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Recombinant Human TrkB Protein (His Tag)

Catalog Number:	633201, 633202
Size:	100 μg, 500 μg
Target Name:	TrkB, Neurotrophic tyrosine kinase receptor type 2, GP145-TrkB
Regulatory Status:	RUO

PRODUCT DETAILS

Application:	Bioassay
Format:	Lyophilized from sterile PBS, pH7.4.
Expression Host:	HEK293
Species:	Human
accession number:	NP_001007098.1
Sources:	A DNA sequence encoding the extracellular domain (Met 1-His 430) of human TrkB (NP_001007098.1) precursor was expressed with a C-terminal polyhistidine tag.
Molecular Weight:	The recombinant human TrkB consists of 410 amino acids and has a predicted molecular mass of 45.7 KDa. In SDS-PAGE under reducing conditions, the apparent molecular mass of rhTrkB is approximately 70-80 KDa due to glycosylation.
Affinity Tag:	C-His
Purity:	≥95% as determined by SDS-PAGE. ≥95% as determined by SEC-HPLC
Endotoxin level:	< 1.0 EU per μg protein
Protein Concentration:	Lyophilized
Storage and Handling:	Samples are stable for up to twelve months from date of receipt at -20℃ to -80℃. Store it under sterile conditions at -20℃ to -80℃. It is recommended that the protein be aliquoted for optimal storage. Avoid repeated freeze-thaw cycles.

BACKGROUND INFORMATION

Neurotrophic tyrosine kinase (NTRK) receptors are a family of receptor tyrosine kinases essential for nervous system development, neuronal survival, differentiation, and synaptic plasticity. The family includes three members: NTRK1 (TrkA), NTRK2 (TrkB), and NTRK3 (TrkC). These receptors regulate neuronal growth, maintain neural connections, and support the function of sensory and central nervous system neurons. Beyond neural biology, NTRK signaling can influence cell proliferation, survival, and differentiation in other tissues.

NTRK receptors are single-pass transmembrane glycoproteins composed of an extracellular ligand-binding region, a transmembrane segment, and an intracellular tyrosine kinase domain. Their extracellular regions contain leucine-rich repeats and immunoglobulin-like domains involved in ligand recognition and receptor interactions. Binding of a neurotrophin promotes receptor dimerization and autophosphorylation, initiating downstream signaling pathways such as RAS–MAPK, PI3K–AKT, and PLCγ. These

pathways regulate gene expression, neuronal survival, differentiation, and synaptic function.

The three receptors have distinct but overlapping ligand preferences. NTRK1 primarily binds nerve growth factor (NGF), NTRK2 binds brain-derived neurotrophic factor (BDNF) and neurotrophin-4 (NT-4), and NTRK3 preferentially binds neurotrophin-3 (NT-3). Ligand specificity is not absolute, as NT-3 can activate other Trk receptors under certain conditions. Neurotrophins may also interact with the p75 neurotrophin receptor, which can modify signaling outcomes.

Altered NTRK signaling is associated with neurological conditions, including sensory and pain disorders, as well as developmental abnormalities. In cancer, chromosomal rearrangements involving NTRK genes can generate oncogenic fusion proteins that retain the kinase domain and drive constitutive signaling. These NTRK fusions occur across multiple tumor types and can promote uncontrolled cell growth and survival.

Therapeutically, selective TRK inhibitors, including larotrectinib and entrectinib, are used to treat eligible cancers harboring NTRK gene fusions. Research also explores modulation of neurotrophin–Trk signaling to support neuronal protection, regeneration, and treatment of neurological disorders.

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