

Table S1. Primer sequences and conditions for genomic amplification of the *PRKCD* gene.

Primer sequence	Conditions
Exon 5-7	38 cycles of 5 s at 95°C, 30 s at 60°C and 30 s at 72°C, using 1X GoTaq reaction buffer (Promega), 1.25U GoTaq DNA polymerase (Promega), 0.2mM for each of the dNTPs (Thermo Fisher) and 0.1µM of each primer.
5'-AGGCTGCCCTCACTGACCTTGTTTC-3'	
5'-TCACGTGCATACACTCTCTGAGCGC-3'	
Exon 8-9	
5'-TGTAGAGGGCTAGACTGGTTCGGCA-3'	
5'-TCATCAGGTCCCTGAATGAGGTGC-3'	
Exon 10	
5'-AACCCAGCAGACCCCACTGAAGC-3'	
5'-TCCCTCTGCTGCCTAAATTCTTGAGTTGCAT-3'	
Exon 13-14	
5'-CTGAGGTGGGAGTCTGTGAATCG-3'	
5'-ATCTCCCCAGTTGCCGCTCTAG-3'	
Exon 16-17	
5'-CAGTCCTGGACTAATACGGCTGAA-3'	
5'-GGGAACATGGTGAGAGAGCAATAC-3'	
Exon 18	
5'-GAGCACTGCAGGCCAGAGTGG-3'	
5'-GTGTGTGCTGAATGACTGAGTCAGGAGG-3'	