

Product datasheet for TA503694S

AIPL1 Mouse Monoclonal Antibody [Clone ID: OTI2D11]

Product data:

Product Type:	Primary Antibodies
Clone Name:	OTI2D11
Applications:	FC, IF, WB
Recommended Dilution:	WB 1:2000, 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:	PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.
Concentration:	0.37 mg/ml
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>



This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2023 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US

OriGene Technologies, Inc.

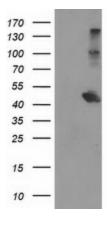
9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Image: Synonyms:AIPL1 Mouse Monoclonal Antibody [Clone ID: OTI2D11] - TA503694SBackground: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

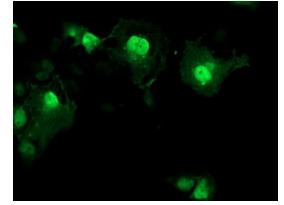
Protein Families:

Druggable Genome

Product images:

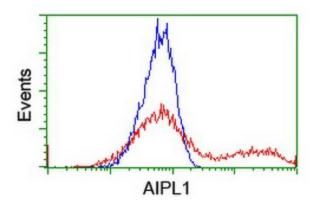


HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY AIPL1 ([RC204079], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-AIPL1. Positive lysates [LY415353] (100ug) and [LC415353] (20ug) can be purchased separately from OriGene.



Anti-AIPL1 mouse monoclonal antibody ([TA503694]) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY AIPL1 ([RC204079]).

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2023 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US



HEK293T cells transfected with either [RC204079] overexpress plasmid (Red) or empty vector control plasmid (Blue) were immunostained by anti-AlPL1 antibody ([TA503694]), and then analyzed by flow cytometry.

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2023 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US