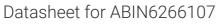
antibodies -online.com





anti-ZIP2 antibody (N-Term)

2 Images



Go to Product page

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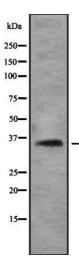
Overview	
Quantity:	100 μL
Target:	ZIP2 (Slc39a2)
Binding Specificity:	N-Term
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This ZIP2 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)
Product Details	
Immunogen:	A synthesized peptide derived from human ZIP2, corresponding to a region within N-terminal amino acids.
Isotype:	IgG
Specificity:	ZIP2 Antibody detects endogenous levels of total ZIP2.
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).
Target Details	
Target:	ZIP2 (Slc39a2)
Alternative Name:	SLC39A2 (Slc39a2 Products)

Target Details

Expiry Date:

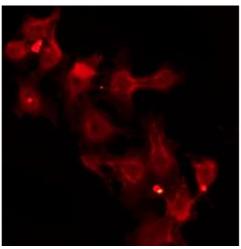
12 months

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Background:	Description: Mediates zinc uptake. Zinc uptake may be mediated by a Zn2+-HCO3- symport mechanism and can function in the presence of albumin. May also transport other divalent cations. May be important in contact inhibition of normal epithelial cells and loss of its expression may play a role in tumorigenesis. Gene: SLC39A2
Molecular Weight:	36kDa
Gene ID:	29986
UniProt:	Q9NP94
Pathways:	Autophagy
Application Details	
Application Notes:	WB 1:1000, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
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:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.



Western Blotting

Image 1. Western blot analysis of ZIP2 using Jurkat whole cell lysates



Immunofluorescence (fixed cells)

Image 2. ABIN6272466 staining Hela by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibody.