

# Molecular characterization and expression analysis of Lily-type lectin (*SmLTL*) in turbot *Scophthalmus maximus*, and its response to *Vibrio anguillarum*\*

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**Abstract** A full-length lily-type lectin (*SmLTL*) was identified from turbot (*Scophthalmus maximus*) in this study. By searching database for protein identification and function prediction, *SmLTL* were confirmed. The full-length cDNA of *SmLTL* is composed of 569 bp and contains a 339 bp ORF that encodes 112 amino acid residues. The *SmLTL* peptide is characterized by a specific  $\beta$ -prism architecture and contains three mannose binding sites in a three-fold internal repeat between amino acids 30–99; two of the repeats share the classical mannose binding domain (QxDxNxVxY) while the third binding site was similar to other fish-specific binding motifs (TxTxGxRxV). The primary, secondary, and tertiary structures of *SmLTL* were predicted and analyzed, indicating that the *SmLTL* protein was hydrophilic, contained 5.36%  $\alpha$ -helices, 39.29% extended strands, 16.07%  $\beta$ -folds, and 39.29% random coils, and three  $\beta$ -folds. Quantitative real-time polymerase chain reaction (qPCR) analysis revealed that the *SmLTL* mRNA was abundantly expressed in skin, gill, and intestine. Low levels of *SmLTL* expression were observed in other tissues. The expression of *SmLTL* in gill, skin and intestine increased at mRNA level after stimulation of *Vibrio anguillarum*, our results suggest that *SmLTL* serve as the first line of defence against microbial infections and play a pivotal role in the innate mucosal immune system. The current study indicates that *SmLTL* is a member of the lily-type lectin family and the information reported here will provide an important foundation for future research on the role of this protein.

**Keyword:** lily-type lectin; full-length of cDNA; quantitative real-time polymerase chain reaction; protein structure; *Vibrio anguillarum*

## 1 INTRODUCTION

The skin of fishes is covered with mucus (Suzuki et al., 2003). The mucosal layer is the first line of defense against the invasion of foreign substances and pathogenic microorganisms, and serves as a main organ of defense (Suzuki et al., 2003). The main constituent of this barrier is a mucous gel that forms a layer which covers the epithelial cells (Van der Marel et al., 2010). The mucous layer is secreted by various

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epidermal or epithelial mucus cells, such as goblet cells (Shephard, 1994; Spitzer and Koch, 1998). The skin mucus layer is mainly composed of water and mucins, which are glycoproteins that contain high molecular weight oligosaccharides. Skin mucus is involved in fish respiration, osmoregulation, reproduction, locomotion, defense against microbial infection, disease resistance, excretion, and communication (Shephard, 1994; Khong et al., 2009). One of the most interesting functions of fish mucus is related to its role in the immune response and disease resistance (Guardiola et al., 2014); however, further characterization of this role is needed.

Lectins are not enzymes but able to bind to carbohydrate, are present in the mucus (Ingram, 1980; Alexander and Ingram, 1992), and are capable of defending against pathogens (Suzuki et al., 2003). The biological effects of lectins primarily occur via the binding of the active site of the lectin chain to carbohydrates (Ke et al., 2005). Fish lectins are mediators of non-self recognition in a variety of biological processes. Specifically, fish lectins are involved in the identification and stimulation of pathogen uptake by phagocytes, the facilitation of innate complement-mediated cell lysis, and the enhancement of natural killer cell activity (Hoffmann et al., 1999; Sharon and Lis, 2004; Kim et al., 2011). There are already several researches of lectin in different fishes, including *Conger myriaster* (Kamiya et al., 1988; Shiomi et al., 1989), *Repomucenus richardsonii* (Shiomi et al., 1990), *Misgurnus anguillicaudatus* (Goto-Nance et al., 1995), *Genypterus capensis* (Toda et al., 1996), *Anguilla japonica* (Tasumi et al., 2002), and *Channa striata* (Arasu et al., 2013).

Based on their distinct structures and functions, lectins have been classified as galectins, C-type lectins, lily-type lectins, and rhamnose-binding lectins (Kim et al., 2011; Suzuki et al., 2003). Pufflectin, a mannose-specific lectin purified from the skin mucus of pufffish by Tsutsui (2003), was named lily-type lectin (LTL). While LTL shares no significant sequence similarity with any known animal lectins, it surprisingly shares sequence homology with mannose-binding lectins of monocotyledonous plants (Suzuki et al., 2003) such as *Galanthus nivalis* (Van Damme et al., 1991), *Clivia miniata* (Van Damme et al., 1994), *Allium prorrhum* (Van Damme et al., 1993), and *Allium sativum* (Smeets et al., 1997). The lectins of these monocotyledonous plants all contain a specific mannose-binding domain (QxDxNxVxY). Interestingly, the amino acid sequence

of pufflectin contains two of these characteristic motifs (Tsutsui et al., 2003). *Channa striata* lily-type lectin (CsLTL-1) was demonstrated to contain two QxDxNxVxY mannose binding sites (Abirami Arasu, 2013). Other studies have also reported the existence of QxDxNxVxY mannose binding sites in fish lectins (Chandra, 1999; Tsutsui, 2003; Tsutsui et al., 2006). These previous reports provide an important basis for the study of *Scophthalmus maximus* lily-type lectin (*SmLTL*).

Turbot (*Scophthalmus maximus*) is an important farmed fish species with high commercial value in northern China and Europe. Since turbot was introduced to China in 1992, breeding scope has been constantly expanding (Huang et al., 2011). One of the most important factors influencing turbot growth is the variety of pathogens (fungi, bacteria, viruses, and parasites) which cause adverse developmental effects and high rates of mortality in this species. Vibriosis is one of the most disturbing bacterial diseases in turbot aquaculture, which have caused great damage and economic loss in aquaculture production among the world (Saulnier, 2000). *Vibrio anguillarum* is one of the major pathogens causing vibriosis (Toranzo, 1997). The anti-*V. anguillarum* response deserves more attention in order to develop more effective methods of preventing *V. anguillarum*. In order to efficiently manage disease and provide theory support for the enhancement of aquaculture production, immune mechanism should be thoroughly studied in turbot. Previous proteomic studies in this laboratory revealed that *SmLTL* protein expression was significantly changed following high temperature stress in turbot (Ma et al., 2013). In this study, the corresponding proteome maps were constructed by two-dimensional gel electrophoresis (2-DE), from which the peptide mass map with matrix-assisted laser desorption / ionization tandem time-of-flight (MALDI-TOF-TOF) was obtained, and *SmLTL* protein was identified by database retrieval. The partial protein sequence of *SmLTL* was also identified by mass spectrometry analysis. The full-length sequence, expression, and structure prediction analysis of *SmLTL* has not been previously published. In the current study, *SmLTL* was cloned, spatially analyzed, and its tertiary structure predicted for the first time. Furthermore, in turbot (*Scophthalmus maximus*) aquaculture, vibriosis is one of the most disturbing bacterial diseases which have caused great damage and economic loss in aquaculture production among the world (Saulnier, 2000). *Vibrio anguillarum*

**Table 1 Primers used in this study**

Primer name	Sequence (5'→3')
L-R-S-5	GACACAAGGACCCGCCACAAGTGAATC
L-R-AS-297	CGCATCCTTTGATTGGCTTGGTTTGAG
L-YGDL-S-117	CGAGTTCAAAGCCATCTTCC
L-YGDL-AS-117	TTATCCGGTTGCAGAAGGAC
β-actin-F	GTAGGTGATGAAGCCCAGAGC
β-actin-R	CTGGGTCATCTTCTCCCTGT
L-QC-S	CTTCTCAGACACAAGGACCCGCC
L-QC-AS	TCTTTATTTCAATCTCAGAGCAAACCGT

is one of the major pathogens causing vibriosis (Toranzo et al., 1997). Antibiotics are used as traditional strategy for fish disease, while it could due to the development and spread of antibiotic resistant pathogens which would have negative impacts on environment and human health (Chen et al., 2016). So we need an efficient and safe method to solve this problem. Recently many innate immune actors in turbot have been characterized, such as Stomatin-like protein2 (Chi, 2016), chemokines (Meng et al., 2013; Chen, 2015), MyD88 (Lin et al., 2015) and lysozyme (Gao et al., 2016). *SmLTL* was one kind of innate immune actors, could be one of the most important players on the mucus for host protection.

## 2 MATERIAL AND METHOD

### 2.1 Fish

Healthy turbot (90±10.2) g were obtained from the Tianyuan Fisheries Co. Ltd. (Yantai, China). Tissue samples, including head-kidney, kidney, liver, spleen, intestine, muscle, gill, and skin, were dissected from euthanized fish and immediately frozen in RNA holder (Tiangen Biotech Co. Ltd., Beijing, China) and stored at -80°C until use.

### 2.2 Cloning and sequencing of *SmLTL*

According to the *SmLTL* protein (Ma et al., 2013), we designed primers (L-R-S-5 and L-R-AS-297) for rapid-amplification of cDNA ends (RACE) (Table 1). Total RNA was isolated from fish skin using an RNAPrep Tissue Kit (Tiangen). The first-strand cDNA was synthesized from total skin RNA using the SMART RACE cDNA amplification kit (Clontech, Mountain View, CA, USA) for 3'-RACE with primers L-R-S-5 and L-R-AS-297. The cDNA was stored at -20°C prior to further analysis. The polymerase chain reactions (PCR) consisted of denaturation at 94°C for 2 min, 35 cycles of amplification (94°C for 30s, 63°C

for 30 s, 72°C for 30 s), and a final extension at 72°C for 2 min. The same procedures were followed for cDNA production for 5'-RACE. All samples were analyzed in triplicate. Cloning and sequencing was performed by Sangon Biotech Co. Ltd. (Shanghai, China) after agarose gel electrophoresis separation and recovery of products by TIANGel Midi Purification Kit (Tiangen). Lasergene Seqman software (DNASTAR, Madison, WI, USA) was used for sequence assembly of the full-length sequence from the 5' and 3' terminals. With primer L-QC-S and L-QC-AS (Table 1) to PCR, after agarose gel electrophoresis detection and recovery agarose gel (Tiangen), cloned and sequenced by Sangon Biotech Co., Ltd. of China to confirm the full-length sequence.

### 2.3 Bioinformatic analysis of *SmLTL*

The full-length *SmLTL* sequence was compared with other sequences available in the NCBI database (<http://blast.ncbi.nlm.nih.gov/Blast>) and the similarities were analyzed. The open reading frame (ORF) and amino acid sequence of *SmLTL* was obtained using NCBI. The hydrophilicity of the *SmLTL* protein was analyzed using ProtScale (<http://web.expasy.org/protscale/>). Sequence identity, similarity and gap percentages were calculated using the FASTA program (<http://fasta.bioch.virginia.edu/fastawww2/fastawww.cgi>). Signal peptide analysis was performed using SignalP (<http://www.cbs.dtu.dk>). The domains and motifs were analyzed using ProtScale (<http://web.expasy.org/protscale/>). Secondary structure was predicted and analyzed using Jpred4 (<http://www.compbio.dundee.ac.uk/jpred/>). The deduced amino acid sequences were submitted to multiple alignment using DNAMAN version 8.0 (Lynnon Biosoft, San Ramon, CA, USA). A phylogenetic tree was constructed using the Neighbor Joining method, considering 1 000 bootstrap hits in DNAMAN. Protein tertiary structure was predicted and inspected using PDBsum Generate (<http://www.ebi.ac.uk/thornton-srv/databases/cgi-bin/pdbsum/GetPage.pl?pdbcode=index.html>). The predicted protein model was checked using PROCHECK (<http://www.ebi.ac.uk/thornton-srv/software/PROCHECK/>).

### 2.4 Tissue-specific expression of *SmLTL*

Total RNA was isolated using an RNAPrep Tissue Kit followed by cDNA synthesis using 0.05–5 µg of total RNA. Total RNA was mixed with random primers and RNase-free d H<sub>2</sub>O, heated to 65°C for 5 min, placed on ice for 5 min, followed by the

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58      ACATGGGGGCCCCCTCTTCAGCTGCTTCTCAGACACAAGGACCCGCCACAACCTGAATC
118     ATGAACAGAACTCCCATCAGCACGGACCGAGAGCTGCGTAAGGGAGAGTTCTCATGAGT
      M N R N S I S T D Q E L R K G E F L M S
178     GTGAATGGCGAGTTCAAAGCCATCTTCCAGGATGACGGCAACTTTGTCTATCTACAAATGG
      V N G E F K A I F Q D D G N F V I Y K W
238     TCTCCCATTTGGGATACTAAGACATGTGGGAAAAACCCATTTGAGTCCTTCTGCAACCG
      S P I W D T K T C G K N P F R V L L Q P
298     GATAACAACCTGGTTATGTACGACAAATTTGTCTAAACAGTTTGGGCCACTGGCACCAC
      D N N L V M Y D K L S K P V W A T G T H
358     TCAAACCAAGCCAATCAAAGGATGCGCCTGACCTTGACTGATGGAGGTCGGCTGGTTCTT
      S N Q A N Q R M R L T L T D G G R L V L
418     GATAAAGATGGAGGTGAAATTTGGGGTCTGGAGGATAATCTTAGCAATGAAAAGTGCTC
      D K D G G E I W G A G G *
478     TATAAATGAAATTTATTATCATTATTATTACTAAATAGATTCCAGAATTCAAACAGAATT
538     CATGATGTGGCAACACACTGTACACGGTTTGCTCTGAGATTGAAATAAGAGCTCTGATG
569     TTTCGAAAAAAAAAAAAAAAAAAAAAAAAA

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**Fig.1 The complete cDNA and deduced amino acid sequence of *SmLTL***

The three specific mannose binding motifs are highlighted in grey.

addition of 2x TS Reaction Mix and RI Enzyme Mix (TransGen Biotech). The mixture was then incubated at 42°C for 30 min and then heated to 85°C for 5 min. The cDNA was stored at -20°C prior to further analysis. Expression analysis of *SmLTL* was conducted using qPCR) with the L-YGDL-S-117 / L-YGDL-AS-117 primers using a SYBR Premix Ex Taq Kit (TaKaRa, Dalian, China) according to the manufacturer's instructions.  $\beta$ -actin was amplified with specific F and R primers (Table 1) for use as a reference gene. The PCR consisted of denaturation at 94°C for 2 min, 35 cycles of amplification (94°C for 30 s, 63°C for 30 s, 72°C for 30 s), and a final extension at 72°C for 2 min. The reactions were performed using an ABI 7500 Real-time Detection System (Applied Biosystems, Foster City, CA, USA). All samples were analyzed in triplicate.

## 2.5 Bacterial challenge

*V. anguillarum* was conserved in our laboratory in Yellow Sea Fisheries Research Institute, Chinese Academy of Fishery Sciences. *V. anguillarum* challenge was carried out as previously reported (Ma et al., 2014). *V. anguillarum* were inoculated on the sterilized TSB, 28°C incubated for about 24 h and were harvested in the logarithmic phase of growth, which was monitored by the optical density assay. *V. anguillarum* cells were washed, resuspended, and diluted to  $10^9$  CFU/mL in sterile PBS. Fish were challenged by intraperitoneal injection with nine concentrations live *V. anguillarum* (dose=experimental fish body weight (g)×1  $\mu$ L/g bacteria) per fish, and

PBS alone was used as a control. Each group had five fish in each of three replicates (i.e. five per tank and 15 in total). At 0, 2, 4, 6, 8, 16, 24 and 24 h post infection (hpi), the gill, intestine and skin were collected and preserved at -80°C until subsequent use. The method of tissue-specific expression of *V. anguillarum* challenge reference 2.4.

## 2.6 Statistical analysis

The data were expressed as mean±SD and subjected to ANOVA (two-way analysis of variance) to determine differences among treatments. The differences were considered as significant at  $P<0.05$ . All the Statistical analysis was performed using SPSS V 19.0 for windows.

## 3 RESULT

### 3.1 Cloning and sequencing of turbot *SmLTL* cDNA

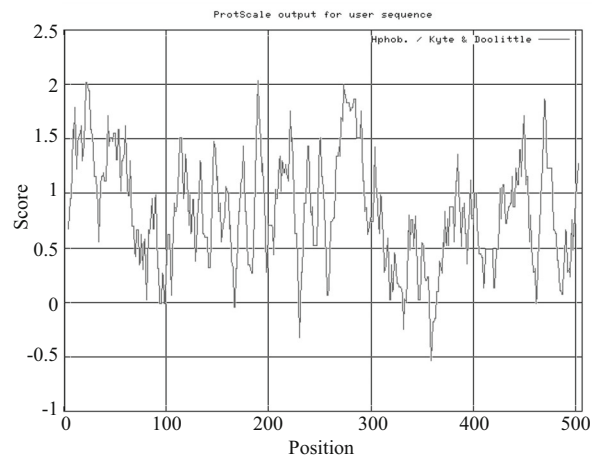
The full-length *SmLTL* cDNA was 569 bp long, included 506 bp coding sequence (CDS) (GenBank accession No. KU199003) and contained an ORF of 339 bp that encoded 112 amino acids (Fig.1). The predicted *SmLTL* peptide has a theoretical molecular mass of 12.652 3 kDa, an isoelectric point (pI) of 7.86, fat factor of 71.34, and an average hydrophilicity of -0.517. (Fig.2).

### 3.2 Structure prediction and analysis of *SmLTL* protein

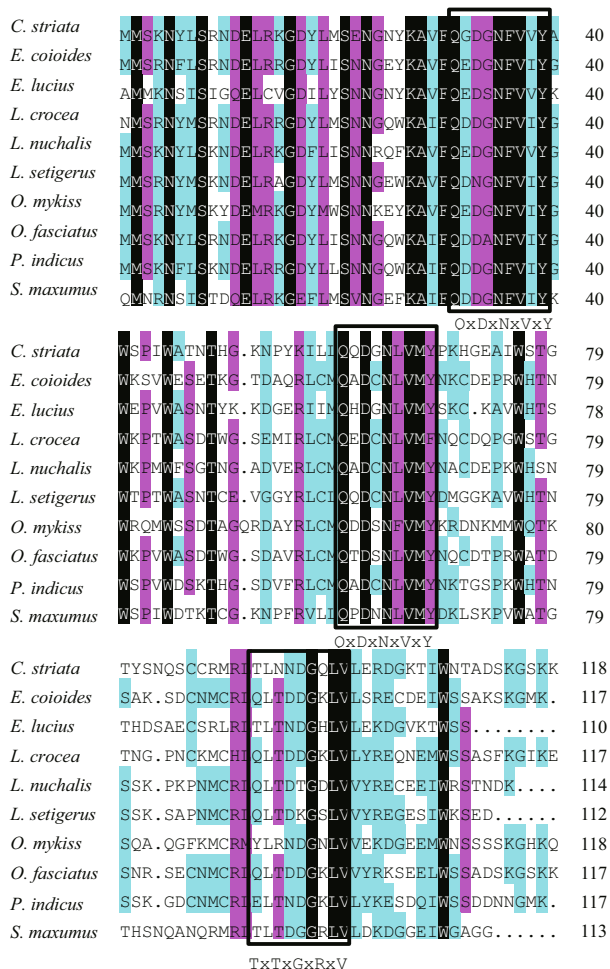
#### 3.2.1 Secondary Structure

The secondary structure analysis of *SmLTL* revealed



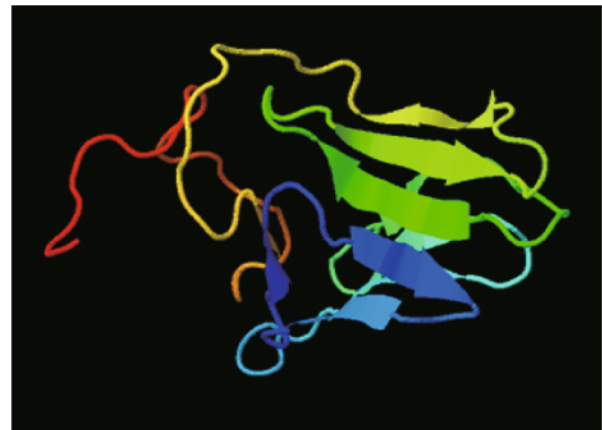


**Fig.2** Hydrophilicity analysis of *SmLTL* using the method of Kyte and Doolittle (1982)



**Fig.3** Multiple sequence alignment of *SmLTL* with other homologous genes: lily-type lectin from *Channa striata*, *Epinephelus coioides*, *Esox lucius*, *Larimichthys crocea*, *Leiognathus nuchalis*, *Lophiomus setigerus*, *Oncorhynchus mykiss*, *Oplegnathus fasciatus*, and *Platycephalus indicus*

The GenBank accession numbers of the homologous are given in supplementary material. The three D-mannose binding sites are highlighted.



**Fig.4** The three-dimensional structure of *SmLTL*

$\beta$ -sheets are indicated by patches. Turns and loops are indicated by lines.

that the protein contains 5.36% alpha helices, 39.29% extended strands, 16.07% beta turns, and 39.29% random coils. The SignalP analysis of the *SmLTL* amino acid sequence did not reveal the presence of a signal peptide or transmembrane region. The *SmLTL* protein contains a bulb-type mannose binding lectin ( $\beta$ -lectin) domain between amino acids 3 and 112 (a total of 109 amino acids). Within the  $\beta$ -lectin domain, two mannose binding sites were found between amino acids 30 and 99 with the specific motif of QxDxNxVxY in a three-fold reversed internal repeat ( $\beta$ -prism architecture) (Fig.1). The first repeat was located at Gln30-Asp32-Asn34-Val36-Tyr38 and the second repeat was located at Gln59-Asp61-Asn63-Val65-Tyr67. The third repeat (TxTxGxRxV) was located at Thr91-Thr93-Gly95-Arg97-Val99. Differences in the amino acid sequence between species were observed within the third repeat region (Fig.3).

The sequence identity of *SmLTL* was compared with other lectin superfamily members, including lily-type lectin, skin mucus lectin, and mannose binding lectin, from different fishes. The *SmLTL* was found to have highest identity with the lily-type lectin from *C. striata* (60%), *Lophiomus setigerus* (53%), *Platycephalus indicus* (52%), and *Larimichthys crocea* (50%). The length of the *SmLTL* sequence was similar to other species, and conserved motifs were observed among the sequences used in the analysis, thus confirming the identity of the gene as *SmLTL* (Fig.3).

### 3.2.2 Tertiary structure

Protein tertiary structure was predicted and inspected by PDBsum Generate (Fig.4). The overall folding of *SmLTL* consisted of three anti-folding  $\beta$ -sheets, comprised of 10  $\beta$ -strands, 14  $\beta$ -turns, and 5

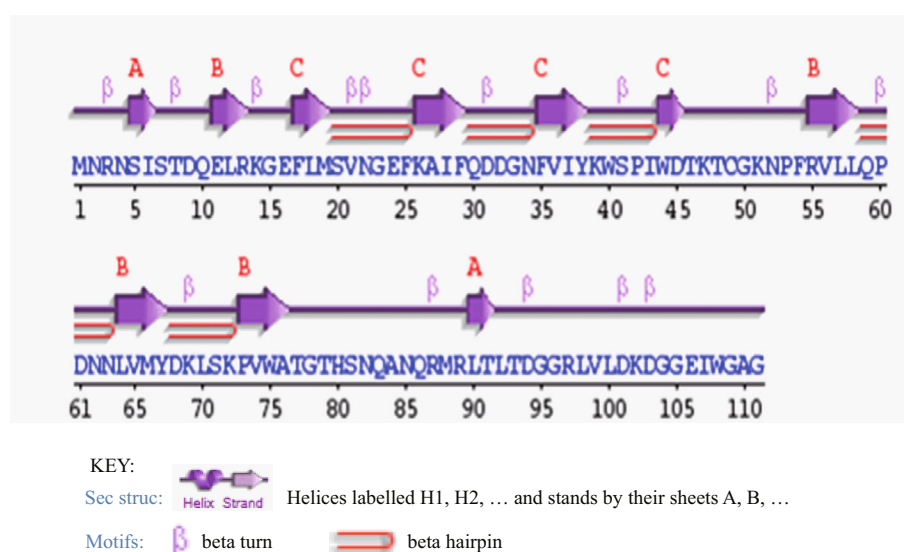


Fig.5 Secondary structure of *SmLTL* contained 10  $\beta$ -strands, 14  $\beta$ -turns, and 5  $\beta$ -hairpins

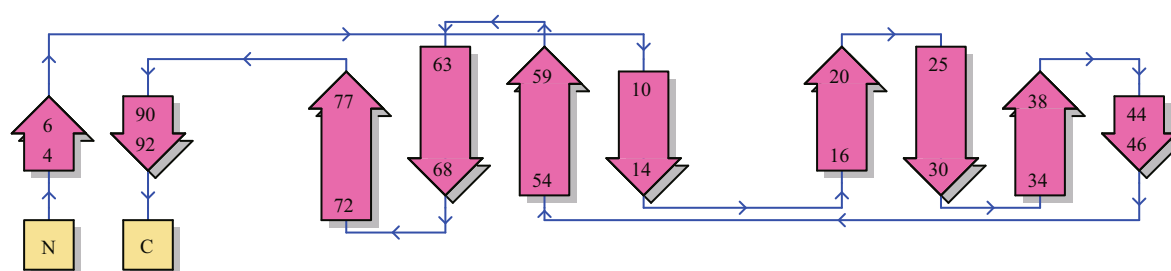


Fig.6 Secondary topology of *SmLTL*

The five beta hairpins are located at amino acids 20–25, 30–34, 38–40, 59–63, and 68–72.

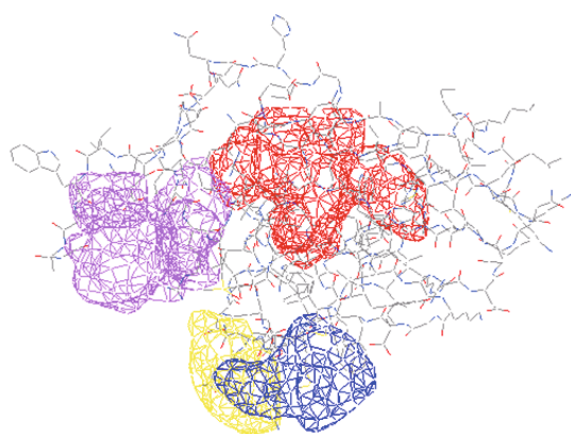


Fig.7 Schematic of the four predicted carbohydrate binding sites of *SmLTL*

Only three of the binding sites are predicted to be functional.

$\beta$ -hairpins (Figs.5, 6), and contained four protein binding sites (Fig.7). The mannose-binding sites were located in the clefts formed by the three bundles of  $\beta$ -sheets.

### 3.3 Tissue distribution of *SmLTL* mRNA

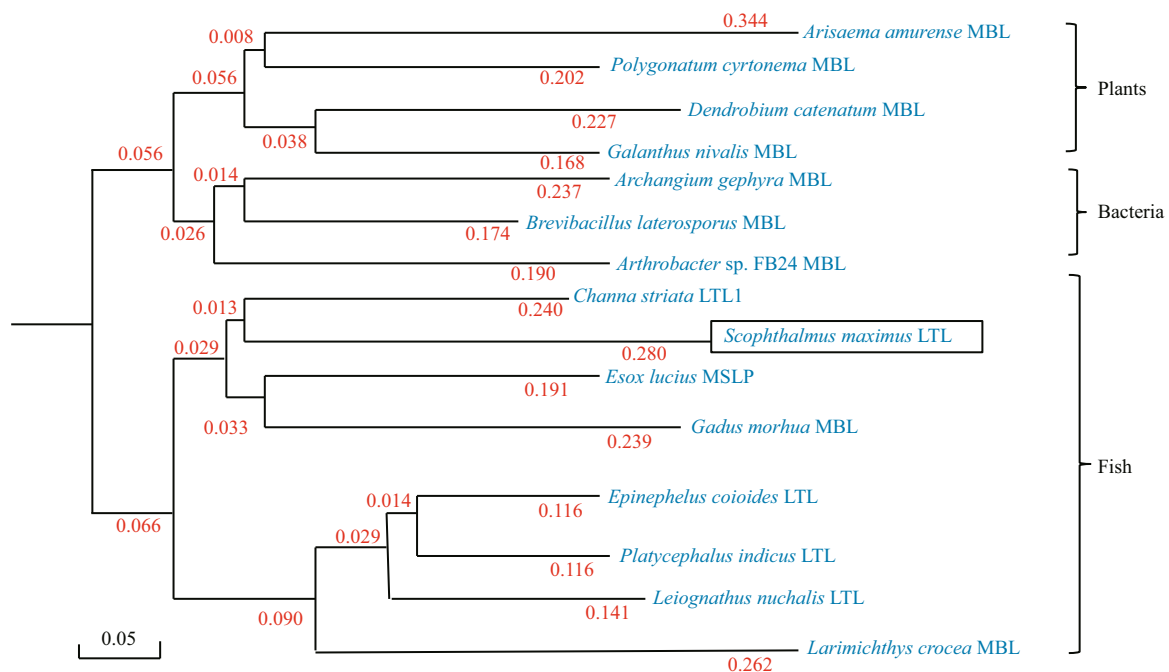
Transcripts for *SmLTL* were abundant ( $P < 0.01$ ) in the skin, intestine, and gill. Lower levels of *SmLTL* transcripts were observed in the liver, head-kidney, spleen, and muscle (Fig.9).

### 3.4 Tissue-specific expression of *V. anguillarum* challenge

Choosing the gill, intestine, and skin, which was abundant ( $P < 0.01$ ) in tissue distribution of *SmLTL* mRNA to challenge *V. anguillarum*. As shown in Fig.10 the mRNA expression of *SmLTL* were up-regulated after *V. anguillarum* challenge, and reached the highest level at 6 h.

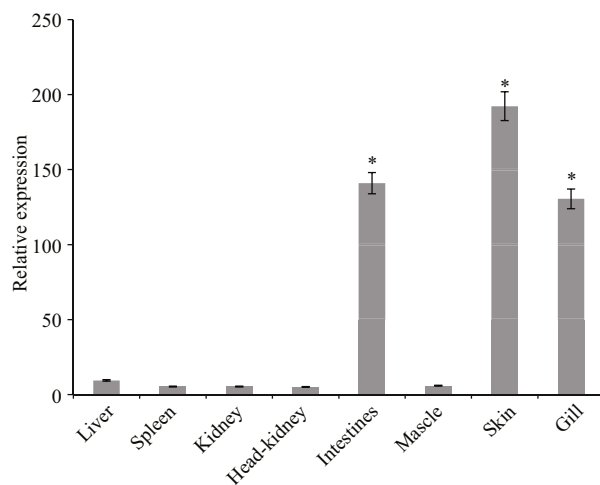
## 4 DISCUSSION

In this study, a novel lectin was isolated from turbot (*S. maximus*) using RACE techniques and was classified as *S. maximus* lily-type lectin (*SmLTL*) based on its structural and functional characteristics. Lily-type lectin is able to bind to specific carbohydrates and



**Fig.8** Phylogenetic tree of lily-type lectins constructed using the Neighbor-Joining method

*SmLTL* is indicated by a black box. The tree is based on an alignment constructed using full-length amino acid sequences using DNAMAN. The numbers at the branches denote bootstrap majority consensus values of 1 000 replicates. The GenBank accession numbers are provided in the supplementary material.



**Fig.9** Tissue distribution of *SmLTL* mRNA detected by quantitative real time reverse transcriptase PCR

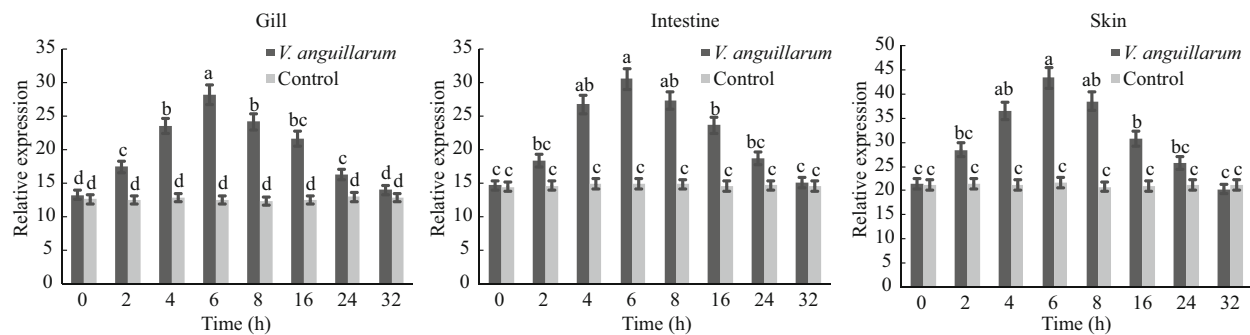
*SmLTL* expression levels in liver, spleen, kidney, head-kidney, intestines, muscle, skin and gill are normalized to that kidney. Vertical bars represent means+SE ( $n=5$ ).

plays key roles in non-self-recognition and clearance of pathogen (Dodd and Drickamer, 2001; Vasta et al., 2004). The study of *SmLTL* could be valuable for further identification of mucus lectin in fish.

The full-length *SmLTL* cDNA that was obtained in this study has a length of 569 bp and contains a 336-bp ORF that encodes a 112-residue protein that lacks a signal peptide. Sequence alignment and phylogenetic analysis (Figs.3, 8) revealed that *SmLTL* had high

similarity with lily-type lectin-1 from *C. striata* (60%), *L. setigerus* (53%), *P. indicus* (52%), *Esox lucius* (52%), *L. crocea* (50%), *Oncorhynchus mykiss* (42%), and other known lectin sequences from bacteria and plants. These data, together with the structural features of *SmLTL*, indicate that it belongs to the lily-type lectin family. The tertiary structure model of the *SmLTL* protein also indicated a high similarity with a variety of  $\beta$ -prism lectins from other species, including *Galanthus nivalis* (30%), *Narcissus pseudonarcissus* (27%), *Galanthus nivalis* (26%), and other monocot plant lectin proteins. The similarities between *SmLTL* and monocot lectins suggest new areas of research regarding the structure and function of *SmLTL*.

The alignment of *SmLTL* with other lily-type lectins revealed that the mannose-binding motifs of some lectins differ slightly from the standard form (QDNVY); however, these amino acid differences are expected to affect their ability to bind mannose (Afolabi-Balogun et al., 2012; Arasu et al., 2013). In particular, repeat three was found to exhibit slight changes in fish based on current reports. For example, repeat three was reported as TxNx DxQxV in *C. striata* (Arasu et al., 2013, Genbank: CCQ25776), TxTx DxHxV in *E. lucius* (Leong et al., 2010, GenBank: ACO14169), and YxRx DxNxV in *O. mykiss* (Berthelot et al., 2014, GenBank:



**Fig.10 *SmLTL* gene expression in turbot gill, intestine and skin at 0–32 h after *V. anguillarum* challenge**

The relative *SmLTL* expression level as exhibited by  $2^{-\Delta\Delta Ct}$  was determined for each group and the values were shown as means $\pm$ S.D ( $n=3$ ). Means with different letters indicate statistical significance ( $P<0.05$ ) between groups.

CDQ78238). However, the other two repeats (one and two) exist in all LTL. These results suggest that these lectin protein motifs are evolutionarily conserved and play an important role in their biological function (Kai et al., 2004). The modification of Arg, Lys, and Ser residues did not modify binding activity, whereas the loss of function following changes in Trp (W) or Asp/Glu (D/E) and Tyr (Y) residues indicated their crucial role in the binding activity of *Colocasia esculenta* lectin (Pereira, 2014). The reason for the presence of amino acid residues that a lack of interaction of lectin with mannose may be due to the substitution, deletion, or insertion of key amino acid residues during evolution (Luo et al., 2007). It was speculated that repeat three (TxTxGxRxV) may lead to changes in binding activity based on species-specific and the standard form (QDNVY) determined them combined with mannose. Further studies are required to investigate the binding of carbohydrates by *SmLTL*.

Three mannose recognition sites were identified in *SmLTL*, in addition to four identical subunits and three anti-folding  $\beta$ -sheets, which were comprised of 10  $\beta$ -strands, 14  $\beta$ -turns, and 5  $\beta$ -hairpins. The secondary structure analysis of *SmLTL* revealed that the protein contains 5.36% alpha helices, 39.29% extended strands, 16.07% beta turns, and 39.29% random coils. The whole folding of *SmLTL*, which typically consists of  $\beta$ -sheets connected by turns and loops, creates a very tight structural scaffold. This is very similar to the 3D structure of other mannose-binding lectins (Barre et al., 2001; Zhao et al., 2003). The monocot  $\beta$ -prism lectin structure also contains three mannose recognition sites and is a homotetrameric protein that is folded in a classic pattern (beta-prism II fold) to form its advanced structure (Hester and Wright, 1996). Molecular characterization of *SmLTL*, such as mannose-binding site analysis, signal cleavage site prediction, and

analyses of secondary and 3D structures, indicated that it shares many Exemplary features with monocot mannose-binding lectins. These similarities signify that *SmLTL* might have similar functions as many other mannose-binding lectins, such as binding to parasites, viruses, and fungi. For example, lectins have been observed to bind *Meloidogyne incognita* (Bhat et al., 2010), HIV (Ding et al., 2008), HSV-II (Luo et al., 2007), and *Rhizoctonia solani* (Tian et al., 2008). The cloning of *SmLTL* performed in the current study will enable further research into its potential functions in disease resistance.

The abundance of *SmLTL* transcripts was highest in the skin, intestine, and gill, but was weak in the liver, head-kidney, spleen, and muscle. The gills in fish are involved in gas exchange and are in continuous contact with the aquatic environment and are, therefore, more susceptible to pathogen infection. The expression of *CsLTL-1* mRNA was significantly higher in the gills, liver, intestine, and skin of *C. striata* (Abirami Arasu, 2013). Similarly, pufferlectin mRNA was also widely expressed in the gills, followed by the oral cavity wall, esophagus, and skin of *Takifugu rubripes* (Suzuki, 2003). Park et al. (2016) also reported that *RbLTL* transcript was abundant in gill and intestinal tissue in rock bream (*Oplegnathus fasciatus*). In addition, fish can absorb environmental antigens into the body via the skin (Moore et al., 1998). The lactose-binding lectin in Japanese eel (AJL-2) was demonstrated to be produced only in the skin (Tasumi et al., 2002). According to Suzuki (2003), the intestinal isoform of pufferlectin was identified in intestine of *T. rubripes*. In teleosts, the gut, skin and gill are the main mucosal surfaces and immune barriers (Goel et al., 2015). The high abundance of *SmLTL* transcripts in the skin, intestine, and gills may reflect the role of this protein in the immune response.



As *V. anguillarum* is one of the important pathogens responsible for major mortalities in turbot fish, the ability of *SmLTL* to inhibit it was of significant importance. Here, *SmLTL* showed the direct activity of facilitating the clearance of *V. anguillarum* in vivo in turbot. The mRNA expression of gill, skin and intestine in *SmLTL* were up-regulated after *V. anguillarum* challenge, and reached the highest level at 6 h. Recently many similar results have showed, such as, a novel C-type lectin (*FcLec4*) in Chinese white shrimp (Wang et al., 2009), a C-type lectin (*AiCTL-3*) in bay scallop (Huang et al., 2013), pathogen recognition receptors TLR2 in turbot (Liu et al., 2016). Obviously, mucosal immune stress response was produced after *V. anguillarum* injection in turbot, and stimulate the secretion of *SmLTL*. Our results suggest that lily-type lectins serve as the first line of defense against microbial infections and play a pivotal role in the innate mucosal immune system. We intend to further investigate the functions of *SmLTL* in the mucosal immune system through comparative pathogens studies.

## 5 CONCLUSION

The *SmLTL* from turbot was identified and characterized in this study. The general characteristics of have been reported here, including protein and cDNA sequences, tissue expression profile, domain architectures and *V. anguillarum* stimulate. The most important result of the present study is that *SmLTL* not only shares similarity with monocotyledonous plant lectins, but also contains identical mannose-binding sites. However, the function of binding site three of *SmLTL* requires further study. The information reported here will be useful for the investigation into the multifaceted functions of *SmLTL*.

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