bs-0707R

[Primary Antibody]

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NF-L Rabbit pAb

DATASHEET -

Host: Rabbit Isotype: IgG

Clonality: Polyclonal

GenelD: 4747 SWISS: P07196

Target: NF-L

Immunogen: KLH conjugated synthetic peptide derived from human NH-L

intermedial: 301-400/543.

Purification: affinity purified by Protein A

Concentration: 1mg/ml

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

Glycerol.

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

freeze/thaw cycles.

Background: Neurofilament light polypeptide also called NF-L; Neurofilament

triplet L protein; 68 kDa neurofilament protein. Neurofilaments usually contain three intermediate filament proteins: L, M, and H which are involved in the maintenance of neuronal caliber. The extra mass and high charge density that distinguish the neurofilament proteins from all other intermediate filament proteins are due to the tailpiece extensions. This region may form a charged scaffolding structure suitable for interaction with other neuronal components or ions. NF-L is the most abundant of the three neurofilament proteins and, as the other nonepithelial intermediate filament proteins, it can form homopolymeric 10-nm

filaments. Belongs to the intermediate filament family.

Applications: WB (1:500-2000)

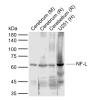
IHC-P (1:100-500) IHC-F (1:100-500) **IF** (1:100-500) Flow-Cyt (1ug/Test)

Reactivity: Human, Mouse, Rat

Predicted MW.: 68 kDa

Subcellular Location: Cytoplasm

VALIDATION IMAGES



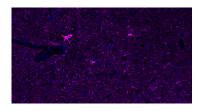
25 ug total protein per lane of various lysates (see on figure) probed with NF-L polyclonal antibody, unconjugated (bs-0707R) at 1:1000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



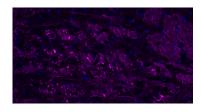
Paraformaldehyde-fixed, paraffin embedded (rat Cerebrum); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (NF-L) Polyclonal Antibody, Unconjugated (bs-0707R) at 1:100 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



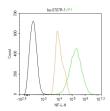
Paraformaldehyde-fixed, paraffin embedded (human cerebellum); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (NF-L) Polyclonal Antibody, Unconjugated (bs-0707R) at 1:100 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded Human Left Parietal Lobe; Antigen retrieval by



Paraformaldehyde-fixed, paraffin embedded Mouse Cerebellum; Antigen retrieval by boiling



The U251 (H) cells were fixed with 4% PFA (10 min at r.t.) and then permeabilized with 90% ice-

boiling in sodium citrate buffer (pH6.0) for 15 min; Antibody incubation with NF-L Polyclonal Antibody, Unconjugated (bs-0707R) at 1:200 overnight at 4°C. Followed by conjugated Goat Anti-Rabbit IgG antibody (Purple, bs-0295D-Cy5), DAPI (blue, C02-04002) was used to stain the cell nuclei.

in sodium citrate buffer (pH6.0) for 15 min; Antibody incubation with NF-L Polyclonal Antibody, Unconjugated (bs-0707R) at 1:200 overnight at 4°C. Followed by conjugated Goat Anti-Rabbit IgG antibody (Purple, bs-0295D-Cy5), DAPI (blue, C02-04002) was used to stain the cell nuclei. cold methanol for 20 min at -20°C, the cells then were incubated in 5%BSA to block non-specific protein-protein interactions (30 min at r.t.). Primary Antibody (green): Rabbit Anti-NF-L antibody (bs-0707R): 1 µg/10^6 cells; Isotype Control (orange): Rabbit IgG (bs-0295P). Blank control (black): PBS. Acquisition of 20,000 events was performed.