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anti-SLC22A17 antibody (Middle Region)



Image



Publication



Go to Product page

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Target:

Quantity:	100 μL
Target:	SLC22A17
Binding Specificity:	Middle Region
Reactivity:	Human, Rat, Mouse, Cow, Dog, Horse, Pig, Guinea Pig, Rabbit
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This SLC22A17 antibody is un-conjugated
Application:	Western Blotting (WB)
Product Details	
Immunogen:	The immunogen is a synthetic peptide directed towards the middle region of human SLC22A17
Sequence:	HCYQPVGGGG SPSDFYLCSL LASGTAALAC VFLGVTVDRF GRRGILLLSM
Predicted Reactivity:	Cow: 100%, Dog: 100%, Guinea Pig: 86%, Horse: 100%, Human: 100%, Mouse: 100%, Pig: 100%, Rabbit: 100%, Rat: 100%
Characteristics:	This is a rabbit polyclonal antibody against SLC22A17. It was validated on Western Blot using a cell lysate as a positive control.
Purification:	Affinity Purified

SLC22A17

Target Details

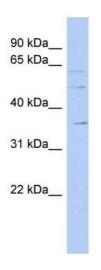
ranget Betane		
Alternative Name:	SLC22A17 (SLC22A17 Products)	
Background:	The specific functin of this protein remains unknown.	
	Alias Symbols: BOCT, BOIT, NGALR, hBOIT, 24p3R, NGALR2, NGALR3	
	Protein Interaction Partner: UBC,	
	Protein Size: 538	
Molecular Weight:	58 kDa	
Gene ID:	51310	
NCBI Accession:	NM_020372, NP_065105	
UniProt:	Q8WUG5	
Pathways:	Transition Metal Ion Homeostasis	
Application Details		
Application Notes:	Optimal working dilutions should be determined experimentally by the investigator.	
Comment:	Antigen size: 538 AA	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Concentration:	Lot specific	
Buffer:	Liquid. Purified antibody supplied in 1x PBS buffer with 0.09 % (w/v) sodium azide and 2 %	
	sucrose.	
Preservative:	Sodium azide	
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which	
	should be handled by trained staff only.	
Handling Advice:	Avoid repeated freeze-thaw cycles.	
Storage:	-20 °C	
Storage Comment:	For short term use, store at 2-8°C up to 1 week. For long term storage, store at -20°C in small	
	aliquots to prevent freeze-thaw cycles.	

Publications

Product cited in:

Rind, Schmeiser, Thiel, Absmanner, Lübbehusen, Hocks, Apeshiotis, Wilichowski, Lehle, Körner: "A severe human metabolic disease caused by deficiency of the endoplasmatic mannosyltransferase hALG11 leads to congenital disorder of glycosylation-lp." in: **Human molecular genetics**, Vol. 19, Issue 8, pp. 1413-24, (2010) (PubMed).

Images



Western Blotting

Image 1. WB Suggested Anti-SLC22A17 Antibody Titration: 0.2-1 ug/ml ELISA Titer: 1:1562500 Positive Control: DU145 cell lysate