SH2-B (E-8): sc-393395



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BACKGROUND

SH2-B, also known as SH2B1 or PSM, is a 756 amino acid protein that is a component of the signaling network and is involved in the regulation of cell shape and movement. SH2-B is related to the APS (adapter molecule containing PH and SH2 domains) family of adapter proteins, which characteristically contain a pleckstrin homology (PH) domain, an SH2 domain and a tyrosine phosphorylation site. SH2-B is alternatively spliced to generate three distinct isoforms, SH2-B $\alpha,\,\beta,$ and $\gamma,$ that share an identical N-terminal sequence, including the PH domain, the SH2 domain, and multiple proline-rich motifs. Containing a PH domain and a SH2 domain, SH2-B shuttles between the nuclues and the cytoplasm. SH2-B is widely expressed with highest expression in skeletal muscle and ovary. SH2-B is phosphorylated on tyrosine residues in response to receptor kinase stimulation.

REFERENCES

- Frank, S.J., et al. 1995. Regions of the JAK2 tyrosine kinase required for coupling to the growth hormone receptor. J. Biol. Chem. 270: 14776-14785.
- Rui, L., et al. 1997. Identification of SH2-Bβ as a substrate of the tyrosine kinase JAK2 involved in growth hormone signaling. Mol. Cell. Biol. 17: 6633-6644.
- Rui, L., et al. 1998. Platelet-derived growth factor (PDGF) stimulates the association of SH2-Bβ with PDGF receptor and phosphorylation of SH2-Bβ.
 J. Biol. Chem. 273: 21239-21245.
- Rui, L., et al. 1999. Identification of SH2-Bβ as a potent cytoplasmic activator of the tyrosine kinase Janus kinase 2. Proc. Natl. Acad. Sci. USA 96: 7172-7177.
- Rui, L., et al. 1999. SH2-B is required for nerve growth factor-induced neuronal differentiation. J. Biol. Chem. 274: 10590-10594.

CHROMOSOMAL LOCATION

Genetic locus: SH2B1 (human) mapping to 16p11.2.

SOURCE

SH2-B (E-8) is a mouse monoclonal antibody raised against amino acids 334-422 mapping within an internal region of SH2-B of human origin.

PRODUCT

Each vial contains 200 μg lgG_1 kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

SH2-B (E-8) is available conjugated to agarose (sc-393395 AC), 500 μ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-393395 HRP), 200 μ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-393395 PE), fluorescein (sc-393395 FITC), Alexa Fluor® 488 (sc-393395 AF488), Alexa Fluor® 546 (sc-393395 AF546), Alexa Fluor® 594 (sc-393395 AF594) or Alexa Fluor® 647 (sc-393395 AF647), 200 μ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-393395 AF680) or Alexa Fluor® 790 (sc-393395 AF790), 200 μ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

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APPLICATIONS

SH2-B (E-8) is recommended for detection of SH2-B of human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for SH2-B siRNA (h): sc-44095, SH2-B shRNA Plasmid (h): sc-44095-SH and SH2-B shRNA (h) Lentiviral Particles: sc-44095-V.

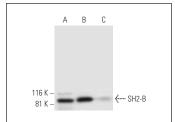
Molecular Weight of SH2-B isoforms: 80-90 kDa.

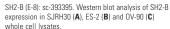
Positive Controls: ES-2 cell lysate: sc-24674, SJRH30 cell lysate: sc-2287 or OV-90 whole cell lysate: sc-364191.

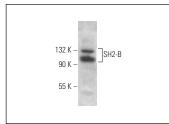
RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-lgG κ BP-HRP: sc-516102 or m-lgG κ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz MarkerTM Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-lgG κ BP-FITC: sc-516140 or m-lgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

DATA







SH2-B (E-8): sc-393395. Western blot analysis of SH2-B expression in A-673 whole cell lysate.

SELECT PRODUCT CITATIONS

 Lundby, A., et al. 2019. Oncogenic mutations rewire signaling pathways by switching protein recruitment to phosphotyrosine sites. Cell 179: 543-560.e26.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

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