



Catenin-γ Monoclonal Antibody

Catalog No	YP-mAb-16942
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	JUP
Immunogen	The antiserum was produced against synthesized peptide derived from human Catenin-gamma. AA range:696-745
Specificity	Catenin-γ Monoclonal Antibody detects endogenous levels of Catenin-γ protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	Beta catenin;Beta-catenin;Cadherin associated protein;Catenin (cadherin associated protein);beta 1;88kDa;Catenin beta 1;Catenin beta-1;CATNB;CHBCAT;CTNB1_HUMAN;CTNNB;CTNNB1;DKFZp686D2253;FLJ2566;FLJ37923;OTTHUMP16282;OTTHUMP165222;OTTHUMP165223;OTTHUMP29288;OTTHUMP29289;Catenin-γ;JUP;Junction plakoglobin;CTNNG;DP3;Catenin gamma;Desmoplakin III;Desmoplakin-3
Observed Band	82kD
Calculated Molecular Weight	82kD
Cell Pathway	Cell junction, adherens junction . Cell junction, desmosome . Cytoplasm, cytoskeleton . Membrane ; Peripheral membrane protein . Cytoplasmic in a soluble and membrane-associated form.
Tissue Specificity	Cervix carcinoma,Epidermal carcinoma,Epithelium,Leukocyte,Lung,Lung carcinoma,Place
Function	disease:Defects in JUP are the cause of familial arrhythmogenic right ventricular dysplasia type 12 (ARVD12) [MIM:611528]; also called arrhythmogenic right ventricular cardiomyopathy type 12 (ARVC12). 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.,disease:Defects in JUP are the cause of Naxos disease (NXD) [MIM:601214]. NXD is an autosomal recessive disorder combining diffuse non-epidermolytic palmoplantar keratoderma with arrhythmogenic right ventricular dysplasia/cardiomyopathy and woolly hair., function:Common junctional plaque protein. The membrane-assoc

Background

This gene encodes a major cytoplasmic protein which is the only known constituent common to submembranous plaques of both desmosomes and intermediate junctions. This protein forms distinct complexes with cadherins and desmosomal cadherins and is a member of the catenin family since it contains a distinct repeating amino acid motif called the armadillo repeat. Mutation in this gene has been associated with Naxos disease. Alternative splicing occurs in this gene; however, not all transcripts have been fully described. [provided by RefSeq, Jul 2008],

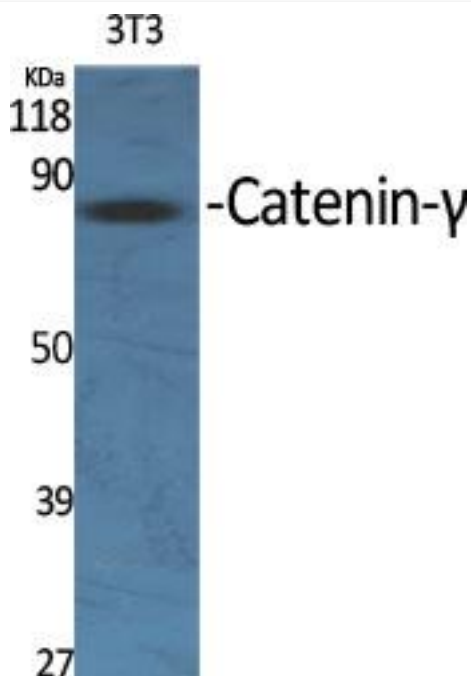
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

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

Products Images



Western Blot analysis of various cells using Catenin-γ Monoclonal Antibody