

## **Product datasheet for TA322663**

**ABCD2 Rabbit Polyclonal Antibody** 

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**Product data:** 

**Product Type:** Primary Antibodies

Applications: IHC

Recommended Dilution: IHC: 25-100

Positive control: Human ovarian cancer

Predicted cell location: Cytoplasm

Reactivity: Human, Mouse, Rat

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

**Immunogen:** Synthetic peptide corresponding to a region derived from 274-288 amino acids of human

ATP-binding cassette, sub-family D (ALD), member 2

Formulation: PBS pH7.3, 0.05% NaN3, 50% glycerol

**Concentration:** lot specific

**Purification:** Antigen affinity purification

**Conjugation:** Unconjugated

**Storage:** Store at -20°C as received.

Stability: Stable for 12 months from date of receipt.

Gene Name: ATP binding cassette subfamily D member 2

Database Link: NP 005155

Entrez Gene 26874 MouseEntrez Gene 84356 RatEntrez Gene 225 Human

Q9UB<sub>12</sub>



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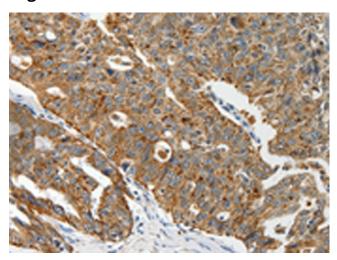
Background:

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.

Synonyms: ABC39; ALDL1; ALDR; ALDRP; hALDR

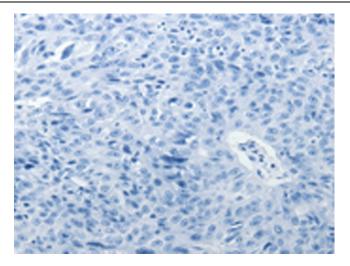
Protein Families: Druggable Genome
Protein Pathways: ABC transporters

## **Product images:**

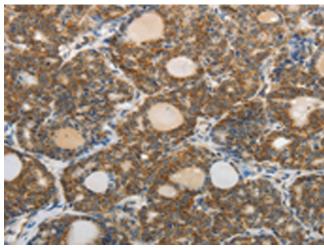


Immunohistochemistry of paraffin-embedded Human ovarian cancer tissue using TA322663 (ABCD2 Antibody) at dilution 1/20 (Original magnification: ×200)

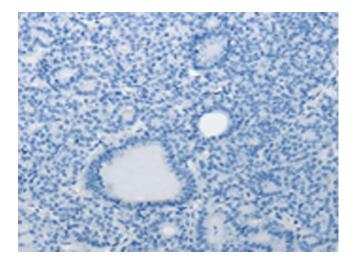




Immunohistochemistry of paraffin-embedded Human ovarian cancer tissue using TA322663 (ABCD2 Antibody) at dilution 1/20, treated with synthetic peptide. (Original magnification: ×200)



Immunohistochemistry of paraffin-embedded Human thyroid cancer tissue using TA322663 (ABCD2 Antibody) at dilution 1/20 (Original magnification: ×200)



Immunohistochemistry of paraffin-embedded Human thyroid cancer tissue using TA322663 (ABCD2 Antibody) at dilution 1/20, treated with synthetic peptide. (Original magnification: ×200)