

ECEL1 Antibody (Center) Blocking Peptide Synthetic peptide

Catalog # BP18053c

Specification

ECEL1 Antibody (Center) Blocking Peptide - Product Information

Primary Accession

<u>095672</u>

ECEL1 Antibody (Center) Blocking Peptide - Additional Information

Gene ID 9427

Other Names Endothelin-converting enzyme-like 1, 3424-, Xce protein, ECEL1, XCE

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.

ECEL1 Antibody (Center) Blocking Peptide - Protein Information

Name ECEL1

Synonyms XCE

Function

May contribute to the degradation of peptide hormones and be involved in the inactivation of neuronal peptides.

Cellular Location Membrane; Single-pass type II membrane protein

Tissue Location

Highly expressed in the CNS, in particular in putamen, spinal cord, medulla and subthalamic nucleus. A strong signal was also detected in uterine subepithelial cells and around renal blood vessels. Detected at lower levels in amygdala, caudate, thalamus, pancreas and skeletal muscle. Detected at very low levels in substantia nigra, cerebellum, cortex, corpus callosum and hippocampus

ECEL1 Antibody (Center) Blocking Peptide - Protocols



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

Blocking Peptides

ECEL1 Antibody (Center) Blocking Peptide - Images

ECEL1 Antibody (Center) Blocking Peptide - Background

This gene encodes a member of the M13 family ofendopeptidases. In general, M13 family members are zinc-containingtype II integral-membrane proteins that are important regulators of neuropeptide and peptide hormone activity. This gene is expressed specifically in the central nervous system and its proteinlocalizes predominately to the endoplasmic reticulum or, in traceamounts, to the cell surface. Disruption of this gene in mouseembryonic stem cells results in neonatal lethality due torespiratory failure shortly after birth. Based on the specificexpression of this gene and the phenotype of the gene deficiency inmouse embryos, it is suggested that this protein plays a criticalrole in neural regulation of the respiratory system. This gene hasmultiple pseudogenes.

ECEL1 Antibody (Center) Blocking Peptide - References

Kiryu-Seo, S., et al. J. Biol. Chem. 283(11):6988-6996(2008)Hillier, L.W., et al. Nature 434(7034):724-731(2005)Benoit, A., et al. Biochem. J. 380 (PT 3), 881-888 (2004) :Clark, H.F., et al. Genome Res. 13(10):2265-2270(2003)Kawamoto, T., et al. Int. J. Oncol. 22(4):815-822(2003)