

Product datasheet for RC203504L3V

OriGene Technologies, Inc.

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SSX2 (NM_175698) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: SSX2 (NM 175698) Human Tagged ORF Clone Lentiviral Particle

Symbol: SSX2

Synonyms: CT5.2; CT5.2A; HD21; HOM-MEL-40; SSX

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK
ACCN: NM 175698

ORF Size: 564 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC203504).

Sequence:
OTI Disclaimer:

The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

RefSeq: <u>NM 175698.1</u>

 RefSeq Size:
 1348 bp

 RefSeq ORF:
 567 bp

 Locus ID:
 6757

 UniProt ID:
 Q16385

 Cytogenetics:
 Xp11.22

Protein Families: Druggable Genome, Transcription Factors

MW: 21.6 kDa







Gene Summary:

The product of this gene belongs to the family of highly homologous synovial sarcoma X (SSX) breakpoint proteins. These proteins may function as transcriptional repressors. They are also capable of eliciting spontaneous humoral and cellular immune responses in cancer patients, and are potentially useful targets in cancer vaccine-based immunotherapy. This gene, and also the SSX1 and SSX4 family members, have been involved in t(X;18)(p11.2;q11.2) translocations that are characteristically found in all synovial sarcomas. This translocation results in the fusion of the synovial sarcoma translocation gene on chromosome 18 to one of the SSX genes on chromosome X. The encoded hybrid proteins are likely responsible for transforming activity. Alternative splicing of this gene results in multiple transcript variants. This gene also has an identical duplicate, GeneID: 727837, located about 45 kb downstream in the opposite orientation on chromosome X. [provided by RefSeq, Jul 2013]