

Datasheet for ABIN7602812

anti-Arylsulfatase E antibody (C-Term)



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Quantity:	100 μg
Target:	Arylsulfatase E (ARSE)
Binding Specificity:	C-Term
Reactivity:	Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This Arylsulfatase E antibody is un-conjugated
Application:	Western Blotting (WB)
Product Details	
Purpose:	Anti-ARSE/ASE Antibody Picoband®
Immunogen:	A synthetic peptide corresponding to a sequence at the C-terminus of rat ARSE/ASE, which shares 77.8% amino acid (aa) sequence identity with human ARSE/ASE.
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Isotype:	IgG
Isotype: Cross-Reactivity (Details):	IgG No cross-reactivity with other proteins.

Target Details

Target:	Arylsulfatase E (ARSE)	
Alternative Name:	Arse (ARSE Products)	
Background:	Synonyms: Pre T-cell antigen receptor alpha, pT-alpha, pTa, pT-alpha-TCR, PTCRA	
	Tissue Specificity: Expressed in immature but not mature T-cells. Also found in CD34+ cells	
	from peripheral blood, CD34+ precursors from umbilical cord blood and adult bone marrow.	
	Background: Arylsulfatase E, also known as ARSE, is an enzyme that, in humans, is encoded by	
	the ARSE gene. Arylsulfatase E is a member of the sulfatase family. It is glycosylated	
	postranslationally and localized to the golgi apparatus. Sulfatases are essential for the correct	
	composition of bone and cartilage matrix. X-linked chondrodysplasia punctata, a disease	
	characterized by abnormalities in cartilage and bone development, has been linked to	
	mutations in this gene. Alternative splicing results in multiple transcript variants. A pseudogene	
	related to this gene is located on the Y chromosome.	
Molecular Weight:	70 kDa	
Gene ID:	310326	
Application Details		
Application Notes:	Western blot, 0.25-0.5 μg/mL, Rat	
	1. Brunetti-Pierri, N., Andreucci, M. V., Tuzzi, R., Vega, G. R., Gray, G., McKeown, C., Ballabio, A.,	
	Andria, G., Meroni, G., Parenti, G. X-linked recessive chondrodysplasia punctata: spectrum of	
	arylsulfatase E gene mutations and expanded clinical variability. Am. J. Med. Genet. 117A: 164-	
	168, 2003. 2. Daniele, A., Parenti, G., d'Addio, M., Andria, G., Ballabio, A., Meroni, G. Biochemical	
	characterization of arylsulfatase E and functional analysis of mutations found in patients with	
	X-linked chondrodysplasia punctata. Am. J. Hum. Genet. 62: 562-572, 1998. 3. Franco, B.,	
	Meroni, G., Parenti, G., Levilliers, J., Bernard, L., Gebbia, M., Cox, L., Maroteaux, P., Sheffield, L.,	
	Rappold, G. A., Andria, G., Petit, C., Ballabio, A. A cluster of sulfatase genes on Xp22.3: mutations	
	in chondrodysplasia punctata (CDPX) and implications for warfarin embryopathy. Cell 81: 15-	
	25, 1995.	
Restrictions:	For Research Use only	
Handling		
Format:	Lyophilized	
Reconstitution:	Add 0.2 mL of distilled water will yield a concentration of 500 µg/mL.	

Handling

Concentration:	500 μg/mL
Buffer:	Each vial contains 4 mg Trehalose, 0.9 mg NaCl, 0.2 mg Na2HPO4, 0.01 mg Sodium azide.
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:	4 °C,-20 °C
Storage Comment:	Store 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 freeze-thaw cycles.