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## Datasheet for ABIN6257103 anti-SLC44A1 antibody (Internal Region)

2 Images



#### Overview

Quantity:	100 µL	
Target:	SLC44A1	
Binding Specificity:	Internal Region	
Reactivity:	Human, Mouse, Rat	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This SLC44A1 antibody is un-conjugated	
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunocytochemistry (ICC), Immunofluorescence (IF)	

## Product Details

Immunogen:	A synthesized peptide derived from human SLC44A1, corresponding to a region within the internal amino acids.
Isotype:	lgG
Specificity:	SLC44A1 Antibody detects endogenous levels of total SLC44A1.
Predicted Reactivity:	Pig,Horse,Sheep,Rabbit,Dog,Chicken
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).

#### Target Details

Т	ar	a	et:

SLC44A1

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#### Target Details

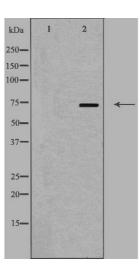
Alternative Name:	SLC44A1 (SLC44A1 Products)
Background:	Description: Choline transporter. May be involved in membrane synthesis and myelin production. Gene: SLC44A1
Molecular Weight:	73 kDa
Gene ID:	23446
UniProt:	Q8WWI5

## Application Details

Application Notes:	WB 1:500-1:1000, IF/ICC 1:100-1:500, IHC 1:50-1:200, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only

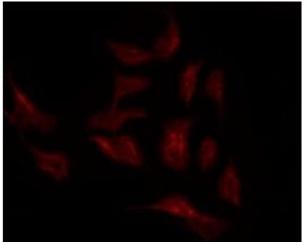
# Handling

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.
Expiry Date:	12 months





**Image 1.** Western blot analysis of extracts from K562 cells, using SLC44A1 antibody.The lane on the left is treated with the antigen-specific peptide.



#### Immunofluorescence (fixed cells)

**Image 2.** ABIN6275869 staining K562 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 25jãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37jãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibod

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