SUPPLEMENTARY TABLE 1: Primers used for qRT-PCR

Gene[1]	Forward[2]	Reverse[2]
EMR1	GCATAATCGCTGCTGGTTGA	CCAGGCAAGGAGGACAGAGTT
MCP-1	GCTGGAGAGCTACAAGAGGATCA	CCTCTCTCTTGAGCTTGGTGACA
TNF-α	AGACCCTCACACTCAGATCATCTTC	CCTCCACTTGGTGGTTTGCT
IL-6	AACGATGATGCACTTGCAGA	CCAGAGGAAATTTTCAATAGGC
IGF-I	CAGTTCGTGTGTGGACCGAG	GCTCCGGAAGCAACACTCAT
IGFBP5	ATACAACCCAGAACGCCAGCT	ACCTGGGCTATGCACTTGATG
IGFBP3	CCAGAACTTCTCCTCCGAGTCTAAG	CTCAGCACATTGAGGAACTTCAGAT
K18	CAAGTCTGCCGAAATCAGGGAC	TCCAAGTTGATGTTCTGGTTT
18S	GTGGGCCTGCGGCTTAAT	GCGAGAGTCTCGTTCGTTATC
FASN	AGAGATCCCGAGACGCTTCT	GCCTGGTAGGCATTCTGTAGT
S14	TGAGAACGACGCTGCTGAAAC	AGGTGGGTAAGGATGTGATGGAG
SREBP-1c	GGAGCCATGGATTGCACATT	GCTTCCAGAGAGGAGGCC
ERα	TGCACCATTGACAAGAACCGGA	AGCACCCATTTCATTTCGGCCT
ARG	AACTCCGCTGCTACCGCTGG	TCCACCGGCACTGTGGTCCC
PR	CAGAAGCCAGCCAGAGCC	CACAGGTAAGCACGCCATAGT
RANK	TTGTGGAATTGGGTCAATGAT	ACCGTCTTCTGGAACCATCTT
RANKL	GGCCACAGCGCTTCTCAG	GAGTGACTTTATGGGAACCCGAT

[1]Primers were designed to measure the abundance of egf-like module containing mucin-like, hormone receptor-like sequence 1 (EMR1), monocyte chemoattractant protein-1 (MCP-1), TNF-α, IL-6, IGF-I, IGF binding protein 5 (IGFBP5), IGF binding protein 3 (IGFBP3), keratin 18 (K18), 18S ribosomal RNA (18S), fatty acid synthase (FASN), thyroid hormone responsive spot 14 (S14), sterol response element binding protein-1c (SREBP-1c), estrogen receptor-α (ERα), amphiregulin (ARG), progesterone receptor (PR), receptor of activated NF-κB (RANK), and RANK ligand (RANKL) transcripts.

[2]Primer sequences are shown in the 5' to 3' orientation