bs-1500R

[Primary Antibody]

Cathepsin B Rabbit pAb



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– DATASHEI	FT		400-901-9800
Host	: Rabbit	Isotype: IgG	Applications: WB (1:500-2000)
Clonality	: Polyclonal		IHC-P (1:100-500)
GenelD	:1508	SWISS: P07858	IF (1:100-500)
Target	: Cathepsin B		Flow-Cyt (1µg/Test)
Immunogen	: KLH conjugated s B heavy chain: 25	synthetic peptide derived from human Cath 51-339/339.	hepsin Reactivity: Human, Mouse, Rat
Purification	: affinity purified b	by Protein A	
Concentration	:1mg/ml		Predicted
Storage	: 0.01M TBS (pH7.4 Glycerol. Shipped at 4°C. S freeze/thaw cycle	4) with 1% BSA, 0.02% Proclin300 and 50% Store at -20°C for one year. Avoid repeated es.	Subcellular Location: Cytoplasm
Background	The protein enco proteinase comp chains, both prote known as amyloi the proteolytic p Incomplete prote a causative facto dementia. Overes member of the p esophageal aden transcript variant this gene. [provid	ided by this gene is a lysosomal cysteine iosed of a dimer of disulfide-linked heavy ar duced from a single protein precursor. It is a d precursor protein secretase and is involve rocessing of amyloid precursor protein (APF eolytic processing of APP has been suggester in Alzheimer disease, the most common co xpression of the encoded protein, which is a eptidase C1 family, has been associated wit nocarcinoma and other tumors. At least five ts encoding the same protein have been fou ded by RefSeq, Jul 2008]	nd light also ed in P). ed to be cause of a th e und for
- VALIDATIO	ON IMAGES —		



Sample: Spleen (Mouse) Lysate at 40 ug Primary: Anti-Cathepsin B (bs-1500R) at 1/300 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 46/28 kD Observed band size: 30 kD



Tissue/cell: Human esophageal carcinoma; 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-Cathepsin B Polyclonal Antibody, Unconjugated(bs-1500R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Blank control: RSC96(blue). Primary Antibody:Rabbit Anti- Cathepsin B antibody(bs-1500R), Dilution: 1µg in 100 µL 1X PBS containing 0.5% BSA; Isotype Control Antibody: Rabbit IgG(orange) ,used under the same conditions); Secondary Antibody: Goat anti-rabbit IgG-PE(white blue), Dilution: 1:200 in 1 X PBS containing 0.5% BSA. Protocol The cells were fixed with 2% paraformaldehyde (10 min) . Antibody (bs-1500R, $5\mu g$ /1x10^6 cells) were incubated for 30 min on the ice, followed by 1 X PBS containing 0.5% BSA + 1 0% goat serum (15 min) to block non-specific protein-protein interactions. Then the Goat Anti-rabbit IgG/PE antibody was added into the blocking buffer mentioned above to react with the primary antibody of bs-1500R at 1/200 dilution for 30 min on ice. Acquisition of 20,000 events was performed.

- SELECTED CITATIONS -

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- [IF=4.372] Lin Tianji. et al. Methylmercury induces lysosomal membrane permeabilization through JNK-activated Bax lysosomal translocation in neuronal cells. Toxicol Lett. 2022 Jan;: IF ;Human. 34999165
- [IF=2.438] Liru Li. et al. Rapamycin Pretreatment Alleviates Cerebral Ischemia/Reperfusion Injury in Dose-Response Manner Through Inhibition of the Autophagy and NFkB Pathways in Rats:. Dose-Response. 2020;18(3): WB,IF ;Rat. 32874166
- [IF=0.18] Liu, B., et al. "Autophagy activation aggravates neuronal injury in the hippocampus of vascular dementia rats." Neural Regeneration Research 9.13 (2014): 1288. IHC ;="Rat". 25221581