

Text S1. Detailed molecular methods

Tissue samples were screened for the presence of 59 infectious agent taxa (Table S1), using HT-qPCR on the Fluidigm Biomark Dynamic Array™ microfluidics platform (Fluidigm, San Francisco, CA, USA) at the Pacific Biological Station, Nanaimo, British Columbia, Canada. This platform has recently been analytically validated for quantitative infectious agent profiling in salmon tissue (Miller et al. 2016) and applied to multiple studies of Pacific salmon (Di Cicco et al. 2017; Miller et al. 2017; Thakur et al. 2018). Infectious agent taxa were chosen based on knowledge of their presence in Canada or evidence of their association with disease worldwide (Miller et al. 2016). Assays utilizing Taqman probes (Table S1) were designed to target both RNA and DNA. Not all of the same assays were used over the course of the qPCR runs, as some new assays were developed (Mordecai et al. 2019) (107 dynamic arrays run over the course of four years).

Total RNA and DNA were extracted using methods previously described in (Miller et al. 2016; Thakur et al. 2018). Briefly, tissues were homogenized separately in TRI-reagent™ (Ambion Inc., Austin, TX, USA). Next, 1-bromo-3-chloropropane was added to the homogenate, and equal volumes of both the aqueous phase (RNA) and the organic/interphase (DNA) from each tissue type were combined for extraction. RNA extractions were carried out using MagMAX™-96 for Microarrays Total RNA Isolation Kits (Ambion Inc.) with a Biomek NXP™ automated liquid-handling instrument. RNA quantity and purity was assessed by measuring the A260/A280 ratio using a Beckman Coulter DTX 880 Multimode Spectrophotometer (Brea, CA, USA). DNA was extracted using the TNE5-6U method following the Qiagen BioSprint protocol.

Normalized RNA (1µg) was reverse transcribed to cDNA using the SuperScript VILO MasterMix Kit (Invitrogen, Carlsbad, CA) following the manufacturer's instructions. DNA and cDNA were then mixed in equal proportions. The assay volume used for qPCR on the BioMark is small (7nL) and therefore a pre-amplification step is recommended by the manufacturer. Thus, 0.2µmol/L of the cDNA/DNA mix from each sample was pre-amplified with primer pairs corresponding to all assays (microbes and 3 reference genes) in a 5µL reaction volume using 1X TaqMan Preamp Master Mix (Applied Biosystems, Foster City, California) according to the BioMark protocol. Unincorporated primers were removed using ExoSAP-IT™ (Affymetrix, Santa Clara, California), and samples were diluted 1:5 in DNA Suspension Buffer (Teknova, Hollister, California).

Artificial positive constructs (APC clones) corresponding to all assays were run in six serial dilutions on the dynamic array to construct a standard curve and calculate efficiency for each assay and estimate RNA copy number for each positive sample. The APC clones contained an additional probe labelled with NED™ reporter dye (Life Technologies) that allowed for the detection of vector contamination (see Miller et al. 2016).

A 5µL sample mix was prepared containing 1X TaqMan Universal Master-Mix (Life Technologies), 1X GE Sample Loading Reagent (Fluidigm PN 85000746), and amplified cDNA/DNA, which was added to each assay inlet of the array following the manufacturer's recommendations. All assays were run in duplicate. Five µL of assay mix was prepared containing 10µM primers (infectious agent in FAM-MGB and APC in NED-MGB) and 3µM probes for the TaqMan assays. After loading the assays and samples into the chip using an IFC controller HX (Fluidigm), PCR was performed with the following conditions: 50°C for 2 min, 95°C for 10 min, followed by 40 cycles of 95°C for 15 s and 60°C for 1 min.

Cycle threshold was determined using the BioMark Real-Time PCR analysis software. Reaction curves for each positive sample-assay combination were visually evaluated for abnormal curve shapes, close correspondence between duplicates, and presence of APC contamination as indicated by NED positives. Using scripts created in R statistical software (R Core Team 2019), we calculated efficiency for each assay (standard curve method (Larionov et al. 2005)), omitted results where only one duplicate was positive for a sample-assay combination, removed NED positive samples, and averaged duplicates. Limit of detection (LOD) is defined as the estimated cycle threshold (Ct) number under which true positive results are expected 95% of the time for a given assay (Miller et al. 2016). Because LOD was established

for maximum compliance with OIE standards but limits the sensitivity of the BioMark to detect low-level infection, we present data exceeding the LOD. Note that we only included detections beyond the LOD for infectious agents that were also detected within the LOD whereas infectious agents only detected beyond the LOD were considered to be false positives.

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Table S1. Taqman assays run for 59 infectious agents and 3 host reference genes (Ref) in Chinook salmon mixed-tissue samples (2008 2018) using the Fluidigm Biomark HT-qRT-PCR platform (DFO Pacific Biological Station, Nanaimo, BC). Below the limit of detection (LOD) Ct value, positive samples are detected 95% of the time.

Scientific name	Abbrev.	LOD (ct)	Forward primer sequence (5'-3')	Reverse primer sequence (5'-3')	Probe sequence (FAM-5'-3'-MGB)
<i>Aeromonas hydrophila</i>	ae_hyd	28.7	ACCGTGCTCATTACTCTGATG	CCAACCCAGACGGGAAGAA	TGATGGTGAGCTGGTTG
<i>Aeromonas salmonicida</i>	ae_sal	25.6	TAAAGCACTGTCTGTACC	GCTACTTCACCTGATTGG	ACATCAGCAGGCTTCAGAGTCACTG
<i>Candidatus Branchiomonas cysticola</i>	c_b_cys	25.7	AATACATCGGAACGTGTCTAGTG	GCCATCAGCCGCTCATGTG	CTCGGTCCAGGCTTCTCTCCCA
<i>Flavobacterium psychrophilum</i>	fl_psy	29.5	GATCCTTATTCTCACAGTACCGTCAA	TGTAAACTGCTTTTGACAGGAA	AAACTCGGTCTGTGACC
<i>Moritella viscosa</i>	mo_vis		CGTTGCGAATGCAGAGGT	AGGCATTGCTTGCTGGTTA	TGCAGGCAAGCCAACTTCGACA
<i>Candidatus Piscichlamydia salmonis</i>	pch_sal	23.3	TCACCCCGAGGCTGCTT	GAATTCATTTCCTCTTG	CAAACTGCTAGACTAGAGT
<i>Piscirickettsia salmonis</i>	pisck_sal	23.3	TCTGGGAAGTGTGGCGATAGA	TCCCGACCTACTCTTGTTCATC	TGATAGCCCCGTACACGAAACGGCATA
<i>Renibacterium salmoninarum</i>	re_sal	25.9	CAACAGGGTGGTTATTCTGCTTTC	CTATAAGAGCCACCAGCTGCAA	CTCCAGCGCCGAGGAGGAC
Rickettsia-like organism	rlo	25.2	GGCTCAACCCAAGAAGTCTT	GTGCAACAGCGTCAGTACT	CCCAGATAACCGCTTCGCTCCG
<i>Candidatus Syngnamydia salmonis</i>	sch	27.9	GGGTAGCCCGATATCTTCAAAGT	CCCATGAGCCGCTCTCTCT	TCCTTCGGGACCTTAC
<i>Tenacibaculum maritimum</i>	te_mar		TGCCTTCTACAGAGGGATAGCC	CTATCGTTGCCATGGTAAGCCG	CACTTTGGAATGGCATCG
<i>Vibrio anguillarum</i>	vi_ang	26.4	CCGTCATGCTATCTAGAGATGTATTTGA	CCATACGCAGCCAAAATCA	TCATTTGACGAGCGTCTTGTTCAGC
<i>Vibrio salmonicida</i>	vi_sal	25.8	GTGTGATGACCGTTCCATATTT	GCTATTGTACTACTCTGTTTCTT	TCGCTTCATGTTGTGTAATTAGGAGCGA
<i>Yersinia ruckeri</i>	ye_ruc	25.8	TGCCGCGTGTGTAAGAA	ACGGAGTTAGCCGGTGCTT	AATAGCACTGAACATTGAC
<i>Dermocystidium salmonis</i>	de_sal	25.5	CAGCCAATCCTTCGCTTCT	GACGGACGCACACCACAGT	AAGCGGCGTGTGCC
<i>Ichthyophonus</i> spp.	ic_hof	24.2	GTCTGTACTGGTACGGCAGTTTC	TCCCGAACTCAGTAGACTCAAA	TAAGAGCACCCACTGCCTTCGAGAAGA
<i>Ceratonova shasta</i>	ce_sha	28.5	CCAGCTTGAGATTAGCTCGGTAA	CCCCGGAACCCGAAAG	CGAGCCAAGTTGGTCTCTCCGTGAAAAC
<i>Sphaerothecum destruens</i>	sp_des	26.5	GGGTATCCTTCTCTCGAAATTG	CCCAAACCTGACGCACACT	CGTGTGCGCTTAAT
<i>Facilispora margolisi</i>	fa_mar	30.6	AGGAAGGAGCACGCAAGAAC	CGCGTGCAGCCAGTAC	TCAGTGATGCCCTCAGA

Scientific name	Abbrev.	LOD (ct)	Forward primer sequence (5'-3')	Reverse primer sequence (5'-3')	Probe sequence (FAM-5'-3'-MGB)
<i>Loma</i> spp.	lo_sal	25.4	GGAGTCGCAGCGAAGATAGC	CTTTTCCTCCCTTACTCATATGCTT	TGCCTGAAATCACGAGAGTGAGACTACCC
<i>Nucleospora salmonis</i>	nuc_sal	26.1	GCCGCAGATCATTACTAAAAACCT	CGATCGCCGCATCTAAACA	CCCCGCGCATCCAGAAATACGC
<i>Paranucleospora theridion</i> (syn. <i>Desmozoon lepeophtherii</i>)	pa_ther	28.2	CGGACAGGGAGCATGGTATAG	GGTCCAGGTTGGGTCTTGAG	TTGGCGAAGAATGAAA
<i>Kudoa thyristes</i>	ku_thy	26.2	TGGCGGCCAAATCTAGGTT	GACCGCACACAAGAAGTTAATCC	TATCGCGAGAGCCGC
<i>Myxobolus arcticus</i>	my_arc	26.8	TGGTAGATACTGAATATCCGGGTTT	AACTGCGCGGTCAAAGTTG	CGTTGATTGTGAGGTTGG
<i>Myxobolus cerebalis</i>	my_cer	26.2	GCCATTGAATTTGACTTTGGATTA	ACCATTGATGTAAGCCCGAACT	TCGAAGCCTTGACCATCTTTTGCC
<i>Myxobolus insidiosus</i>	my_ins	26.4	CCAATTTGGGAGCGTCAAA	CGATCGGCAAAGTTATCTAGATTCA	CTCTCAAGGCATTAT
<i>Parvicapsula kabatai</i>	pa_kab	25.6	CGACCATCTGCACGGTACTG	ACACCACAACCTGCCTTCCA	CTTCGGGTAGGTCCGG
<i>Parvicapsula minibicornis</i>	pa_min	29.6	AATAGTTGTTTGTCTGCACTCTGT	CCGATAGGCTATCCAGTACCTAGTAAG	TGTCCACCTAGTAAGGC
<i>Parvicapsula pseudobranchicola</i>	pa_pse	25.2	CAGCTCCAGTAGTGATTTCA	TTGAGCACTCTGCTTTATCAA	CGTATTGCTGCTTTGACATGCAGT
<i>Tetracapsuloides bryosalmonae</i>	te_bry	25.0	GCGAGATTTGTTGCATTTAAAAAG	GCACATGCAGTGTCCAATCG	CAAATTGTGGAACCGTCCGACTACGA
<i>Gyrodactylus salaris</i>	gy_sal	26.4	CGATCGTCACTCGGAATCG	GGTGGCGCACCTATTCTACA	TCTTATTAACCACTTCTGC
<i>Nanophyetus salmincola</i>	na_sal	24.3	GATCTGCATTTGGTTCTGTAACA	CCAACGCCACAATGATAGCTATAC	TGAGGCGTGTTTTATG
<i>Cryptobia salmositica</i>	cr_sal	24.3	TCAGTGCCTTTCCAGGACATC	GAGGCATCCACTCCAATAGAC	AGGAGGACATGGCAGCCTTTGTAT
<i>Ichthyophthirius multifiliis</i>	ic_mul	23.7	AAATGGGCATACGTTTGCAAA	AACCTGCCTGAAACTCTA	ACTCGGCCTTCACTGGTTCGACTTGGATTTT
<i>Neoparamoeba perurans</i>	ne_per	25.4	GTTCTTTCGGGAGCTGGGAG	GAATATCGCCGGCACAAAAG	CAATGCCATTCTTTTCGGA
<i>Spironucleus salmonicida</i>	sp_sal	26.1	GCAGCCGCGTAATTCC	CGAACTTTTAACTGCAGCAACA	ACACGGAGAGTATTCT
Atlantic salmon calicivirus virus	ascv		ACCGACTGCCGGTTGT	CTTAGGGTTAAAGCAGTCG	CTCCGATTGCCTGTGATAATACC
Atlantic salmon paramyxovirus	Aspv	26.2	CCCATATTAGCAAATGAGCTCTATCTT	CGTTAAGGAACTCATCATTG	AGCCCTTTTGTCTGCAGCTT
Chinook aquareovirus	Reov		AACTTCGGCTTTCTGCTATGC	GAGGACAAGGGTCTCCATCTGA	TTAATTGCGGTACTGCTC
Cutthroat trout virus 2	ctv		CCACTTGTGCTACGATGAAAC	ATGCCGGGCATC	CGCCTCCTTTGCCCTTCTC
Erythrocytic necrosis virus	ven	24.9	CGTAGGGCCCAATAGTTTCT	GGAGGAAATGCAGACAAGATTG	TCTTGCCGTTATTTCCAGCACCCG

Scientific name	Abbrev.	LOD (ct)	Forward primer sequence (5'-3')	Reverse primer sequence (5'-3')	Probe sequence (FAM-5'-3'-MGB)
Infectious hematopoietic necrosis virus	ihnv	27.6	AGAGCCAAGGCACTGTGCG	TTCTTTGCGGCTTGGTTGA	TGAGACTGAGCGGGACA
Infectious pancreatic necrosis virus	ipnv	27.6	GCAACTTACTTGAGATCCATTATGCT	AGACCTCTAAGTTGTATGACGAGGTCTCT	CGAGAATGGGCCAGCAAGCA
Infectious salmon anemia virus 7	isa7	27.0	TGGGATCATGTGTTTCTGCTA	GAAAATCCATGTTCTCAGATG-CAA	CACATGACCCCTCGTC
Infectious salmon anemia virus 8	isa8	26.1	TGGGCAATGGTGTATGGTATGA	GAAGTCGATGAACTGCAGCGA	CAGGATGCAGATGTATGC
Pacific salmon nidovirus	cov		GGATAATCCCAACCGAAAAGTTT	GCATGAAATGTTGTCTCGGTTTAA	CGATCCCATTATC
Pacific salmon parvovirus	pspv	26.4	CCCTCAGGCTCCGATTTTAT	CGAAGACAACATGGAGGTGACA	CAATTGGAGGCAACTGTA
Piscine myocarditis virus	pmcv	26.3	TTCCAAACAATTCGAGAAGCG	ACCTGCCATTTTCCCCTCTT	CCGGGTAAAGTATTTGCGTC
Piscine orthoreovirus	PRv	26.1	TGCTAACACTCCAGGAGTCATTG	TGAATCCGCTGCAGATGAGTA	CGCCGGTAGCTCT
Putative narna-like virus	pnarna		TGTCCCTGAAGATTCATTTTCA	TCCTAGGTGATGATATAAT	CTATGTAAAGCCTCGTCGGTGAT
Putative RNA virus 1	smallUK		GTACCTAATTTAACTGGAACAGTAGAC	TGCAACAGGCAAGTGATATGCTTGA	CGTTCAGTAACACAAGTATCCAAA
Putative toti-like virus	toti		TCTGCGCGCTGCACCTA	CAAGTGCTACTGCG	ATGCGGAGGAACTCACACACT
Rainbow trout orthomyxovirus	ortho		GGAAGCAGTGGACGCTAACC	TCGCGAAGGTCTCTCAATGTC	ATTCTTCTCATCAAAGGCA
Salmon alphavirus	sav	26.3	CCGGCCCTGAACCAGTT	GTAGCCAAGTGGGAGAAAGCT	TCGAAGTGGTGGCCAG
Salmon gill pox virus	sgpx		ATCCAAAATACGGAACATAAGCAAT	CAACGACAAGGAGATCAACGC	CTCAGAACTTCAAAGGA
Salmonid herpesvirus	shv	26.6	GCCTGGACCACAATCTCAATG	CGAGACAGTGTGGCAAGACAAC	CCAACAGGATGGTCATTA
Salmon pescarenavirus 1	arena1		CCTGCCTCTTTGCTCATTGTG	AGAAAAAGCTGTGGTACTTTAGAAAGC	ATCCGCCTAACGGTTGG
Salmon pescarenavirus 2	arena2		AACATGAAGGCGGATTTCGTT	CAGCCCCTGACTGAGT	CAAGTGATGTAAGCTTG
Viral encephalopathy and retinopathy virus	ver	26.2	TTCCAGCGATACGCTGTTGA	CACCGCCGTGTTTGC	AAATTCAGCCAATGTGCCCC
Viral hemorrhagic septicemia virus	vhsv	26.9	ATGAGGCAGGTGTCGGAGG	TGTAGTAGGACTCTCCAGCATCC	TACGCCATCATGATGAGT
78d16.1	Ref	NA	GTCAAGACTGGAGGCTCAGAG	GATCAAGCCCCAGAAGTGTGTTG	AAGGTGATCCCTCGCCGTCCGA
COIL-P84-2	Ref	NA	GCTCATTTGAGGAGAAGGAGGATG	CTGGCGATGCTGTTCTGAG	TTATCAAGCAGCAAGCC
MRPL40	Ref	NA	CCCAGTATGAGGCACCTGAAGG	GTTAATGCTGCCACCCTCTCAC	ACAACAACATCACCA

Table S2: Groupings of infectious agents used in spatial models for hypothesis testing via meta-analysis (excluding distance to aquaculture, see Table S3). References are provided as evidence for groupings.

Taxa	Transmission environment [§]	Temperature preference [£]	Transmission complexity [¥]
<i>Ca. Branchiomonas cysticola</i>	<i>freshwater</i> ^{1,2}	unknown	freshwater-simple ²
<i>Flavobacterium psychrophilum</i>	freshwater ^{3,4}	cold ⁵	freshwater-simple
<i>Ca. Syngnamydia salmonis</i>	saltwater ¹	unknown	saltwater-simple
<i>Piscirickettsia salmonis</i>	saltwater ⁶	cold ⁶	saltwater-simple ⁷
<i>Renibacterium salmoninarum</i>	<i>freshwater</i>	warm ⁸	freshwater-simple
Rickettsia-like organism	freshwater ⁹	warm ¹⁰	freshwater-simple
<i>Tenacibaculum maritimum</i>	saltwater ¹¹	warm ¹¹	saltwater-simple
<i>Ichthyophonus hoferi</i>	<i>saltwater</i> ^{1,12}	warm ¹³	saltwater-complex ¹²
<i>Sphaerothecum destruens</i>	<i>saltwater</i> [*]	warm ¹⁴	saltwater-simple
<i>Facilispora margolisi</i>	saltwater ¹⁵	unknown	saltwater-complex
<i>Loma salmonae</i>	<i>saltwater</i> ^{16*}	warm ^{16,17}	saltwater-simple ¹⁸
<i>Paranucleospora theridion</i>	saltwater ¹⁹	warm ^{19,20}	saltwater-complex ²¹
<i>Ceratonova shasta</i>	freshwater ²²	warm ^{10,23}	freshwater-complex ²²
<i>Kudoa thyrsites</i>	saltwater	warm ¹⁰	saltwater-complex
<i>Myxobolus arcticus</i>	freshwater ²⁴	unknown	freshwater-complex ²⁴
<i>Parvicapsula kabatai</i>	saltwater ¹	unknown	saltwater-complex
<i>Parvicapsula minibicornis</i>	freshwater ²⁵	warm ^{10,26}	freshwater-complex ²⁵
<i>Parvicapsula pseudobranchicola</i>	saltwater ²⁷	unknown	saltwater-complex
<i>Tetracapsuloides bryosalmonae</i>	freshwater	warm ^{28,29}	freshwater-complex ³⁰
<i>Cryptobia salmositica</i>	freshwater ³¹	cold ³¹	Freshwater-complex ³¹
<i>Nanophyetus salmincola</i>	freshwater	unknown	freshwater-complex
<i>Ichthyophthirius multifiliis</i>	freshwater ³²	warm ^{32,33}	freshwater-simple
Pacific salmon arena virus 1	saltwater [*]	cold ³⁴	saltwater-simple
Pacific salmon nidovirus	freshwater [*]	cold ³⁴	freshwater-simple
Piscine orthoreovirus	<i>saltwater</i> [*]	cold ³⁴	saltwater-simple
Erythrocytic necrosis virus	<i>saltwater</i> ³⁵	cold ³⁴	saltwater-simple
Putative RNA virus 1 ³⁶	saltwater [*]	cold ³⁴	saltwater-complex ³⁶

§ Pathogens in italics can be transmitted in both environments. We chose the environment where transmission is most likely to first occur for juvenile Chinook salmon in the study area.

* If evidence from published studies was insufficient to categorize pathogen, we relied upon unpublished observations from the SSHI dataset

£ Given that viability of most fish viruses decline as water temperature increases³⁴, we categorized all viruses, including those recently discovered, as cold-associated.

¥ For myxozoans where life cycle is unknown, transmission is assumed to involve an invertebrate alternate host

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Table S3. Grouping of infectious agents for the “distance to aquaculture” hypothesis meta-analysis. References are provided to support categorization rationale.

Taxa	Ranking	Rationale
<i>Cryptobia salmositica</i>	Moderate	Outbreaks in BC Pacific salmon in marine netpens ¹ Transmission without alternate host requires physical contact ²
<i>Ca. Branchiomonas cysticola</i>	Moderate	PCR detections in BC aquaculture ^{3,4} Frequently the most common pathogen in studies of Pacific wild salmon ^{5,6}
<i>Flavobacterium psychrophilum</i>	High	Major pathogen in FW aquaculture but little known in SW ⁷ PCR detections in BC aquaculture ^{3,4} Elevated eDNA at active BC aquaculture sites relative to inactive ⁸
<i>Ca. Syngnamydia salmonis</i>	Moderate	Present in Norwegian and BC aquaculture ^{4,9}
<i>Piscirickettsia salmonis</i>	High	Frequent outbreaks in Chile ¹⁰ PCR detections in BC aquaculture ⁴ Elevated eDNA at active BC aquaculture sites relative to inactive ⁸
<i>Renibacterium salmoninarum</i>	High	Outbreaks and transmission in SW for BC aquaculture ¹¹
Rickettsia-like organism	Low	No evidence of presence in BC aquaculture or evidence of SW aquaculture presence elsewhere
<i>Tenacibaculum maritimum</i>	High	Outbreaks in BC aquaculture ¹² Potential for transmission to BC wild salmon from aquaculture ¹³ Elevated eDNA at active BC aquaculture sites relative to inactive ⁸
<i>Ichthyophonus hoferi</i>	Moderate	Potential to persist in feeds that contain herring ¹⁴ Presence in first summer <i>S. salar</i> in BC marine netpens ¹⁵
<i>Sphaerothecum destruens</i>	Moderate	Outbreaks in netpen Pacific salmon in WA, USA ¹⁶
<i>Facilispora margolisi</i>	Moderate	PCR detections in BC aquaculture ³
<i>Loma salmonae</i>	Moderate	Outbreaks in netpen Pacific salmon ¹⁷
<i>Paranucleospora theridion</i>	High	High prevalence in Norwegian aquaculture ¹⁸ High prevalence in BC aquaculture ⁴
<i>Ceratonova shasta</i>	Low	Freshwater pathogen with no evidence of presence in marine aquaculture
<i>Kudoa thyrsites</i>	High	Common in BC aquaculture ¹⁹ Elevated eDNA at active BC aquaculture sites relative to inactive ⁸
<i>Myxobolus arcticus</i>	Low	Freshwater pathogen with no evidence of presence in marine aquaculture
<i>Parvicapsula kabatai</i>	Moderate	PCR detections in BC aquaculture ⁴
<i>Parvicapsula minibicornis</i>	Low	Freshwater pathogen with no evidence of presence in marine aquaculture
<i>Parvicapsula pseudobranchicola</i>	High	High prevalence in Norwegian aquaculture ²⁰ Elevated eDNA at active BC aquaculture sites relative to inactive ⁸
<i>Tetracapsuloides bryosalmonae</i>	Low	Freshwater pathogen with no evidence of presence in marine aquaculture
<i>Nanophyetus salmincola</i>	Low	Freshwater pathogen with no evidence of presence in marine aquaculture
<i>Ichthyophthirius multifiliis</i>	Moderate	Outbreaks in FW salmonids ²¹ PCR detections in BC aquaculture ³

Taxa	Ranking	Rationale
Pacific salmon arena virus 1	Moderate	PCR detections in Pacific salmon in BC aquaculture ²²
Pacific salmon nidovirus	Low	Little known virus with majority of detections in freshwater ²³
Piscine orthoreovirus	High	High prevalence in BC aquaculture ^{3,24} Association between likelihood of infection and distance to aquaculture ²⁴
Erythrocytic necrosis virus	Moderate	Infections in Pacific and Atlantic salmon in BC aquaculture ¹⁵ PCR detections in BC aquaculture ⁴
Putative RNA virus 1	Low	No evidence of presence in marine aquaculture

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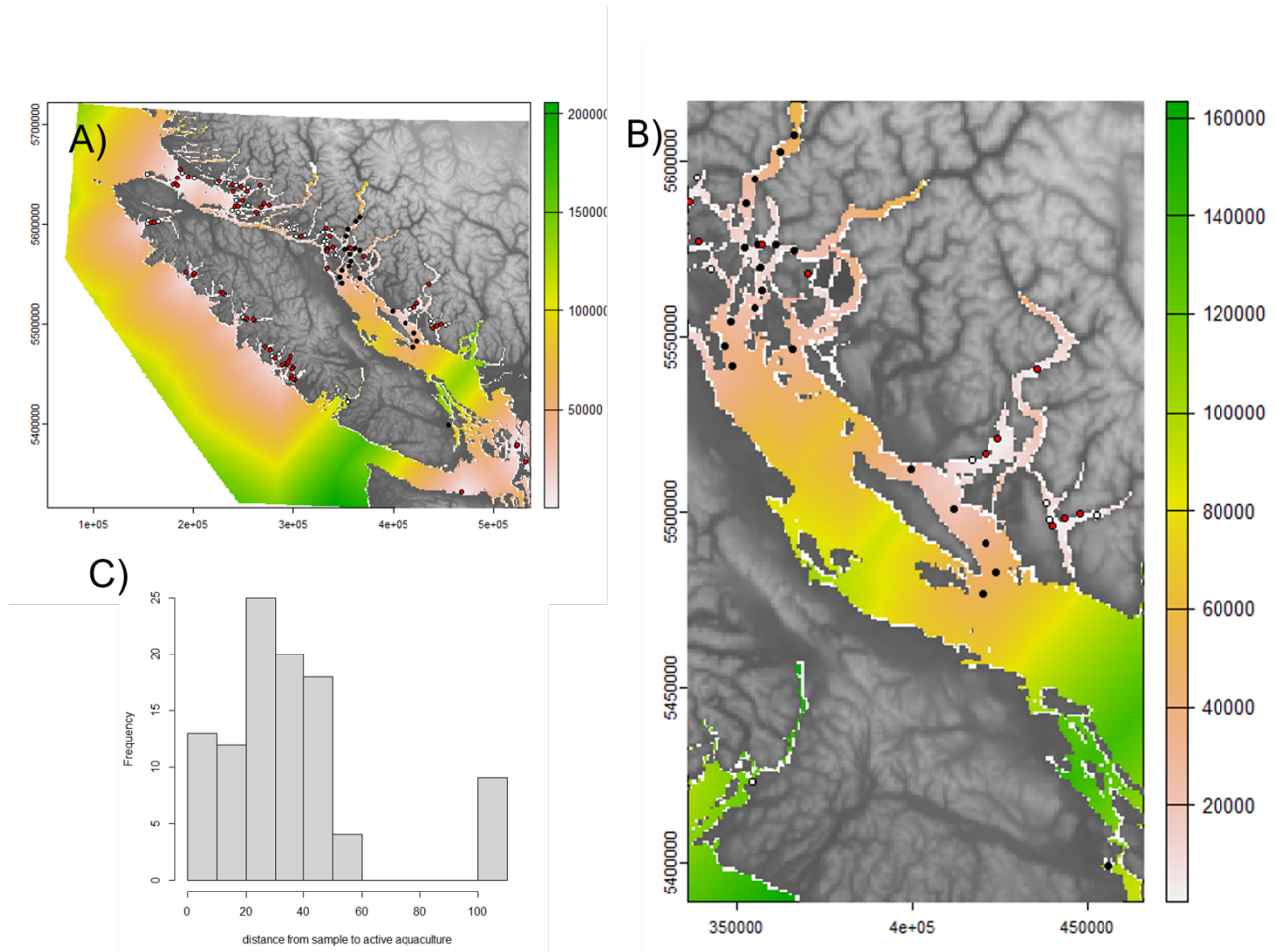


Figure S1. Example of seaway distance from active aquaculture data. Panel A shows the seaway distance raster from September, 2014. Cell color of raster over sea surface (1 km resolution) indicates distance from nearest active finfish aquaculture netpen facilities (red points) in meters. White points are aquaculture facilities that were inactive in September, 2014. Black points are locations where Chinook salmon were collected in September, 2014. Panel B is zoomed in to the extent of Chinook salmon collections in September, 2014. Panel C shows a histogram of seaway distances (km) to active aquaculture for Chinook collected in September, 2014 (corresponding to points in B).

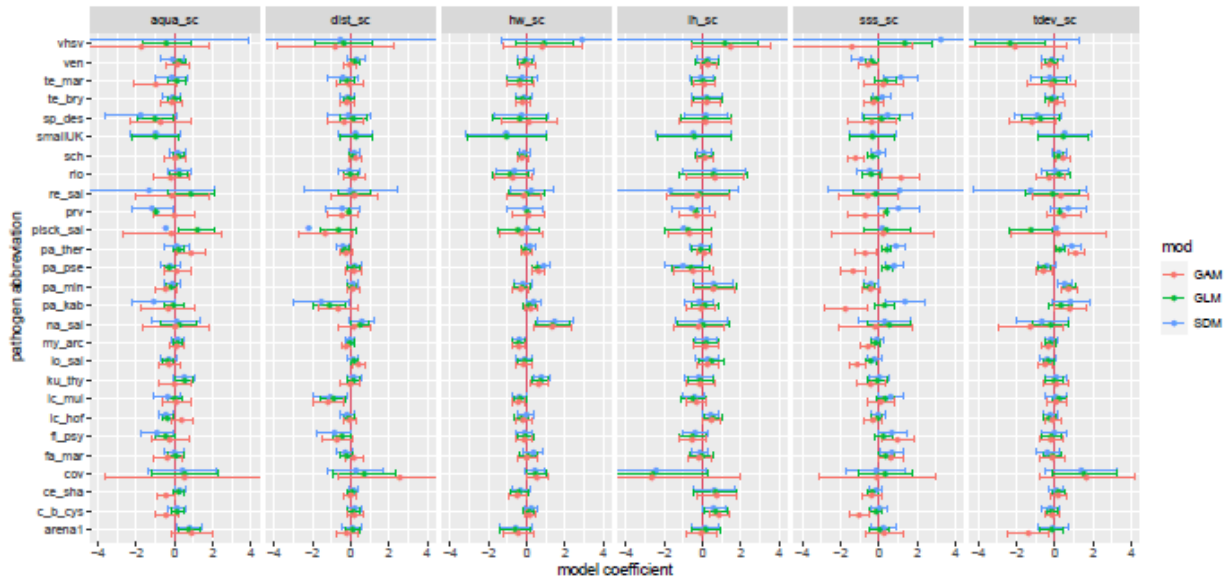


Figure S2. Plots of all Chinook model coefficients for three spatial model types demonstrating the consistency between geostatistical modeling approaches. Error bars represent 99% confidence intervals. Full names corresponding to pathogen abbreviations can be found in Table S1. Model variables at top of plots represent: distance to active aquaculture (aqua), distance to nearest shoreline (dist), finclip status (hw), age at ocean entry (lh), sea surface salinity (SSS), and sea surface temperature deviation (tdev).

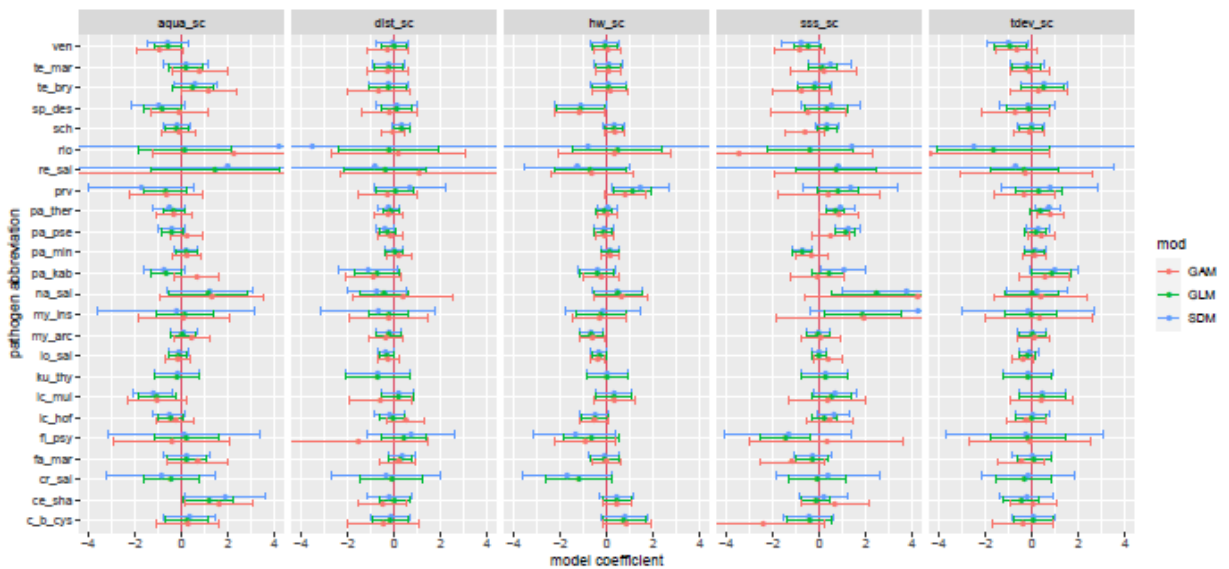


Figure S3. Plots of all Coho model coefficients for three spatial model types demonstrating the consistency between geostatistical modeling approaches. Error bars represent 99% confidence intervals. Full names corresponding to pathogen abbreviations can be found in Table S1. Model variables at top of plots represent: distance to active aquaculture (aqua), distance to nearest shoreline (dist), finclip status (hw), age at ocean entry (lh), sea surface salinity (SSS), and sea surface temperature deviation (tdev).

Figures S4–S13. Forest plots for hypothesis tests using sdmTMB model coefficients (see Tables 2 & 3). Squares are model estimates (orange = Chinook, blue = Coho) with 99% confidence intervals. Diamonds represent meta-analytical means with 95% confidence intervals. The bottom diamond and dotted red vertical line represent the mean coefficient estimate across all models. Square size is proportional to weight, which represents a given coefficient estimates relative contribution to the group and overall means.

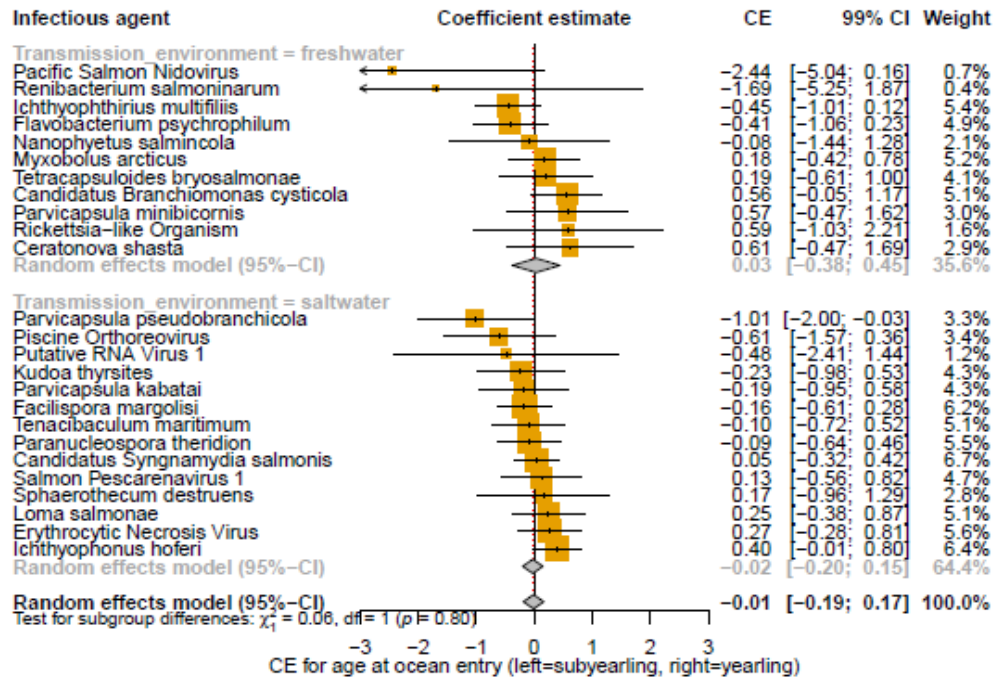


Figure S4

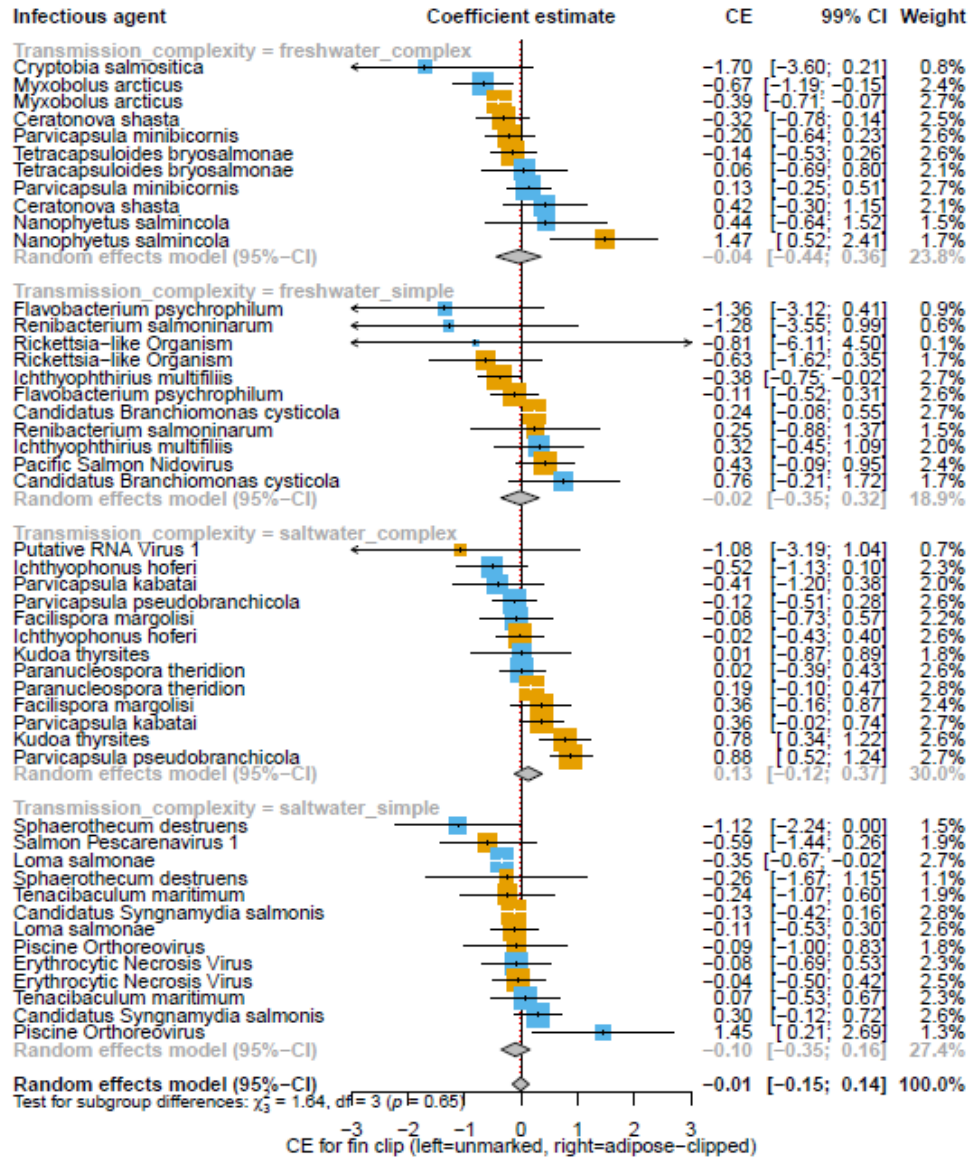


Figure S5

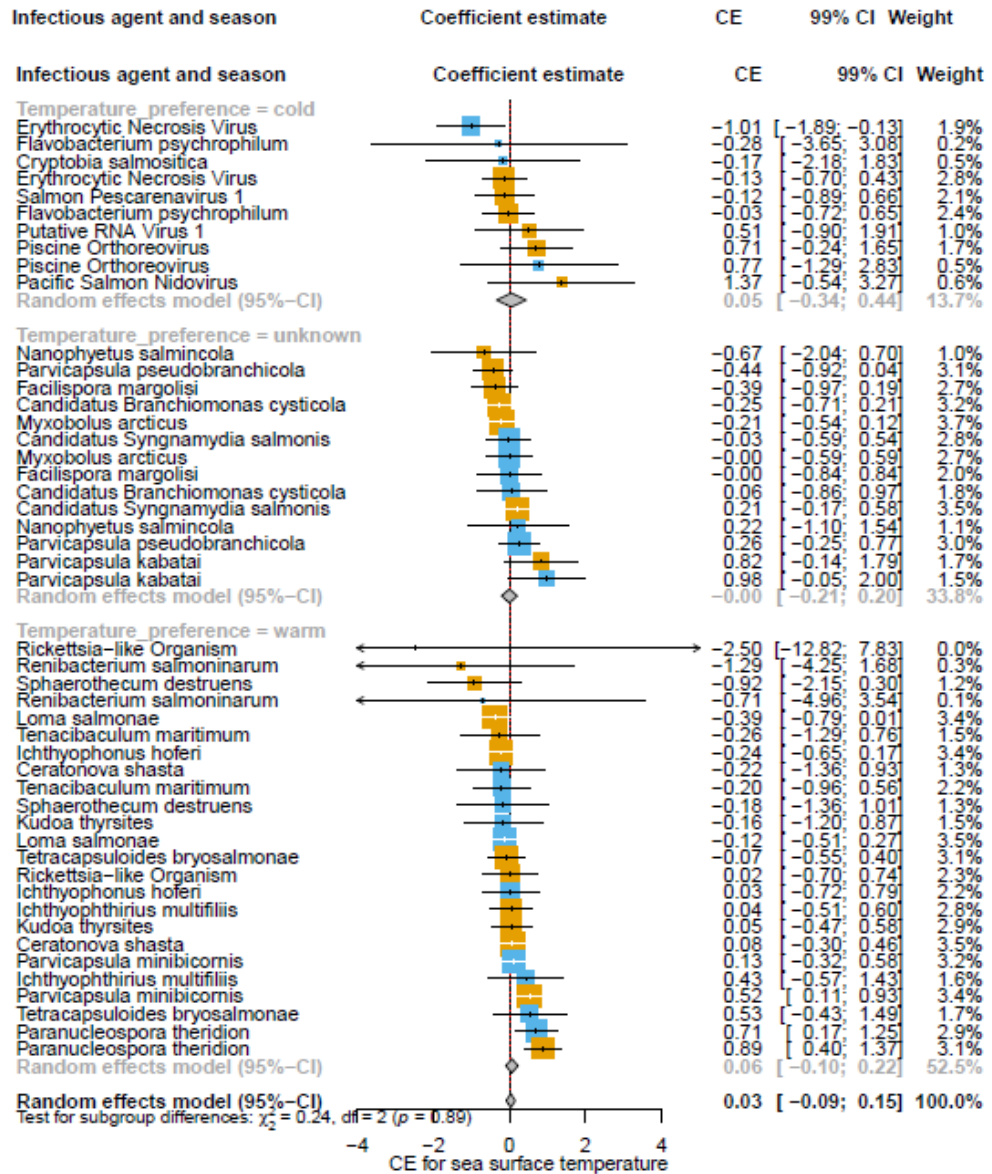


Figure S6

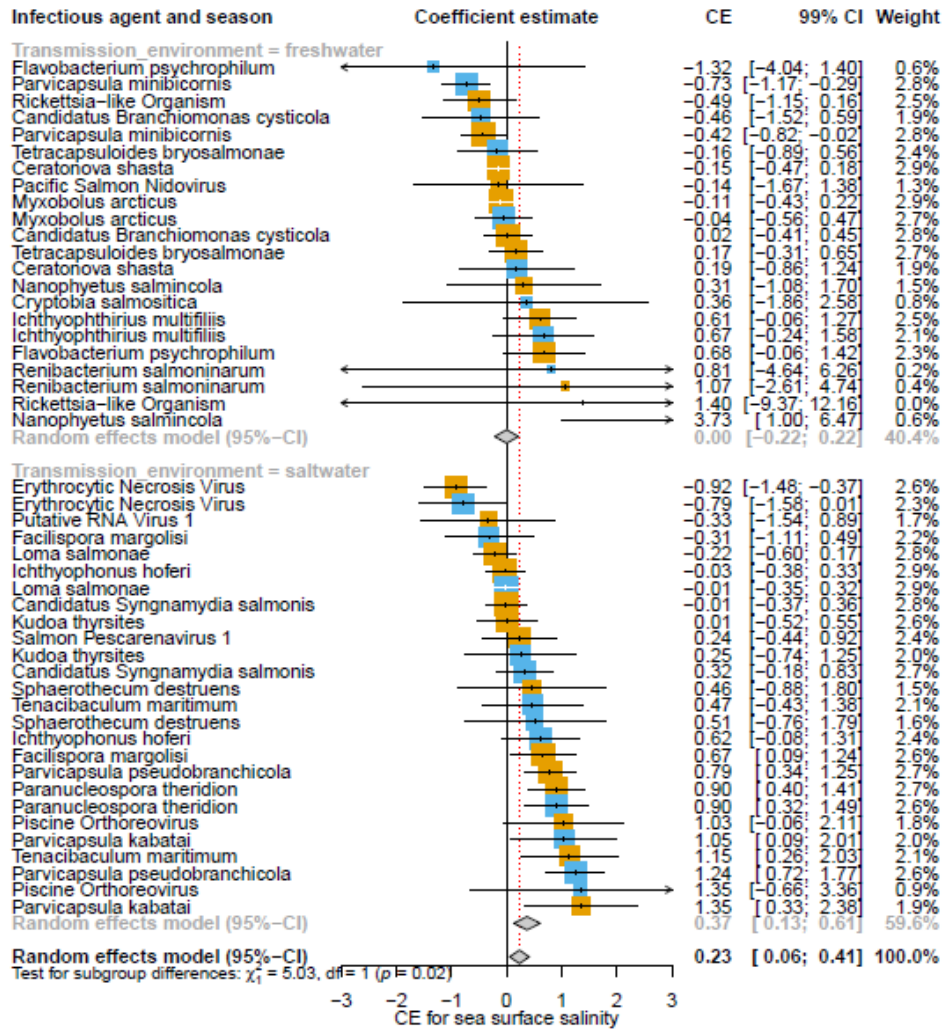


Figure S7

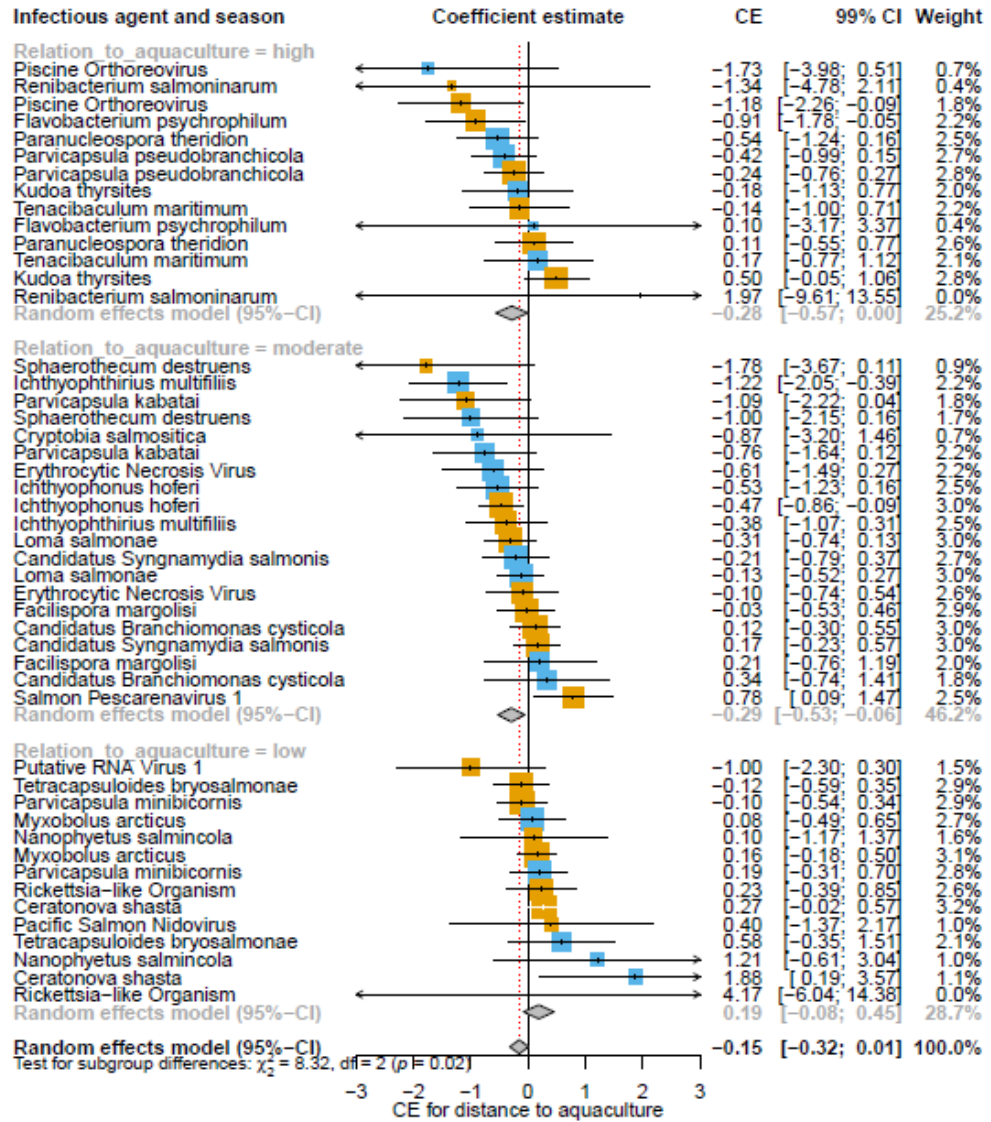


Figure S8

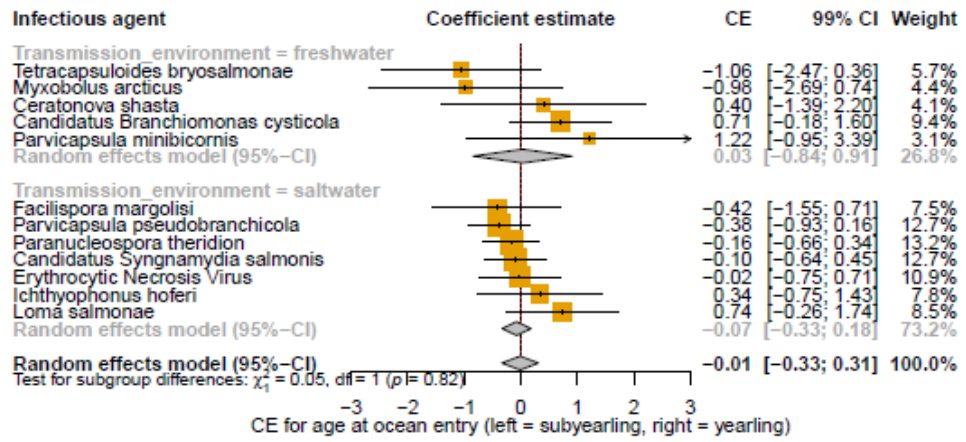


Figure S9

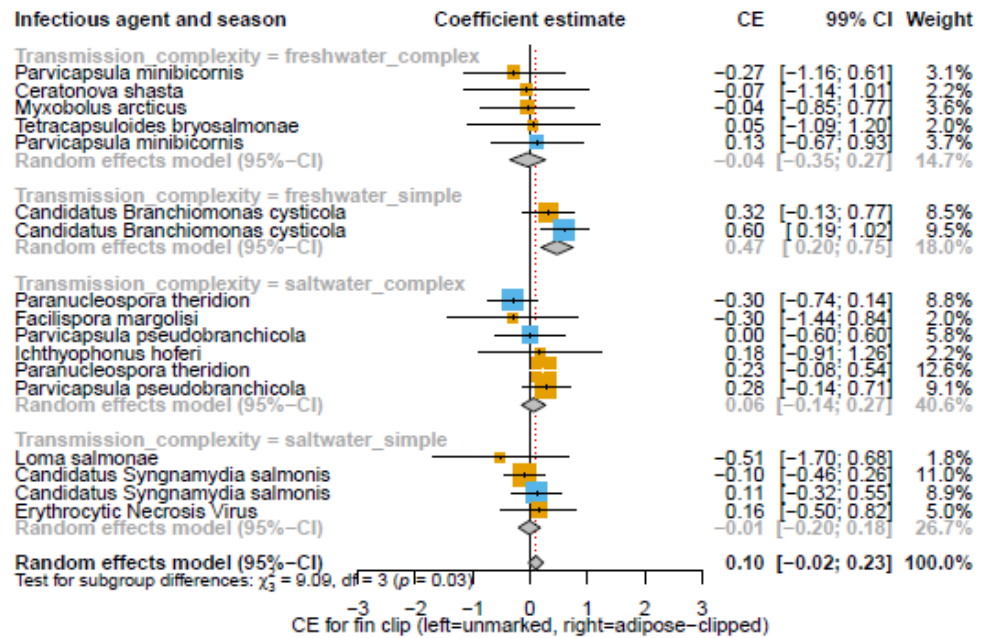


Figure S10

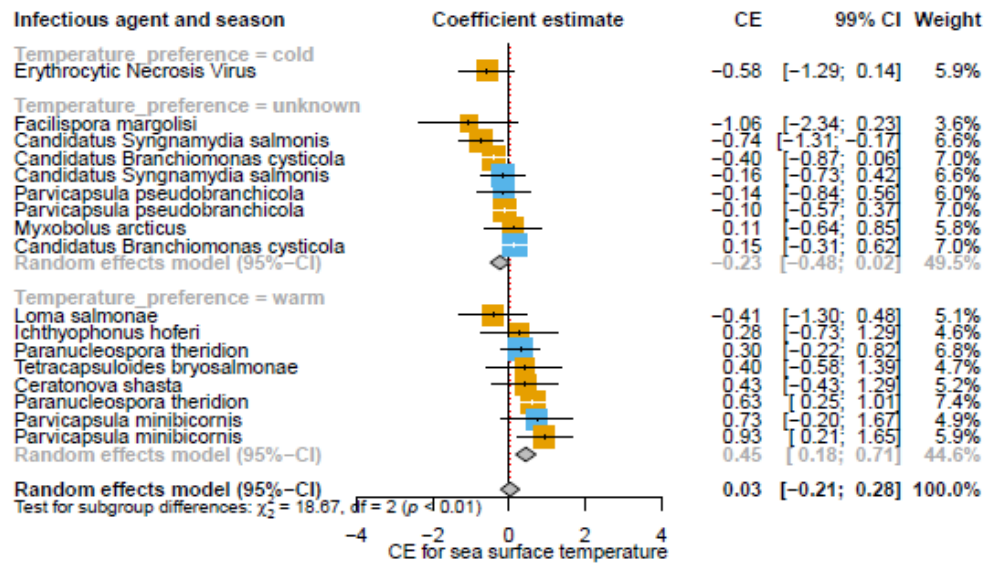


Figure S11

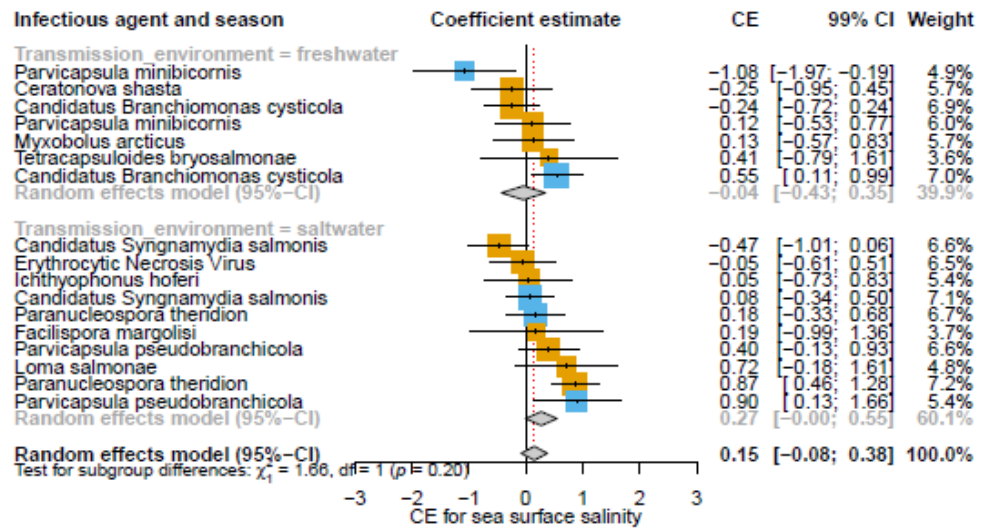


Figure S12

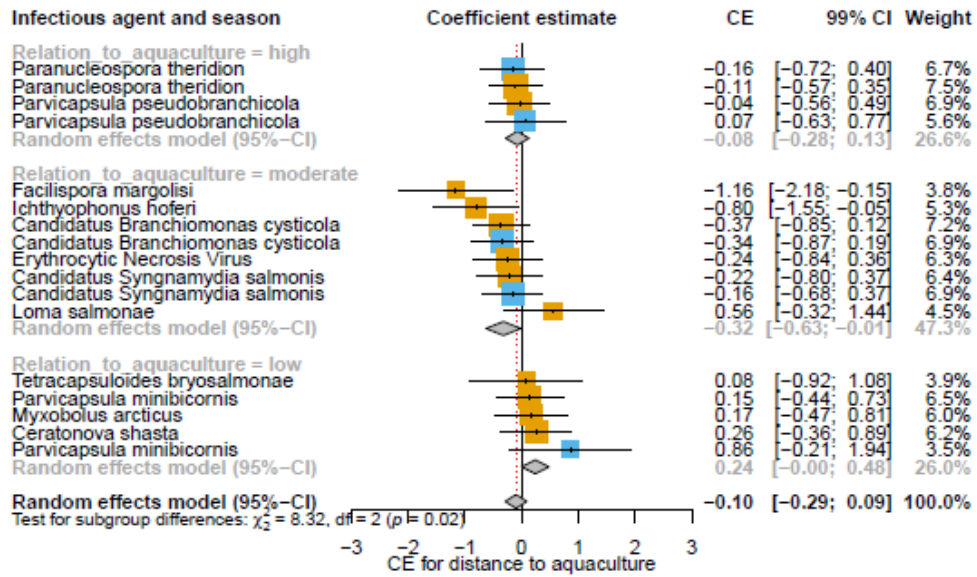


Figure S13