ALDH18A1 Rabbit Polyclonal Antibody

CAB12114



Product Information Size: 20uL, 50uL, 100uL, 200uL Observed MW:	Protein Background This gene is a member of the aldehyde dehydrogenase family and encodes a bifunctional ATP- and NADPH-dependent mitochondrial enzyme with both gamma-glutamyl kinase and gamma- glutamyl phosphate reductase activities. The encoded protein catalyzes the reduction of glutamate to delta1-pyrroline-5-carboxylate, a critical step in the de novo biosynthesis of proline, ornithine and arginine. Mutations in this gene lead to hyperammonemia,		
		Refer to figures	hypoornithinemia, hypocitrullinemia, hypoargininemia and hypoprolinemia and may be associated with neurodegeneration, cataracts and connective tissue diseases. Alternatively
		Calculated MW:	spliced transcript variants, encoding different isoforms, have been described for this gene.
		87kDa	Immunogen information
Applications:	Gene ID:		
WB	5832		
Reactivity:	Uniprot		
Human	P54886		
	Synonyms:		
Antibody Information	ALDH18A1; ADCL3; ARCL3A; GSAS; P5CS; PYCS; SPG9; SPG9A; SPG9B		
Recommended dilutions: WB 1:500 - 1:2000			
	Immunogen:		
Source: Rabbit	Recombinant fusion protein containing a sequence corresponding to amino acids 1-240 of human ALDH18A1 (NP_002851.2).		
lsotype:	Storage:		
IgG	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide, 50% glycerol, pH7.3.		

Purification: Affinity purification