

Datasheet for ABIN7601544 anti-SGCE antibody (AA 38-407)



Overview

Characteristics:

Purification:

Overview		
Quantity:	100 μg	
Target:	SGCE	
Binding Specificity:	AA 38-407	
Reactivity:	Human, Mouse, Rat	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This SGCE antibody is un-conjugated	
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Flow Cytometry (FACS)	
Product Details		
Purpose:	Anti-SGCE Antibody Picoband®	
Immunogen:	E.coli-derived human SGCE recombinant protein (Position: Y38-D407).	
Isotype:	IgG	
Cross-Reactivity (Details):	No cross-reactivity with other proteins.	

Immunogen affinity purified.

designated as Picoband, ensuring unmatched performance.

Anti-SGCE Antibody Picoband® (ABIN7601544). Tested in ELISA, Flow Cytometry, IHC, WB

applications. This antibody reacts with Human, Mouse, Rat. The brand Picoband indicates this

is a premium antibody that guarantees superior quality, high affinity, and strong signals with

minimal background in Western blot applications. Only our best-performing antibodies are

Target Details

Target:	SGCE
Alternative Name:	SGCE (SGCE Products)
Background:	Synonyms: Tumor necrosis factor receptor superfamily member 4, MRC 0X40, 0X40 antigen
	OX40L receptor, CD134, Tnfrsf4, Ox40, Txgp1I
	Background: Epsilon-sarcoglycan is a protein that in humans is encoded by the SGCE gene.
	This gene encodes the epsilon member of the sarcoglycan family. Sarcoglycans are
	transmembrane proteins that are components of the dystrophin-glycoprotein complex, which
	link the actin cytoskeleton to the extracellular matrix. Unlike other family members which are
	predominantly expressed in striated muscle, the epsilon sarcoglycan is more broadly
	expressed. Mutations in this gene are associated with myoclonus-dystonia syndrome. This
	gene is imprinted, with preferential expression from the paternal allele. Alternatively spliced
	transcript variants encoding different isoforms have been found for this gene. A pseudogene
	associated with this gene is located on chromosome 2.
Molecular Weight:	50 kDa
Gene ID:	8910
UniProt:	O43556
Application Details	

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Western blot, 0.25-0.5 μ g/mL, Human, Mouse, Rat Immunohistochemistry(Paraffin-embedded Section), 2-5 μ g/mL, Human, Mouse, Rat Flow Cytometry (Fixed), 1-3 μ g/1x10⁶ cells, Human ELISA, 0.1-0.5 μ g/mL, -

1. Asmus, F., Salih, F., Hjermind, L. E., Ostergaard, K., Munz, M., Kuhn, A. A., Dupont, E., Kupsch, A., Gasser, T. Myoclonus-dystonia due to genomic deletions in the epsilon-sarcoglycan gene. Ann. Neurol. 58: 792-797, 2005. 2. Asmus, F., Zimprich, A., Tezenas du Montcel, S., Kabus, C., Deuschl, G., Kupsch, A., Ziemann, U., Castro, M., Kuhn, A. A., Strom, T. M., Vidailhet, M., Bhatia, K. P., Durr, A., Wood, N. W., Brice, A., Gasser, T. Myoclonus-dystonia syndrome: epsilon-sarcoglycan mutations and phenotype. Ann. Neurol. 52: 489-492, 2002. 3. DeBerardinis, R. J., Conforto, D., Russell, K., Kaplan, J., Kollros, P. R., Zackai, E. H., Emanuel, B. S. Myoclonus in a patient with a deletion of the epsilon-sarcoglycan locus on chromosome 7q21. Am. J. Med. Genet. 121A: 31-36, 2003.

Restrictions:

For Research Use only

Handling

Format:	Lyophilized	
Reconstitution:	Adding 0.2 mL of distilled water will yield a concentration of 500 µg/mL.	
Concentration:	500 μg/mL	
Buffer:	Each vial contains 4 mg Trehalose, 0.9 mg NaCl, 0.2 mg Na2HPO4.	
Storage:	4 °C,-20 °C	
Storage Comment:	At -20°C for one year from date of receipt. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for six months. Avoid repeated freezing and thawing.	