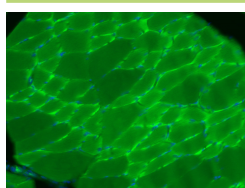


Desmuslin Rabbit pAb

- Catalog Number: bs-8555R
Target Protein: Desmuslin
Concentration: 1mg/ml
Form: Liquid
Host: Rabbit
Clonality: Polyclonal
Isotype: IgG
Applications: IHC-P (1:100-500), IHC-F (1:100-500), IF (1:200-800)
Reactivity: Rat (predicted:Human, Mouse, Rabbit, Pig, Cow)
Predicted MW: 172 kDa
Subcellular Locations: Cell membrane ,Nucleus
Entrez Gene: 23336
Swiss Prot: O15061
Source: KLH conjugated synthetic peptide derived from human Desmuslin: 501-650/1565.
Purification: affinity purified by Protein A
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: The protein encoded by this gene is an intermediate filament (IF) family member. IF proteins are cytoskeletal proteins that confer resistance to mechanical stress and are encoded by a dispersed multigene family. This protein has been found to form a linkage between desmin, which is a subunit of the IF network, and the extracellular matrix, and provides an important structural support in muscle. Two alternatively spliced variants encoding different isoforms have been described for this gene. [provided by RefSeq, Jul 2008].

VALIDATION IMAGES



Paraformaldehyde-fixed, paraffin embedded (Rat 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 (Desmuslin) Polyclonal Antibody, Unconjugated (bs-8555R) at 1:400 overnight at 4°C, followed by a conjugated Goat Anti-Rabbit IgG antibody (bs-0295G-FITC) for 90 minutes, and DAPI for nuclei staining.

PRODUCT SPECIFIC PUBLICATIONS

[IF=15.887] Weihl, Conrad C.. et al. Loss of function variants in DNAJB4 cause a myopathy with early respiratory failure. ACTA NEUROPATHOL. 2022 Oct;;1-17 WB ; Mouse . 36264506

[IF=6.39] Bengoechea, Rocio, et al. "Myofibrillar disruption and RNA binding protein aggregation in a mouse model of limb girdle muscular dystrophy 1D." Human Molecular Genetics (2015): ddv363. WB ; ="Mouse" . 26362252

[IF=6.183] Jipeng Jiang. et al. 3D printing collagen/heparin sulfate scaffolds boost neural network reconstruction and motor function recovery after traumatic brain injury in canine. Biomater Sci-Uk. 2020 Nov;8(22):6362-6374 IF ; Dog . 33026366

[IF=4.902] Bengoechea et al. Myofibrillar disruption and RNA-binding protein aggregation in a mouse model of limb-girdle muscular dystrophy 1D. (2015) Hum.Mol.Genet. 24:6588-602 WB ; Mouse . 26362252