

Science, Service, Stewardship



Transmission of the pathogen
Renibacterium salmoninarum
under different laboratory-
controlled barging conditions

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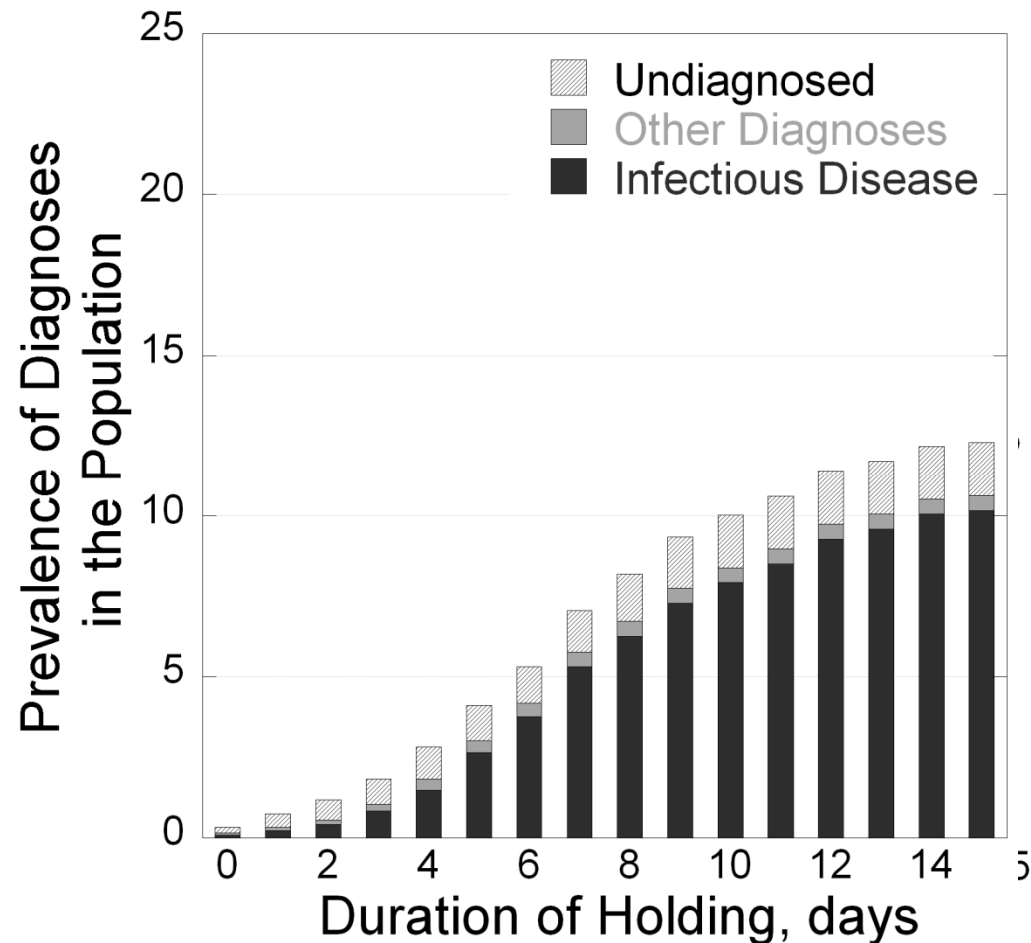


Differential mortality of transported and in-river outmigrants

Annually – delayed mortality **greater** in barged fish than in-river fish

Estuary net pen observations

- Greater barged fish mortality within 6 days of holding
 - Majority of mortalities within 14 days
- >80% of mortalities associated with infectious disease

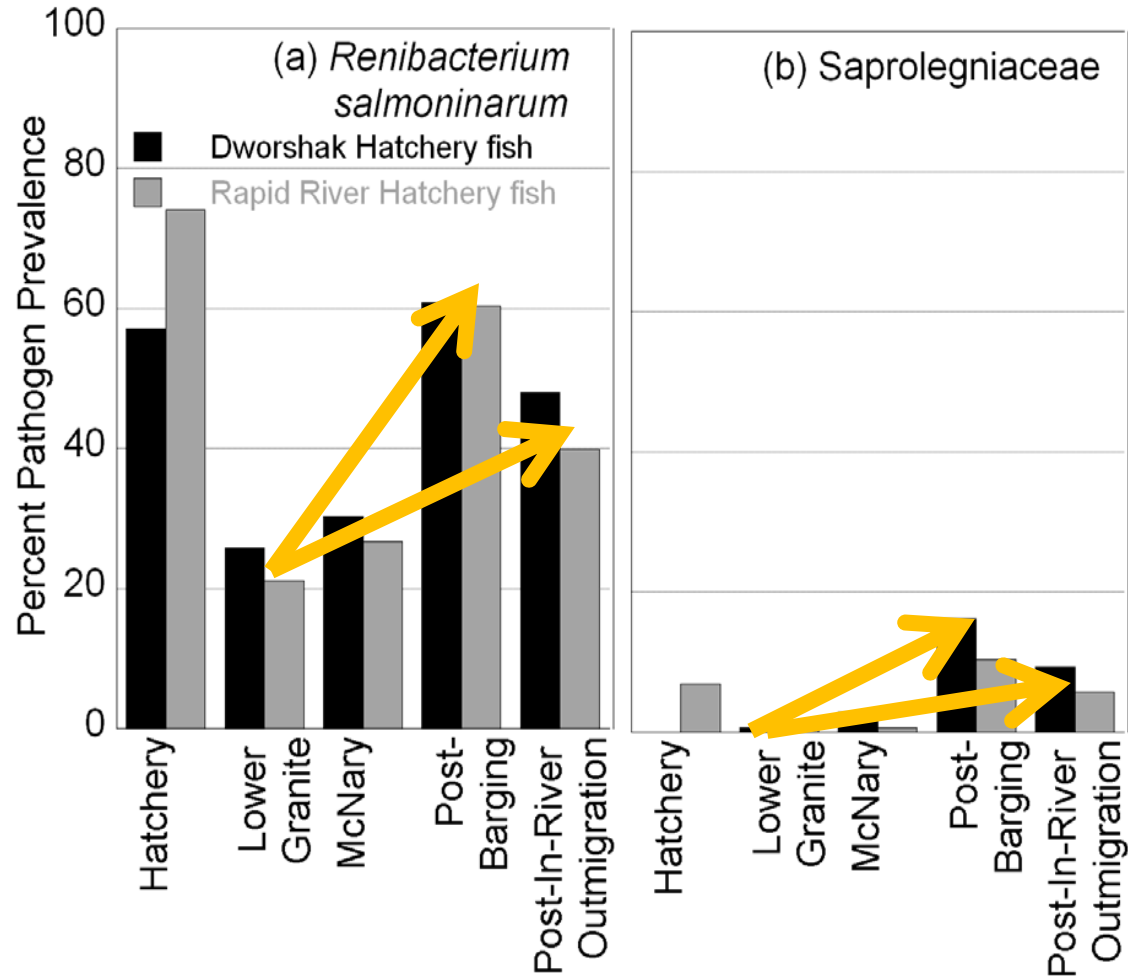




2007 Pathogen Prevalence

Pathogen prevalence increased during outmigration

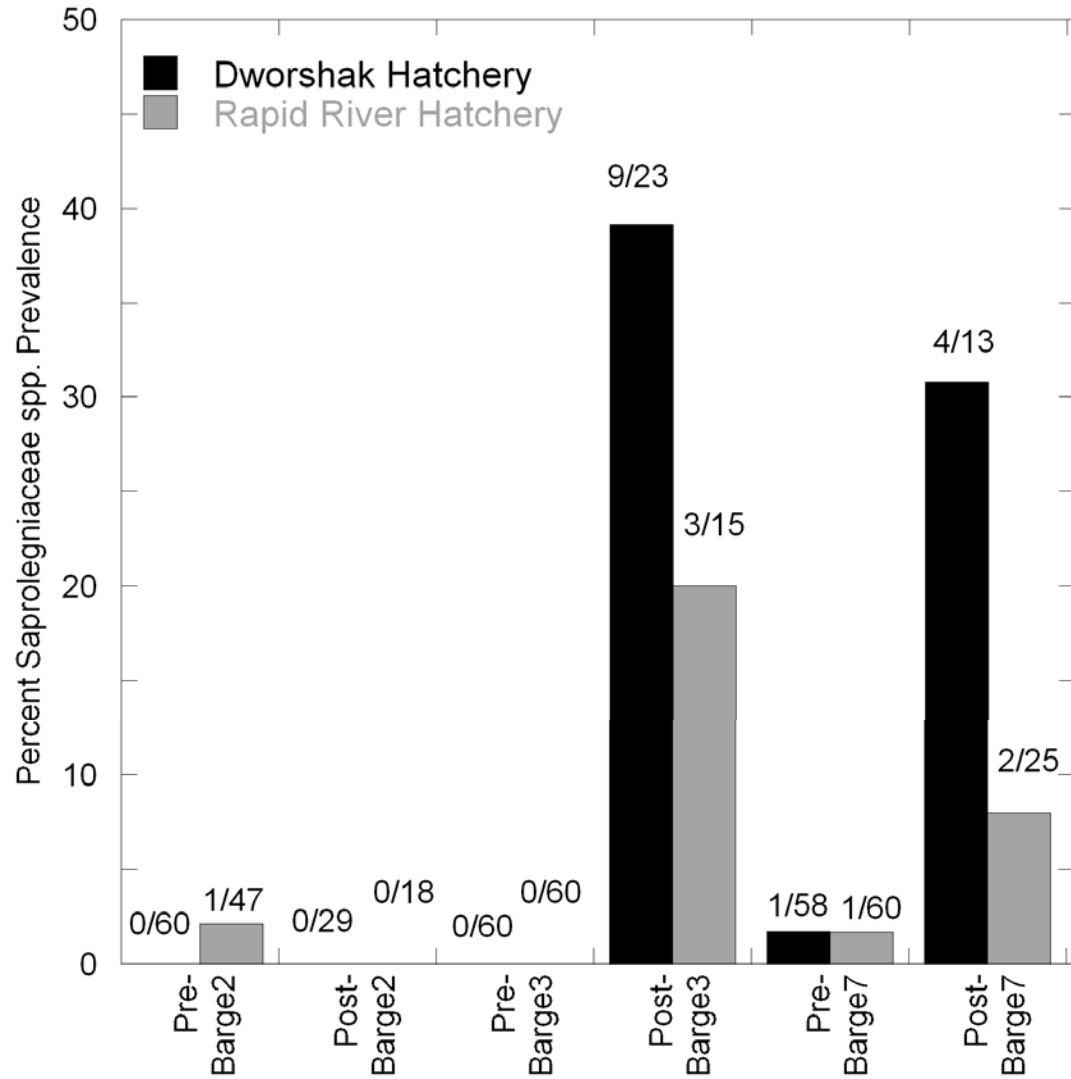
Increased pathogens detected in barged fish relative to in-river





Preliminary Transmission

- Pathogen transmission during barging
 - Individual barges
- Live-box experiments
 - Pascho and Elliot, 1989
 - Elliot and Pascho, 1991





Objective

To identify operating conditions that reduce disease transmission and subsequent delayed mortality in transported Snake River spring/summer Chinook salmon.

- Determine the impacts of holding density and flow exchange rates have on *Renibacterium salmoninarum* transmission under laboratory conditions



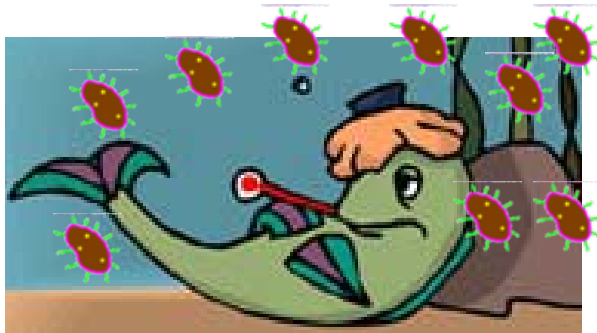
Disease Transmission

Vertical Transmission

Infected parent passes pathogen to their progeny

Horizontal transmission

Infected fish infects non-progeny fish due to shared contact or environment



Donor



Susceptible



Transport Operations from Lower Granite Dam

Transport Timing

Raceway- up to 24 hours

Barges- ~36 hours

Fish Density

Raceway- 0.01 – 0.40 lb/gal

Barges- 0.01 – 0.26 lb/gal

Barge Hold Exchange Rates

1 engine (*<0.17lb fish/gal*) – 1.8 exchanges/hr

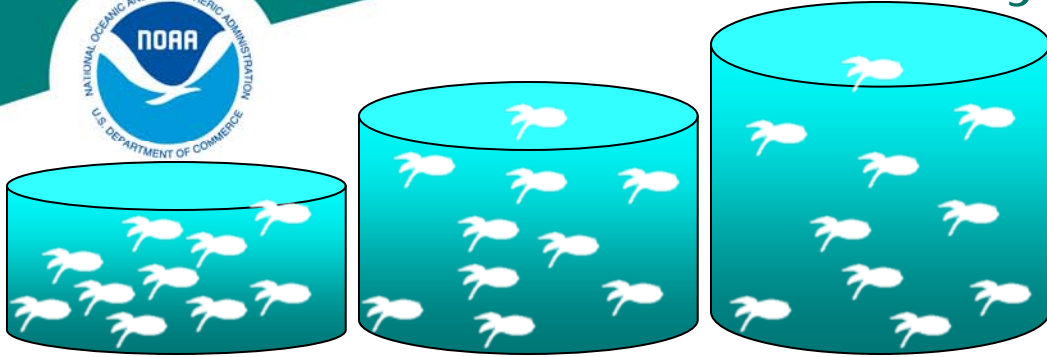
2 engines (*0.17-0.33lb fish/gal*)– 3.7 exchanges/hr

3 engines (*0.33-0.42lb fish/gal*) – 5.5 exchanges/hr





Laboratory disease transmission



Densities

0.05, 0.18, & 0.5 lb/gal

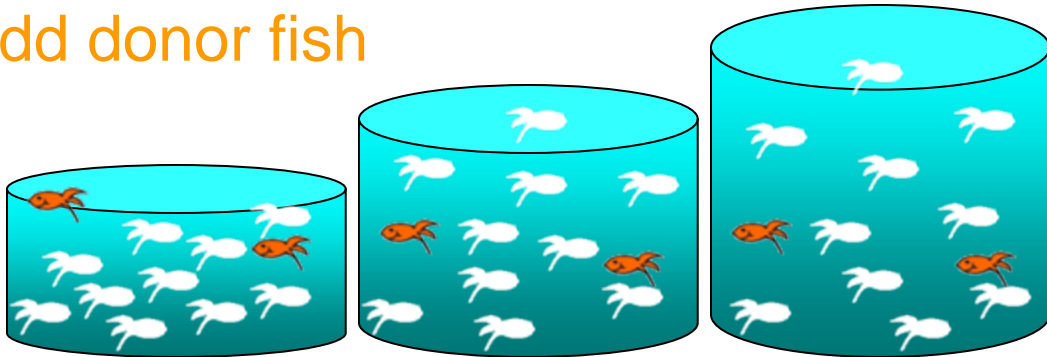
Exchange Rates

2.0 and 5.7 ex/hr

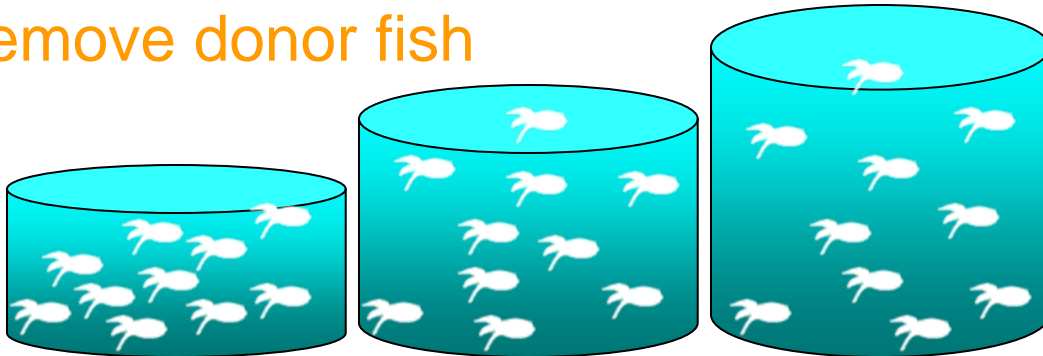
Exposure Period

Raceway + Barge
= 60 hours

add donor fish



remove donor fish



Monitor for pathogen

After 0, 35, and 75 days

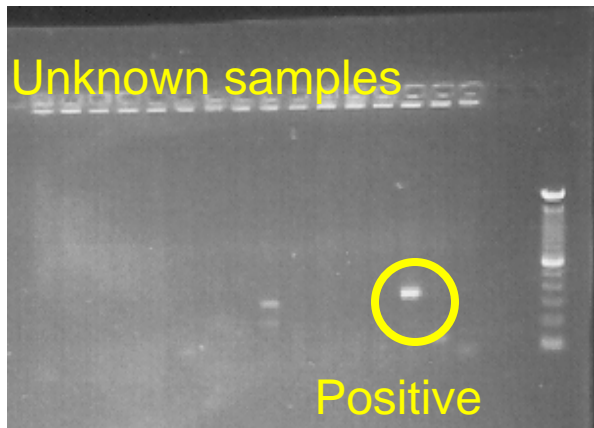


Monitor Pathogen DNA by PCR

Nested PCR

Conventional
presence/absence PCR
Highly sensitive detection
of DNA

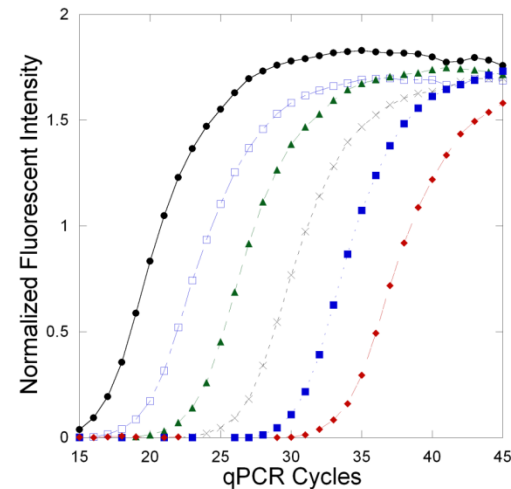
Cannot indicate severity



Quantitative PCR

Real-time tracking of PCR
amplification
Quantitative output indicative
of infection severity

Cannot detect small amounts





Work In Progress

Nested and quantitative PCR sample analysis

Kidney and gills of susceptible fish

0, 35, and 75-days post-exposure

Preliminary experiments

Characterizing donor fish

Pathogen shedding

Donor prevalence

Transmission confirmation



Characterizing Donor Fish: Pathogen Shedding

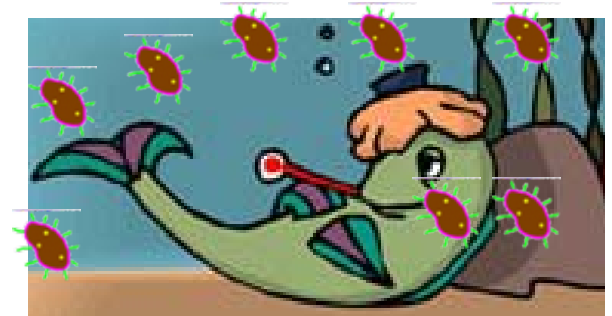
Aims:

Determine concentration of
pathogens shed

Determine the time course of
shedding

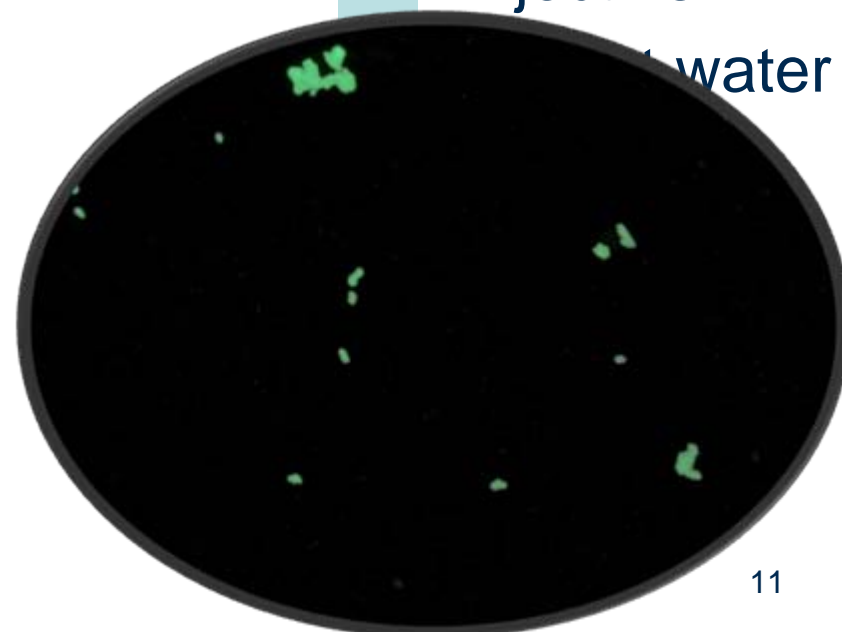
Method^a:

MFAT = membrane filtration
fluorescent antibody technique



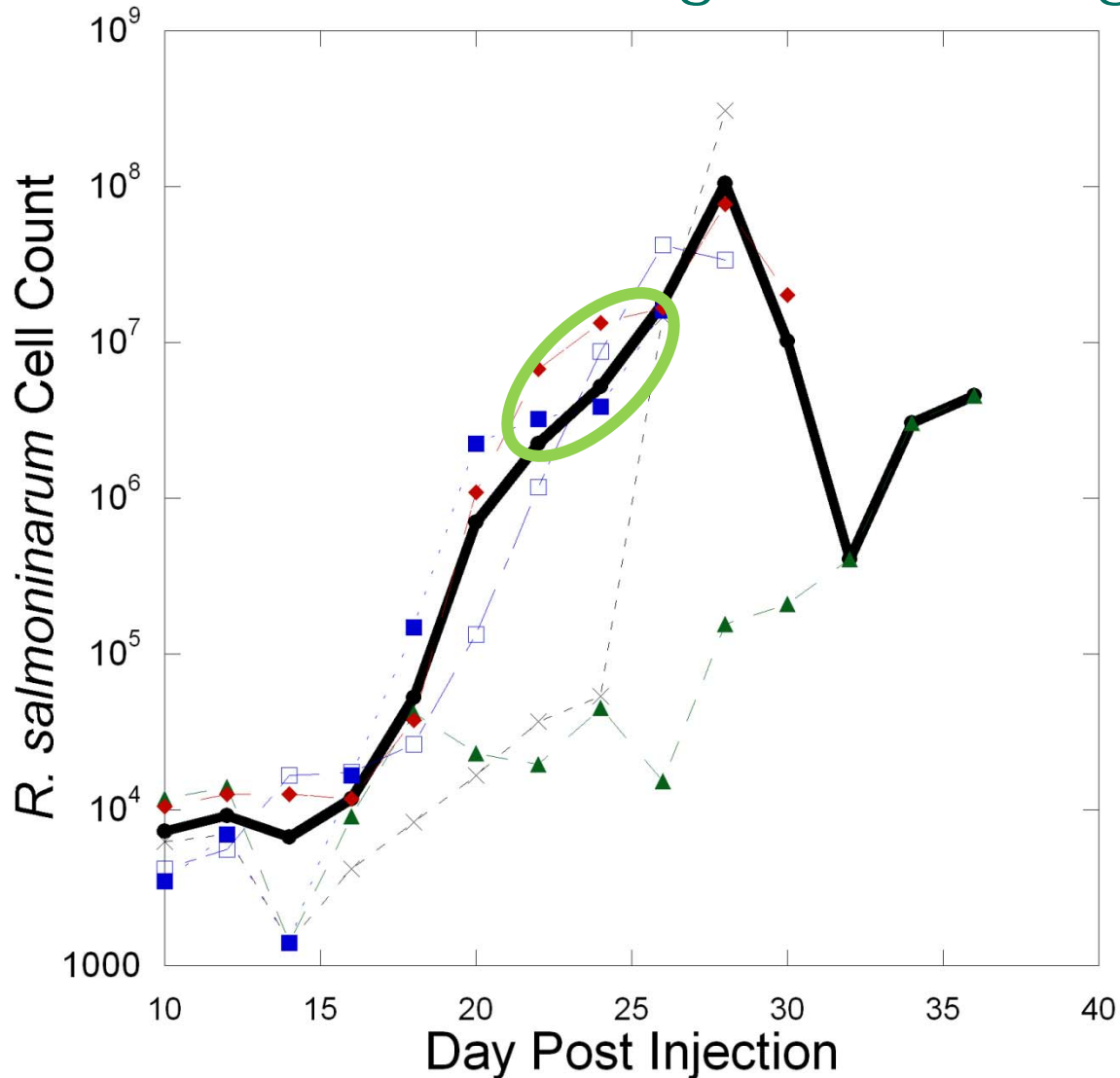
Inject fish

water





Characterizing Donor Fish: Pathogen Shedding





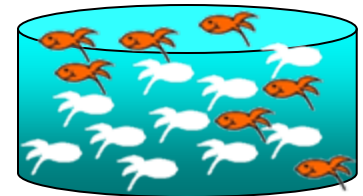
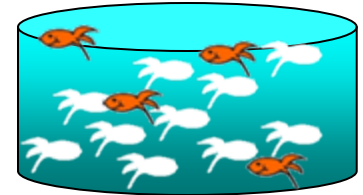
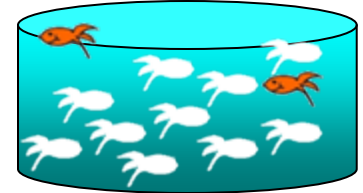
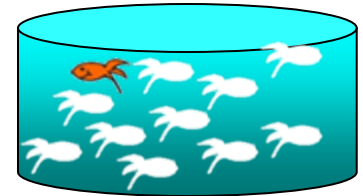
Characterizing Donor Fish: Quantity

Aim:

Evaluate pathogen transmission and variability with multiple donors

Method:

- Prevalence of donor fish (🐟) among susceptible fish was varied for 60 hours (8%, 17%, 27%, 40%)
- PCR detection of *R. salmoninarum* in donors (kidneys) and susceptibles (kidneys + gills)



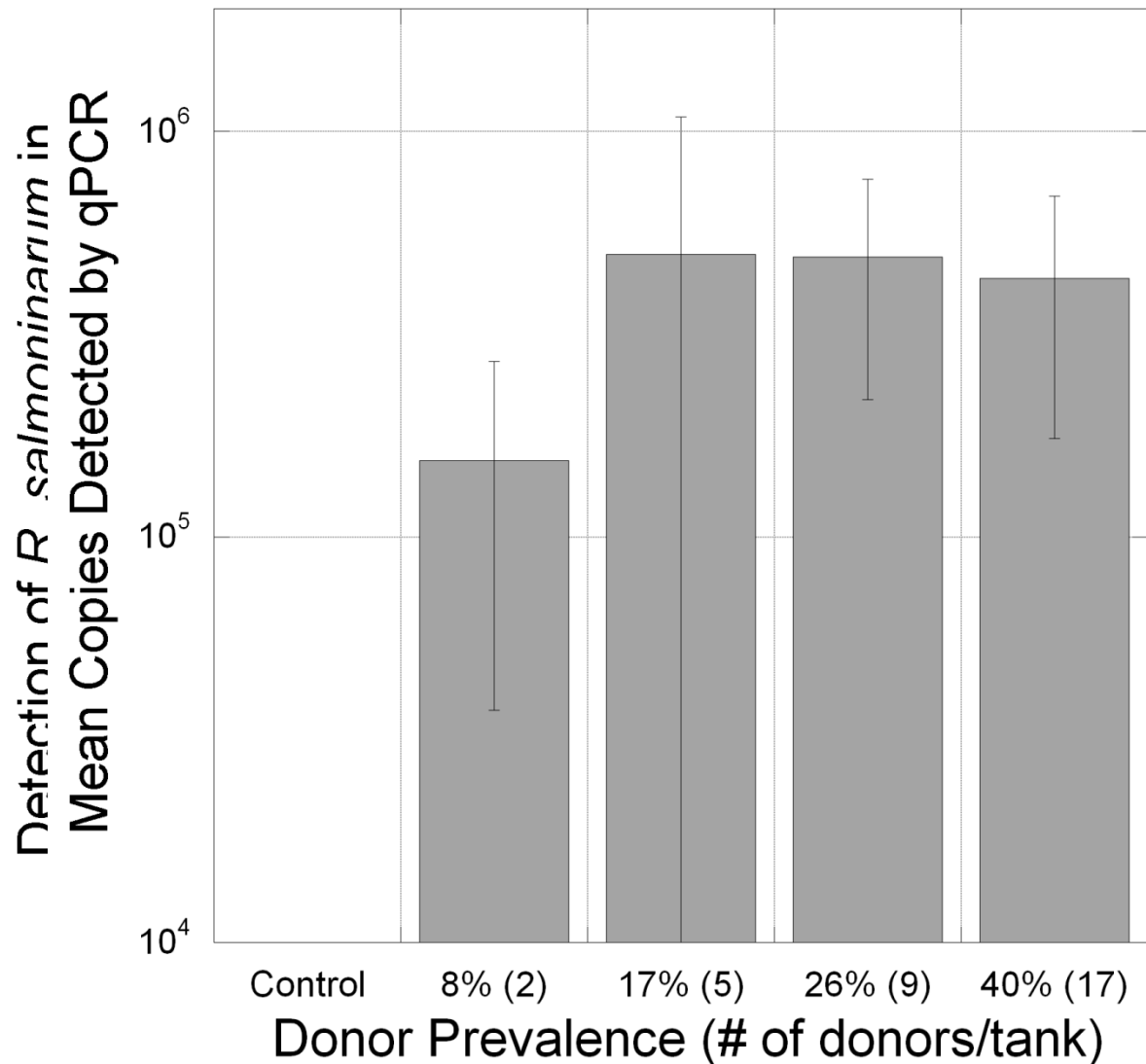


Characterizing Donor Fish: Quantity

Nested PCR is more sensitive

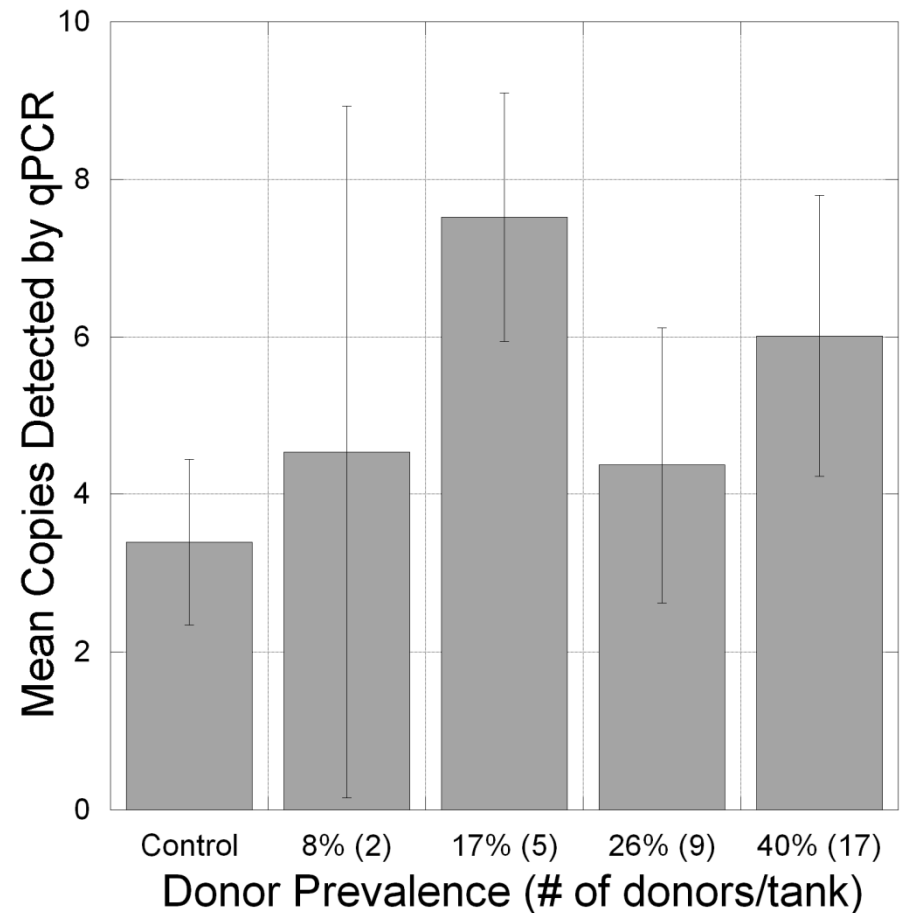
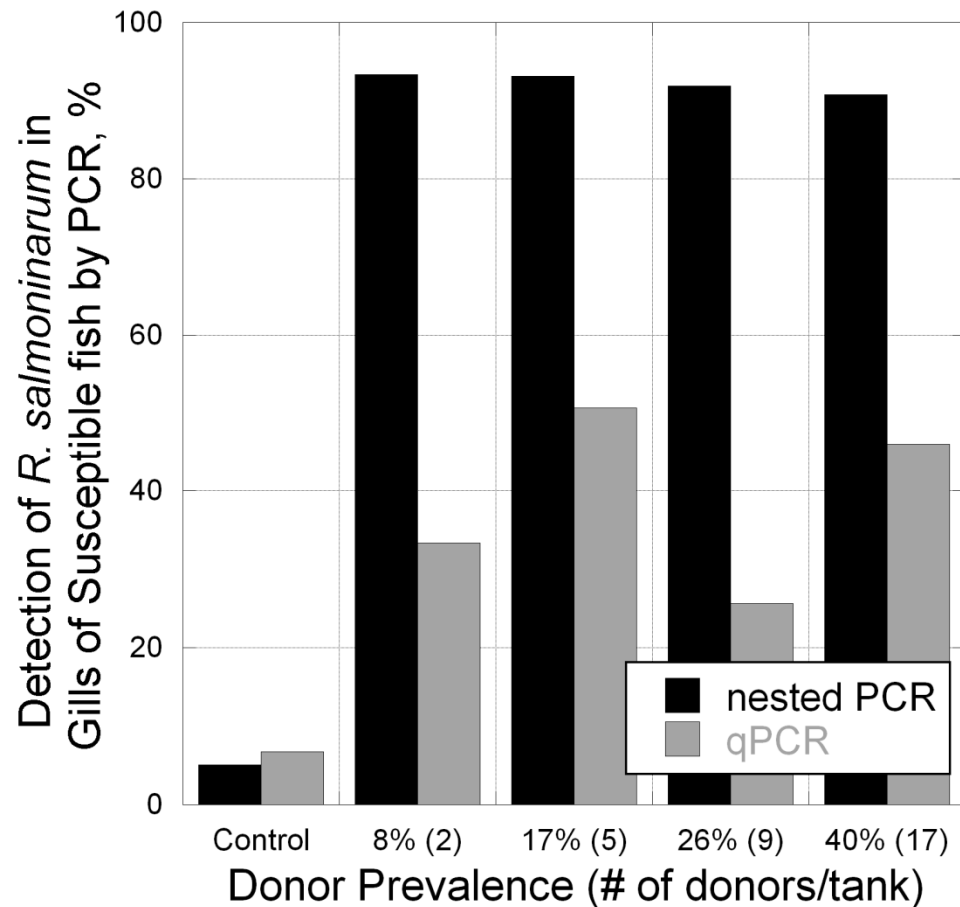
qPCR indicates:

- Control positives are negligible
- Stable mean copy number with >5 donors





Transmission of *R. salmoninarum* to Gills of Susceptible Fish





Preliminary Conclusions

- Lab conditions are close surrogates for field transport conditions
- Up to 615 million cells shed per severely infected fish per day
- Transmission to gills occurred in over 90% of susceptible fish in 60 hours



Acknowledgements

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UC Davis

Don Thompson



Citations

Elliott, D. G. and T. Y. Barila. 1987. Membrane filtration - fluorescent antibody staining procedure for detecting and quantifying *Renibacterium salmoninarum* in coelomic fluid of chinook salmon (*Oncorhynchus tshawytscha*). Can. J. Fish. Aquatic Sci. 44(1): 206-210.

Elliott DG, Pascho RJ. 1991. Juvenile fish transportation: Impact of bacterial kidney disease on survival of spring/summer Chinook salmon stocks, 1989. Annual Report, 1989 (Contract E86880047) prepared by the U.S. Fish and Wildlife Service, Seattle, WA. for the U. S. Army Corps of Engineers, Walla Walla, WA.

Pascho RJ, Elliott DG. 1989. Juvenile fish transportation: Impact of bacterial kidney disease on survival of spring/summer Chinook salmon stocks, 1988. Annual Report, 1988 (Contract E86880047) prepared by the U.S. Fish and Wildlife Service, Seattle, WA. for the U.S. Army Corps of Engineers, Walla Walla, Wa.

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Field Transport Conditions Vs. Lab

	Field Conditions	Lab Conditions
Exposure time, hrs	~60	60
Density, lb/gal	0.01 – 0.4	0.05, 0.18, 0.5
Exchanges, ex/hr	1.8, 3.7, 5.5	2.0, 5.7
Donor prevalence	20-35%	37%