bs-2591R

- DATASHEET -

[Primary Antibody]

Fbx32 Rabbit pAb



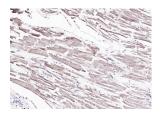
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| DATASITLET | | |
|--|--|--|
| Host: Rabbit | Isotype: IgG | Applications: IHC-P (1:100-500) |
| Clonality: Polyclonal | | IHC-F (1:100-500) IF (1:100-500) |
| GenelD: 114907 | SWISS: Q969P5 | Flow-Cyt (0.2ug/test) |
| Target: Fbx32 | | Reactivity: Human, Rat |
| Immunogen: KLH conjugated synthetic peptide derived from human MAFbx: 31-130/355. | | (predicted: Mouse, Rabbit, Pig, Cow, Dog, Horse) |
| Purification: affinity purified by | Protein A | |
| Concentration: 1mg/ml | | Predicted MW.: ^{42 kDa} |
| 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: ^{Cytoplasm} ,Nucleus |
| ubiquitin-mediate highly expressed o this gene were fou recognize and bind | quitin ligase that initiates ATP dependent d proteolysis and promotes muscle atrophy. It is luring muscle atrophy, whereas mice deficient ir nd to be resistant to atrophy. It is also thought t d to some phosphorylated proteins and promote n and degradation during skeletal muscle | D |

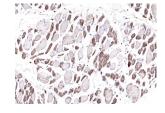
atrophy. Fbx32 interacts with MyoD by ubiquitination via a sequence found in transcriptional coactivators and therefore may play an important role in the course of muscle differentiation by

determining the abundance of MyoD.

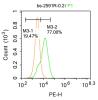
- VALIDATION IMAGES



Paraformaldehyde-fixed, paraffin embedded (human heart); 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 (Fbx32) Polyclonal Antibody, Unconjugated (bs-2591R) at 1:50 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Paraformaldehyde-fixed, paraffin embedded (human skeletal muscle); 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 (Fbx32) Polyclonal Antibody, Unconjugated (bs-2591R) at 1:50 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Blank control: Hela. Primary Antibody (green line): Rabbit Anti-Fbx32 antibody (bs-2591R) Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-PE Dilution: 1µg /test. Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at-20°C. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

- SELECTED CITATIONS -

- [IF=6.684] Ma X. et al. Endoplasmic Reticulum Stress Is Involved in Muscular Pathogenesis in Idiopathic Inflammatory Myopathies.. Front Cell Dev Biol. 2022 Feb;10:791986-791986 IHC ;Human. 35237595
- [IF=5.62] Lin, Xuzhu, et al. "Hindlimb Immobilization, but not Castration, Induces Reduction of Undercarboxylated

Osteocalcin Associated with Muscle Atrophy in Rats." Journal of Bone and Mineral Research (2016). WB ;="Rat". 27291707

• [IF=3.412] Zhang Quan-Bing. et al. Effect of Electrical Stimulation on Disuse Muscular Atrophy Induced by Immobilization: Correlation with Upregulation of PERK Signal and Parkin-Mediated Mitophagy. AM J PHYS MED REHAB. 2023 Jan;:10.1097/PHM.00000000002182 IHC ;Rabbit. 36630294