

## SH2-B $\alpha/\beta/\gamma$ (E-20): sc-10827

### BACKGROUND

SH2- $\beta$  is a component of the signaling network involved in the regulation of cell shape and movement. SH2- $\beta$  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- $\beta$  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. The isoform SH2- $\beta$ b contributes to the regulation of the actin cytoskeleton as it associates with various tyrosine kinases in response to growth factor stimulation. Following PDGF stimulation, SH2- $\beta$ b can directly interact with the PDGF receptor (PDGFR) where it is phosphorylated on tyrosine residues and functions as a signaling protein for the PDGFR pathway. In addition, SH2- $\beta$ b is also a substrate for JAK2 and, thereby, mediates the cytoskeletal reorganization that is induced by the signaling pathways of various growth factors.

### CHROMOSOMAL LOCATION

Genetic locus: SH2B1 (human) mapping to 16p11.2; Sh2b1 (mouse) mapping to 7 F3.

### SOURCE

SH2-B  $\alpha/\beta/\gamma$  (E-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of SH2-B  $\beta$  of rat origin.

### PRODUCT

Each vial contains 200  $\mu$ g IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-10827 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

### RESEARCH USE

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

### APPLICATIONS

SH2-B  $\alpha/\beta/\gamma$  (E-20) is recommended for detection of SH2-B isoforms  $\alpha$ ,  $\beta$  and  $\gamma$  of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

SH2-B  $\alpha/\beta/\gamma$  (E-20) is also recommended for detection of SH2-B  $\alpha$ ,  $\beta$  and  $\gamma$  in additional species, including equine, canine, bovine and porcine.

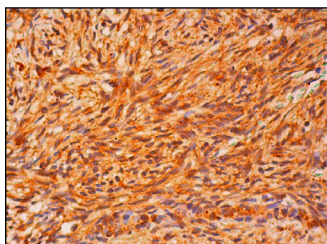
Suitable for use as control antibody for SH2-B siRNA (h): sc-44095, SH2-B siRNA (m): sc-40333, SH2-B shRNA Plasmid (h): sc-44095-SH, SH2-B shRNA Plasmid (m): sc-40333-SH, SH2-B shRNA (h) Lentiviral Particles: sc-44095-V and SH2-B shRNA (m) Lentiviral Particles: sc-40333-V.

Molecular Weight of SH2-B  $\alpha/\beta/\gamma$  isoforms: 70-95 kDa.

### RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941. 3) Immunohistochemistry: use ImmunoCruz™: sc-2053 or ABC: sc-2023 goat IgG Staining Systems.

### DATA



SH2-B  $\alpha/\beta/\gamma$  (E-20): sc-10827. Immunoperoxidase staining of formalin fixed, paraffin-embedded human ovary tissue showing cytoplasmic and nuclear staining of ovarian stroma cells.

### SELECT PRODUCT CITATIONS

- Miquet, J.G., et al. 2005. Desensitization of the JAK2/Stat5 GH signaling pathway associated with increased CIS protein content in liver of pregnant mice. *Am. J. Physiol. Endocrinol. Metab.* 289: E600-E607.
- Miquet, J.G., et al. 2005. Increased SH2-B  $\beta$  content and membrane association in transgenic mice overexpressing GH. *J. Endocrinol.* 185: 301-306.
- Miquet, J.G., et al. 2005. Increased sensitivity to GH in liver of Ames dwarf (Prop1<sup>df</sup>/Prop1<sup>df</sup>) mice related to diminished CIS abundance. *J. Endocrinol.* 187: 387-397.

### STORAGE

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

### PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) or our catalog for detailed protocols and support products.

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Satisfaction  
Guaranteed

Try **SH2-B (E-8): sc-393395** or **SH2-B  $\alpha/\beta/\gamma/\delta$  (C-11): sc-514142**, our highly recommended monoclonal alternatives to SH2-B  $\alpha/\beta/\gamma$  (E-20).