

order:order@affbiotech.com

BRAF Ab

Cat.#: DF7134 Size: 100ul,200ul	Concn.: 1mg/ml Source: Rabbit	Mol.Wt.: 84kDa Clonality: Polyclonal
Application:	WB 1:500-1:2000 IHC 1:50-1:200	
Reactivity:	Human,Mouse,Rat	
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink™ Coupling Resin (Thermo Fisher Scientific).	
Specificity:	BRAF Ab detects endogenous levels of total BRAF.	
Immunogen:	A synthetic peptide of human BRAF.	
Uniprot:	P15056	
Description:	BRAF: v-raf murine sarcoma viral oncogene homolog B1, also known as BRAF1; RAFB1; B-RAF1; FLJ95109. Entrez Protein NP_004324. It is the main effectors recruited by GTP- bound Ras to activate the MEK-MAP kinase pathway. B-Raf contains three consensus Akt phosphorylationsites (Ser364, Ser428, and Thr439). B-Raf is a key regulatory molecule of the mitogen-activated protein kinase kinase (MEK), it has a long amino-terminal region,the region is essential for homo- dimerization of B-Raf and hetero-dimerization of B-Raf and c- Raf at the plasma membrane, followed by phosphorylation of Thr118 in the amino-terminal B-Raf-specific region. Notably, in calcium ionophore-stimulated HeLa cells, B-Raf could propagate signals to MEK under the basal level of GTP- Ras. Expression of Raf-B is highly restricted with highestlevels in the cerebrum and testes and defects in braf are involved in a wide range of cancers. The BRAF gene mutation is frequently detected in papillary thyroid carcinoma,melanocytic nevi, primary cutaneous melanomas and colorectal cancers.	
Subcellular Location:	Nucleus. Cytoplasm. Cell memb and RAF1 in both the cytoplasm	
Tissue Specificity:	Brain and testis.	
Similarity:	Belongs to the protein kinase superfamily. TKL Ser/Thr protein kinase family. RAF subfamily.	
Storage Condition and Buffer:	Rabbit IgG in phosphate buffere NaCl, 0.02% sodium azide and °C.Stable for 12 months from d	50% glycerol.Store at -20





Western blot analysis of Hela whole cell lysates, using BRAF Ab. The lane on the left was treated with the antigen-specific peptide.



DF7134 at 1/100 staining Rat liver tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit Ab was used as the secondary.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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