

OriGene Technologies, Inc.

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Product datasheet for TA500444AM

B Raf (BRAF) Mouse Monoclonal Antibody (Biotin conjugated) [Clone ID: OTI4C5]

Product data:

Product Type:	Primary Antibodies	
Clone Name:	OTI4C5	
Applications:	IP, WB	
Recommended Dilution:	WB 1:2000 IHC 1:50	
Reactivity:	Human, Mouse, Rat	
Host:	Mouse	
lsotype:	lgG1	
Clonality:	Monoclonal	
Immunogen:	Full-length protein expressed in 293T cell transfected with human BRAF expression vector	
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:	84.4 kDa	
Gene Name:	B-Raf proto-oncogene, serine/threonine kinase	
Database Link:	<u>NP_004324</u> <u>Entrez Gene 109880 MouseEntrez Gene 114486 RatEntrez Gene 673 Human</u> <u>P15056</u>	



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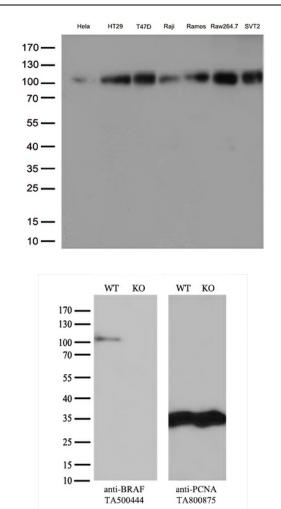
	af (BRAF) Mouse Monoclonal Antibody (Biotin conjugated) [Clone ID: OTI4C5] – TA500444AM	
Background:	This gene encodes a protein belonging to the raf/mil family of serine/threonine protein kinases. This protein plays a role in regulating the MAP kinase/ERKs signaling pathway, which affects cell division, differentiation, and secretion. Mutations in this gene are associated with cardiofaciocutaneous syndrome, a disease characterized by heart defects, mental retardation and a distinctive facial appearance. Mutations in this gene have also been associated with various cancers, including non-Hodgkin lymphoma, colorectal cancer, malignant melanoma, thyroid carcinoma, non-small cell lung carcinoma, and adenocarcinoma of lung. A pseudogene, which is located on chromosome X, has been identified for this gene. [provided by RefSeq]	
Synonyms:	B-raf; B-RAF1; BRAF1; NS7; RAFB1	
Protein Families:	Druggable Genome, Protein Kinase	
Protein Pathways:	Acute myeloid leukemia, Bladder cancer, Chemokine signaling pathway, Chronic myeloid leukemia, Colorectal cancer, Endometrial cancer, ErbB signaling pathway, Focal adhesion, Glioma, Insulin signaling pathway, Long-term depression, Long-term potentiation, MAPK signaling pathway, Melanoma, mTOR signaling pathway, Natural killer cell mediated cytotoxicity, Neurotrophin signaling pathway, Non-small cell lung cancer, Pancreatic cancer, Pathways in cancer, Progesterone-mediated oocyte maturation, Prostate cancer, Regulation of actin cytoskeleton, Renal cell carcinoma, Thyroid cancer, Vascular smooth muscle contraction	

Product images:

170 —	
130 —	
100 —	
70 —	
55 —	
40 —	
35 —	
25 —	
15 —	
10 —	

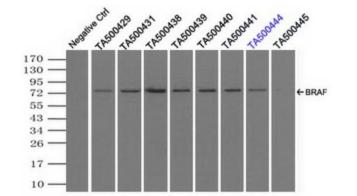
HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY BRAF ([RC211013], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-BRAF (1:500).

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Western blot analysis of extracts (35ug) from 7 cell lines lysates by using anti-BRAF monoclonal antibody (1:500).

Equivalent amounts of cell lysates (10 ug per lane) of wild-type HeLa cells (WT, Cat# LC810HELA) and BRAF-Knockout HeLa cells (KO, Cat# [LC835315]) were separated by SDS-PAGE and immunoblotted with anti-BRAF monoclonal antibody [TA500444] (1:500). Then the blotted membrane was stripped and reprobed with anti-PCNA antibody as a loading control.



Immunoprecipitation (IP) of BRAF by using TrueMab monoclonal anti-BRAF antibodies (Negative control: IP without adding anti-BRAF antibody.). For each experiment, 500ul of DDK tagged BRAF overexpression lysates (at 1:5 dilution with HEK293T lysate), 2ug of anti-BRAF antibody and 20ul (0.1mg) of goat anti-mouse conjugated magnetic beads were mixed and incubated overnight. After extensive wash to remove any non-specific binding, the immunoprecipitated products were analyzed with rabbit anti-DDK polyclonal antibody.

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