

### **CLN8 Antibody (C-term) Blocking Peptide**

Synthetic peptide Catalog # BP18729b

### **Specification**

### CLN8 Antibody (C-term) Blocking Peptide - Product Information

Primary Accession

Q9UBY8

# CLN8 Antibody (C-term) Blocking Peptide - Additional Information

**Gene ID 2055** 

#### **Other Names**

Protein CLN8, CLN8, C8orf61

### **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.

### CLN8 Antibody (C-term) Blocking Peptide - Protein Information

Name CLN8

Synonyms C8orf61

#### **Function**

Could play a role in cell proliferation during neuronal differentiation and in protection against cell death.

### **Cellular Location**

Endoplasmic reticulum membrane; Multi-pass membrane protein. Endoplasmic reticulum-Golgi intermediate compartment membrane; Multi-pass membrane protein. Endoplasmic reticulum

# CLN8 Antibody (C-term) Blocking Peptide - Protocols

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

### • Blocking Peptides

## CLN8 Antibody (C-term) Blocking Peptide - Images



# CLN8 Antibody (C-term) Blocking Peptide - Background

This gene encodes a transmembrane protein belonging to afamily of proteins containing TLC domains, which are postulated tofunction in lipid synthesis, transport, or sensing. The proteinlocalizes to the endoplasmic reticulum (ER), and may recyclebetween the ER and ER-Golgi intermediate compartment. Mutations in this gene are associated with progressive epilepsy with mentalretardation (EMPR), which is a subtype of neuronal ceroidlipofuscinoses (NCL). Patients with mutations in this gene havealtered levels of sphingolipid and phospholipids in the brain.

# CLN8 Antibody (C-term) Blocking Peptide - References

Reinhardt, K., et al. Clin. Genet. 77(1):79-85(2010)Vantaggiato, C., et al. Hum. Mutat. 30(7):1104-1116(2009)Kousi, M., et al. Brain 132 (PT 3), 810-819 (2009) :Striano, P., et al. Epilepsy Behav 10(1):187-191(2007)Hermansson, M., et al. J. Neurochem. 95(3):609-617(2005)