

Product datasheet for CF503693

AIPL1 Mouse Monoclonal Antibody [Clone ID: OTI3B4]

Product data:

Product Type:	Primary Antibodies		
Clone Name:	OTI3B4		
Applications:	FC, IF, IHC, WB		
Recommended Dilution:	WB 1:500~2000, IHC 1:150, IF 1:100, FLOW 1:100		
Reactivity:	Human, Mouse, Rat		
Host:	Mouse		
lsotype:	lgG1		
Clonality:	Monoclonal		
Immunogen:	Full length human recombinant protein of human AIPL1(NP_055151) produced in HEK293T cell.		
Formulation:	Lyophilized powder (original buffer 1X PBS, pH 7.3, 8% trehalose)		
Reconstitution Method:	For reconstitution, we recommend adding 100uL distilled water to a final antibody concentration of about 1 mg/mL. To use this carrier-free antibody for conjugation experiment, we strongly recommend performing another round of desalting process. (OriGene recommends Zeba Spin Desalting Columns, 7KMWCO from Thermo Scientific)		
Purification:	Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G)		
Conjugation:	Unconjugated		
Storage:	Store at -20°C as received.		
Stability:	Stable for 12 months from date of receipt.		
Predicted Protein Size:	43.7 kDa		
Gene Name:	aryl hydrocarbon receptor interacting protein like 1		
Database Link:	<u>NP_055151</u> <u>Entrez Gene 59110 RatEntrez Gene 114230 MouseEntrez Gene 23746 Human</u> <u>Q9NZN9</u>		



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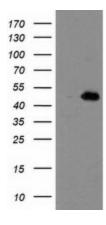
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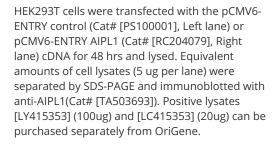
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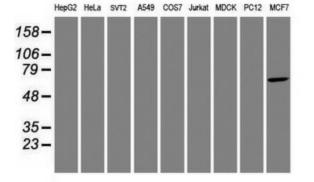
	AIPL1 Mouse Monoclonal Antibody [Clone ID: OTI3B4] – CF503693
Background:	Leber congenital amaurosis (LCA) accounts for at least 5% of all inherited retinal disease and is the most severe inherited retinopathy with the earliest age of onset. Individuals affected with LCA are diagnosed at birth or in the first few months of life with severely impaired vision or blindness, nystagmus and an abnormal or flat electroretinogram. The photoreceptor/pineal -expressed gene, AIPL1, encoding aryl-hydrocarbon interacting protein- like 1, was mapped within the LCA4 candidate region. The protein contains three tetratricopeptide motifs, consistent with nuclear transport or chaperone activity. AIPL1 mutations may cause approximately 20% of recessive LCA. [provided by RefSeq]
Synonyms:	AIPL2; LCA4

Protein Families:	Druggable Genome

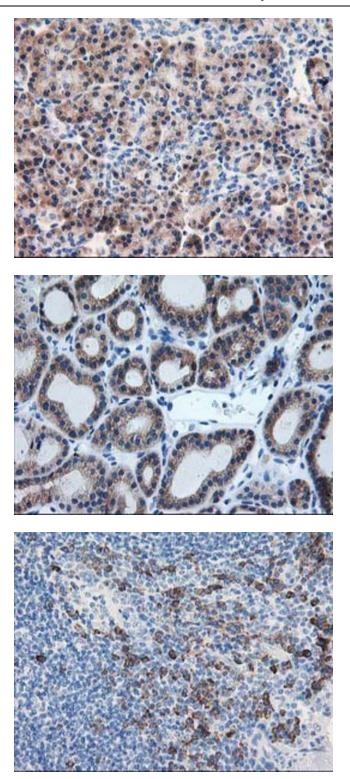
Product images:







Western blot analysis of extracts (35ug) from 9 different cell lines by using anti-AIPL1 monoclonal antibody.

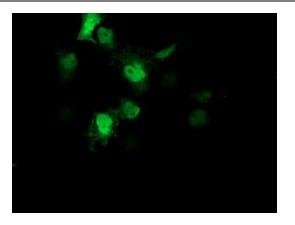
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Immunohistochemical staining of paraffinembedded Human pancreas tissue within the normal limits using anti-AIPL1 mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

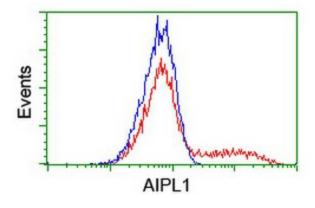
Immunohistochemical staining of paraffinembedded Carcinoma of Human thyroid tissue using anti-AIPL1 mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

Immunohistochemical staining of paraffinembedded Human lymph node tissue within the normal limits using anti-AIPL1 mouse monoclonal antibody. Heat-induced epitope retrieval by EDTA solution buffer pH 8.0 at 120°C for 3 min.

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Anti-AlPL1 mouse monoclonal antibody ([TA503693]) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY AIPL1 ([RC204079]).



HEK293T cells transfected with either [RC204079] overexpress plasmid (Red) or empty vector control plasmid (Blue) were immunostained by anti-AlPL1 antibody ([TA503693]), and then analyzed by flow cytometry.

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