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# Datasheet for ABIN6256944 anti-SLC16A3 antibody (Internal Region)

2 Images



#### Overview

Quantity:	100 µL
Target:	SLC16A3
Binding Specificity:	Internal Region
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This SLC16A3 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)

## Product Details

Immunogen:	A synthesized peptide derived from human SLC16A3, corresponding to a region within the internal amino acids.
lsotype:	lgG
Specificity:	SLC16A3 Antibody detects endogenous levels of total SLC16A3.
Predicted Reactivity:	Pig,Rabbit,Xenopus
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).

Target Details

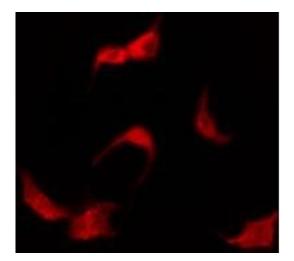
Target:

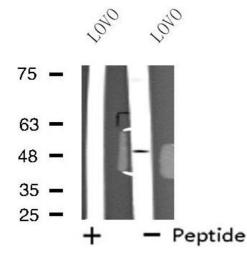
SLC16A3

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Target Details	
Alternative Name:	SLC16A3 (SLC16A3 Products)
Background:	Description: Proton-linked monocarboxylate transporter. Catalyzes the rapid transport across the plasma membrane of many monocarboxylates such as lactate, pyruvate, branched-chain oxo acids derived from leucine, valine and isoleucine, and the ketone bodies acetoacetate, beta- hydroxybutyrate and acetate (By similarity). Gene: SLC16A3
Molecular Weight:	49 kDa
Gene ID:	9123
UniProt:	015427
Pathways:	Warburg Effect
Application Details	
Application Notes:	WB 1:500-1:1000, 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.
Expiry Date:	12 months

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### Immunofluorescence (fixed cells)

**Image 1.** ABIN6275250 staining LOVO 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 25jaC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37jaC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibod

### Western Blotting

**Image 2.** Western blot analysis of extracts from LOVO cells, using MOT4 antibody.

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