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## TRKβ, Active

Recombinant protein expressed in Sf9 cells

# Catalog # 7758-5

## **Product Description**

Recombinant human TRKB (455-end) was expressed by baculovirus in \$9 insect cells using an N-terminal GST tag. The gene accession number is NM 001018064.

#### **Gene Aliases**

NTRK2, GP145-TrkB

#### **Formulation**

Recombinant protein stored in 50mM Tris-HCI, pH 7.5, 150mM NaCl, 0.25mM DTT, 0.1mM EGTA, 0.1mM EDTA, 0.1mM PMSF, 25% glycerol.

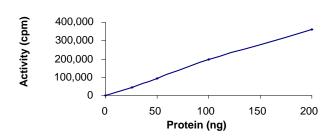
## Storage and Stability

Store product at  $-70^{\circ}$ C. For optimal storage, aliquot target into smaller quantities after centrifugation and store at recommended temperature. For most favorable performance, avoid repeated handling and multiple freeze/thaw cycles.

#### Scientific Background

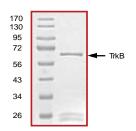
TRKB is a member of the neurotrophic tyrosine receptor kinase (NTRK) family. TRKB is the high affinity catalytic receptor for several "neurotrophins", which are small protein growth factors that induce the survival and differentiation of distinct cell populations (1). TRKB is a membrane-bound receptor that, upon neurotrophin binding, phosphorylates itself and members of the MAPK pathway (2). Signalling through TRKB leads to cell differentiation. Mutations in the TRKB gene have been

## **Specific Activity**



The specific activity of TRKB was determined to be **74 nmol/min/mg** asper activity assay protocol.

## **Purity**



The purity of TRKB was determined to be >80% by densitometry, approx. MW 67kDa.

## TRKB, Active

Full-length recombinant protein expressed in Sf9 cells

Catalog Number
Quantity
Specific Activity
Lot Specific Number

7758-5 5μg 74 nmol

Purity

>80%



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- Gregory, T. et al: Signal Transduction Mediated by the Truncated trkB Receptor Isoforms, trkB.T1 and trkB.T2. J. Neurosci. 1997; 17(8); 2683-2690.
- Pearse, R N. et al: A neurotrophin axis in myeloma: TrkB and BDNF promote tumor-cell survival. Blood. 2005;105(11):4429-36.

quantities after certifity and store at recommended temperature. For most favorable performance, avoid repeated handling and multiple freeze/thaw cycles. Product shipped on dry ice.

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## **Activity Assay Protocol**

#### **Reaction Components**

#### **Active Kinase** (Catalog # 7758-5)

Active TrkB  $(0.1\mu g/\mu l)$  diluted with Kinase Dilution Buffer and assayed as outlined in sample activity plot. (Note: these are suggested working dilutions and it is recommended that the researcher perform a serial dilution of Active SYK for optimal results).

#### Kinase Dilution Buffer, pH 7.2

Kinase Assay Buffer II diluted at a 1:4 ratio (5X dilution) with 50 ng/ $\mu$ l BSA solution.

### Kinase Assay Buffer II, pH 7.2

Buffer components: 25mM MOPS pH 7.2, 12.5mM  $\beta$ -glycerol-phosphate, 20mM MgC1 $_2$ , 12.5mM MnC1 $_2$ , 5mM EGTA, 2mM EDTA. Add 0.25mM DTT to Kinase Assay Buffer prior to use.

## [32P]-ATP Assay Cocktail

Prepare 250 $\mu$ M [ $^{32}$ P]-ATP Assay Cocktail in a designated radioactive working area by adding the following components: 150 $\mu$ l of 10mM ATP Stock Solution, 100 $\mu$ l [ $^{32}$ P]-ATP (1mCi/100 $\mu$ l), 5.75ml of Kinase Assay Buffer). Store 1ml aliquots at  $-20^{\circ}$ C.

#### 10mM ATP Stock Solution

Prepare ATP stock solution by dissolving 55mg of ATP in 10ml of Kinase Assay Buffer. Store  $200\mu l$  aliquots at  $-20^{\circ}C$ .

#### **Substrate**

Poly (Glu:Tyr, 4:1) synthetic peptide substrate diluted in distilled  $H_2O$  to a final concentration of 1 mg/ml.

#### **Assay Protocol**

- Step 1. Thaw [32P]-ATP Assay Cocktail in shielded container in a designated radioactive working area.
- Step 2. Thaw the Active TrkB, Kinase Assay Buffer, Substrate and Enzyme Dilution Buffer on ice.
- Step 3. In a pre-cooled microfuge tube, add the following reaction components bringing the initial reaction volume up to 20μl:

Component 1. 10μl of diluted Active TrkB (Catalog # 7758-5).

Component 2. 5µl of 1 mg/ml stock solution of substrate.

Component 3. 5µl distilled H,O (4°C)

- Step 4. Set up the blank control as outlined in step 3, excluding the addition of the substrate. Replace the substrate with an equal volume of distilled H<sub>2</sub>O.
- Step 5. Initiate the reaction by the addition of 5μl [32P]-ATP Assay Cocktail bringing the final volume up to 25μl and incubate the mixture in a water bath at 30°C for 15 minutes.
- Step 6. After the 15 minute incubation period, terminate the reaction by spotting 20μl of the reaction mixture onto individual pre-cut strips of phosphocellulose P81 paper.
- Step 7. Air dry the pre-cut P81 strip and sequentially wash in a 1% phosphoric acid solution (dilute 10ml of phosphoric acid and make a 1L solution with distilled H<sub>2</sub>O) with constant gentle stirring. It is recommended that the strips be washed a total of 3 intervals for approximately 10 minutes each.
- Step 8. Count the radioactivity on the P81 paper in the presence of scintillation fluid in a scintillation counter.
- Step 9. Determine the corrected cpm by removing the blank control value (see Step 4) for each sample and calculate the kinase specific activity as outlined below.

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### Kinase Specific Activity (SA) (pmol/min/μg or nmol/min/mg)

Corrected cpm from reaction / [(SA of <sup>32</sup>P-ATP in cpm/pmol)\*(Reaction time in min)\*(Enzyme amount in μg or mg)]\*[(Reaction Volume) / (Spot Volume)]

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## MATERIAL SAFETY DATA SHEET

### Article 1 - Product Identification and Use

Product Name: Active TRKB

## Catalog # 7758-5

This product is sold only for research use by qualified laboratory personnel, and is not to be used as a drug, medical device, food additive, cosmetic, nor household chemical. It is not to be used in diagnostic, therapeutic, consumer, agricultural, nor pesticidal applications.

Seller's Name: BioVision, Inc.

Street Address: 980 Linda Vista Avenue
City, Prov. Postal Code: Mountain View, CA 94043

Fa x: 659-428-0336 PHONE: 650-428-0236

## **Article 2 - Hazardous Ingredients**

NOT AVAILABLE. We are not aware of any hazards associated with this product or its ingredients, but the chemical, physical, and toxicological properties of this product have not been investigated thoroughly. Observe normal laboratory precautions.

## **Article 3 - Physical Data**

This product consists of purified protein in Tris-HCI buffer shipped on dry ice. The physical properties of this product have not been investigated thoroughly.

## **Article 4 - Fire and Explosion Hazard**

NOTAPPLICABLE

## **Article 5 - Reactivity Data**

NOTAPPLICABLE

## Article 6 - Toxicologically Data

May be harmful by inhalation, ingestion, or skin absorption. The toxicological properties of this product have not been investigated thoroughly. Exercise due caution.

## **Article 7 - Preventative Measures**

Wear chemical safety goggles and compatible chemical-resistant gloves. Avoid inhalation, contact with eyes, skin or clothing.

#### \*\*\*\*\*MULTIPLE COMPONENT SPILL OR LEAK PROCEDURES\*\*\*\*\*

- Wear protective equipment.
- Absorb on sand or vermiculite and place in closed containers for disposal.
- Observe all federal, state and local environmental regulations.

## **Article 8 - First Aid Measures**

- If swallowed, wash out mouth with water, provided person is conscious. Call a physician.
- In case of skin contact, flush with copious amounts of water for at least 15 minutes. Remove contaminated clothing and shoes. If a rash
  or other irritation develops, call a physician.
- If inhaled, remove to fresh air. If breathing becomes difficult, call a physician.
- In case of eye contact, flush with copious amounts of water for at least 15 minutes while separating the eyelids with fingers. Call a
  physician.

The above information is believed to be correct but does not purport to be all-inclusive and shall be used only as a guide. Bio Vision shall not be held liable for any damage resulting handling or from contact with the above product. See the Technical Specification. Packing Sign. Invoice, and Product Catalogue for additional terms and conditions of sale.

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