the Australian Government and the fishing industry. If I sent this e-mail message to you in error, please accept my apologies. I would appreciate a return e-mail or telephone call (02 6285 0400) so that I can prevent the error from happening again. I ask that you do not distribute or print any part of a wrongly sent e-mail message, or take any action as a result of knowing its contents, but that you destroy all copies and any attachment(s).  
Your cooperation is appreciated.

## **RESUME: Anna Mouton**

### **A. Personal details**

Title : Dr

Full names : Anna-Louise Mouton

Address : PO Box 967  
Stanford  
7210 South Africa

Contact number : +27 28 341 0391

Nationality : South African

### **B. History**

I obtained a BVSc from the University of Pretoria in 1995 and a BSc from the University of South Africa in 2002. After I qualified as a veterinarian, I initially worked in private practice, both in England and South Africa. From September 1996 to the end of January 2001, I was with the Onderstepoort Veterinary Institute, primarily to do diagnostic bacteriology and was the deputy head of the reference veterinary bacteriology laboratory for the country. My involvement in aquatic diagnostics began at Onderstepoort and I initiated a health management program for the abalone industry. During this time, I spent a month training under Carolyn Friedman at the Bodega Marine Laboratory of the University of California. After Onderstepoort, I was with the Department of Agriculture in the Western Cape, based at their regional veterinary laboratory in Stellenbosch. I was responsible for all aquatic diagnostic services, including investigation of disease and mortalities in fish and shellfish, running health management programs for abalone and koi production units, consulting to the commercial aquaculture sector, and advising government bodies on aquaculture policy. During this time, I developed a vaccine against streptococcosis in trout which is still successfully used by the Mpumalanga trout industry. In addition, I was responsible for diagnostic bacteriology and food hygiene services. Since October 2002, I have been directly employed by the Abalone Farmers' Association of South Africa under an industry government partnership in conjunction with Marine and Coastal Management of the Department of Environmental Affairs and Tourism.

### C. Papers presented

□ **The occurrence of bacterial fish diseases in South Africa**

Fourth Congress of the Aquaculture Association of South Africa  
Stellenbosch, South Africa, 1997

repeated by invitation  
Regional Workshop on Aquaculture  
Lilongwe, Malawi, 1997

□ **Health management and disease surveillance in abalone, *Haliotis midae*, in South Africa**

Fourth International Abalone Symposium  
Cape Town, South Africa, 2000

□ **Undesirable aliens: the importation of fish diseases**

Fifth Congress of the Aquaculture Association of South Africa  
Pretoria, South Africa, 2000

□ **The current status of abalone health management in South Africa**

and

**Histological changes associated with stress in intensively cultured South African abalone, *Haliotis midae***

Fifth International Abalone Symposium  
Qingdao, China, 2003

repeated by invitation  
Tenth Annual Abalone Aquaculture Workshop  
Port Lincoln, Australia, 2003

□ I was invited as guest speaker to the Abalone Health Workshop held in conjunction with the Australasian Aquaculture Conference in 2004. I gave twelve talks on various issues related to abalone health.

□ **Factors affecting the prevalence of gut associated parasites in the South African abalone, *Haliotis midae***

Sixth International Abalone Symposium  
Puerto Varas, Chile, 2006

#### **D. Research interests**

I am currently involved in the projects listed below.

□ **The epidemiology of parasites infecting South African abalone, *Haliotis midae*, in western Cape aquaculture facilities**

principal investigator

□ **A national survey of abalone (*Haliotis midae*) health**

principal investigator

□ **Oxygen and ammonia as limiting factors in abalone (*Haliotis midae*) aquaculture systems: husbandry, handling protocols, energetics and growth**

coresearcher

□ **The characterization of an intracellular bacterium infecting the digestive gland of the South African abalone, *Haliotis midae***

cosupervisor

□ **The characterisation of an intracellular protozoan parasite infecting the digestive gland of abalone, *Haliotis midae***

cosupervisor

#### **E. Present responsibilities**

I am currently responsible for all aspects of the abalone health management program. The program includes routine herd health, diagnostic examinations and disease surveillance. The primary aim of the program is to optimise abalone health for production purposes. Abalone movement within South Africa is not officially regulated, but the Abalone Farmers' Association of South Africa has adopted a voluntary stock movement protocol. I developed this protocol and am responsible for its implementation, including all health examinations of abalone. In addition, I act as advisor to Marine and Coastal Management, the official regulatory body, on importation of live abalone. Other responsibilities include compiling the annual industry survey, which is a survey of the technical aspects of abalone production. Results are presented on a comparative basis, with each participating unit receiving their figures relative to the industry as a whole.

# Dr Anna Mouton

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**SATURDAY, 23 SEPTEMBER 2006**

TBA                        Return Flight to South Africa



---

**From:** Crane, Mark (LI, Geelong)  
**Sent:** Friday, 8 September 2006 3:34 PM  
**To:** 'Mehdi.Doroudi@dpi.vic.gov.au'  
**Subject:** RE: Abalone Virus - Scientific Forum - 21 September 2006

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Hi Mehdi,

Thank you for the invitation which I accept and I look forward to participating in the Scientific Forum.

Cheers

Mark

MARK CRANE Ph.D.  
Project Leader  
AAHL Fish Diseases Laboratory  
Australian Animal Health Laboratory  
CSIRO Livestock Industries  
Private Bag 24  
Geelong Vic 3220

International Phone: +61 3 52 275118  
International Fax: +61 3 52 275555  
email: [mark.crane@csiro.au](mailto:mark.crane@csiro.au)

-----Original Message-----

**From:** [Caroline.McGowan@dpi.vic.gov.au](mailto:Caroline.McGowan@dpi.vic.gov.au) [<mailto:Caroline.McGowan@dpi.vic.gov.au>] On Behalf Of [Mehdi.Doroudi@dpi.vic.gov.au](mailto:Mehdi.Doroudi@dpi.vic.gov.au)  
**Sent:** Friday, 8 September 2006 13:58  
**Subject:** Abalone Virus - Scientific Forum - 21 September 2006  
**Importance:** High

Dear All,

The outcomes of the recent investigation into unusual mortality of abalone at four Victorian aquaculture farms and wild abalone in close proximity to one of the affected farms have confirmed the presence of a virus as the cause of mortality. The virus causes ganglioneuritis in abalone. An epidemiology study has been undertaken to provide us with a better understanding of the nature of this outbreak. The Department of Primary Industries Victoria and Western Abalone Divers Association have invited international experts including Dr Anna Mouton (from South Africa), Dr Carolyn Friedman (from USA), Dr Mike Hine (from New Zealand) and Dr Tristan Renault (from France) to provide us with an independent review of the epidemiology study undertaken to date and further advice.

To maximise the benefits and to share our current knowledge and information with each other, DPI / WADA would like to extend an invitation to you to participate in an Abalone Virus - Scientific Forum on Thursday, 21 September 2006 from 10:00 am to 3:00 pm at DPI, 1 Spring Street, Melbourne. The purpose of the forum is:

- to provide you with an update on the current status of the outbreak and its management response and
- to identify / discuss the current knowledge gaps and research priorities

required to address future biosecure management issues.

There are limited places available for the forum. Please nominate a representative from your organisation to attend. RSVPs are due by COB, Friday 15 September 2006. Please RSVP to Caroline McGowan either by email at [caroline.mcgowan@dpi.vic.gov.au](mailto:caroline.mcgowan@dpi.vic.gov.au) or by phone on 03 5258 0266. Please note that you are responsible for any costs incurred.

An agenda for the forum will be forwarded closer to the date.

I look forward to seeing you on the 21st September.

Regards,

Mehdi Doroudi - DVM, PhD  
Research Director  
Marine & Freshwater Systems - PIRVic  
Department of Primary Industries

Ph: +613 5258 0272 Fax: +613 5258 0270  
Mobile: +61400 845 406

Email: [mehdi.doroudi@dpi.vic.gov.au](mailto:mehdi.doroudi@dpi.vic.gov.au)  
2A Bellarine Hwy, Queenscliff VIC 3225 Australia  
PO Box 114, Queenscliff VIC 3225 Australia

---

**From:** Corbeil, Serge (LI, Geelong)  
**Sent:** Tuesday, 12 September 2006 11:53 AM  
**To:** Hyatt, Alex (LI, Geelong)  
**Cc:** Crane, Mark (LI, Geelong)  
**Subject:** Herpes virus meeting

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Hi Alex,

This coming Monday 18th September AFDL is hosting a meeting to discuss the herpes-like virus outbreak in abalone. Members of the abalone diving association will be present as well as 3 international experts on mollusc diseases (Tristan Renault from Ifremer, Carolyn Friedman from Washington Uni and Mike Hine from NZ). I thought you could be interested in coming along to have a chat with these experts and show few EM photos of our herpes-virus on Power Point. The meeting will be held in the board room at around 11:30 am (exact time is not available because our guest will be driving from Port Fairie). Let me know if you are interested in attending the meeting so that can arrange lunch for the appropriate number of people.

Serge

---

**From:** Malcolm.Lancaster@dpi.vic.gov.au  
**Sent:** Friday, 15 September 2006 8:05 AM  
**To:** Crane, Mark (LI, Geelong); McColl, Ken (LI, Geelong); Corbeil, Serge (LI, Geelong)  
**Cc:** Slater, Joanne (LI, Geelong)  
**Subject:** abalone matters

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Hi guys:

A number of international (and national) abalone experts are visiting Victoria next week, as you are no doubt aware. On Tuesday 19th September there will be a histopathology session here at Attwood, using our facilities to present microscopic images to a crowd. You are all welcome to participate, bring slides etc. The session kicks off at 10:00 am, and will not be restricted to ganglioneuritis, although that is the major issue.

This international group will also be involved in assessing research requirements for abalone ganglioneuritis. After they have made their recommendations then the various Victorian players will have to decide what they want and what resources will be allocated.

Given that the FRDC deadline is not till 1st Nov, do we have to decide today what shape our proposal needs to take?

Regards,  
Malcolm Lancaster

**From:** Caroline.McGowan@dpi.vic.gov.au on behalf of Mehdi.Doroudi@dpi.vic.gov.au  
**Sent:** Friday, 15 September 2006 12:32 PM  
**To:** peter.appleford@dpi.vic.gov.au; hugh.millar@dpi.vic.gov.au; jon.presser@dpi.vic.gov.au;  
sally.ridge@dpi.vic.gov.au; malcolm.lancaster@dpi.vic.gov.au;  
mehdi.doroudi@dpi.vic.gov.au; anthony.forster@dpi.vic.gov.au; dallas.@dpi.vic.gov.au  
d'silva@dpi.vic.gov.au; stewart.mcglashan@dpi.vic.gov.au; mike.jeffers@dpi.vic.gov.au;  
harry.gorfine@dpi.vic.gov.au; paul.hamer@dpi.vic.gov.au; graeme.hanel@dpi.vic.gov.au;

s22

Mark (LI, Geelong);

Crane,

**DELETION**

s22

**Subject:** Agenda for Abalone Virus - Scientific Forum, 21st September 2006  
**Attachments:** Agenda - Abalone Virus - Scientific Forum 21 September 2006.pdf  
**Importance:** High  
**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Good Afternoon

Please find attached a copy of the agenda for the Abalone Virus - Scientific Forum on Thursday, 21st September 2006.

For those of you who have not done so already, could you please advise of your attendance by COB today, 15 September?

We look forward to seeing you at the forum.

Regards

Mehdi Doroudi - DVM, PhD  
Research Director  
Marine & Freshwater Systems - PIRVic  
Department of Primary Industries

Ph: +613 5258 0272 Fax: +613 5258 0270  
Mobile: +61400 845 406

Email: [mehdi.doroudi@dpi.vic.gov.au](mailto:mehdi.doroudi@dpi.vic.gov.au)  
2A Bellarine Hwy, Queenscliff VIC 3225 Australia  
PO Box 114, Queenscliff VIC 3225 Australia

(See attached file: Agenda - Abalone Virus - Scientific Forum 21 September 2006.pdf)

## ABALONE VIRUS - SCIENTIFIC FORUM

### AGENDA

Thursday, 21 September 2006  
Conference Room, Ground Level  
1 Spring Street, Melbourne

- 09.30 Registration**
- 10.00 Welcome, Introduction & Background**  
Chair - Dr Peter Appleford, Executive Director Fisheries Victoria
- 10.30 DPI's Response**  
Dr Hugh Millar, Chief Veterinary Officer, Biosecurity Victoria
- 11.00 Epidemiological aspects**  
Dr Paul Hardy-Smith, Managing Director, Panaquatic Health Solutions Pty Ltd
- 1130 Open Forum with Panel Experts**  
Dr Peter Appleford - Chair  
Dr Carolyn Friedman, United States of America  
Dr Judith Handlinger, Tasmania  
Dr Mike Hine, New Zealand  
Dr Anna Mouton, South Africa  
Dr Tristan Renault, France
- 12.30 Lunch**
- 13.15 Research undertaken to date**  
Dr Mark Crane, Project Leader, AAHL Fish Diseases Laboratory, CSIRO
- 1345 Current research proposals**  
Dr Mehdi Doroudi, Research Director, Marine & Freshwater Systems, PIRVic
- 1415 Future directions**  
Dr Peter Appleford, Executive Director Fisheries Victoria
- 1430 Open Forum with Panel Experts**  
Dr Peter Appleford - Chair  
Dr Carolyn Friedman, United States of America  
Dr Judith Handlinger, Tasmania  
Dr Mike Hine, New Zealand  
Dr Anna Mouton, South Africa
- 15.00 Close**

---

**From:** Crane, Mark (LI, Geelong)  
**Sent:** Tuesday, 19 September 2006 1:32 PM  
**To:** 'Mehdi.Doroudi@dpi.vic.gov.au'  
**Subject:** RE: Agenda for Abalone Virus - Scientific Forum, 21st September 2006

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Hi Caroline,

I did get the copy of the epidemiology report - thanks.

Also - my powerpoint presentation for the scientific forum is too big to send as an attachment - would it be OK to bring it on CD or on a USB memory stick on the day?

Mark

MARK CRANE Ph.D.  
Project Leader  
AAHL Fish Diseases Laboratory  
Australian Animal Health Laboratory  
CSIRO Livestock Industries  
Private Bag 24  
Geelong Vic 3220

International Phone: +61 3 52 275118  
International Fax: +61 3 52 275555  
email: mark.crane@csiro.au

-----Original Message-----

**From:** Caroline.McGowan@dpi.vic.gov.au [mailto:Caroline.McGowan@dpi.vic.gov.au] On Behalf Of Mehdi.Doroudi@dpi.vic.gov.au  
**Sent:** Tuesday, 19 September 2006 10:16  
**To:** Crane, Mark (LI, Geelong)  
**Subject:** RE: Agenda for Abalone Virus - Scientific Forum, 21st September 2006

Wonderful - thanks Mark. Hope you had a great time away from the office.

Did you get the copy of the epidemiology report I sent you by express post?

Kind Regards

Caroline

Mehdi Doroudi - DVM, PhD  
Research Director  
Marine & Freshwater Systems - PIRVic  
Department of Primary Industries

Ph: +613 5258 0272 Fax: +613 5258 0270  
Mobile: +61400 845 406

Email: mehdi.doroudi@dpi.vic.gov.au  
2A Bellarine Hwy, Queenscliff VIC 3225 Australia  
PO Box 114, Queenscliff VIC 3225 Australia

Mark.Crane@csiro.  
au

19/09/2006 10:13  
Scientific Forum, 21st September 2006  
AM

To: Mehdi.Doroudi@dpi.vic.gov.au  
cc:  
Subject: RE: Agenda for Abalone Virus -

Hi Caroline/Mehdi,

As discussed with Mehdi last week - I will be participating. I have only just got back from a few days away and will get my presentation to you asap.

Cheers

Mark

MARK CRANE Ph.D.  
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Australian Animal Health Laboratory  
CSIRO Livestock Industries  
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Geelong Vic 3220

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**From:** Crane, Mark (LI, Geelong)  
**Sent:** Tuesday, 19 September 2006 2:21 PM  
**To:** 'Caroline.McGowan@dpi.vic.gov.au'  
**Subject:** Powerpoint presentation part 1

**Follow Up Flag:** Follow up  
**Flag Status:** Completed




Crane Scientific  
Forum part 1....

**MARK CRANE Ph.D.**  
**Project Leader**  
**AAHL Fish Diseases Laboratory**  
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**Private Bag 24**  
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Abalone Herpesvirus – Results of current research

Mark Crane<sup>2</sup>, Malcolm Lancaster<sup>1</sup> & Serge Corbeil<sup>2</sup> & Ken McColl<sup>2</sup>

<sup>1</sup>PIRVic Attwood, Department of Primary Industries, Attwood Victoria 3049  
<sup>2</sup>CSIRO Livestock Industries, Geelong Victoria 3220

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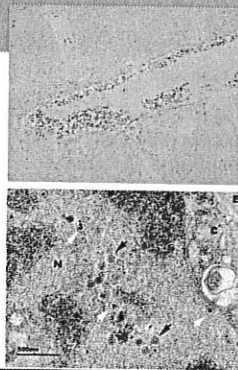
Victoria

**Abalone herpesvirus: Background**

Histopathological examination on moribund animals indicated a ganglioneuritis

Examination by electron microscopy revealed the presence of a herpes-like virus in the pleuropedal ganglion

A similar case was reported in cultured abalone, *Haliotis diversicolor supertexta*, in Taiwan (Jan 2003) (Chang et al., *Dis Aquat Org* 65: 23-27 2005)



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**Immediate Research Need**

As discussed with DPI Victoria: Information on the virus transmissibility was required in order to assist implementation of effective management strategies to control the disease and/or eradicate the virus

Two infection trials were planned and undertaken

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**Herpes-like virus transmission trial 1**

**Objectives:**

- Confirm that the virus isolated from sick abalone can cause disease in healthy abalone
- Confirm Henle-Koch's postulates
- Determine if moribund abalone (virus infected) can transmit the virus to healthy uninfected abalone through the water column

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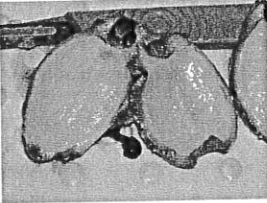
Victoria

**Herpes-like virus transmission trial 1**

**Method:**

Inocula were prepared from 6 frozen (infected) and 9 moribund abalone

Ganglia were dissected, pooled (1 frozen group and 1 fresh group), homogenised, centrifuged, supernatant filtered and kept at 4°C until inoculation into healthy abalone



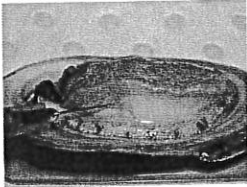
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The State of Victoria

**Herpes-like virus transmission trial 1**

**Treatment Groups (12-15 abalone/group)**

- 1) Co-habitation with sick abalone
- 2) Injected with frozen virus (100uL)
- 3) Injected with fresh virus (100uL)
- 4) Injected with DMEM only (100uL)

Experiments were carried out in 100 L plastic tanks containing 80 L aerated salt water (sea). Water flow rate through the system was 3 Litres/hour



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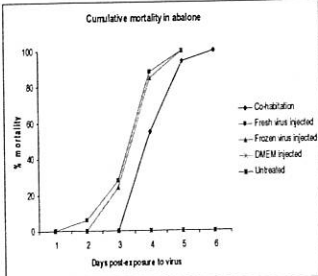
**Herpes-like virus transmission trial 1**

- An extra tank contained healthy naïve abalone as negative (no treatment) control
- Health status and mortality was recorded daily
- Histological examination and electron microscope analyses were performed on some moribund animals

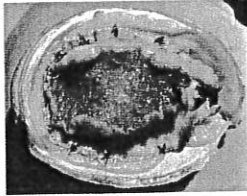
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The State of Victoria

**Herpes-like virus transmission trial 1: Results**


Cumulative mortality in abalone



Days post-exposure to virus	Cohabitation	Fresh virus injected	Frozen virus injected	Unbelled
1	0	0	0	0
2	0	0	0	0
3	25	25	0	0
4	85	85	20	0
5	95	95	80	10
6	100	100	90	80




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### Conclusions

1. The virus is transmitted through the water column from sick abalone to the healthy abalone
2. The virus is highly pathogenic, killing abalone within a few days of infection
3. Injection of the virus in the abalone foot causes disease and mortality
4. The virus remains virulent and pathogenic after being frozen at -80°C

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**From:** Crane, Mark (LI, Geelong)  
**Sent:** Tuesday, 19 September 2006 2:26 PM  
**To:** 'Caroline.McGowan@dpi.vic.gov.au'  
**Subject:** Part 2


**Follow Up Flag:** Follow up  
**Flag Status:** Completed



Crane Scientific  
Forum part 2....


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 **Herpes-like virus transmission trial 2:  
Water dilution and virus transmission**

**Objective:**

To determine if dilution of contaminated water is a sufficient means of disease control

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 **Herpes-like virus transmission trial 2:  
Water dilution and virus transmission**


**Method:**


Sample 200L of farm water (on-going abalone mortalities) and transport to AAHL

Expose healthy abalone to various dilution levels (100%, 10%, 1%, 0.01%, 0.001%) of farm water for a 48 hour period

Record morbidity and mortality


Perform histological examination of moribund animals


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 **Herpes-like virus transmission trial 2:  
Water dilution and virus transmission**

**Results:**


- No disease or mortality in any of the experimental groups
- Positive control group (virus-injected abalone) demonstrated typical clinical signs and mortality

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 **Herpes-like virus transmission trial 2:  
Water dilution and virus transmission**

**Conclusions:**

- 1) There was no (or little) virus present in the water at the time of sampling (virus titres may fluctuate in a farm setting, particularly in flow-through systems)  
and/or
- 2) The transportation of the water to AAHL affected the viability of the virus present in the water  
and/or
- 3) The virus in the water was present at a titre that was too low to cause a productive infection in the abalone

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**Herpes-like virus transmission trial 2:  
Water dilution and virus transmission**


**Control Experiment:**

Co-habitation of healthy abalone with diseased animals (held in basket) until approx. 50% mortality

Expose healthy abalone to various dilution levels (100%, 10%, 1%, 0.01%, 0.001%) of simulated farm water from the co-habitation group for a 48 hour period

Record morbidity and mortality

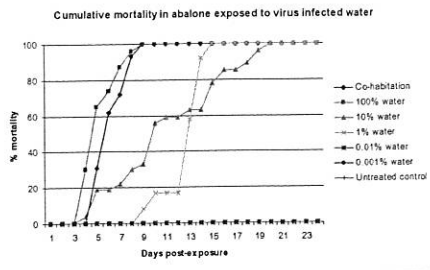
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
**Herpes-like virus transmission trial 2:  
Water dilution and virus transmission**

**Results**

Cumulative mortality in abalone exposed to virus infected water



Days post-exposure	Co-habitation	100% water	10% water	1% water	0.01% water	0.001% water	Untreated control
1	0	0	0	0	0	0	0
3	0	0	0	0	0	0	0
5	65	0	0	0	0	0	0
7	100	0	0	0	0	0	0
9	100	0	0	0	0	0	0
11	100	0	0	0	0	0	0
13	100	0	0	0	0	0	0
15	100	0	0	0	0	0	0
17	100	0	0	0	0	0	0
19	100	0	0	0	0	0	0
21	100	0	0	0	0	0	0
23	100	0	0	0	0	0	0


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**Herpes-like virus transmission trial 2:  
Water dilution and virus transmission**

**Conclusions**

The virus remained infectious to animals even after a 1 in 100 dilution (although only one of the duplicate tanks was affected at this dilution compared with both tanks affected at 100% and 10% levels)


Even though the dilution factor at the outlet of the farm is greater than 1 in 100 and while dilution reduces infectious dose, it remains possible that wild mollusc species could become infected by virus released into the environment


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**Abalone Herpes-like virus: Summary**

**What we know:**


- The disease can be transmitted horizontally
- The virus is transmitted through the water column – direct contact is not required for transmission
- The virus is highly pathogenic, killing abalone within a few days of infection
- Injection of the virus in the abalone foot causes disease and mortality
- The virus remains virulent and pathogenic after being frozen at -80°C
- Cannot be grown in fish cell lines
- Oyster herpesvirus PCR negative

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 **Abalone Herpes-like virus  
Research Needs**

**What we need to know:**  
Geographic range  
Is it the same as the Taiwanese virus?  
Host range  
Distribution and prevalence  
Infectious dose  
Sensitivity to physico-chemical conditions  
Stability in the environment  
Routes of transmission  
Tissue distribution  
Mechanisms of resistance, if any  
Control methods

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Research & Innovation

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**From:** Crane, Mark (LI, Geelong)  
**Sent:** Tuesday, 19 September 2006 3:29 PM  
**To:** 'Caroline.McGowan@dpi.vic.gov.au'  
**Subject:** Part 1

**Follow Up Flag:** Follow up  
**Flag Status:** Completed


**Categories:** NQS

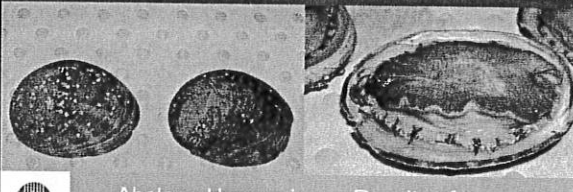



Crane Scientific  
Forum part 11...

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**AAHL Fish Diseases Laboratory**  
**Australian Animal Health Laboratory**  
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 **Abalone Herpesvirus – Results of current research**

Mark Crane<sup>2</sup>, Malcolm Lancaster<sup>1</sup> & Serge Corbeil<sup>2</sup> & Ken McColl<sup>2</sup>


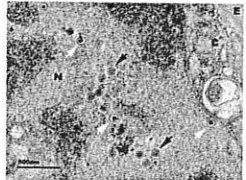
<sup>1</sup>PIRVic Attwood, Department of Primary Industries, Attwood Victoria 3049  
<sup>2</sup>CSIRO Livestock Industries, Geelong Victoria 3220

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 **Abalone herpesvirus: Background**


Histopathological examination on moribund animals indicated a ganglioneuritis

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A similar case was reported in cultured abalone, *Haliotis diversicolor supertexta*, in Taiwan (Jan 2003) (Chang et al., *Dis Aquat Org* 65: 23-27 2005)


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 **Immediate Research Need**

As discussed with DPI Victoria: Information on the virus transmissibility was required in order to assist implementation of effective management strategies to control the disease and/or eradicate the virus


Two infection trials were planned and undertaken

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 **Herpes-like virus transmission trial 1**

**Objectives:**

- Confirm that the virus isolated from sick abalone can cause disease in healthy abalone
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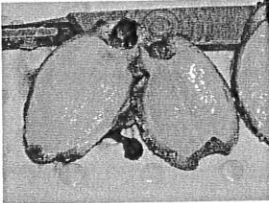
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**Herpes-like virus transmission trial 1**

**Method:**

Inocula were prepared from 6 frozen (infected) and 9 moribund abalone

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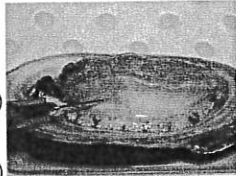
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**Herpes-like virus transmission trial 1**

**Treatment Groups (12-15 abalone/group)**

- 1) Co-habitation with sick abalone
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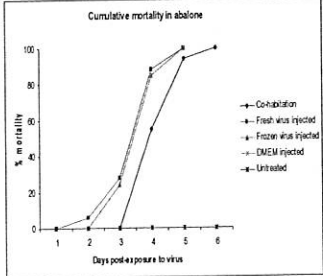
**Herpes-like virus transmission trial 1**

- An extra tank contained healthy naïve abalone as negative (no treatment) control
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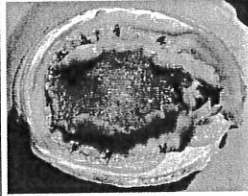
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**Herpes-like virus transmission trial 1: Results**


**Cumulative mortality in abalone**



Days post-exposure to virus	Co-habitation	Fresh virus injected	Frozen virus injected	DMEM injected	Untreated
1	0	0	0	0	0
2	0	0	0	0	0
3	0	0	0	0	0
4	~85	~45	~10	0	0
5	~95	~75	~15	0	0
6	100	85	10	0	0




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### Conclusions

1. The virus is transmitted through the water column from sick abalone to the healthy abalone
2. The virus is highly pathogenic, killing abalone within a few days of infection
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4. The virus remains virulent and pathogenic after being frozen at  $-80^{\circ}\text{C}$

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**From:** Caroline.McGowan@dpi.vic.gov.au  
**Sent:** Tuesday, 19 September 2006 4:54 PM  
**To:** Crane, Mark (LI, Geelong)  
**Subject:** Re: Powerpoint presentation  
**Attachments:** Mark Crane Presentation - Scientific Forum.pdf

**Follow Up Flag:** Follow up  
**Flag Status:** Completed

Mark

I've made a couple of format changes (bullet points etc - hope that is ok). I've saved it as a pdf version should you want to forward it electronically to anyone.

Please let me know if the attached is ok?

Kind Regards

(See attached file: Mark Crane Presentation - Scientific Forum.pdf)

Caroline McGowan  
 EA to Dr Mehdi Doroudi DVM, Phd  
 Research Director  
 Marine and Freshwater Systems  
 PIRVic  
 Department of Primary Industries

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[Mark.Crane@csiro.](mailto:Mark.Crane@csiro.au)  
 au

19/09/2006 03:31  
 PM

**To:** [Caroline.McGowan@dpi.vic.gov.au](mailto:Caroline.McGowan@dpi.vic.gov.au)  
**cc:**  
**Subject:** Powerpoint presentation

Hi Caroline,

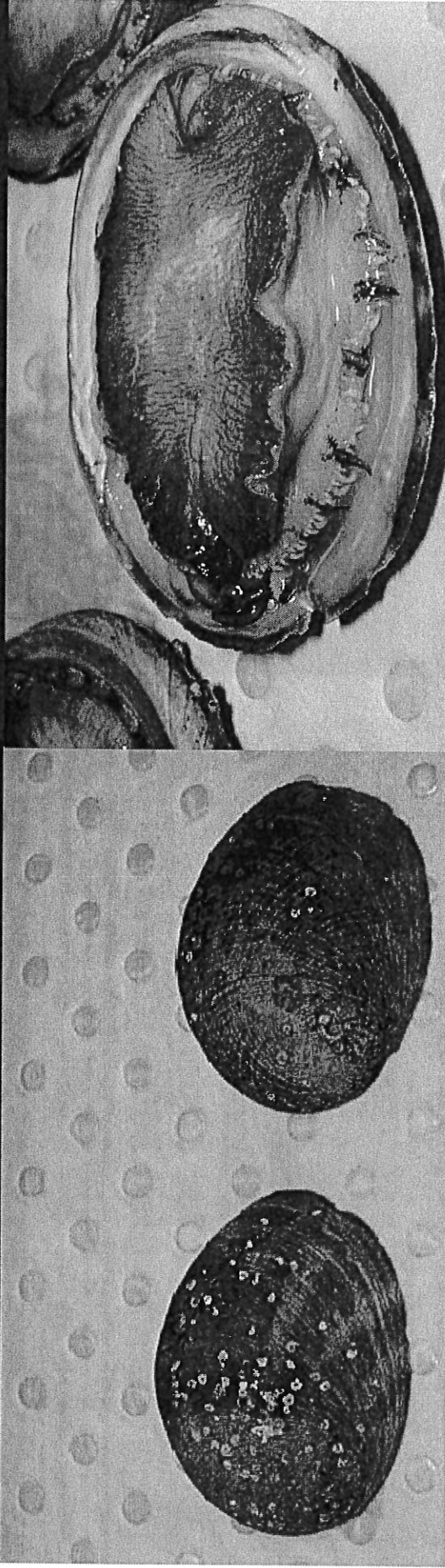
You should have rec'd 2 emails with my presentation in 2 parts; part 1 (labelled ...part 11) is the first half and part 2 is the remainder. If you could let me know that you have rec'd these OK and you can put these together that would be great.

Cheers

Mark

MARK CRANE Ph.D.  
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Australian Animal Health Laboratory  
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## Abalone Herpesvirus – Results of current research

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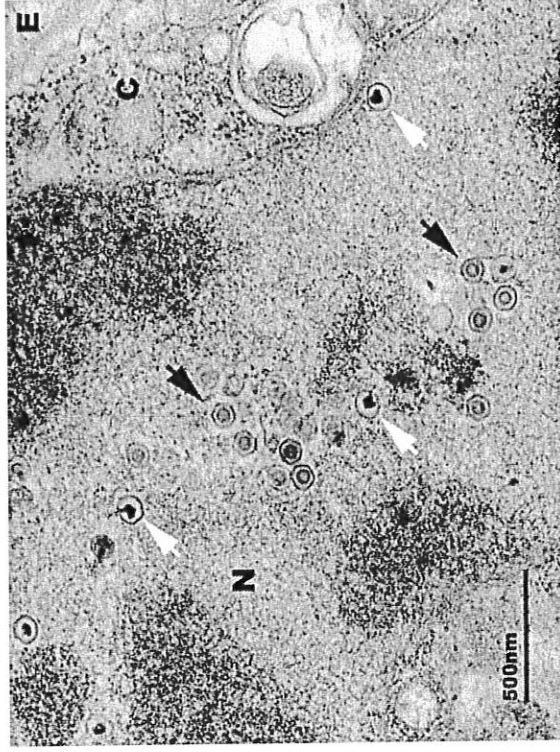
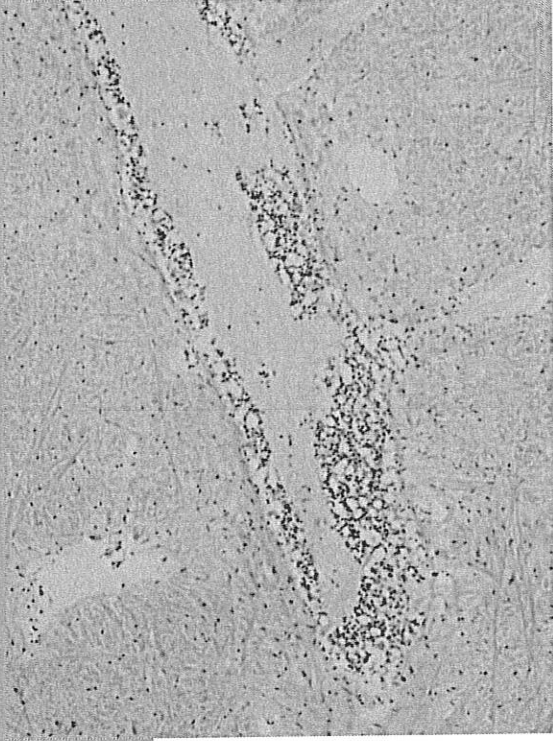
<sup>1</sup>PIRVic Attwood, Department of Primary Industries, Attwood Victoria 3049

<sup>2</sup>CSIRO Livestock Industries, Geelong Victoria 3220



## Abalone herpesvirus: Background

- Histopathological examination on moribund animals indicated a ganglioneuritis
- Examination by electron microscopy revealed the presence of a herpes-like virus in the pleuropedal ganglion
- A similar case was reported in cultured abalone, *Haliotis diversicolor supertexta*, in Taiwan (Jan 2003) (Chang et al., *Dis Aquat Org* 65: 23-27 2005)





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- As discussed with DPI Victoria: Information on the virus transmissibility was required in order to assist implementation of effective management strategies to control the disease and/or eradicate the virus
- Two infection trials were planned and undertaken



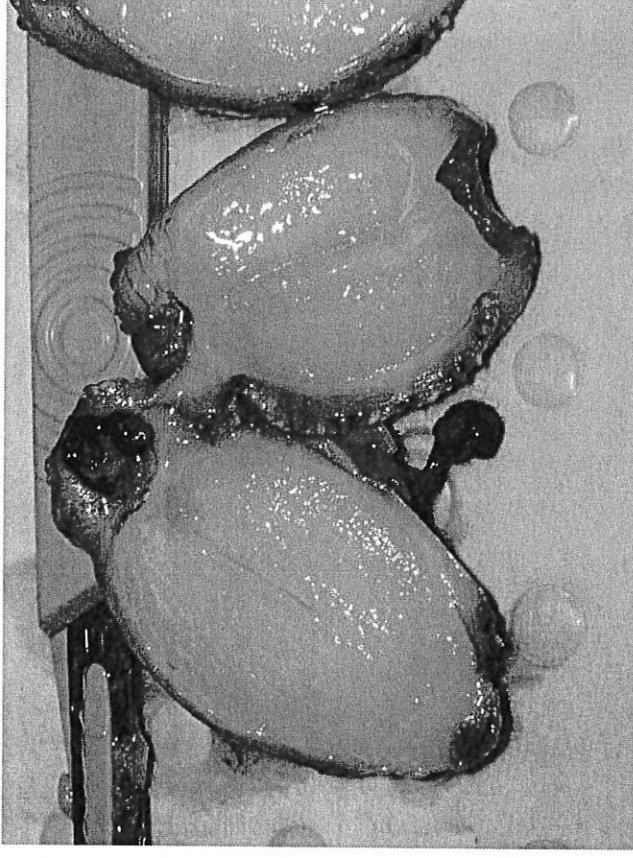
# Herpes-like virus transmission trial 1

## Objectives:

- Confirm that the virus isolated from sick abalone can cause disease in healthy abalone
- Confirm Henle-Koch's postulates
- Determine if moribund abalone (virus infected) can transmit the virus to healthy uninfected abalone through the water column

## Method:

- Inocula were prepared from 6 frozen (infected) and 9 moribund abalone
- Ganglia were dissected, pooled (1 frozen group and 1 fresh group), homogenised, centrifuged, supernatant filtered and kept at 4°C until inoculation into healthy abalone

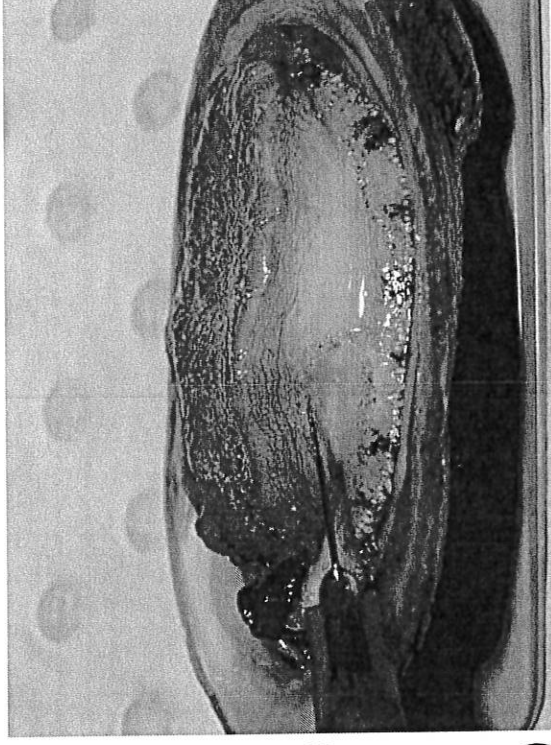




# Herpes-like virus transmission trial 1

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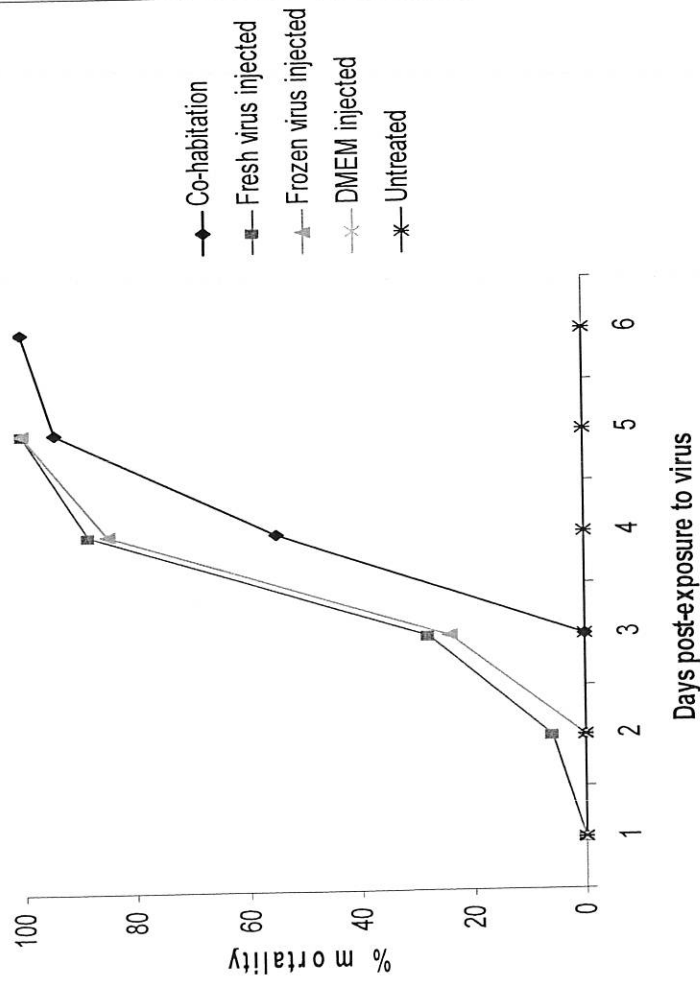
## Herpes-like virus transmission trial 1

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- Histological examination and electron microscope analyses were performed on some moribund animals



# Herpes-like virus transmission trial 1: Results

Cumulative mortality in abalone



## Conclusions

- The virus is transmitted through the water column from sick abalone to the healthy abalone
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- Injection of the virus in the abalone foot causes disease and mortality
- The virus remains virulent and pathogenic after being frozen at -80°C



## Herpes-like virus transmission trial 2: Water dilution and virus transmission

### **Objective:**

To determine if dilution of contaminated water is a sufficient means of disease control



## Herpes-like virus transmission trial 2: Water dilution and virus transmission

### **Method:**

- Sample 200L of farm water (on-going abalone mortalities) and transport to AAHL
- Expose healthy abalone to various dilution levels (100%, 10%, 1%, 0.01%, 0.001%) of farm water for a 48 hour period
- Record morbidity and mortality
- Perform histological examination of moribund animals



## Herpes-like virus transmission trial 2: Water dilution and virus transmission

### Results:

- No disease or mortality in any of the experimental groups
- Positive control group (virus-injected abalone) demonstrated typical clinical signs and mortality



## Herpes-like virus transmission trial 2: Water dilution and virus transmission

### Conclusions:

- 1) There was no (or little) virus present in the water at the time of sampling (virus titres may fluctuate in a farm setting, particularly in flow-through systems)  
and/or
- 2) The transportation of the water to AAHL affected the viability of the virus present in the water  
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## Herpes-like virus transmission trial 2: Water dilution and virus transmission

### Control Experiment:

- Co-habitation of healthy abalone with diseased animals (held in basket) until approx. 50% mortality
- Expose healthy abalone to various dilution levels (100%, 10%, 1%, 0.01%, 0.001%) of simulated farm water from the co-habitation group for a 48 hour period
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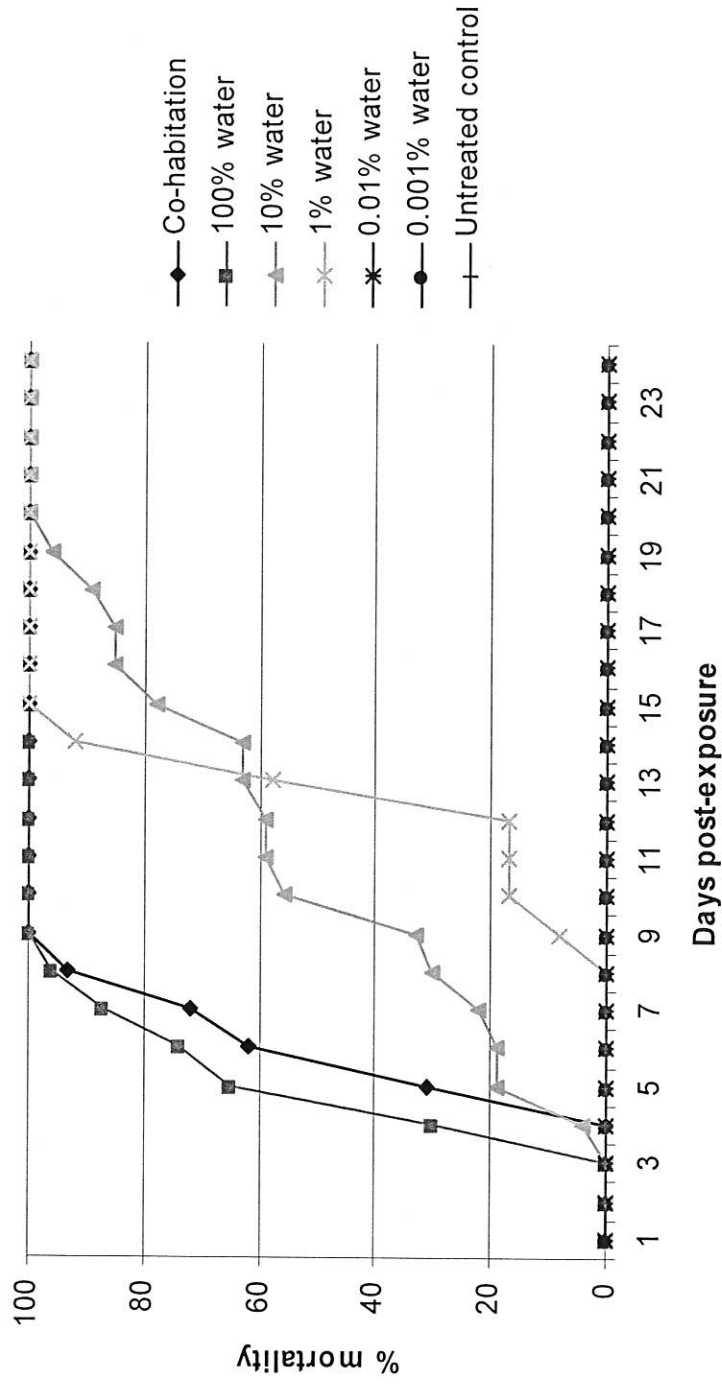




# Herpes-like virus transmission trial 2: Water dilution and virus transmission

## Results

Cumulative mortality in abalone exposed to virus infected water





## Herpes-like virus transmission trial 2: Water dilution and virus transmission

### Conclusions

The virus remained infectious to animals even after a 1 in 100 dilution (although only one of the duplicate tanks was affected at this dilution compared with both tanks affected at 100% and 10% levels)

Even though the dilution factor at the outlet of the farm is greater than 1 in 100 and while dilution reduces infectious dose, it remains possible that wild mollusc species could become infected by virus released into the environment



# Abalone Herpes-like virus: Summary

## What we know:

- The disease can be transmitted horizontally
- The virus is transmitted through the water column – direct contact is not required for transmission
- The virus is highly pathogenic, killing abalone within a few days of infection
- Injection of the virus in the abalone foot causes disease and mortality
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- Cannot be grown in fish cell lines
- Oyster herpes virus PCR negative



# Abalone Herpes-like virus Research Needs

## What we need to know:

- Geographic range
- Is it the same as the Taiwanese virus?
- Host range
- Distribution and prevalence
- Infectious dose
- Sensitivity to physico-chemical conditions
- Stability in the environment
- Routes of transmission
- Tissue distribution
- Mechanisms of resistance, if any
- Control methods

