

Table S1. All the sequences of pair of primers used in the current study

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Assay	Primer names	Primer Sequence	Restriction Site
Gene clon	AvHKT1(F)	ATGCCCAAGGACTTGGACTT	
	AvHKT1(R)	TCACATCATCACAACGAACACCA	
transformation	AvHKT1(F)	gagaacacgggggactctagaATGCCCAAGGACTTGGACTTG	<i>XbaI, BamH1</i>
	AvHKT1(R)	gcccttgctaccatggatccCATCATCACAACGAACACCACC	
transgenic line detection	35S-30(F)	AACCTCCTCGGATTCCATTGCC	
	AvHKT1(R)	TCACATCATCACAACGAACACCA	
ACTIN	F	TGCATGAGCGATCAAGTTTCAAG	
	R	TGTCCCATGTCTGGTTGATGACT	
qRT-PCR	YHKT (F)	CCTCGTGCCTACACCTTGTA	
	YHKT (R)	TGGCTGAAACGCACGAAGAA	
subcellular localization	pAN580-GFP-f	aagtccggagctagctctagatgcccaaggacttgactgtttttcac	<i>XbaI, BamH1</i>
	pAN580-GFP-r	agcgccgctgtacaggatccatcatcacaacgaacaccaccaatatagc	
Promoter clon	Promoter-F	TTCCCCATTACGCTGCTGAG	
	Promoter-R	CTTCCCCACCCACCAACATC	