

Supplementary Table 1. PCR-RFLP condition for microRNA machinery genes polymorphism

SNP	Ref. gene	Polymorphism	Forward primer (5'-3')	Reverse primer (5'-3')	Restriction enzyme*
rs13078	<i>DICER</i>	A>T	5'-CTA GTT TTC CTG CAG ACA ATG CA-3'	5'-GTA ATG CAC ATT CAC CAA AGT CA-3'	<i>Bcc I</i>
rs3742330		A>G	5'- GGT CTC AGT TTG GTG GCT TC -3'	5'- CCT GCC TTG ACA ACA TGA AA -3'	<i>Ban II</i>
rs10719	<i>DROSHA</i>	T>C	5'-CTA GTT TTC CTG CAG ACA ATG CA-3'	5'-GTA ATG CAC ATT CAC CAA AGT CA-3'	<i>Dra III</i>
rs6877842		G>C	5'-GGG CGC AAA AAC ATG AGT GAC-3'	5'-TCC TCT CCA CAG CAA CGG AAT A-3'	<i>Sau 96I</i>
rs14035	<i>RAN</i>	C>T	5'-GAA GCA CTT GCT CAA AAT CTG TGA C-3'	5'- TGC CAT CCA CTG ATG TTC CAT C-3'	<i>Bsi I</i>
rs11077	<i>XPO5</i>	A>C	5'-TGC TTT GGG CAA GAA TCT GGT CAC-3'	5'-TAA AGG GGA TGT TAG CAC TAA AGA AT -3'	<i>Bsm I</i>

PCR-RFLP, polymerase chain reaction–restriction fragment length polymorphism; RAN, Ran GTPase; XPO5, exportin 5.

*All of the restriction enzymes were available from New England Biolabs (Ipswich, MA, USA) and the reaction conditions recommended by the instructions were used.