

EST and Mitochondrial DNA Sequences Support a Distinct Pacific Form of Salmon Louse, *Lepeophtheirus salmonis*

Ryosuke Yazawa · Motoshige Yasuike · Jong Leong ·
Kristian R. von Schalburg · Glenn A. Cooper ·
Marianne Beetz-Sargent · Adrienne Robb ·
William S. Davidson · Simon R. M. Jones · Ben F. Koop

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Abstract Nuclear deoxyribonucleic acid sequences from approximately 15,000 salmon louse expressed sequence tags (ESTs), the complete mitochondrial genome (16,148bp) of salmon louse, and 16S ribosomal ribonucleic acid (rRNA) and cytochrome oxidase subunit I (*COI*) genes from 68 salmon lice collected from Japan, Alaska, and western Canada support a Pacific lineage of *Lepeophtheirus salmonis* that is distinct from that occurring in the Atlantic Ocean. On average, nuclear genes are 3.2% different, the complete mitochondrial genome is 7.1% different, and 16S rRNA and *COI* genes are 4.2% and 6.1% different, respectively. Reduced genetic diversity within the Pacific form of *L. salmonis* is consistent with an introduction into the Pacific from the Atlantic Ocean. The level of divergence is consistent with the hypothesis that the Pacific form of *L. salmonis* coevolved with Pacific salmon (*Onchorhynchus* spp.) and the Atlantic form coevolved with Atlantic salmo-

nids (*Salmo* spp.) independently for the last 2.5–11 million years. The level of genetic divergence coincides with the opportunity for migration of fish between the Atlantic and Pacific Ocean basins via the Arctic Ocean with the opening of the Bering Strait, approximately 5 million years ago. The genetic differences may help explain apparent differences in pathogenicity and environmental sensitivity documented for the Atlantic and Pacific forms of *L. salmonis*.

Keywords Salmon lice · *Lepeophtheirus salmonis* · Expressed sequence tags (ESTs) · Mitochondrial genome · 16S rRNA · Cytochrome oxidase subunit I (*COI*) gene

Introduction

The salmon louse, *Lepeophtheirus salmonis*, is an economically important ectoparasite of farmed and wild salmon throughout the northern hemisphere (see reviews by Pike and Wadsworth 1999; Johnson and Fast 2004; Johnson et al. 2004; Boxaspen 2006; Costello 2006). Indirect and direct annual losses due to *L. salmonis* in the global salmonid aquaculture industry are estimated to exceed US\$ 100 million (Johnson et al. 2004). In addition, elevated abundances of sea lice on wild salmon smolts in coastal waters occupied by salmon aquaculture have led to the hypothesis that wild populations of Atlantic salmonids have been negatively impacted by parasites derived from farmed salmon (Costello 2006). Uncertainty concerning the transmission of *L. salmonis* between farmed and wild salmon populations in British Columbia, Canada, has led to considerable research effort and scientific debate.

The development of the parasite includes two nonparasitic nauplii stages that facilitate dispersal in the plankton,

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R. Yazawa · M. Yasuike · J. Leong · K. R. von Schalburg ·
G. A. Cooper · M. Beetz-Sargent · A. Robb · B. F. Koop (✉)
Centre for Biomedical Research, University of Victoria,
P.O. Box 3020 STN CSC, Victoria, BC V8W 3N5, Canada
e-mail: bkoop@uvic.ca

W. S. Davidson
Department of Molecular Biology and Biochemistry,
Simon Fraser University,
8888 University Drive,
Burnaby, BC V5A1S6, Canada

S. R. M. Jones (✉)
Pacific Biological Station, Fisheries and Oceans Canada,
3190 Hammond Bay Road,
Nanaimo, BC V9T 6N7, Canada
e-mail: simon.jones@dfo-mpo.gc.ca

an infective copepodid stage, four chalimus stages that are tethered to the host by a frontal filament, two preadult stages, and one adult stage. Preadults and adults are not tethered and are mobile on the surface of the fish. Following mating, adult females produce eggs that hatch to complete the life cycle (see reviews by Johnson and Fast 2004; Boxaspen 2006; Costello 2006). *L. salmonis* has become an important model for the study of ectoparasitic infestations on salmon. Disease due to *L. salmonis* on Atlantic salmon (*Salmo salar*) and sea trout (*S. trutta*) results from the feeding behavior and the secretion of bioactive compounds by the parasite (Pike and Wadsworth 1999; Dawson et al. 1997; Fast et al. 2007). The parasite feeds on mucus, epidermal cells, and underlying tissues causing physical damage, changes in the composition of blood electrolytes, physiological stress, immune dysfunction, impairment of swimming ability, and possibly death (see reviews by Johnson and Fast 2004; Boxaspen 2006; Costello 2006; Tully and Nolan 2002). However, physiological and immunological studies of *L. salmonis* remain limited (see review by Wagner et al. 2008).

Innate resistance to the salmon louse varies among the various species of salmon and trout (Jones 2001; Johnson and Albright 1992; Fast et al. 2003, 2006). Laboratory studies show that the heaviest infestations and greatest impacts are observed on sea trout (*S. trutta*) and Atlantic salmon (*S. salar*) followed by rainbow trout (*Oncorhynchus mykiss*), chinook (*O. tshawytscha*), and coho salmon (*O. kisutch*; Dawson et al. 1997; Johnson and Albright 1992; Fast et al. 2002). More recently, pink salmon (*O. gorbuscha*) were shown to rapidly reject *L. salmonis* and avoid the clinical consequences of infestation (Jones et al. 2007). Morphological and protein data suggest that the development of an inflammatory reaction, both systemically and at the site of parasite attachment, is a distinguishing feature of *Oncorhynchus* spp. that is not observed in the more susceptible Atlantic salmonids (e.g., *S. salar*; Fast et al. 2002). The kinetics of these inflammatory processes suggests they play a role in parasite rejection.

Oncorhynchus and *Salmo* species have been geographically isolated since the Miocene, approximately 18 to 30 million years ago (Devlin 1993; McKay et al. 1996). In light of differential responses of salmon species to lice, the question arises as to whether Atlantic and Pacific parasites such as *L. salmonis* have coevolved with Atlantic salmon and Pacific salmon, respectively, as distinct populations. Earlier *L. salmonis* microsatellite data based on six loci identified significant differentiation (fixation index = 0.0595) between one Pacific population and Atlantic forms but noted that only 6% of the overall variation was across oceans (Todd et al. 2004). In addition, a study of four mitochondrial genes noted clear differences between samples from a Japanese population and six Atlantic popula-

tions but excluded analysis and reporting of the Japanese data because of reduced length and numbers of sequences (Tjensvoll et al. 2006). In the present study, we examine the mitochondrial genome of the Pacific *L. salmonis* and compare it to the Atlantic form.

Genomic characterization of Atlantic salmon and rainbow trout (Rexroad et al. 2003; Rise et al. 2004; Govoroun et al. 2006; Adzhubei et al. 2007, Wynne et al. 2008) has enabled an expanded capacity for exploring the salmonid response to infectious disease and other environmental impacts (Rise et al. 2004; von Schalburg et al. 2005). In contrast, the present availability of sequence data from fewer than 200 salmon louse genes (GenBank: Nov 2007) limits our ability to measure and characterize parasite responses prior to and during infection. In the present study, we report on an expression sequence tag (EST) analysis of *L. salmonis* collected from salmon in the Pacific Ocean as part of a larger effort to improve our understanding of the coincident expression of host and parasite genes during infection.

Materials and Methods

Salmon Lice Samples

Gravid *L. salmonis* were collected from postharvest Atlantic salmon reared in seawater net pens in the Broughton Archipelago region of coastal British Columbia, Canada. Dissected egg strings were incubated in four 3-L beakers to permit egg hatching as described earlier (Jones et al. 2007). After 8 days of incubation, copepodids comprised approximately 40% of the developing lice, and separate suspensions of copepodids and nauplii were obtained in ice-cold filtered seawater by pipetting from a series of 5-mL samples (total volume sampled was 50 mL). The copepodid and nauplii suspensions were concentrated onto 0.45- μ m nitrocellulose membrane filters. The membranes were immediately placed in separate 1.5-mL capped centrifuge tubes, frozen in liquid nitrogen, and stored at -80°C . The remaining viable copepodids were used to infect salmon. Sixty naive Atlantic salmon (75 g) were maintained in 100-L tanks of filtered seawater and exposed to *L. salmonis* at a rate of approximately 288 copepodids per fish using the methods of Jones et al. (2007). Parasitic developmental stages of *L. salmonis* were obtained from exposed salmon 9 (n =ten fish; copepodid, chalimus I), 19 (n =10 fish, chalimus II, III, IV), and 38 days (n =31 fish; preadult, adult) after exposure. Immediately following removal from the fish, the specimens belonging to each stage were placed separately in 1.5-mL capped tubes and frozen and stored as described above. For the Pacific salmon louse distribution study, 68 samples of *L. salmonis* were obtained from

widely separated locations along the British Columbia coast (Broughton Archipelago: LsBa; Sidney: LsSi; Sooke: LsSo, and Ucluelet: LsUc), Alaska (Port Moller: LsPm; Port of Kodiak: LsPk, and Juneau: LsJu), the mid Bering Sea (180° W, between 55° and 57° N: LsBs) and Japan (Yoichi, Hokkaido: LsJp; Fig. 1).

mRNA Isolation and Construction of cDNA Libraries

Total ribonucleic acid (RNA) was extracted from frozen samples using TRIzol reagent (Invitrogen) and Poly(A)+RNA purified by using Poly(A) Purist™ (Ambion). The non-normalized complementary deoxyribonucleic acid (cDNA) libraries for different developmental stages (copepodid, chalimus I, III, and IV, preadult male, preadult female, adult male, and adult female) were constructed using pBluescript II XR cDNA Library Construction Kits (Stratagene). To obtain enough RNA, particularly from the early stages, several hundreds of individuals were pooled. Poly(A)+RNAs of 2.5 to 5 µg were used for each cDNA library following methods previously described (Rise et al. 2004). A normalized library containing equal amounts of RNA from all eight developmental stages was also constructed (Evrogen).

Sequencing, Sequence Analysis, and Contig Assembly

cDNA libraries were manually arrayed in 384-well microtiter plates, and glycerol stocks of overnight cultures were prepared. Plasmid DNAs were extracted and sequenced on an ABI 3730 DNA analyzer (Applied Biosystems) with M13 Reverse or Forward primers. The resulting ESTs were assembled with CAP3 (Huang and Madan 1999) with

default parameters. The assembled total contigs (clusters+singletons) were annotated using RPS-BLAST or BLASTX comparisons with the Conserved Domain Database (www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=cdd) or SWISSProt (Bairoch and Boeckmann 1992). The best BLAST match (E value threshold of $1e^{-10}$) was used to identify contigs. Contigs not meeting this threshold were annotated as unknown.

Genomic Characterization and Annotation for the Pacific Sea Lice mtDNA

The total genomic DNAs were extracted from an adult male *L. salmonis* collected from Broughton Archipelago (British Columbia, Canada) using the following method: A sample placed in 5% Chelex-100 resin (Sigma) solution (5% Chelex-100 resin, 0.2% sodium dodecyl sulfate in Tris-ethylenediamine tetraacetic acid, with proteinase K [100 µg/mL]) was incubated for 30 min at 55°C, and the proteinase K was then inactivated for 10 min at 90°C. The polymerase chain reaction (PCR) primer sets were designed for 12 fragments of the Pacific salmon louse (Fig. 2 and Supplemental Table 1) based on the EST sequences encoding mitochondrial DNA (mtDNA) genes and the complete mtDNA genomic sequence previously reported by Tjensvoll et al. (2005). PCR amplification was performed using 1.0 µL of extracted total genomic DNA of *L. salmonis* with an initial denaturation step of 2 min at 95°C and then 30 cycles as follows: 30 s of denaturation at 95°C, 30 s of annealing at 55°C, and 3 min of extension at 72°C. PCR products were cloned into pCR2.1 vector (TA Cloning Kit, Invitrogen) with the manufacturer's protocol, and each positive PCR product was sequenced as described above.

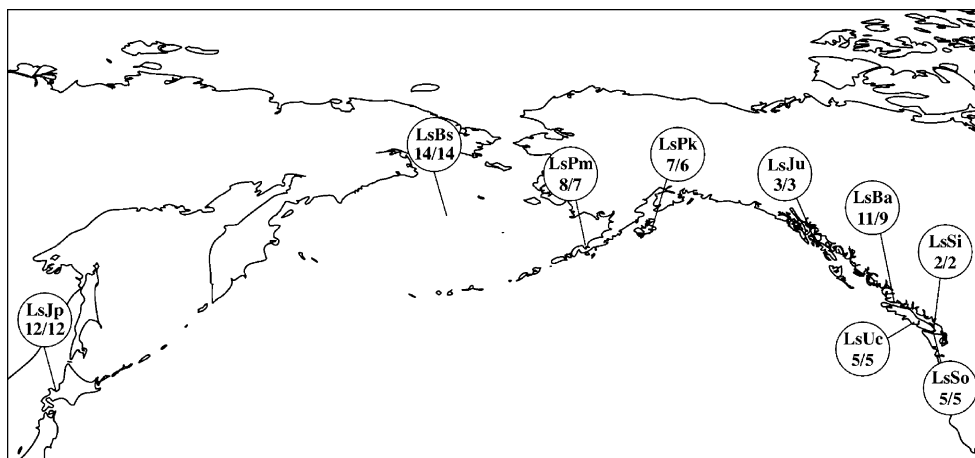


Fig. 1 Map of the northern hemisphere showing the locations Pacific form *L. salmonis* sample site. The sampling locations are shown with sample name (from British Columbia coast, Broughton Archipelago: LsBa; Sidney: LsSi; Sooke: LsSo, and Ucluelet: LsUc; from Bering Sea, midway between the Aleutians and Kamchatka: LsBs; Port

Moller: LsPm; Port of Kodiak: LsPk, and Juneau area: LsJu, and from Japan, Yoichi, Hokkaido: LsJp) and the number of samples sequenced (left: 16S rRNA; right: COI gene) for each location in the circles, respectively

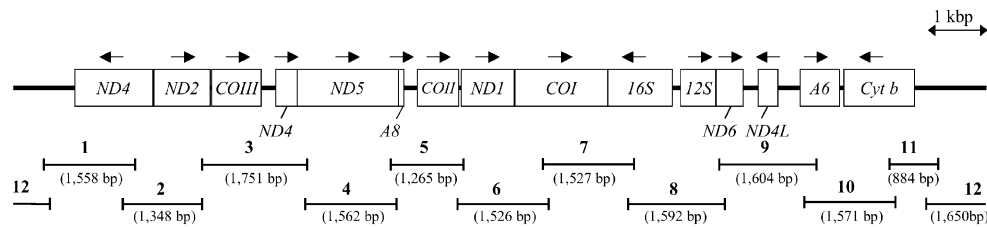


Fig. 2 Genomic organization of the Pacific form *L. salmonis* mtDNA (16,148bp). Boxes represent each mtDNA gene, and transcription directions are shown by arrowheads. *16S* and *12S* refer to 16S and 12S rRNA; *COI*, *COII*, and *COIII* refer to cytochrome oxidase subunit I, II, and III; *Cyt b* refers to cytochrome *b*; *ND1–6* and *ND4L* refer to

NADH dehydrogenase subunits 1–6 and 4L, and *A6* and *A8* refer to ATP synthase subunits 6 and 8, respectively. The bars below the diagram show overlaps of PCR amplicons, and their sizes are indicated in parentheses

The sequenced clones were assembled using PHRED (Ewing and Green 1998), PHRAP (Ewing et al. 1998), and Consed (Gordon et al. 1998) and annotated using Sequin (Altschul et al. 1990). Dotter (Sonnhammer and Durbin 1995) was used to compare the D-loop region sequences of the Atlantic and Pacific forms.

The Screening for Sequence Variation in *COI* and 16S Genes

Total DNA extracts were obtained from fresh or ethanol-fixed samples as described above. The partial gene sequences of the 16S ribosomal RNA (rRNA) and cytochrome oxidase subunit I (*COI*) genes were amplified with the following primer sets: the 16S rRNA, LsPc-16S-F, and LsPc-16S-R; *COI* gene, LsPc-COI-F and LsPc-COI-R. PCR amplification was performed using 1.0 μ L of *L. salmonis* genomic DNA with an initial denaturation step of 5min at 95°C and then 40 cycles as follows: 30s of denaturation at 95°C, 30s of annealing at 55°C, and 2min of extension at 72°C. The PCR products were purified with QIAquick PCR purification kit (Qiagen) and directly sequenced with the internal sequencing primers, respectively. All primers used in this study are shown in Supplemental Table 1.

Phylogenetic Analysis

The partial sequences of the Pacific *L. salmonis* 16S rRNA (total of 67 samples) and *COI* gene (total of 63 samples) were obtained by the PCRs described above (for 16S rRNA: LsBa; 11 samples, LsSi; two samples, LsSo; five samples, LsUc; five samples, LsBs; 14 samples, LsJu; three samples, LsPm; eight samples, LsPk; seven samples, LsJp; 12 samples. For *COI* gene: LsBa; nine samples, LsSi; two samples, LsSo; five samples, LsUc; five samples, LsBs; 14 samples, LsJu; three samples, LsPm; seven samples, LsPk; six samples, LsJp; 12 samples; Fig. 1). The 16S rRNA and *COI* gene sequences that were originally identified by Tjensvoll et al. (2006) were used for the Atlantic form of *L. salmonis* sequences (16S rRNA [GenBank: AY602770-AY602949] and *COI* gene

[GenBank:AY602587-AY602766]). All sequences were trimmed to the same length (16S rRNA; 796bp, *COI* gene; 1,300bp) and aligned using CLUSTALW (Higgins and Sharp 1988). Distance matrices (Kimura two-parameter) and data for the phylogenetic tree were generated by the PHYLIP program package (Felsenstein 1989) using the neighbor-joining and unweighted pair group method with arithmetic mean (UPGMA) methods. For simplifying the phylogenetic trees, identical sequences were grouped into clusters. The phylogenetic trees were generated by NJplot software (Perrine and Gouy 1996).

Results and Discussion

cDNA Libraries and ESTs

Independent cDNA libraries were constructed for copepodids, chalimus I, III, and IV stages, male and female preadult, and male and female adult stages of *L. salmonis*. Several hundreds of individual copepodid and chalimus stage individuals were pooled to obtain sufficient RNA quantities. One hundred to 900 sequence reads were obtained from each library and assembled into contiguous sequences (contigs). An analysis of these contigs showed that more than 30% of the sequences were rRNA gene transcripts indicating very active protein translation. There was also a very high level of transcript redundancy making random sequencing strategies far too inefficient. To obtain the broadest possible representation of genes, equal amounts of messenger RNA from the different life stages listed above were combined, and a normalized cDNA library was constructed. Inserts of 5,760 random clones from this normalized library were sequenced from both the 5' and 3' ends resulting in 11,252 total sequences. In the normalized library, the average contig had 1.33 sequences with the largest contig consisting of ten individual sequence reads. A combined total of 14,994 EST sequences from all of the libraries were assembled into 5,256 unique contigs, of which 1,407 were composed of single sequences, 3,849 composed of two or more sequences, and 1,326 of three or

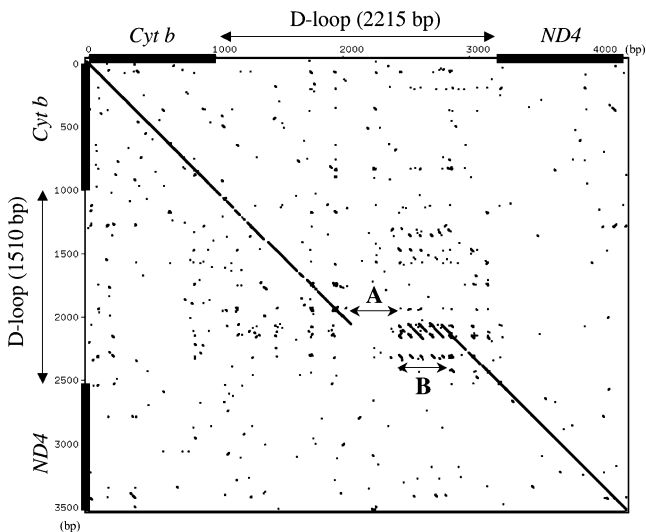


Fig. 3 Dot-matrix analysis of Pacific form D-loop region sequence (horizontal) versus the Atlantic form D-loop region sequence (vertical). Black boxes on the X- and Y-axes represent genes on the mtDNA genome. *Cyt b* and *ND4* refer to the genes cytochrome *b* and NADH dehydrogenase subunit 4, respectively. Region A shows missing region in D-loop region of Atlantic form mtDNA. Region B indicated by arrow shows repeat region in Pacific form mtDNA

more sequences. Contigs were annotated by RPS-BLAST or BLASTX comparisons to known protein domain profiles and protein entries in public databases (Conserved Domain Database; www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=cdd, and SWISS-PROT; Bairoch and Boeckmann 1992). Of the 5,256 contigs, 2,557 matched at least one entry in the databases, and the others remain unidentified. EST sequences are available in GenBank (EX475086–

EX486337) and contigs along with their proposed annotation are available through the cGRASP website (www.uvic.ca/cbr/grasp). The identification of 5,256 unique contigs provides a novel resource with which to study sea louse biology, as well as serving as the basis for a cDNA microarray. Efforts are currently underway to build a sea louse microarray that will complement existing salmonid microarrays (Rise et al. 2004; von Schalburg et al. 2005) to enable profiling of both host and parasite gene expression during infection.

Comparison of Atlantic and Pacific Form *L. salmonis* Genes

A total of 155 of the 5,256 contigs from Pacific *L. salmonis* matched (BLAST E value $< 1e^{-100}$) at least one of the approximately 200 nuclear gene sequences from the Atlantic form of *L. salmonis* available in the public databases. These comparisons showed an average of 96.8% identity over an average of 765bp (data not shown). The importance of a 3.2% difference is difficult to determine without knowledge of gene duplications or establishing natural population variation for each gene, and as contig comparisons include 5' (presumably genic) and 3' (3' untranslated region) sequences, they provide only a very rough estimate of overall sequence similarity. However, nuclear gene sequence comparisons do show clear genetic differences between Atlantic and Pacific forms of *L. salmonis*.

Nineteen of the 5,256 EST contigs were identified as mitochondrial sequences and spanned approximately 80%

Table 1 The summary of nucleotide and protein differences between Pacific and Atlantic form

Genes	In nucleic sequence		In deduced amino acid sequence	
	Identities (bp)	Percent	Identities (amino acid residues)	Percent
12S ribosomal RNA	583/590	98.8	N.A.	
16S ribosomal RNA	1,037/1,070	96.9	N.A.	
Similar to ATPase 8	90/93	96.8	N.A.	
ATP synthase F0 subunit 6	601/654	91.9	208/217	95.9
Cytochrome <i>b</i>	1,098/1,172	93.7	383/389	98.5
Cytochrome <i>c</i> oxidase subunit I	1,430/1,539	92.9	508/512	99.2
Cytochrome <i>c</i> oxidase subunit II	645/690	93.5	229/229	100.0
Cytochrome <i>c</i> oxidase subunit III	759/825	92.0	266/271	98.2
NADH dehydrogenase subunit 1	821/885	92.8	285/294	96.9
NADH dehydrogenase subunit 2	845/930	90.9	293/309	94.8
NADH dehydrogenase subunit 3	327/357	91.6	115/118	97.5
NADH dehydrogenase subunit 4	1,182/1,296	91.2	399/431	92.6
NADH dehydrogenase subunit 4L	313/332	94.3	107/110	97.3
NADH dehydrogenase subunit 5	1,524/1,680	90.7	534/559	95.5
NADH dehydrogenase subunit 6	423/451	93.8	146/150	97.3
Average	–	93.4		97.0

of the complete 14.5-kb mtDNA genome of a previously described Atlantic isolate (GenBank: AY625897; Tjensvoll et al. 2005). The EST contigs for Pacific *L. salmonis* mtDNA genes differed from Atlantic mtDNA by an average of 8%. Similar differences in mitochondrial contigs from the Pacific form were also apparent in comparisons to ATPase subunit 6, *COI*, cytochrome *b*, and 16S rRNA mtDNA gene sequences from 180 Atlantic Ocean isolates (Tjensvoll et al. 2006).

Characterization of the Pacific Form *L. salmonis* mtDNA Genome

To determine whether the divergence observed between EST contigs and existing mitochondrial sequences was due to transcriptional or genomic differences, oligonucleotide primers based on conserved sequences within the EST contigs were designed and used to amplify and sequence the complete mtDNA genome of a single Pacific *L. salmonis* individual. Primers based on the EST contigs were effective in amplifying 12 overlapping segments of genomic mtDNA from which we successfully characterized the complete mtDNA genomic sequence (16,148bp; Fig. 2). Structurally, the Atlantic (Tjensvoll et al. 2005) and Pacific mtDNA genomes are identical in gene organization, direction of transcription, and transfer RNA locations (Fig. 2). However, they differ substantially in the D-loop region. The Pacific form D-loop region has a 350-bp insert with respect to the Atlantic form and differs in the number of repeated 70-bp sequences (Fig. 3). Sequence divergence in the D-loop region, not including repeats, is very high (22.7%). Among mitochondrial genic regions, Pacific and Atlantic forms showed an average of 7.1% difference (combined 13 mtDNA genes and two rRNA DNA; 11,680/12,570bp). Individual gene and protein similarities are shown in Table 1.

Distribution of Pacific Form of *L. salmonis*

To examine the distribution of the Pacific mitochondrial form, we amplified and sequenced the 16S rRNA and *COI* gene from 68 specimens obtained from widely separated locations along the British Columbia coast (Broughton Archipelago, Ucluelet, Sidney, and Sooke), Alaska (Port Moller, Port of Kodiak, and Juneau), the mid Bering Sea (180° W, between 55° and 57° N), and Japan (Yoichi, Hokkaido; Fig. 1). These sequences were compared to the 16S rRNA and *COI* gene sequences previously described from six locations in the Atlantic Ocean (Scotland, Russia, Eastern Canada, and three locations in Norway; Tjensvoll et al. 2006). All Pacific and Atlantic 16S rRNA and *COI* gene sequences were aligned and phylogenetic analyses performed (Fig. 4). All individual sequences of *L. salmonis*

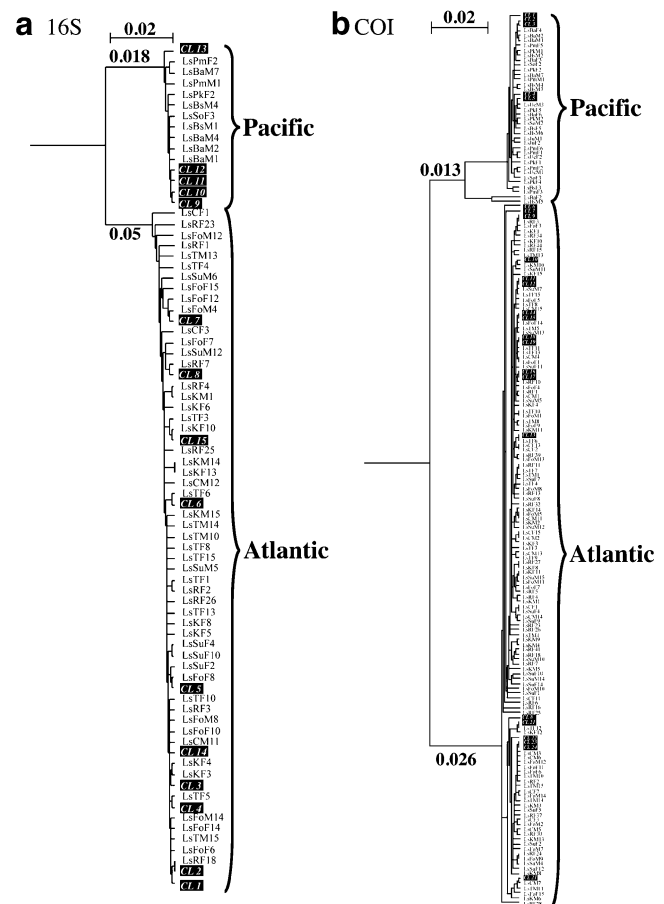


Fig. 4 Molecular phylogeny of Pacific and Atlantic form *L. salmonis* inferred from a UPGMA analysis of nucleotide sequence of **a** 16S rRNA and **b** *COI* mtDNA genes. *CL* refers to the clusters that contain 2–90 identical sequences and are shown by white letters in black boxes. All clusters consist of only Pacific ($n=67, 63$) or Atlantic ($n=180, 180$) isolates. The numbers of sequences in each cluster are described below. For 16S rRNA, CL1: 90 sequences (sq); CL2, 4, 5, 7, 9, 10, 12, and 13: 2 sq; CL3: 13 sq; CL6 and 8: 3 sq; CL11: 11sq. For *COI* gene, CL1–7, 10, 13–20, 23–25, 27, 28, 30, and 31: 2 sq; CL8, 9, 11, and 26: 3 sq; CL12: 4 sq; CL21: 9 sq; CL22: 6 sq; CL29: 12 sq. The partial sequences of *Caligus elongatus* (16S rRNA [GenBank: AY660020] and *COI* gene [GenBank: AY861371]) were used to root the neighbor-joining trees (not shown in this figure). The numbers are branch lengths from UPGMA analysis. The same major branches were obtained from a neighbor-joining analysis. *M* or *F* in the sample names are for the male or female

from the Pacific closely resemble and confirm the Pacific mitochondrial genome sequence and form a single branch, distinct from a branch with representatives from the Atlantic Ocean. Samples from the mid Bering Sea, the closest Pacific location to the Atlantic Ocean, clearly grouped with the Pacific forms. Atlantic and Pacific forms could be uniquely identified by fixed differences at 13 (100%) and 22 (>95%) aligned nucleotide positions in the 16S rRNA sequence (in 796bp) and at 26 (100%) and 67 (>95%) positions in the *COI* gene (in 1,300bp). The large number of fixed differences between Atlantic and Pacific populations strongly supports a long period of isolation.

Table 2 The divergence of 16S rRNA and *COI* genes in and between Pacific and Atlantic forms of *L. salmonis*

	16S rRNA (%)			<i>COI</i> (%)		
	In Pacific	In Atlantic	Atl vs. Pac	In Pacific	In Atlantic	Atl vs. Pac
Average	0.14	0.26	4.16	0.62	0.76	6.06
Max	1.01	1.91	5.48	4.83	2.73	7.89
Min	0.00	0.00	2.95	0.00	0.00	5.06
Stdev	0.00169	0.00267	0.00226	0.00400	0.00770	0.00286

“Average” indicates average for Kimura two-parameter distance (K2P) of all samples in each group. “Max” and “Min” refer to the maximum and minimum values of K2P in each group. “Stdev” refers to standard deviation of K2P values in each group. “Atl vs. Pac” indicates the values for K2P and Stdev between the Atlantic and Pacific isolates

The average difference between the Pacific and Atlantic *L. salmonis* forms is 4.2% at the 16S rRNA locus and 6.1% at the *COI* gene locus (Table 2). These differences are greater than those found between different species of *Drosophila* (Kopp and True 2002; Kastanis et al. 2003). Moreover, the 6.1% divergence between the *COI* gene of the Pacific and Atlantic forms is much greater than the 3% differences between lepidopteran and vertebrate species in the Barcoding programs (Hebert et al. 2003). These results strongly support distinct Pacific and Atlantic forms of *L. salmonis*. Whether these forms constitute separate species will require additional study as Margolis (1958) found Atlantic and Pacific forms morphologically identical.

The 68 Pacific *L. salmonis* samples precluded a robust analysis of population structure in the biogeographical distribution of alleles. The evidence for structure among samples collected from the Atlantic was not found in an earlier study (Tjensvoll et al. 2006). The overall average divergence between individuals within the Pacific population is 0.14% in the 16S rRNA locus and 0.62% in the *COI* gene locus. The intraspecific divergence values from the Pacific samples are consistently lower than those seen in the Atlantic form (Table 2) and indicate lower genetic variability within the Pacific salmon louse population. In general, these values are consistent with intraspecific

variation found in many other species (Hebert et al. 2003). It is interesting to note that the two most distinct individuals were found at distant locations in the Pacific (LsBs: the mid-Bering Sea and LsBa; Broughton Archipelago, British Columbia; Fig. 4). These two Pacific sea lice individuals differ by 1.8% from each other and by 3.4% from the other 61 Pacific isolates at the *COI* locus, which indicates the possibility of population structure. A more extensive sampling from various Pacific locations is required to determine the existence of population structure.

Conclusions

These results suggest that the Pacific salmon (*Oncorhynchus* spp.) and the Pacific salmon louse have coevolved over an extended period of time, in parallel to the coevolution of Atlantic salmonids (*Salmo* spp.) and the Atlantic salmon louse. On the basis of fossil specimens found in Idaho, pink, chum, and sockeye salmon (*Oncorhynchus* spp) have been separated into distinct species for at least 6 million years (Smith 1992). Moreover, estimations of the length of separation between *Salmo* and *Oncorhynchus* range from 18 to 30 million years (Devlin 1993; McKay et al. 1996) and correspond to 4.3% and 14.5% divergences for 16S

Table 3 Ranges of divergence based on Kimura two-parameter distance and crustacean molecular-clock calibrations

	16S rRNA				<i>COI</i>			
	Distance (K2P, %)	Divergence range (Myr)				Distance (K2P, %)	Divergence range (Myr)	
		Ano	Fid	Gra (low)	Gra (high)		Alp	Gra
Mean	4.16	10.92	3.74	6.40	4.73	6.06	2.53	3.65
Max	5.48	14.37	4.37	8.42	6.22	7.89	3.29	4.75
Min	2.95	7.75	2.66	4.54	3.36	5.06	2.11	3.05

The values for “Distance” are the median of the Kimura two-parameter (K2P) distance between the Atlantic and the Pacific form *L. salmonis* in 16S rRNA and *COI* genes. Rates of molecular evolution used for the 16S rRNA gene include 0.38% K2P/million year (Myr) for anomurans: Ano (Cunningham et al. 1992); 0.90% K2P/Myr for fiddler crabs: Fid (Sturmbauer et al. 1996), and 0.65 (low)–0.88% (high) K2P/Myr obtained from grapsid crabs: Gra (Schubart et al. 1998). *COI* calibrations were calculated with 2.4% K2P/Myr obtained from alpheid shrimps: Alp (Knowlton et al. 1993), and 1.66% K2P/Myr obtained from grapsid crabs: Gra (Schubart et al. 1998)

rRNA and *COI* genes. Atlantic and Pacific salmon lice differ by 4.2% and 6.1% for 16S rRNA and *COI* genes (Table 2), respectively. These values are not easily comparable since rates of evolution vary tremendously among taxa. However, the divergence between Pacific and Atlantic salmon lice is substantially lower than between Pacific and Atlantic salmon hosts even though the evolutionary rates of mitochondrial genes in copepods are thought to be faster than for vertebrates. Sequence divergences, in conjunction with a molecular clock calibrated by previous copepod studies (Rocha-Olivares et al. 2001), suggest that the Pacific and Atlantic forms of *L. salmonis* diverged 2.5 to 11 million years ago (Table 3), roughly corresponding with the first continuous connection between the Pacific and Atlantic oceans through the Bering Strait (5 million years ago; Marincovich and Gladenkov 1999). These estimates are very broad, but it appears that the Atlantic and Pacific forms of the salmon louse arose well after the separation of Atlantic and Pacific salmonids and about the time of the opening of the Bering Strait.

The level of separation between Pacific and Atlantic salmon lice mitochondrial genomes, the estimated time of Pacific and Atlantic salmon louse separation coinciding with the opening of the Bering Strait, and the reduced overall variation found within the 16S rRNA and *COI* genes from Pacific salmon lice all support an Atlantic Ocean origin of *L. salmonis* followed by a limited introduction into the Pacific Ocean coincident with the opening of the Bering Strait, approximately 5 million years ago. Parallel coevolution of salmon lice on their respective hosts in the Pacific and Atlantic Oceans has resulted in nuclear and mitochondrial genetic changes that may help to explain apparent phenotypic differences observed between these forms. In recent work using Scottish *L. salmonis* specimens, Bricknell et al. (2006) provided evidence of reduced tolerance of copepodids for low salinity in comparison to similar studies using lice specimens from British Columbia (Johnson and Albright 1992). Similarly, Saksida et al. (2007) documented a lower incidence of disease and a reduced need to treat farmed *S. salar* for *L. salmonis* in British Columbia compared with farmed *S. salar* in Scotland and Norway. More research is required to test these hypotheses. The high level of sequence divergence between the Pacific and Atlantic *L. salmonis* indicates that a taxonomic revision of these forms may be warranted.

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