

Product datasheet for TA504364S

SH2D2A Mouse Monoclonal Antibody [Clone ID: OTI4F3]

Product data:

Product Type:	Primary Antibodies
Clone Name:	OTI4F3
Applications:	FC, IF, IHC, WB
Recommended Dilution:	WB 1:2000, IHC 1:150, IF 1:100, FLOW 1:100
Reactivity:	Human
Host:	Mouse
lsotype:	lgG1
Clonality:	Monoclonal
Immunogen:	Full length human recombinant protein of human SH2D2A(NP_003966) produced in HEK293T cell.
Formulation:	PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.
Concentration:	0.71 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:	42.8 kDa
Gene Name:	SH2 domain containing 2A
Database Link:	<u>NP_003966</u> <u>Entrez Gene 9047 Human</u> <u>Q9NP31</u>
Background:	This gene encodes an adaptor protein thought to function in T-cell signal transduction. A related protein in mouse is responsible for the activation of lymphocyte-specific protein-tyrosine kinase and functions in downstream signaling. Alternative splicing results in multiple transcript variants. [provided by RefSeq]
Synonyms:	F2771; SCAP; TSAD; VRAP



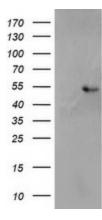
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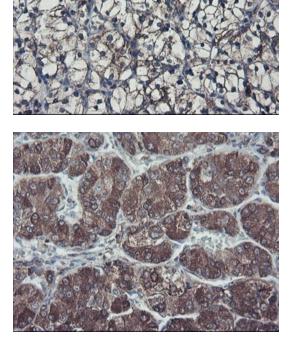
Protein Pathways: VEGF signaling pathway

Product images:



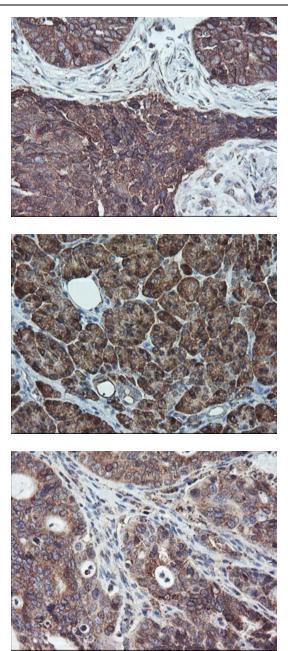
HEK293T cells were transfected with the pCMV6-ENTRY control (Cat# [PS100001], Left lane) or pCMV6-ENTRY SH2D2A (Cat# [RC204162], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-SH2D2A(Cat# [TA504364]). Positive lysates [LY418322] (100ug) and [LC418322] (20ug) can be purchased separately from OriGene.

Immunohistochemical staining of paraffinembedded Carcinoma of Human kidney tissue using anti-SH2D2A mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA504364])



Immunohistochemical staining of paraffinembedded Carcinoma of Human liver tissue using anti-SH2D2A mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA504364])

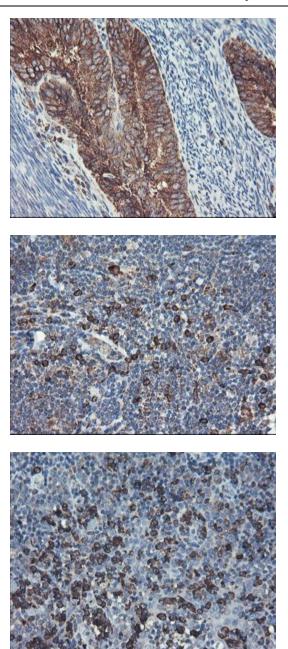
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Immunohistochemical staining of paraffinembedded Carcinoma of Human lung tissue using anti-SH2D2A mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA504364])

Immunohistochemical staining of paraffinembedded Human pancreas tissue within the normal limits using anti-SH2D2A mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA504364])

Immunohistochemical staining of paraffinembedded Carcinoma of Human pancreas tissue using anti-SH2D2A mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA504364])

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Immunohistochemical staining of paraffinembedded Adenocarcinoma of Human endometrium tissue using anti-SH2D2A mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA504364])

Immunohistochemical staining of paraffinembedded Human lymphoma tissue using anti-SH2D2A mouse monoclonal antibody. (Heatinduced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA504364])

Immunohistochemical staining of paraffinembedded Human tonsil within the normal limits using anti-SH2D2A mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA504364])

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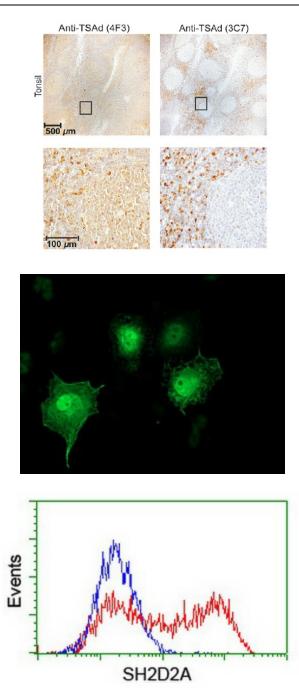


Figure from citation: Comparison of IHC on tonsil with clone 4F3 ([TA504364]) and clone 3C7 ([TA504372]) SH2D2A mAb and HRP-conjugated secondary antibodies. <u>View Citation</u>

Anti-SH2D2A mouse monoclonal antibody ([TA504364]) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY SH2D2A ([RC204162]).

HEK293T cells transfected with either [RC204162] overexpress plasmid (Red) or empty vector control plasmid (Blue) were immunostained by anti-SH2D2A antibody ([TA504364]), and then analyzed by flow cytometry.

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