



Molecular cloning, characterization, and expression profiles of androgen receptors in spotted scat (*Scatophagus argus*)

H.P. Chen*, S.P. Deng*, M.L. Dai, C.H. Zhu and G.L. Li

Key Laboratory of Aquaculture in South China Sea for Aquatic Economic,
Animal of Guangdong Higher Education Institutes, Fisheries College,
Guangdong Ocean University, Zhanjiang, China

*These authors contributed equally to this study.

Corresponding author: G.L. Li

E-mail: guangli211@163.com

Genet. Mol. Res. 15 (2): gmr.15027838

Received October 10, 2015

Accepted December 29, 2015

Published April 7, 2016

DOI <http://dx.doi.org/10.4238/gmr.15027838>

ABSTRACT. Androgen plays critical roles in vertebrate reproductive systems via androgen receptors (ARs). In the present study, the full-length spotted scat (*Scatophagus argus*) androgen receptor (sAR) cDNA sequence was cloned from testis. The sAR cDNA measured 2448 bp in length with an open-reading frame of 2289 bp, encoding 763 amino acids. Amino acid alignment analyses showed that the sARs exhibited highly evolutionary conserved functional domains. Phylogenetically, the sARs clustered within the AR β common vertebrate group. Real-time polymerase chain reaction (RT-PCR) revealed that sAR expression varied in level and distribution throughout the tissues of both females and males. sAR expression was detected during testicular development by quantitative RT-PCR. The results showed that the highest transcription of sARs was observed in the mid-testicular stage, and remained at a high expression level until the late-testicular stage. In addition, the effects of 17 α -methyltestosterone (MT) and estrogen (E₂) on the expression of sARs in ovaries were determined using quantitative RT-PCR. sAR expression increased at 12 and 24 h post-

MT treatment and decreased with E₂ treatment. The present study provides preliminary evidence indicating gonadal plasticity of spotted scat under exogenous steroidal hormone treatments. It also provides a theoretical basis for sex reversal and production of artificial pseudo-males for female monosex breeding.

Key words: *Sarcophagus argus*; Androgen receptor; Steroid hormones; Testicular development