



Identification and assessment of differentially expressed genes involved in growth regulation in *Apostichopus japonicus*

L. Zhu¹, C.H. Li¹, X.R. Su¹, C.Y. Guo¹, Z. Wang¹, C.H. Jin¹, Y. Li¹ and T.W. Li^{1,2}

¹School of Marine Sciences, Ningbo University, Ningbo, Zhejiang Province, China

²Ningbo City College of Vocational Technology, Ningbo, Zhejiang Province, China

Corresponding author: C.H. Li
E-mail: chli@yic.ac.cn

Genet. Mol. Res. 12 (3): 3028-3037 (2013)

Received January 7, 2013

Accepted June 18, 2013

Published August 20, 2013

DOI <http://dx.doi.org/10.4238/2013.August.20.4>

ABSTRACT. Rapid and efficient growth is a major consideration and challenge for global mariculture. The differential growth rate of the sea cucumber, *Apostichopus japonicus*, has significantly hampered the total production of the industry. In the present study, forward and reverse suppression subtractive hybridization libraries were constructed and sequenced from a fast-growth group and a slow-growth group of the sea cucumber. A total of 142 differentially expressed sequence tags (ESTs) with insertions longer than 150 bp were identified and further analyzed. Fifty-seven of these ESTs (approximately 40%) were functionally annotated for cell structure, energy metabolism, immunity response, and growth factor categories. Six candidate genes, arginine kinase, cytochrome *c* oxidase subunit I, HSP70, β -actin, ferritin, and the ADP-ribosylation factor, were further validated by quantitative PCR. Significant differences were found between the fast- and slow-growth groups ($P < 0.05$) for the expression levels of arginine kinase, cytochrome

c oxidase, HSP70, the ADP-ribosylation factor, and β -actin. However, no significant difference was observed for ferritin. Our results provide promising candidate gene markers for practical size screening, and also further promote marker-assisted selective breeding of this species.

Key words: *Apostichopus japonicus*; Growth rate; qPCR; Suppression subtractive hybridization