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anti-VAMP8 antibody (N-Term)

2 Images



Go to Product page

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Overview		
Quantity:	400 μL	
Target:	VAMP8	
Binding Specificity:	N-Term	
Reactivity:	Human	
Host:	Mouse	
Clonality:	Monoclonal	
Conjugate:	This VAMP8 antibody is un-conjugated	
Application:	Western Blotting (WB), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p))	
Product Details		
Immunogen:	This VAMP8 antibody is generated from a mouse immunized with a KLH conjugated synthetic peptide between 2~24 amino acids from the N-terminal region of human VAMP8.	
Clone:	1414CT354-12-23-93	
Isotype:	IgG1 kappa	
Purification:	This antibody is purified through a protein G column, followed by dialysis against PBS.	
Target Details		
Target:	VAMP8	
Alternative Name:	VAMP8 (VAMP8 Products)	
Background:	SNAREs, Soluble N-ethylmaleimide-sensitive factor- attachment protein receptors, are essential	

Target Details

proteins for fusion of cellular membranes. SNAREs localized on opposing membranes assemble to form a trans-SNARE complex, an extended, parallel four alpha-helical bundle that drives membrane fusion. VAMP8 is a SNARE involved in autophagy through the direct control of autophagosome membrane fusion with the lysososome membrane. Also required for densegranule secretion in platelets. Plays also a role in regulated enzyme secretion in pancreatic acinar cells. Involved in the abscission of the midbody during cell division, which leads to completely separate daughter cells. Involved in the homotypic fusion of early and late endosomes.

Molecular Weight:

11438

UniProt:

Q9BV40

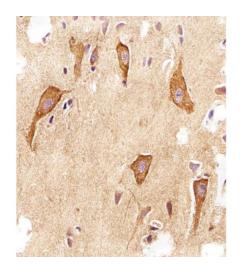
Application Details

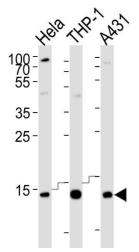
Application Notes: WB: 1:1000-1:2000. IHC-P: 1:25

Restrictions: For Research Use only

Handling

Format:	Liquid	
Buffer:	Purified monoclonal antibody supplied in PBS with 0.09 % (W/V) sodium azide.	
Preservative:	Sodium azide	
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.	
Storage:	4 °C,-20 °C	
Expiry Date:	6 months	





Immunohistochemistry (Paraffin-embedded Sections)

Image 1. Immunohistochemical analysis of paraffinembedded H. brain section using VP8 Antibody (N-term) 8440a. 8440a was diluted at 1:25 dilution. A undiluted biotinylated goat polyvalent antibody was used as the secondary, followed by DAB staining.

Western Blotting

Image 2. Western blot analysis of lysates from Hela, THP-1, A431 cell line (from left to right), using VP8 Antibody (Nterm) 8440a. 8440a was diluted at 1:1000 at each lane. A goat anti-mouse IgG H&L(HRP) at 1:10000 dilution was used as the secondary antibody. Lysates at 20 µg per lane.