



Dsg2 Polyclonal Antibody

Function	disease:Defects in DSG2 are the cause of familial arrhythmogenic right ventricular dysplasia 10 (ARVD10) [MIM:610193]; also known as arrhythmogenic right ventricular cardiomyopathy 10 (ARVC10). ARVD is an autosomal dominant disease characterized by partial degeneration of the myocardium of the right ventricle, electrical instability, and sudden death. It is clinically defined by electrocardiographic and angiographic criteria; pathologic findings, replacement of ventricular myocardium with fatty and fibrous elements, preferentially involve the right ventricular free wall.,domain:Calcium may be bound by the cadherin-like repeats .,function:Component of intercellular desmosome junctions. Involved in the interaction of plaque proteins and intermediate filaments mediating cell-cell adhesion.,similarity:Contains 4 cadherin domains.,tissue specificity:All of the
Tissue Specificity	All of the tissues tested and carcinomas.
Cell Pathway	Cell membrane ; Single-pass type I membrane protein. Cell junction, desmosome.
Observed Band	140kD
Synonyms	DSG2; CDHF5; Desmoglein-2; Cadherin family member 5; HDGC
Storage Stability	-20°C/1 year
Purity	≥90%
Concentration	1 mg/ml
Dilution	Western Blot: 1/500 - 1/2000. ELISA: 1/40000. Not yet tested in other applications.
Purification	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
Source	Polyclonal, Rabbit,IgG
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Specificity	Dsg2 Polyclonal Antibody detects endogenous levels of Dsg2 protein.
Immunogen	The antiserum was produced against synthesized peptide derived from human DSG2. AA range:401-450
Protein Name	Desmoglein-2
Gene Name	DSG2
Applications	WB;ELISA
Reactivity	Human;Rat;Mouse;
Isotype	IgG
Catalog No	BYab-17016

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tissues tested and carcinomas.,

This gene encodes a member of the desmoglein family and cadherin cell adhesion molecule superfamily of proteins. Desmogleins are calcium-binding transmembrane glycoprotein components of desmosomes, cell-cell junctions between epithelial, myocardial, and other cell types. The encoded preproprotein is proteolytically processed to generate the mature glycoprotein. This gene is present in a gene cluster with other desmoglein gene family members on chromosome 18. Mutations in this gene have been associated with arrhythmogenic right ventricular dysplasia, familial, 10. [provided by RefSeq, Jan 2016],

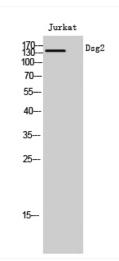
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

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

Products Images



Western Blot analysis of Jurkat cells using Dsg2 Polyclonal Antibody

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