

Datasheet for ABIN7602709 anti-SERAC1 antibody (AA 97-623)



Overview

Overview	
Quantity:	100 μg
Target:	SERAC1
Binding Specificity:	AA 97-623
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This SERAC1 antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (IHC), ELISA, Flow Cytometry (FACS)
Product Details	
Purpose:	Anti-SERAC1 Antibody Picoband®
Immunogen:	E.coli-derived human SERAC1 recombinant protein (Position: E97-H623).
Isotype:	IgG
Cross-Reactivity (Details):	No cross-reactivity with other proteins.
Characteristics:	Anti-SERAC1 Antibody Picoband® (ABIN7602709). 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 designated as Picoband, ensuring unmatched performance.
Purification:	Immunogen affinity purified.

Target Details

Target:	SERAC1
Alternative Name:	SERAC1 (SERAC1 Products)
Background:	Synonyms: FERM domain-containing protein 6, Willin, FRMD6, C14orf31
	Tissue Specificity: Expressed in fetal and adult brain. Also detected in fetal liver and skeletal
	muscle, but not in their adult counterparts.
	Background: Serine active site-containing protein 1, or Protein SERAC1 is a protein in humans
	that is encoded by the SERAC1 gene. The protein encoded by this gene is a
	phosphatidylglycerol remodeling protein found at the interface of mitochondria and
	endoplasmic reticula, where it mediates phospholipid exchange. The encoded protein plays a
	major role in mitochondrial function and intracellular cholesterol trafficking. Defects in this gen
	are a cause of 3-methylglutaconic aciduria with deafness, encephalopathy, and Leigh-like
	syndrome (MEGDEL). Two transcript variants, one protein-coding and the other non-protein
	coding, have been found for this gene.
Molecular Weight:	74, 20 kDa
Gene ID:	84947
Gene id.	0.15.17
Pathways:	Inositol Metabolic Process
Pathways:	
Pathways: Application Details	Inositol Metabolic Process
Pathways: Application Details	Inositol Metabolic Process Western blot, 0.25-0.5 μg/mL, Human
Pathways: Application Details	Inositol Metabolic Process Western blot, 0.25-0.5 μg/mL, Human Immunohistochemistry(Paraffin-embedded Section), 2-5 μg/mL, Human, Mouse, Rat
Pathways: Application Details	Inositol Metabolic Process Western blot, 0.25-0.5 μg/mL, Human Immunohistochemistry(Paraffin-embedded Section), 2-5 μg/mL, Human, Mouse, Rat Flow Cytometry (Fixed), 1-3 μg/1x10 ⁶ cells, Human
Pathways: Application Details	Western blot, 0.25-0.5 μg/mL, Human Immunohistochemistry(Paraffin-embedded Section), 2-5 μg/mL, Human, Mouse, Rat Flow Cytometry (Fixed), 1-3 μg/1x10 ⁶ cells, Human ELISA, 0.1-0.5 μg/mL, -
Pathways: Application Details	Western blot, 0.25-0.5 μg/mL, Human Immunohistochemistry(Paraffin-embedded Section), 2-5 μg/mL, Human, Mouse, Rat Flow Cytometry (Fixed), 1-3 μg/1x10 ⁶ cells, Human ELISA, 0.1-0.5 μg/mL, - 1. Hartz, P. A. Personal Communication. Baltimore, Md. 7/17/2012. 2. Lumish, H. S., Yang, Y.,
Pathways: Application Details	Western blot, 0.25-0.5 μg/mL, Human Immunohistochemistry(Paraffin-embedded Section), 2-5 μg/mL, Human, Mouse, Rat Flow Cytometry (Fixed), 1-3 μg/1x10 ⁶ cells, Human ELISA, 0.1-0.5 μg/mL, - 1. Hartz, P. A. Personal Communication. Baltimore, Md. 7/17/2012. 2. Lumish, H. S., Yang, Y., Xia, F., Wilson, A., Chung, W. K. The expanding MEGDEL phenotype: optic nerve atrophy,
Pathways: Application Details	Western blot, 0.25-0.5 μg/mL, Human Immunohistochemistry(Paraffin-embedded Section), 2-5 μg/mL, Human, Mouse, Rat Flow Cytometry (Fixed), 1-3 μg/1x10 ⁶ cells, Human ELISA, 0.1-0.5 μg/mL, - 1. Hartz, P. A. Personal Communication. Baltimore, Md. 7/17/2012. 2. Lumish, H. S., Yang, Y., Xia, F., Wilson, A., Chung, W. K. The expanding MEGDEL phenotype: optic nerve atrophy, microcephaly, and myoclonic epilepsy in a child with SERAC1 mutations. JIMD Rep. 16: 75-79,
Pathways: Application Details	Western blot, 0.25-0.5 μg/mL, Human Immunohistochemistry(Paraffin-embedded Section), 2-5 μg/mL, Human, Mouse, Rat Flow Cytometry (Fixed), 1-3 μg/1x10 ⁶ cells, Human ELISA, 0.1-0.5 μg/mL, - 1. Hartz, P. A. Personal Communication. Baltimore, Md. 7/17/2012. 2. Lumish, H. S., Yang, Y., Xia, F., Wilson, A., Chung, W. K. The expanding MEGDEL phenotype: optic nerve atrophy, microcephaly, and myoclonic epilepsy in a child with SERAC1 mutations. JIMD Rep. 16: 75-79, 2014. 3. Maas, R. R., Iwanicka-Pronicka, K., Kalkan Ucar, S., Alhaddad, B., AlSayed, M., Al-Owain
Pathways: Application Details	Western blot, 0.25-0.5 μg/mL, Human Immunohistochemistry(Paraffin-embedded Section), 2-5 μg/mL, Human, Mouse, Rat Flow Cytometry (Fixed), 1-3 μg/1x10 ⁶ cells, Human ELISA, 0.1-0.5 μg/mL, - 1. Hartz, P. A. Personal Communication. Baltimore, Md. 7/17/2012. 2. Lumish, H. S., Yang, Y., Xia, F., Wilson, A., Chung, W. K. The expanding MEGDEL phenotype: optic nerve atrophy, microcephaly, and myoclonic epilepsy in a child with SERAC1 mutations. JIMD Rep. 16: 75-79, 2014. 3. Maas, R. R., Iwanicka-Pronicka, K., Kalkan Ucar, S., Alhaddad, B., AlSayed, M., Al-Owain M. A., Al-Zaidan, H. I., Balasubramaniam, S., Baric, I., Bubshait, D. K., Burlina, A., Christodoulou,
Pathways: Application Details	Western blot, 0.25-0.5 μg/mL, Human Immunohistochemistry(Paraffin-embedded Section), 2-5 μg/mL, Human, Mouse, Rat Flow Cytometry (Fixed), 1-3 μg/1x10 ⁶ cells, Human ELISA, 0.1-0.5 μg/mL, - 1. Hartz, P. A. Personal Communication. Baltimore, Md. 7/17/2012. 2. Lumish, H. S., Yang, Y., Xia, F., Wilson, A., Chung, W. K. The expanding MEGDEL phenotype: optic nerve atrophy, microcephaly, and myoclonic epilepsy in a child with SERAC1 mutations. JIMD Rep. 16: 75-79, 2014. 3. Maas, R. R., Iwanicka-Pronicka, K., Kalkan Ucar, S., Alhaddad, B., AlSayed, M., Al-Owain M. A., Al-Zaidan, H. I., Balasubramaniam, S., Baric, I., Bubshait, D. K., Burlina, A., Christodoulou, J., and 46 others. Progressive deafness-dystonia due to SERAC1 mutations: a study of 67
Pathways: Application Details Application Notes:	Western blot, 0.25-0.5 μg/mL, Human Immunohistochemistry(Paraffin-embedded Section), 2-5 μg/mL, Human, Mouse, Rat Flow Cytometry (Fixed), 1-3 μg/1x10 ⁶ cells, Human ELISA, 0.1-0.5 μg/mL, - 1. Hartz, P. A. Personal Communication. Baltimore, Md. 7/17/2012. 2. Lumish, H. S., Yang, Y., Xia, F., Wilson, A., Chung, W. K. The expanding MEGDEL phenotype: optic nerve atrophy, microcephaly, and myoclonic epilepsy in a child with SERAC1 mutations. JIMD Rep. 16: 75-79, 2014. 3. Maas, R. R., Iwanicka-Pronicka, K., Kalkan Ucar, S., Alhaddad, B., AlSayed, M., Al-Owain M. A., Al-Zaidan, H. I., Balasubramaniam, S., Baric, I., Bubshait, D. K., Burlina, A., Christodoulou, J., and 46 others. Progressive deafness-dystonia due to SERAC1 mutations: a study of 67 cases. Ann. Neurol. 82: 1004-1015, 2017.

Handling

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.