

Table 1: Sequences of primers used for characterization of the labial gland enzymes. Primers were used in rapid amplification of cDNA ends (RACE), confirming the open reading frame (ORF), synthesis of RNA probes for in situ hybridization (IS) and double stranded RNA for RNA interference (RNAi), and for real time RT-PCR both target genes (qPCR) and reference genes (qPCR ref). Gene name abbreviations: Lepeophtheirus salmonis (Ls), labial gland protein (LGP), labial gland astacin (LGA), labial gland serine protease 1 (LGSP1), labial gland apyrase (LsLGAp1), Trypsin 1 (Trp1), elongation factor (EF), adenine nucleotide translocator (ADT), Salmo salar (Ss), interleukin (IL), matrix metalloproteinase 13 (MMP13), tumor necrosis factor alpha (TNF α), non-spesific cytotoxic cell reseptor P1 (NCCRP1), interferon gamma (IFN γ), T-cell reseptor beta (TCR β), cluster of differentiation (CD), major histocompatibility complex II (MHC2), immunoglobulin (Ig), tripartatiae motif-containing protein 16 (TRIM16).

Gene	Forward	Reverse	Usage
LsLGP1L	AAACAATGGCTATAAAATGAATAATCCCATT	GAATAACCTCGTCATTCAAAGATAACACCCT	qPCR
LsLGP1	AAAACGAAGAGACAGTTGTCCTCAAATT	CGTCATTCAAAGATAACATCGTTTTCTC	qPCR
LsLGP2	GCTCTCCTTAGTATCTTATTGGGCTCC	ACACGTAAGCACAGCTGTGGATGC	qPCR
LsLGP3	TCAAGAAGGATTGAAAAGTTGACGAATG	TGCCGATTGTTCCATAATTCCGTT	qPCR
LsLGP4	TGTTTTCAAGATCGACTCTTCCAATGG	AGCACACCTTCTTTGCAAGTTCCATA	qPCR
LsLGSP1	GGGACAAACCTCGGTAGCGAT	CGCACATTGCTTATCCATGGTGGTTCCCA	RACE
LsLGSP1	TCAGTAAAAGCTGTTAGGTGAACG	ATTCCACATATGCCACAAGGAT	ORF
LsLGSP1	GGGACAAACCTCGGTAGCGAT	CCTGGTCGCACTTAACCTCATGCC	IS/RNAi
LsLGSP1	TCTTAATACTGTTATGCCTGGAGTCTTGG	TGCATTGGGATCAATTTCAGCAATT	qPCR
LsLGA1	TCCATTATAACAAAATCAATGCAAGACTAA	ATCTTGATAAATTGGTAGTCTGCATTGA	RACE
LsLGA1	GGCAAGTCCACAGCTGAGCATTGCG	TCTACAAGTTAACATTACATCCTTTGTAATGGC	ORF
LsLGA1	GACTTGGATGTGATTAAATC	GCATAATAGATCCATAGTCA	IS/RNAi
LsLGA1	CTCACTTCCATGATGTCTATTCAAGCG	CCTTGAAGCCATTTCCTGTCTGTGA	qPCR
LsLGA2	ATTAACAAGTTATTGCGAAAGCCGCCAGA	CATTCTCCCCACAAAGATGGCCAATTGG	RACE
LsLGA2	CAAGCGTCGTATTGACCGTGGCA	TCCACACGTTAACATTACAACATTGTAATTTC	ORF
LsLGA2	GAAGTGAGTGTGATTAAATC	GCATGATGGATCCATAGTCA	IS/RNAi
LsLGA2	CTCCTGACTCTCATGATGTCTATTCAAGTG	TGGAAGCCATTTCCTGCCTGTGA	qPCR
LsLGA3	CAAGTTATTGCAAAGCAATCCATCAGTCA	CATTCTCCCCACAAAGATGGCCAATTGG	RACE
LsLGA3	CGCCATTCCGCAGCTGAGCATTGCA	TCTACAAGTTAACATTACAGCTTGTAACTCC	ORF
LsLGA3	GATGTGGATGTGATTAAATC	GCATGATAGATCCATAGTCA	IS/RNAi
LsLGA3	CCTGACTTTCCAGATGTCTATTCAAGCG	ATTCCATTGAAGCAATTTCCTGTCTATGG	qPCR
LsLGA4	TTTATACAAAAAAATTGCGAAAGCCATCA	AACCTTACAACCATTTGATAGATTGATG	RACE
LsLGA4	CAAAAGTCCACAGCTGAACATTGCA	TCTACAAGTTAACATTACAGTCTTGTAAATGGT	ORF
LsLGA4	GATGTGGATGTGATTAAATC	GCATGATAGATCCATAGTCA	IS/RNAi
LsLGA4	CCTGACTTTCCATGATGTTATTCAAGCA	AAATTCTGAAGCCATTTCCTGTTGTAA	qPCR
LsLGA5	TGATCGATGTTGGCTGGATACCG	CCGTTGATTAGTTCGTGGGATCCAT	RACE
LsLGA5	CCTTGGATTGCAATATGACTACC	AGACATACTTCGATGGCCTTCAT	RACEn
LsLGA5	CATCAGTTGAATAACAAAAATGATGA	AATTACAATGACTTCAGTTATCACAAA	ORF
LsLGA5	CGGCAAGACCTGCTTAGGAAT	TTTGTGCTCATCGTAGGGTATCCC	IS/RNAi

Gene	Forward	Reverse	Usage
LsLGA6	GCACCTACTACCACAACCCCAGCAT	GCCAATATTAGCGAAGCATTCCGTCT	RACE
LsLGA6	AGCGACTATAAGCATCTTGTGGAA	ATGTTGATCCGAAATGTGCTTA	RACEn
LsLGA6	TCATTAGGTTTTAAAGATGGCAA	TTTCTTGATTGCTCCAACGAA	ORF
LsLGA6	ATGCGTCACATATGTTGAAAGAAC	CCACAAAGATGCTTATAGTCGCTACA	IS/RNAi
LsLGA6	GAATGAACCTGCTGCTATTCCA	TGAAAAGTATCGAAGTCCGTAGTGC	qPCR
LsLGA7	GGTCATGAGCAACACGACCTGACCG	CTGGGTCTGCATTGGGTGGTACGAT	RACE
LsLGA7		TAATACCACTATCGTACCAACCCCAATGCAG	RACEn
LsLGA7	AGCATATTCCGAACTGAGCCAT	CCACAAAGTGGCTTACAATTCTC	ORF
LsLGA7	TTCCGAACTGAGCCATGCTCAA	CGAATTCCAAACGGTACAGGTCTCAG	IS/RNAi
LsLGA7	ATGGTGCCCCTTAAATATACTAGC	GTCAAAGCAATCTGAAGTGGTTGTA	qPCR
LsLGA8	CCAAAAATGTGATCCATCCAGA	CCATGTTCCCCTCCTGTT	IS/RNAi
LsLGSP1	TCTTAATACTGTTATGCCTGGGAGTCTTGG	CGCACATTGCTTATCCATGGTGGTTCCA	RACE
LsLGSP1	TCAGTAAAAGCTGTTGAGGTGAACG	ATTCACATATGCCACAAGGAT	ORF
LsLGSP1	GGGACAAACCTCGGTCAAGCGAT	CCTGGTCGCACTTAACTTCATGCC	IS/RNAi
LsLGSP1	TCTTAATACTGTTATGCCTGGGAGTCTTGG	TGCATTGGGATCAATTTCAGCAATT	qPCR
LsLGAp1	TGTGAAACTCTGGACGCTCACTCATCAA	TGTGTCGTCGGGTTGTGAGGTGTACTCAG	RACE
LsLGAp1	ATAGTCACGCCCTTCAGCTA	GATCCGGCCATCCCCTTAG	ORF
LsLGSP1	TCTTAATACTGTTATGCCTGGGAGTCTTGG	TGCATTGGGATCAATTTCAGCAATT	qPCR
LsLGAp1	TGTGAAACTCTGGACGCTCACTCATCAA	TGTGTCGTCGGGTTGTGAGGTGTACTCAG	RACE
LsLGAp1	ATAGTCACGCCCTTCAGCTA	GATCCGGCCATCCCCTTAG	ORF
LsLGAp1	AAATGGAAGGCAGTGACGCCAT	GACCTCCGACAACGACATCA	IS/RNAi
LsLGAp1	ATAGTCACGCCCTTCAGCTA	GCATCTCGAAGAAGAACGCG	qPCR
LsTrypt1	CACCTTCTCCAGTTCTAAAGCTGTT	AGATCATGGTCTCATCAATAGATCCA	qPCR
LsEF1 α	GGTCGACAGACGTACTGGTAAATCC	TGCGGCCTTGGTGGTGGTC	qPCRref (ref)
LsADT3	CTGGAGAGGAATTGGCTAACGTG	GACCCTGGACACCGTCAGACTTCA	qPCRref (ref)
SsIL1 β	GCTGGAGAGTGTGGAGAAGA	TGCTCCCTCCTGCTCGTAG	qPCR
SsIL8	GCATCAGAATGTCAGCCAGCC	ACGCCTCTCAGACTCATCCC	qPCR
SsIL6	ACCAACAGTTGTGGAGGAGTTCCAGAAC	CCTGCAGACATGCCTCCTGTTG	qPCR
SsMMP13	ACTCTTGCCAATATGCCACCCA	TGGGCCCTCGTTGAACGCA	qPCR
SsTNF α	CACTGCCACCAAGAGCCAAG	CGCCAGTTGTACCGCATACC	qPCR
SsNCCR1P1	AATCCTGCGCCTCACGGTGTGAGTC	GCGAGGAGGTCTTCTGGTGGAAAC	qPCR
SsIL10	ATGAGGCTAATGACGAGCTGGAGA	GGTGTAGAATGCCTCGTCCAACA	qPCR
SsIFN γ	ATGGATGTGTTATCAAGGGCTGTGATGTG	CAGCTGGCCTTGGAGATCTTATAGTGGAC	qPCR
SsIL4/13a	CGTACCGGCAGCATAAAATCACCATCC	CCTTGCATTTGTGGTGGTCCA	qPCR
SsMHCII	GGACGTGAGGTGAAGTCTGATGTGACC	CTGATGTGCTCCACCATGCAGGA	qPCR
SsIgM	TGAGGAGAACTGTGGCTACACT	TGTTAATGACCACTGAATGTGCAT	qPCR
SsIgT	GGTGGTCATGGACGTACTATT	CCTGTGCAGGCTCATATCTT	qPCR
SsIgD	CACCAGGAGGAAAGTTGGCATCA	CCCCAAGGAGCTCTGGTTGGA	qPCR

Gene	Forward	Reverse	Usage
SsCD4	GAGTACACCTGCGCTGTGGAAT	GGTTGACCTCCTGACCTACAAAGG	qPCR
SsCD8α	TAGAGTGCAAGACAACGCTGGAATGGA	TCTCGAGCCTTTGAAAGCCTTCAG	qPCR
SsEF1α	CACCACCGGCCATCTGATCTACAA	TCAGCAGCCTCCTCTCGAACTTC	qPCR (ref)
SsTRIM16	TTACTGTAGGAGCTGTATTGAGGGCTGCTG	TTCTCCACCAGCTCAGCCAACATG	qPCR (ref)

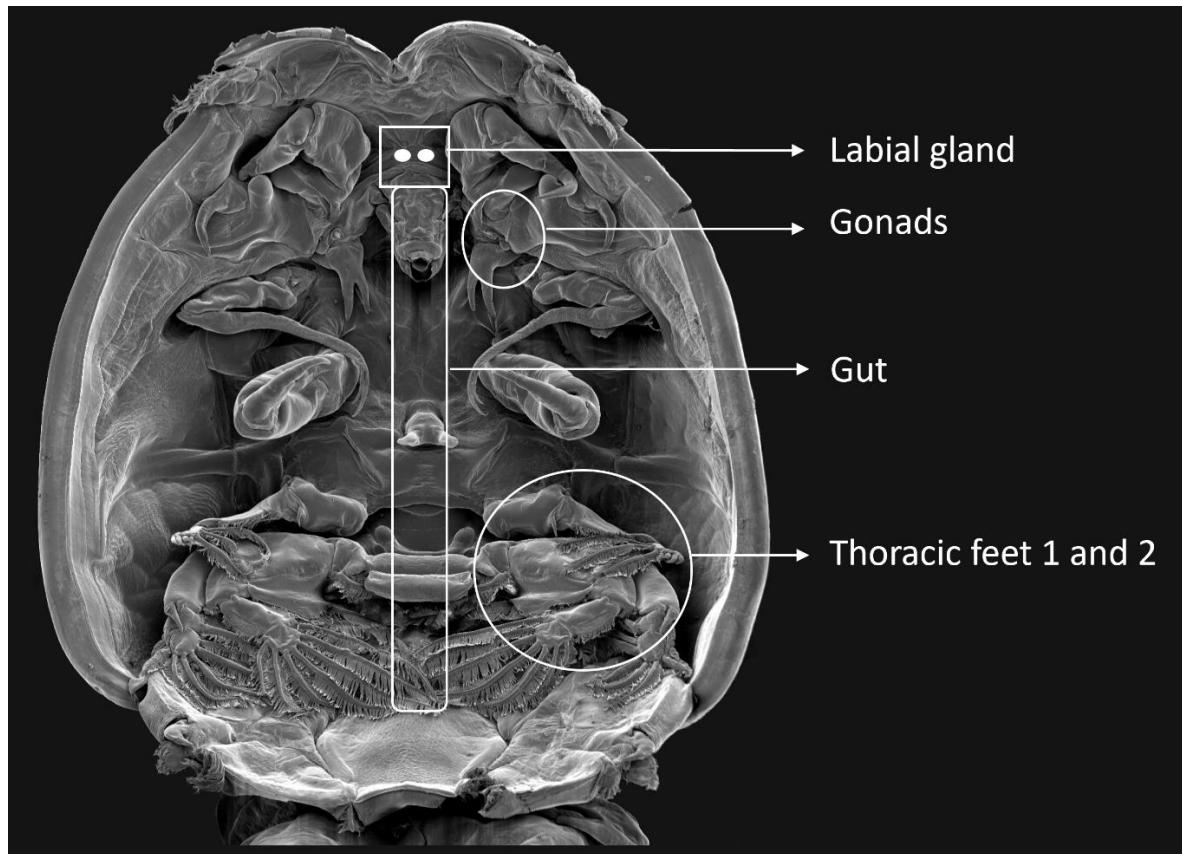


Figure 1. An overview of where the sample high in labial gland were dissected from, and the other areas used to compare the transcriptome of the labial gland sample to enable the identification of labial gland expressed genes. If the genes were high in the labial gland sample, but not in gut, gonads and the thoracic feet that holds all the other gland types identified in the lice, the gene were analyzed further if it was mainly expressed in parasitic lice stages.