

Table S1. Primers used for qRT-PCR and cloning

Genes	Forward primer sequences:	Reverse primer sequences
<i>Runx2</i>	TCCTGTAGATCCGAGCACCA	CTGCTGCTGTTGTTGCTGTT
<i>Alp</i>	CCAGAAAGACACCTTGACTGTGG	TCTTGTCCGTGTCGCTCACCAT
<i>Osterix</i>	GGCTTTTCTGCGGCAAGAGGTT	CGCTGATGTTTGCTCAAGTGGTC
<i>Bsp</i>	AATGGAGACGGCGATAGTTCCG	GGAAAGTGTGGAGTTCTCTGCC
<i>Opn</i>	GCTTGGCTTATGGACTGAGGTC	CCTTAGACTCACCGCTCTTCATG
<i>β-actin</i>	AAGACCTCTATGCCAACACAG	GGAGGAGCAATGATCTTGATC
<i>MTSS1</i>	CACCTGAAGCTGCCAACCAAGTT	AGCGTAGTCAGGAAGGTGGACA
<i>Src</i>	GTTGCTTCGGAGAGGTGTGGAT	CACCAGTTTCTCGTGCCTCAGT
<i>Src full-length cloning</i>	TTGGTACCGAGCTCGGATCCGCCACC ATGGGCAGCAACAAGAGCAA	GCTGGATATCTGCAGAATTCCTAT AGGTTCTCCCCGGGCTGGTACT
<i>MTSS1</i> shRNA oligo pair	GATCCGTAGACAACCTGGTTGGCAGC TTCAAGAGAGCTGCCAACCAAGTTGT CTATTTTTTACGCGTG	AATTCACGCGTAAAAAATAGACAAC TGGTTGGCAGCTCTCTTGAAGCTG CCAACCAGTTGTCTACG

Table S2. siRNA sequences used for gene silencing

Genes	Sense sequences	Antisense sequences
MTSS1 siRNA-1	GCAAUAUACCCUCUCCUUTT	AAGGAAGAGGGUAUAUUGCTT
MTSS1 siRNA-2	GCUGCCAACCAGUUGUCUATT	UAGACAACUGGUUGGCAGCTT
