

OriGene Technologies, Inc.

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Product datasheet for TA501362AM

MOB4A (MOB1B) Mouse Monoclonal Antibody (Biotin conjugated) [Clone ID: OTI1F6]

Product data:

Product Type:	Primary Antibodies
Clone Name:	OTI1F6
Applications:	FC, WB
Recommended Dilution:	WB 1:2000, FLOW 1:100
Reactivity:	Human, Mouse, Rat
Host:	Mouse
lsotype:	lgG2b
Clonality:	Monoclonal
Immunogen:	Full length human recombinant protein of human MOBKL1A(NP_775739) produced in HEK293T cell.
Formulation:	PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.
Concentration:	0.5 mg/ml
Purification:	Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G)
Conjugation:	Biotin
Storage:	Store at -20°C as received.
Stability:	Stable for 12 months from date of receipt.
Predicted Protein Size:	24.9 kDa
Gene Name:	MOB kinase activator 1B
Database Link:	<u>NP_775739</u> <u>Entrez Gene 68473 MouseEntrez Gene 360920 RatEntrez Gene 92597 Human</u> <u>Q7L9L4</u>
Background:	The protein encoded by this gene is similar to the yeast Mob1 protein. Yeast Mob1 binds Mps1p, a protein kinase essential for spindle pole body duplication and mitotic checkpoint regulation. [provided by RefSeq]. COMPLETENESS: complete on the 3' end.
Synonyms:	MATS2; MOB4A; MOBKL1A



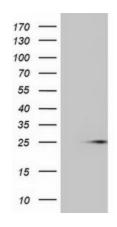
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Product images:

Events

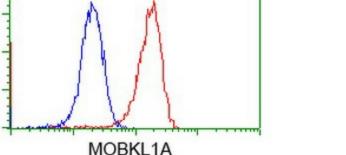
Events



MOBKL1A

HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY MOBKL1A ([RC206337], 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-MOBKL1A. Positive lysates [LY406621] (100ug) and [LC406621] (20ug) can be purchased separately from OriGene.

Flow cytometric Analysis of Hela cells, using anti-MOBKL1A antibody ([TA501362]), (Red), compared to a nonspecific negative control antibody (TA50011), (Blue).



Flow cytometric Analysis of Jurkat cells, using anti-MOBKL1A antibody ([TA501362]), (Red), compared to a nonspecific negative control antibody (TA50011), (Blue).

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