

Additional file:

Fig. S1 Strategy of cloning of PNKL gene.

The CDS sequence without signal peptide of PNKL was cloned, *EcoR* I and *Not*I was insert in 5' and 3' respectively.

***EcoR* I**

GAATTCCCAGGGCTGGCCTTTTCCGGTCTGACCCCTGAGCA
CTCTGCCCTGGCAAGGGCCCACCCATGCGACGGAGAGCAG
TTCTGCCAGAACCTGGCCCCGGAGGACCCCCAGGGTGACC
AGCTGCTCCAAAGAGAGGAGCTGGGCCTCATCTGTGAGTCT
TGTCGGAAGATAATCCAGAAGCTGGAGGACATGGTGGGACC
ACAACCCAACGAGGACACTGTCACCCAGGCAGCCTCCCGG
GTGTGTGACAAGATGAAGATACTGAGAGGTGTGTGCAAGA
AGATCATGAGGACCTTTTCTCCGTCGCATCTCCAAGGACATCC
TGACTGGGAAGAAACCCCAGGCTATCTGTGTGGACATCAAG
ATCTGTAAAGAGAAGACAGGTCTCATCTGA***GCGGCCGC***

***Not* I**

Fig. S2 Results from agarose gel electrophoresis (EcoRI/Not I digestion map).

A Digestion of the recombinant expression vectors. Lane M₁ DNA marker (DL 10 000), Lanes 1 PNKL (without digestion). Lanes 2 PNKL (digestion with EcoRI and NotI) Lane M₂ DNA marker (DL 2000). b RT-PCR of PNKL. Lane M DNA marker (DL 4000), Lanes 1-2 PNKL (production of five tubes of reaction solution).

