

# SPG7 Antibody (Center) Blocking peptide

Synthetic peptide Catalog # BP12476c

# **Specification**

## SPG7 Antibody (Center) Blocking peptide - Product Information

**Primary Accession** 

**Q9UQ90** 

# SPG7 Antibody (Center) Blocking peptide - Additional Information

**Gene ID** 6687

#### **Other Names**

Paraplegin, 3424-, Spastic paraplegia 7 protein, SPG7, CAR, CMAR, PGN

#### **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.

# SPG7 Antibody (Center) Blocking peptide - Protein Information

Name SPG7 (HGNC:11237)

#### **Function**

ATP-dependent zinc metalloprotease. Plays a role in the formation and regulation of the mitochondrial permeability transition pore (mPTP) and its proteolytic activity is dispensable for this function (PubMed:<a href="http://www.uniprot.org/citations/26387735" target="blank">26387735</a>).

#### **Cellular Location**

Mitochondrion inner membrane; Multi-pass membrane protein

## **Tissue Location**

Ubiquitous.

### SPG7 Antibody (Center) Blocking peptide - Protocols

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

• Blocking Peptides



# SPG7 Antibody (Center) Blocking peptide - Images SPG7 Antibody (Center) Blocking peptide - Background

This gene encodes a nuclear-encoded mitochondrialmetalloprotease protein that is a member of the AAA (ATPasesassociated with a variety of cellular activities) protein family. Members of this protein family share an ATPase domain and haveroles in diverse cellular processes including membrane trafficking, intracellular motility, organelle biogenesis, protein folding, and proteolysis. Two transcript variants encoding distinct isoformshave been identified for this gene. Mutations associated with thisgene cause autosomal recessive spastic paraplegia 7. [provided byRefSeq].

# SPG7 Antibody (Center) Blocking peptide - References

Warnecke, T., et al. Mov. Disord. 25(4):413-420(2010) Augustin, S., et al. Mol. Cell 35(5):574-585(2009)Karlberg, T., et al. PLoS ONE 4 (10), E6975 (2009): Brugman, F., et al. Neurology 71(19):1500-1505(2008)Tzoulis, C., et al. J. Neurol. 255(8):1142-1144(2008)