



Cytokeratin 16 Monoclonal Antibody

Catalog No	YP-mAb-03112
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	KRT16
Immunogen	The antiserum was produced against synthesized peptide derived from human Keratin 16. AA range:421-470
Specificity	Cytokeratin 16 Monoclonal Antibody detects endogenous levels of Cytokeratin 16 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	KRT16;Keratin16;keratin 16;Cytokeratin;Keratin;CK-16;Cytokeratin 16;Keratin type I cytoskeletal 16;KRT16A;type I cytoskeletal 16;Cytokeratin-16;Keratin-16; K16
Observed Band	52kD
Calculated Molecular Weight	52kD
Cell Pathway	nucleus,cytoskeleton,intermediate filament,extracellular exosome,
Tissue Specificity	Expressed in the corneal epithelium (at protein level).
Function	disease:Defects in KRT16 are a cause of pachyonychia congenita type 1 (PC1) [MIM:167200]; also known as Jadassohn-Lewandowsky syndrome. PC1 is an autosomal dominant ectodermal dysplasia characterized by hypertrophic nail dystrophy resulting in onychogryposis (thickening and increase in curvature of the nail), palmoplantar keratoderma, follicular hyperkeratosis, and oral leukokeratosis. Hyperhidrosis of the hands and feet is usually present., disease:Defects in KRT16 are a cause of unilateral palmoplantar verrucous nevus (UPVN) [MIM:144200]. UPVN is characterized by a localized thickening of the skin in parts of the right palm and the right sole.,disease:Defects in KRT16



are the cause of palmoplantar keratoderma non-epidermolytic (NEPPK) [MIM:600962]. NEPPK is a dermatological disorder characterized by focal palmoplantar keratoderma with oral, genital, and follicular lesions., disease:KRT16

Background

The protein encoded by this gene is a member of the keratin gene family. The keratins are intermediate filament proteins responsible for the structural integrity of epithelial cells and are subdivided into cytokeratins and hair keratins. Most of the type I cytokeratins consist of acidic proteins which are arranged in pairs of heterotypic keratin chains and are clustered in a region of chromosome 17q12-q21. This keratin has been coexpressed with keratin 14 in a number of epithelial tissues, including esophagus, tongue, and hair follicles. Mutations in this gene are associated with type 1 pachyonychia congenita, non-epidermolytic palmoplantar keratoderma and unilateral palmoplantar verrucous nevus. [provided by RefSeq, Jul 2008],

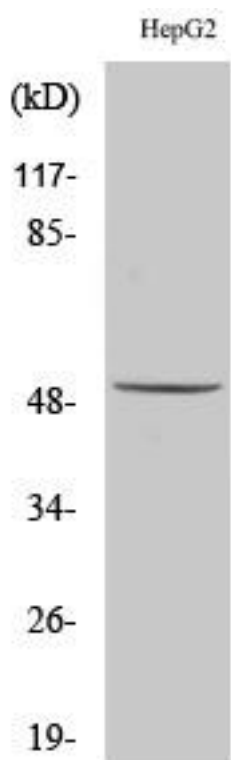
matters needing attention

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 Cytokeratin 16 Monoclonal Antibody