

OriGene Technologies, Inc.

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Product datasheet for TA332070

ABCD2 Rabbit Polyclonal Antibody

Product data:

Product Type:	Primary Antibodies
Applications:	WB
Recommended Dilution:	WB
Reactivity:	Human
Host:	Rabbit
lsotype:	lgG
Clonality:	Polyclonal
Immunogen:	The immunogen for Anti-ABCD2 Antibody: synthetic peptide directed towards the middle region of human ABCD2. Synthetic peptide located within the following region: WRFEQLDTAIRLTLSEEKQKLESQLAGIPKMQQRLNELCKILGEDSVLKT
Formulation:	Liquid. Purified antibody supplied in 1x PBS buffer with 0.09% (w/v) sodium azide and 2% sucrose. Note that this product is shipped as lyophilized powder to China customers.
Conjugation:	Unconjugated
Storage:	Store at -20°C as received.
Stability:	Stable for 12 months from date of receipt.
Predicted Protein Size:	83 kDa
Gene Name:	ATP binding cassette subfamily D member 2
Database Link:	<u>NP_005155</u> <u>Entrez Gene 225 Human</u> <u>Q9UBJ2</u>



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GRIGENE ABCD2 Rabbit Polyclonal Antibody – TA332070

Background:	The protein encoded by this gene is a member of the superfamily of ATP-binding cassette
	(ABC) transporters. ABC proteins transport various molecules across extra- and intra-cellular
	membranes. ABC genes are divided into seven distinct subfamilies (ABC1, MDR/TAP, MRP,
	ALD, OABP, GCN20, White). This protein is a member of the ALD subfamily, which is involved
	in peroxisomal import of fatty acids and/or fatty acyl-CoAs in the organelle. All known
	peroxisomal ABC transporters are half transporters which require a partner half transporter
	molecule to form a functional homodimeric or heterodimeric transporter. The function of this
	peroxisomal membrane protein is unknown; however this protein is speculated to function
	as a dimerization partner of ABCD1 and/or other peroxisomal ABC transporters. Mutations in
	this gene have been observed in patients with adrenoleukodystrophy, a severe demyelinating
	disease. This gene has been identified as a candidate for a modifier gene, accounting for the
	extreme variation among adrenoleukodystrophy phenotypes. This gene is also a candidate
	for a complement group of Zellweger syndrome, a genetically heterogeneous disorder of
	peroxisomal biogenesis. [provided by RefSeq, Jul 2008]
Synonyms:	ABC39; ALDL1; ALDR; ALDRP; hALDR
Note:	Immunogen sequence homology: Dog: 100%; Pig: 100%; Rat: 100%; Horse: 100%; Human:
	100%; Mouse: 100%; Bovine: 100%; Rabbit: 100%; Guinea pig: 100%; Yeast: 85%; Zebrafish:
	79%
Protein Families:	Druggable Genome
Protein Pathways:	ABC transporters
Product images:	

90 kDa_____ 65 kDa____ 40 kDa____ 31 kDa_____ 22 kDa___

WB Suggested Anti-ABCD2 Antibody Titration: 0.2-1 ug/ml; Positive Control: THP-1 cell lysate

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