bs-0044R

DATACHEET

[Primary Antibody]

BNP Rabbit pAb



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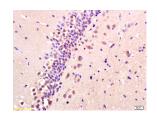
| - DATASHE | | | 1 |
|---|--|--|---|
| | Rabbit | Isotype: IgG | Applications: IHC-P (1:100-500) |
| Clonality: Polyclonal | | | IHC-F (1:100-500) IF (1:100-500) |
| GenelD | 4879 | SWISS: P16860 | |
| Target | BNP | | Reactivity: Mouse, Rat (predicted: Human |
| Immunogen | KLH conjugated synth 103-120/134. | etic peptide derived from human BNP 32: | |
| Purification: affinity purified by Protein A | | | Predicted MW.: ^{3.5 kDa} |
| Concentration | 1mg/ml | | MW.: 000 M2 4 |
| 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. | | | Subcellular Location: Secreted |
| Background | peptide, is a hormone ventricles in response ventricular blood volu The 32-amino acid po 76-amino acid N-term proBNP (BNPT), which binds to and activates and to a lesser extent peptide (ANP) but with life of BNP, however, i | lypeptide BNP is secreted attached to a ninal fragment in the prohormone called NT- n is biologically inactive. Once released, BNP s the atrial natriuretic factor receptor NPRA, NPRB, in a fashion similar to atrial natriuretic h 10-fold lower affinity. The biological half- s twice as long as that of ANP, and that of NT r, making these peptides better targets than | |

— VALIDATION IMAGES -

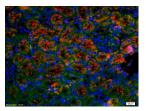


ANP for diagnostic blood testing.

Tissue/cell: rat brain tissue; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-BNP Polyclonal Antibody, Unconjugated(bs-0044R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



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Tissue/cell: mouse kidney tissue;4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-BNP Polyclonal Antibody, Unconjugated(bs-0044R) 1:200, overnight at 4°C; The secondary antibody was Goat Anti-Rabbit IgG, Cy3 conjugated(bs-0295G-Cy3)used at 1:200 dilution for 40 minutes at 37°C. DAPI(5ug/ml,blue,C-0033) was used to stain the cell nuclei