

**Table S1: TaqMan primer/probes designed for specific amplification of KCC isoforms and parameters for standard curves of copy numbers.**

Identification		Sequence	Co-eff <sup>a</sup>	Slope <sup>b</sup>	E <sup>c</sup>
KCC1	F-primer	5'-GGGCTCAGTTGGGTGGAC-3'	0.999	-3.55	1.91
	R-primer	5'-AAGAAAAGGGCTGCTCTCTCTGT-3'			
	Probe	<b>6FAM-TGACTCGGACGGACATG-MGB</b>			
KCC1b	F-primer	5'-GCCAGACCTTGGCTGTCTACA-3'	0.996	-3.53	1.92
	R-primer	same as KCC1 R-primer			
	Probe	<b>TET-TGAGCCCCGGACATG-MGB</b>			
KCC3a <sup>d</sup>	F-primer	5'-CCCAGTGACCGGACTTCTCA-3'	0.995	-3.48	1.94
	R-primer	5'-TCTTATGTCCGTCGTCTAACAGTTG -3'			
	Probe	<b>6FAM-TGTCATCGAGGACCTGAG-MGB</b>			
KCC3b <sup>d</sup>	F-primer	5'GCAGACATAAAAAGCCCGGATT-3'	0.993	-3.58	1.90
	R-primer	same as KCC3a R-primer			
	Probe	<b>TET-CAGATGAACCAGACCTGAG-MGB</b>			
KCC4 <sup>e</sup>	Inventoried	TaqMan Gene Expression Assay ID Hs00383447_m1	1.000	-3.54	1.92
mKCC1	F-primer	5'- TGACAGAGATTCCTGGCATACT-3'	0.998	-3.48	1.94
	R-primer	5'- CGGCAGCCCATGCTTCT -3'			
	Probe	<b>6FAM- CTCCAGGAGAACCTC -MGB</b>			
mKCC3	F-primer	5'- ATCTCAATCCAAGGCATTCCA -3'	0.998	-3.57	1.91
	R-primer	5'- TTGGCTGATGGCTTTTCAATT -3'			
	Probe	<b>6FAM- TCATTACTGAAAATCTTTG -MGB</b>			
mKCC4	F-primer	5'- GCACAGAACAACGTTACTGAGATACA -3'	0.998	-3.52	1.92
	R-primer	5'-CACCTTTCTTTTCCACAAATGC-3'			
	Probe	<b>6FAM- CTCCTGGATAACCTGTGG -MGB</b>			

<sup>a</sup> Correlation coefficient ( $R^2$ ) is listed for isoform-specific standard curves of copy numbers. Data were derived from 2-3 sets of plasmid dilution samples with 6-10 measurements.

<sup>b</sup> The slopes (S) were derived from standard curves of copy numbers as:  $Ct = S * \log_{10} [\text{Copy Number}] + n$ .

<sup>c</sup> The PCR amplification efficiency (E) for each primer/probe set was determined by  $10^{(-1/S)}$ .

<sup>d</sup> The probes for KCC3a and KCC3b crossed exon 1 and 2 junction, and R-primer were designed to cross exon 2 and 3 junction.

<sup>e</sup> The TaqMan Assay was purchased from Applied Biosystems Inc as premade reaction mix.

**Table S2. Forward and reverse primers and restriction sites used to clone myc expression tag into pcDNA3.1 expression vectors coding for full length KCC cDNAs.**

mycKCC1	mycKCC1FW	CAGAATTCCatg <b>GAACAAAAACTCATCTCAGAAGAGGATCTG</b> cctcacttcaccgtggtgcc
	KCC1D-RV	CCTCCAAGGGGGAAAGAAAAG
	Restriction sites	EcoRI and BssHII.
mycKCC3a	mycKCC3aFW	CAGAATTCCatg <b>GAACAAAAACTCATCTCAGAAGAGGATCTG</b> catcctccagaaaccaccac
	KCC3NB2-1	TGTGTTCCCTGTGATGGAGTTC
	Restriction sites	EcoRI and BamHI
mycKCC4	mycKCC4FW	CAGAATTCCatg <b>GAACAAAAACTCATCTCAGAAGAGGATCTG</b> cccaccaacttcaccgtggtgcc
	KCC4B2	ATCGCACTCATGGAAATGGC
	Restriction sites	EcoRI and BsrGI

Each forward primer (FW) contained the myc-tag coding sequence (bold upper case) which was in-frame linked to the KCC-specific coding sequence (lowercase). Reverse primers (RV) were also isoform-specific.