次世代シークエンサーによるライギョダマシ脾臓 RNA のトランスクリプトーム解析

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Transcriptomic analysis of Antarctic toothfish spleen RNA by next generation sequencer

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Notothenioid fish, the dominant species in the Southern Ocean, is one of the most intriguing group of species from evolutionary viewpoint. One major reason of their evolutionary success is the acquisition of anti-freeze glycopeptides (AFGP) by the modification of trypsinogen, which enables them to survive even under harsh environment of the Southern Ocean where temperature can become as low as about -2°C. Another peculiar example of adaptive evolution is nonfunctionalization of hemoglobin genes in icefish. Although loss of hemoglobin is generally lethal for vertebrates, extensive modification of cardiovascular system in icefish has compensated such disadvantage with ensuring sufficient oxygen transport. Recent studies of mitochondrial genes, furthermore, have shown that the evolutionary rates of genes involved in cellular respiration are much higher compared with other teleosts (see also Nagata and Ota in this symposium).

In order to shed light on the molecular evolution of notothenioid genes in larger scale, we are currently investigating the genes expressed in Antarctic toothfish (*Dissostichus mawsoni*) spleen, one of important tissue of hematopoiesis, with taking advantage of technological progress of next generation sequencer. So far 55180 nucleotide sequences were assembled from about 11 million 50-mers by using velvet program (table 1). Subsequent analysis of assembled sequences then revealed that genes highly expressed are those involved in protein synthesis and folding, protein degradation, cellular respiration, cellular movement, innate and adaptive immunity, oxygen transport, iron ion storage and homeostasis, and protection against freezing, as shown in table 2. In the presentation, I am going to further discuss the results obtained in detail from molecular evolutionary perspectives.

Size of sequences (bp)	Number of sequences	Gene
$100 \sim 199$	33253	1. rRNA promoter binding protein
$200 \sim 299$	5714	2. astacin like metalloprotease
$300 \sim 399$	1774	3. β actins
$400 \sim 499$	812	4. complement molecules
$500 \sim 599$	382	5. elongation factors
$500 \sim 699$	201	6. thymosin β
$700 \sim 799$	139	7. immunoglobulins
$800 \sim 899$	67	8. type II antifreeze proteins
$900 \sim 999$	48	9. peptidyl-prolyl cis-trans isomerase
$1000 \sim 1099$	33	10. β_2 -microglobulin
$1100 \sim 1199$	20	11. hemoglobins
$1200 \sim 1299$	19	12. trypsinogens (or chimeric AFGP/ trypsinogen -like protease)
1300 ~ 1399	8	13. ferritins
$1400 \sim 1499$	2	14. heat shock protein 70
$1500 \sim 1599$	4	15. hepcidin
$1600 \sim 1699$	6	16. C-C motif chemokine
$1700 \sim 1799$	4	17. profilin
1800 \sim	4	18. carboxypeptidase

Table 1. Number of assembled nucleotide sequences*

Table 2. Highly expressed genes**

*The largest nucleotide sequence assembled is the elongation factor 2 (2473bp).

** Excluding ribosomal genes, mitochondrial genes and genes whose function were unknown.