# elF2Bβ (h2): 293T Lysate: sc-172832



The Power to Question

## **BACKGROUND**

The initiation of protein synthesis in eukaryotic cells is regulated by interactions between protein initiation factors and RNA molecules. The eukaryotic initiation complex elF2B exists as a five subunit complex composed of elF2 $\alpha$ , elF2 $\beta$ , elF2 $\gamma$ , elF2B $\delta$ , and elF2B $\epsilon$ . The elF2B complex catalyzes the exchange of GDP for GTP on the elF2 complex, following the interaction of elF2/GTP with the 40S ribosomal subunit. Guanine nucleotide exchange factor (GEF) activity is exhibited by the elF2B $\epsilon$  subunit alone, but is greater in the presence of all five elF2B subunits. Phosphorylation of elF2 inhibits GEF activity of elF2B, an inhibition that requires the elF2 $\alpha$  subunit.

## **REFERENCES**

- 1. Trachsel, H. and Staehelin, T. 1978. Binding and release of eukaryotic initiation factor eIF2 and GTP during protein synthesis initiation. Proc. Natl. Acad. Sci. USA 75: 204-208.
- Benne, R., Amesz, H., Hershey, J.W. and Voorma, H.O. 1979. The activity
  of eukaryotic initiation factor eIF2 in ternary complex formation with GTP
  and Met-tRNA. J. Biol. Chem. 254: 3201-3205.
- 3. Ernst, H., Duncan, R.F. and Hershey, J.W. 1987. Cloning and sequencing of complementary DNAs encoding the  $\alpha$ -subunit of translational initiation factor eIF2. Characterization of the protein and its messenger RNA. J. Biol. Chem. 262: 1206-1212.
- Pathak, V.K., Nielsen, P.J., Trachsel, H. and Hershey, J.W. 1988. Structure
  of the β subunit of translational initiation factor eIF2. Cell 54: 633-639.
- Kaufman, R.J., Davies, M.V., Pathak, V.K. and Hershey, J.W. 1989. The phosphorylation state of eucaryotic initiation factor 2 alters translational efficiency of specific mRNAs. Mol. Cell. Biol. 9: 946-958.
- 6. Gaspar, N.J., Kinzy, T.G., Scherer, B.J., Humbelin, M., Hershey, J.W. and Merrick, W.C. 1994. Translation initiation factor eIF2. Cloning and expression of the human cDNA encoding the  $\gamma$  subunit. J. Biol. Chem. 269: 3415-3422.

#### **CHROMOSOMAL LOCATION**

Genetic locus: EIF2B2 (human) mapping to 14q24.3.

## **PRODUCT**

elF2B $\beta$  (h2): 293T Lysate represents a lysate of human elF2B $\beta$  transfected 293T cells and is provided as 100  $\mu$ g protein in 200  $\mu$ l SDS-PAGE buffer.

#### **APPLICATIONS**

elF2B $\beta$  (h2): 293T Lysate is suitable as a Western Blotting positive control for human reactive elF2B $\beta$  antibodies. Recommended use: 10-20  $\mu$ l per lane.

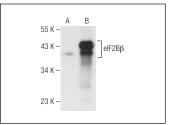
Control 293T Lysate: sc-117752 is available as a Western Blotting negative control lysate derived from non-transfected 293T cells.

elF2B $\beta$  (E-12): sc-376478 is recommended as a positive control antibody for Western Blot analysis of enhanced human elF2B $\beta$  expression in elF2B $\beta$  transfected 293T cells (starting dilution 1:100, dilution range 1:100-1:1,000).

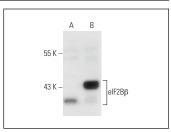
## **RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG $\kappa$  BP-HRP: sc-516102 or m-lgG $\kappa$  BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker<sup>TM</sup> Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048.

#### **DATA**







elF2Bβ (P-4): sc-9979. Western blot analysis of elF2Bβ expression in non-transfected: sc-117752 (**A**) and human elF2Bβ transfected: sc-172832 (**B**) 293T whole cell lysates

#### STORAGE

Store at -20° C. Repeated freezing and thawing should be minimized. Sample vial should be boiled once prior to use. Non-hazardous. No MSDS required.

# **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

## **PROTOCOLS**

See our web site at www.scbt.com for detailed protocols and support products.

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