

Product datasheet for TP700031

OriGene Technologies, Inc.

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B Raf (BRAF) (NM_004333) Mutant (V600E) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Recombinant protein of human v-raf murine sarcoma viral oncogene homolog B1 (BRAF)

V600E mutant, expressed in human cells

Species: Human
Expression Host: HEK293T

Expression cDNA Clone

or AA Sequence:

A DNA sequence from TrueORF clone, RC211013, encoding the V600E mutant form of human

BRA

Tag: C-Myc/DDK

Predicted MW: 84 kDa

Concentration: >0.05 μg/μL as determined by microplate BCA method

Purity: > 80% as determined by SDS-PAGE and Coomassie blue staining

Buffer: 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

Bioactivity: BRAF kinase activity was measured in an HTRF? assay. Varying concentrations of BRAF were

added to a reaction mix containing ATP and a biotinylated kinase substrate (HTRF substrate 2) and was incubated at 37C for phosphorylation. HTRF detection reagents were then added, the reaction was incubated for 30 minutes at room temperature. Time-resolved fluorescent

signal (Delta R) was measured on a Flexstation 3 microplate reader.

Enzyme activity (PMID: 26814611)

Note: For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

RefSeq: <u>NP 004324</u>

Locus ID: 673

 UniProt ID:
 P15056

 RefSeq Size:
 2949

 Cytogenetics:
 7q34





RefSeq ORF: 2298

Synonyms: B-raf; B-RAF1; BRAF1; NS7; RAFB1

Summary: This gene encodes a protein belonging to the RAF family of serine/threonine protein kinases.

This protein plays a role in regulating the MAP kinase/ERK signaling pathway, which affects cell division, differentiation, and secretion. Mutations in this gene, most commonly the V600E mutation, are the most frequently identified cancer-causing mutations in melanoma, and have been identified in various other cancers as well, including non-Hodgkin lymphoma, colorectal cancer, thyroid carcinoma, non-small cell lung carcinoma, hairy cell leukemia and

adenocarcinoma of lung. Mutations in this gene are also associated with

cardiofaciocutaneous, Noonan, and Costello syndromes, which exhibit overlapping

phenotypes. A pseudogene of this gene has been identified on the X chromosome. [provided

by RefSeq, Aug 2017]

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:





