

Product datasheet for TA500844S

OriGene Technologies, Inc.

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B Raf (BRAF) Mouse Monoclonal Antibody [Clone ID: OTI1D2]

Product data:

Product Type: Primary Antibodies

Clone Name: OTI1D2

Applications: FC, IF, IP, WB

Recommended Dilution: WB 1:2000, IF 1:100, Flow 1:100

Reactivity: Human, Mouse, Rat

Host: Mouse Isotype: IgG1

Clonality: Monoclonal

Immunogen: Full length human recombinant protein of human BRAF (NP_004324) produced in HEK293T

cell

Formulation: PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.

Concentration: 1 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: 84.4 kDa

Gene Name: B-Raf proto-oncogene, serine/threonine kinase

Database Link: NP 004324

Entrez Gene 109880 MouseEntrez Gene 114486 RatEntrez Gene 673 Human

P15056





Background:

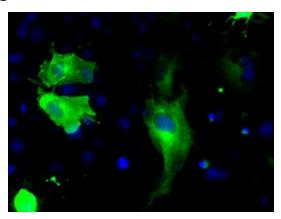
This gene encodes a protein belonging to the raf/mil family of serine/threonine protein kinases. This protein plays a role in regulating the MAP kinase/ERKs signaling pathway, which affects cell division, differentiation, and secretion. Mutations in this gene are associated with cardiofaciocutaneous 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.

Synonyms: B-raf; B-RAF1; BRAF1; NS7; RAFB1
Protein Families: Druggable Genome, Protein Kinase

Protein Pathways: Acute myeloid leukemia, Bladder cancer, C

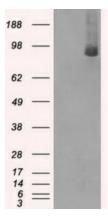
Acute myeloid leukemia, Bladder cancer, Chemokine signaling pathway, Chronic myeloid leukemia, Colorectal cancer, Endometrial cancer, ErbB signaling pathway, Focal adhesion, Glioma, Insulin signaling pathway, Long-term depression, Long-term potentiation, MAPK signaling pathway, Melanoma, mTOR signaling 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

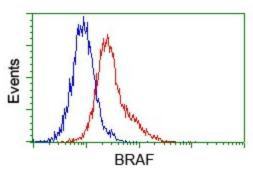
Product images:

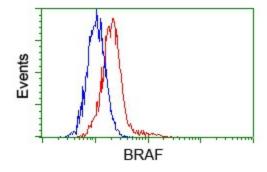


Anti-BRAF mouse monoclonal antibody ([TA500844]) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY BRAF ([RC211013]).







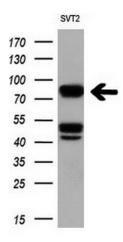


HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY BRAF ([RC211013], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-BRAF. Positive lysates [LY401382] (100ug) and [LC401382] (20ug) can be purchased separately from OriGene.

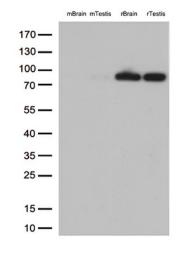
Flow cytometric analysis of Jurkat cells, using anti-BRAF antibody ([TA500844]), (Red) compared to a nonspecific negative control antibody (TA50011) (Blue).

Flow cytometric analysis of Hela cells, using anti-BRAF antibody ([TA500844]), (Red) compared to a nonspecific negative control antibody (TA50011) (Blue).

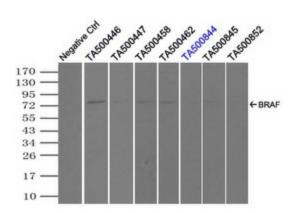




Western blot analysis of extracts (10ug) from 1 cell line by using anti-BRAF monoclonal antibody (1:200).



Western blot analysis of extracts (35ug) from 4 tissue lysates by using anti-BRAF monoclonal antibody (1:500).



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.