

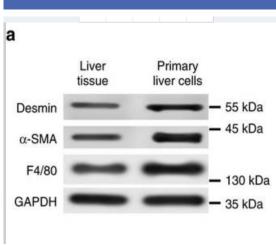
Desmin Polyclonal Antibody

Catalog No :	YT1326		
Reactivity :	Human;Mouse;Rat		
Applications :	WB;IHC;IF;ELISA		
Target :	Desmin		
Fields :	>>Hypertrophic cardiomyopathy;>>Arrhythmogenic right ventricular cardiomyopathy;>>Dilated cardiomyopathy		
Gene Name :	DES		
Protein Name :	Desmin		
Human Gene Id :	1674		
Human Swiss Prot No :	P17661		
Mouse Gene Id :	13346		
Mouse Swiss Prot No :	P31001		
Rat Gene Id :	64362		
Rat Swiss Prot No :	P48675		
Immunogen :	The antiserum was produced against synthesized peptide derived from human Desmin. AA range:421-470		
Specificity :	Desmin Polyclonal Antibody detects endogenous levels of Desmin protein.		
Formulation :	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.		
Source :	Polyclonal, Rabbit,IgG		
Dilution :	WB 1:500 - 1:2000. IHC 1:100 - 1:300. IF 1:200 - 1:1000. ELISA: 1:10000. Not yet tested in other applications.		



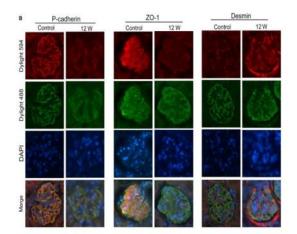
Purification :	The antibody was affinity-purified from rabbit antiserum by affinity- chromatography using epitope-specific immunogen.		
Concentration :	1 mg/ml		
oonochination .			
Storage Stability :	-15°C to -25°C/1 year(Do not lower than -25°C)		
Observed Band :	55kD		
Cell Pathway :	Hypertrophic cardiomyopathy (HCM);Arrhythmogenic right ventricular cardiomyopathy (ARVC);Dilated cardiomyopathy;		
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],		
Function :	disease:Defects in DES are the cause of cardiomyopathy dilated type 11 (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].		
Subcellular	Cytoplasmic		
Location : Expression :	Muscle,Skeletal muscle,		
Tag :	orthogonal,hot		
Sort :	267		
No4 :	1		
Host :	Rabbit		
Modifications :	Unmodified		



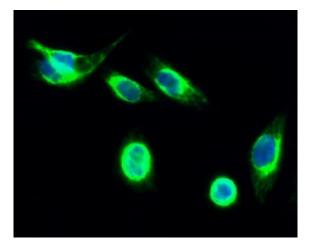


Products Images

Zhang, Haiyang, et al. "Exosome-delivered EGFR regulates liver microenvironment to promote gastric cancer liver metastasis." Nature communications 8 (2017): 15016.

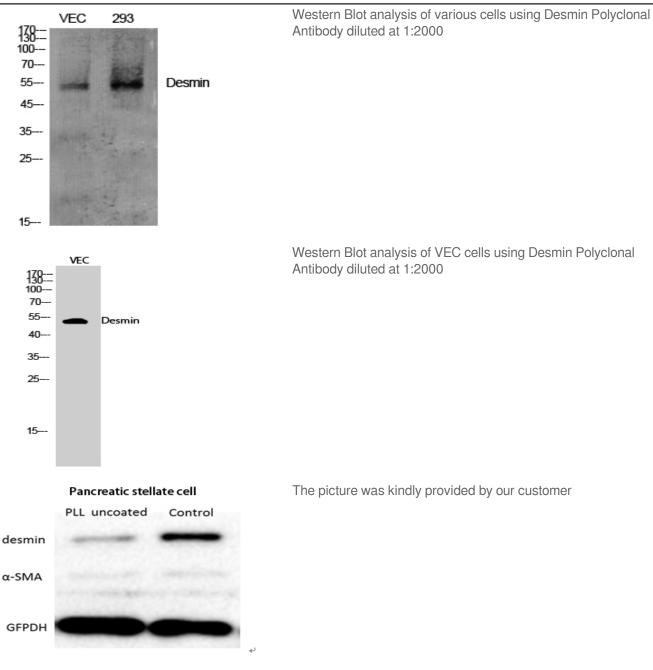


Hu, Mengsi, et al. "Lnc RNA MALAT 1 is dysregulated in diabetic nephropathy and involved in high glucose-induced podocyte injury via its interplay with β -catenin." Journal of cellular and molecular medicine 21.11 (2017): 2732-2747.



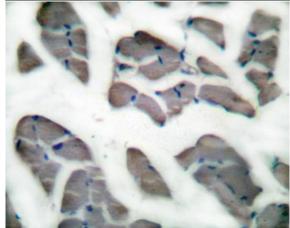
Immunofluorescence analysis of Hela cell. 1,Desmin Polyclonal Antibody(green) was diluted at 1:200(4° overnight). 2, Goat Anti Rabbit Alexa Fluor 488 Catalog:RS3211 was diluted at 1:1000(room temperature, 50min). 3 DAPI(blue) 10min.





Union Hospital of Wuhan





Immunohistochemistry analysis of paraffin-embedded human skeletal muscle tissue, using Desmin Antibody. The picture on the right is blocked with the synthesized peptide.

	K562	
	-	117
	-	85
Desmin Antibody	_	
		48
	-	34
		26
	-	19
		(kD)

Western blot analysis of lysates from K562 cells, using Desmin Antibody. The lane on the right is blocked with the synthesized peptide.