



# Desmin Rabbit mAb

<b>Catalog No</b>	YP-rAb-17506
<b>Isotype</b>	IgG
<b>Reactivity</b>	Human,Mouse,Rat
<b>Applications</b>	WB,IHC,IF,ELISA
<b>Gene Name</b>	DES
<b>Protein Name</b>	CMD1I;CSM1;CSM2;DES;DESM_HUMAN;Desmin;FLJ12025;FLJ39719;FLJ41013;FLJ41793;Intermediate filament protein;OTTHUMP00000064865
<b>Purification Process</b>	Protein A
<b>Specificity</b>	Endogenous
<b>Formulation</b>	PBS, 50% glycerol, 0.05% Proclin 300, 0.05%BSA
<b>Source</b>	Monoclonal, Rabbit,IgG
<b>Dilution</b>	IHC 1:1000-1:5000; WB 1:20000-1:50000; IF 1:200-1:1000; ELISA 1:5000-1:20000; Note: For IHC, we suggest antigen retrieval with TE buffer pH 9.0
<b>Concentration</b>	0.5 mg/ml
<b>Purity</b>	≥90%
<b>Storage Stability</b>	-15° C to -25° C/1 year(Do not lower than -25° C)
<b>Synonyms</b>	CMD1I ; CSM1 ; CSM2 ; DES ; DESM_HUMAN ; Desmin ; FLJ12025 ; FLJ39719 ; FLJ41013 ; FLJ41793 ; Intermediate filament protein ; OTTHUMP00000064865
<b>Observed Band</b>	53kD
<b>Calculated Molecular Weight</b>	53kD
<b>Cell Pathway</b>	Cytoplasmic
<b>Tissue Specificity</b>	Appendix/ Colon
<b>Function</b>	Disease:Defects in DES are the cause of cardiomyopathy dilated type 1I (CMD1I) [MIM:604765]. Dilated cardiomyopathy is a disorder characterized by ventricular dilation and impaired systolic function, resulting in congestive heart failure and arrhythmia. Patients are at risk of premature death.,Disease:Defects in DES are the cause of desmin-related cardio-skeletal myopathy (CSM) [MIM:601419]; also known as desmin-related myopathy (DRM). CSM is characterized by skeletal muscle weakness associated with cardiac conduction blocks, arrhythmias, restrictive heart failure, and by intracytoplasmic accumulation of desmin-reactive deposits in cardiac and skeletal muscle cells. A desmin-related myopathy can have a distal onset, it is then known as hereditary distal myopathy (HDM).,Disease:Defects in DES are the cause of neurogenic scapuloperoneal





syndrome Kaeser type (Kaeser syndrome) [MIM:181400]. Kaeser syndrome is an autosomal dominant disorder with a peculiar scapulo-peroneal distribution of weakness and atrophy. A large clinical variability is observed ranging from scapulo-peroneal, limb girdle and distal phenotypes with variable cardiac or respiratory involvement. Facial weakness, dysphagia and gynecomastia are frequent additional symptoms. Affected men seemingly bear a higher risk of sudden, cardiac death as compared to affected women. Histological and immunohistochemical examination of muscle biopsy specimens reveal a wide spectrum of findings ranging from near normal or unspecific pathology to typical, myofibrillar changes with accumulation of desmin. Function: Desmin are class-III intermediate filaments found in muscle cells. In adult striated muscle they form a fibrous network connecting myofibrils to each other and to the plasma membrane from the periphery of the Z-line structures. online information: Desmin entry, similarity: Belongs to the intermediate filament family, subunit: Homopolymer.

#### Background

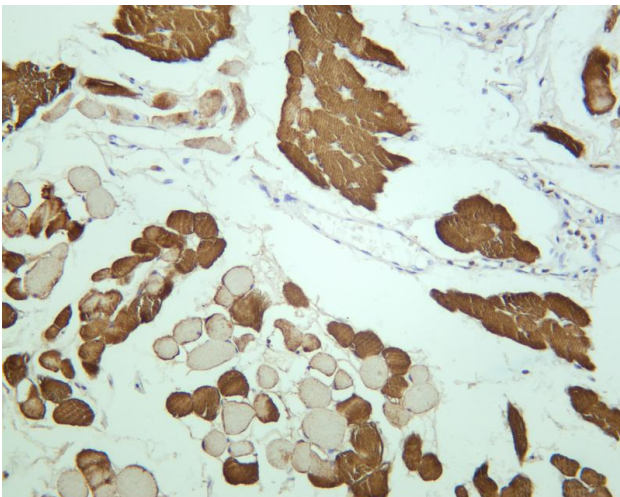
This gene encodes a muscle-specific class III intermediate filament. Homopolymers of this protein form a stable intracytoplasmic filamentous network connecting myofibrils to each other and to the plasma membrane. Mutations in this gene are associated with desmin-related myopathy, a familial cardiac and skeletal myopathy (CSM), and with distal myopathies. [provided by RefSeq, Jul 2008],

#### matters needing attention

Avoid repeated freezing and thawing!

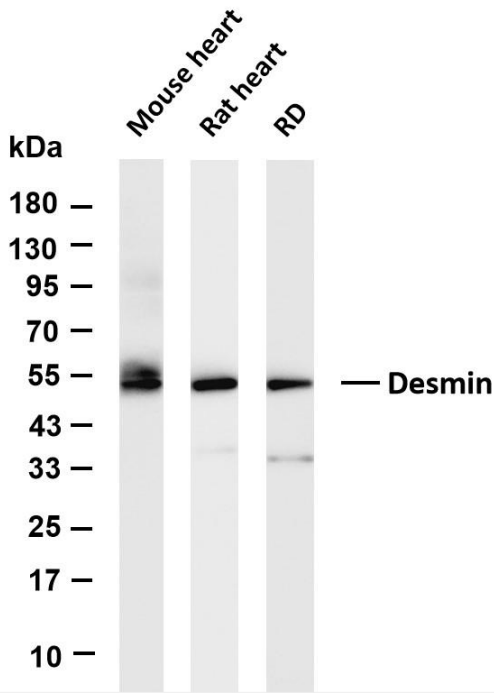
#### Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

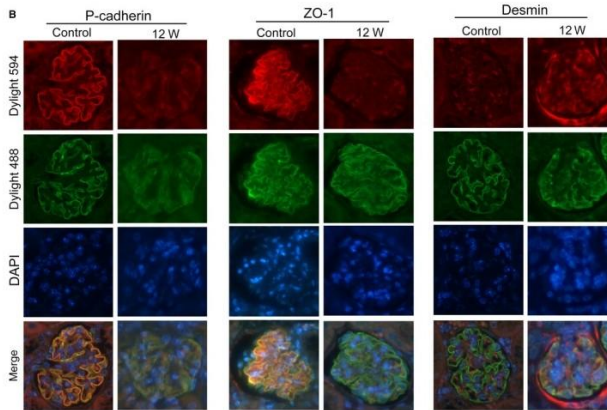


Human skeletal muscle was stained with anti-Desmin rabbit antibody

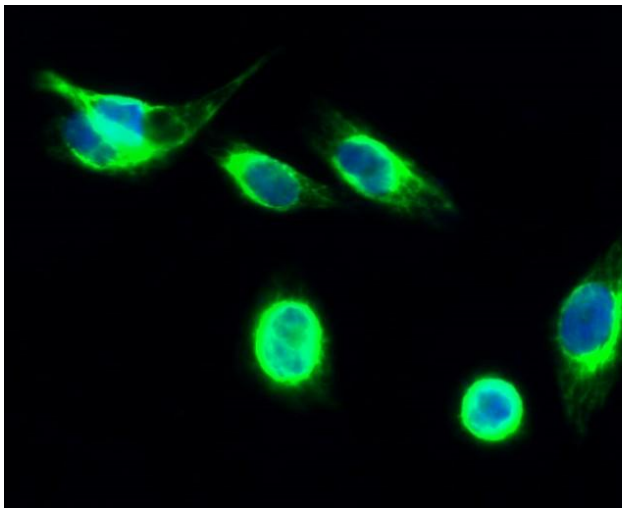




Various whole cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with anti-Desmin antibody. The HRP-conjugated Goat anti-Rabbit IgG(H + L) antibody was used to detect the antibody. Lane 1: Mouse heart Lane 2: Rat heart Lane 3: RD Predicted band size: 53kDa Observed band size: 53kDa



Hu, Mengsi, et al. "Lnc RNA MALAT 1 is dysregulated in diabetic nephropathy and involved in high glucose - induced podocyte injury via its interplay with  $\beta$  - catenin." *Journal of cellular and molecular medicine* 21.11 (2017): 2732-2747.



Immunofluorescence analysis of Hela cell. 1, Desmin Monoclonal Antibody(green) was diluted at 1:200(4° overnight). 2, Goat Anti Rabbit Alexa Fluor 488 Catalog:RS3211 was diluted at 1:1000(room temperature, 50min). 3 DAPI(blue) 10min.

