

SSPN Blocking Peptide (N-term) Synthetic peptide Catalog # BP20496a

Specification

SSPN Blocking Peptide (N-term) - Product Information

Primary Accession

<u>Q14714</u>

SSPN Blocking Peptide (N-term) - Additional Information

Gene ID 8082

Other Names Sarcospan, K-ras oncogene-associated protein, Kirsten-ras-associated protein, SSPN, KRAG

Target/Specificity

The synthetic peptide sequence is selected from aa 21-34 of Human SSPN

Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

Storage Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

Precautions This product is for research use only. Not for use in diagnostic or therapeutic procedures.

SSPN Blocking Peptide (N-term) - Protein Information

Name SSPN

Synonyms KRAG

Function

Component of the dystrophin-glycoprotein complex (DGC), a complex that spans the muscle plasma membrane and forms a link between the F-actin cytoskeleton and the extracellular matrix. Preferentially associates with the sarcoglycan subcomplex of the DGC.

Cellular Location

Cell membrane; Multi-pass membrane protein. Cell membrane, sarcolemma Postsynaptic cell membrane; Multi-pass membrane protein. Note=Also found in myotendinous junctions and in the postsynaptic membrane of neuromuscular junctions.

Tissue Location

Isoform 1 is expressed exclusively in heart and skeletal muscle. Isoform 2 is expressed in heart, skeletal muscle, thymus, prostate, testis, ovary, small intestine, colon and spleen



SSPN Blocking Peptide (N-term) - Protocols

Provided below are standard protocols that you may find useful for product applications.

<u>Blocking Peptides</u>

SSPN Blocking Peptide (N-term) - Images

SSPN Blocking Peptide (N-term) - Background

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SSPN Blocking Peptide (N-term) - References

Crosbie R.H., et al. J. Biol. Chem. 272:31221-31224(1997). Crosbie R.H., et al. Submitted (APR-2000) to the EMBL/GenBank/DDBJ databases. Wiemann S., et al. Genome Res. 11:422-435(2001). Heighway J., et al. Genomics 35:207-214(1996).