bsm-33192M

GAP43 Mouse mAb

[Primary Antibody]

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– DATASHEET –

Host: Mouse Isotype: IgG
Clonality: Monoclonal CloneNo.: 6F11
GeneID: 2596 SWISS: P17677

Target: GAP43

Purification: affinity purified by Protein G

Concentration: 1mg/ml

Storage: Size: 50ul/100ul/200ul

0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50%

Glycerol.

Size: 200ug (PBS only)

0.01M PBS

Shipped at 4°C. Store at -20°C for one year. Avoid repeated

freeze/thaw cycles.

Background: The protein encoded by this gene has been termed a 'growth' or

'plasticity' protein because it is expressed at high levels in neuronal growth cones during development and axonal regeneration. This protein is considered a crucial component of an effective

regenerative response in the nervous system. Alternatively spliced transcript variants encoding distinct isoforms have been found for

this gene. [provided by RefSeq, Jul 2008]

Applications: WB (1:500-2000)

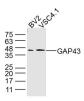
IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500) ICC/IF (1:50-200)

Reactivity: Human, Mouse, Rat

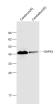
Predicted MW.: 43 kDa

Subcellular Extracellular matrix ,Cell **Location:** membrane ,Cytoplasm

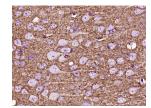
VALIDATION IMAGES



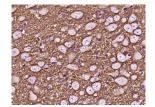
Sample: BV2 (Mouse) Lysate at 40 ug VSC4.1(Rat) Lysate at 40 ug Primary: Anti-GAP43 (bsm-33192M) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Mouse IgG at 1/20000 dilution Predicted band size: 43 kD Observed band size: 43 kD



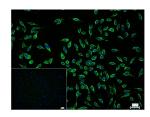
Sample: Cerebrum(Rat) Lysate at 40 ug Cerebellum(Rat) Lysate at 40 ug Primary: Anti-GAP43 (bsm-33192M) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Mouse IgG at 1/20000 dilution Predicted band size: 43-46 kD Observed band size: 43 kD



Paraformaldehyde-fixed, paraffin embedded (Mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (GAP43) Monoclonal Antibody, Unconjugated (bs-33192M-8A8) at 1:400 overnight at 4°C, followed by a conjugated secondary antibody (sp-0023) for 20 minutes and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (Rat brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (GAP43) Monoclonal Antibody,



4% Paraformaldehyde-fixed SH-SY5Y (H) cell; Triton X-100 at r.t. for 20 min; Antibody incubation with (GAP43) monoclonal Antibody, unconjugated (bsm-33192M) 1:100, 90 min at 37°C; followed by conjugated Goat Anti-Mouse IgG antibody (green, bs-40296G-FITC) at 37°C for 90 min, DAPI (blue, C02-04002) was used to stain Unconjugated (bs-33192M-8A8) at 1:400 overnight at 4°C, followed by a conjugated secondary antibody (sp-0023) for 20 minutes and DAB staining.

the cell nuclei. PBS instead of the primary antibody was used as the blank control.

- SELECTED CITATIONS -

- [IF=7.9] Mei-feng Zhang. et al. Catalpol attenuates ischemic stroke by promoting neurogenesis and angiogenesis via the SDF-1α/CXCR4 pathway. PHYTOMEDICINE. 2024 Jan;:155362 WB; Rat. 10.1016/j.phymed.2024.155362
- [IF=3.7] Sun Xuri. et al. Exogenous NT-3 Promotes Phenotype Switch of Resident Macrophages and Improves Sciatic Nerve Injury through AMPK/NF-kB Signaling Pathway. NEUROCHEM RES. 2024 Jun;:1-15 IF,WB;Rat. 38904909