



## Expression profile analysis reveals putative prostate cancer-related microRNAs

H. Song<sup>1\*</sup>, Y. Liu<sup>2\*</sup>, J. Pan<sup>3</sup> and S.T. Zhao<sup>1</sup>

<sup>1</sup>Department of Urology, The Second Hospital of Shandong University, Jinan, China

<sup>2</sup>Department of Urology, General Hospital of Jinan Military Command, Jinan, China

<sup>3</sup>Jinan Municipal Center for Disease Control and Prevention, Jinan, China

\*These authors contributed equally to this study.

Corresponding author: S.T. Zhao

E-mail: zhaoshengtianzhao@hotmail.com

Genet. Mol. Res. 12 (4): 4934-4943 (2013)

Received March 7, 2013

Accepted August 30, 2013

Published October 24, 2013

DOI <http://dx.doi.org/10.4238/2013.October.24.4>

**ABSTRACT.** Annotation of prostate cancer (PC) genomes provides a foundation for discoveries that can improve the understanding and treatment of the disease. Therefore, in the present study, we used the Student *t*-test to identify differentially expressed PC-related mRNAs and microRNAs (miRNAs). Then, we performed interrelated mapping of miRNA target genes between abnormally expressed mRNAs and miRNAs, and explored mRNA-target miRNA interrelated pairs to explain the biological functions of miRNA during the progression of PC, thus revealing the occurrence of miRNA-mediated PC. After Gene Set Functional Similarity analysis, we obtained 20 abnormal PC-related candidate miRNAs, including hsa-miR-26a, hsa-miR-152, hsa-miR-19a, hsa-miR-30c, hsa-miR-19b, and hsa-miR-146b-5p, among others. These results suggest that it may be possible to predict the clinical behavior of prostate cancer based on gene expression analysis.

**Key words:** miRNA; mRNA; Prostate cancer; Target gene; Gene set; Functional similarity