



Keratin(50-60kDa) mouse mAb

Catalog No	BYab-02962
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	krt5
Protein Name	
Immunogen	Purified recombinant human KRT5 beta protein fragments expressed in E.coli.
Specificity	This antibody detects endogenous levels of Keratin(50-60kDa) proteins.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse
Purification	The antibody was affinity-purified from mouse ascites by affinity-chromatography using epitope-specific immunogen.
Dilution	wb dilution 1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	58 kDa cytokeratin; CK 5; CK-5; CK5; Cytokeratin-5; Cytokeratin5; DDD; DDD1; EBS 2; EBS2; epidermolysis bullosa simplex 2 Dowling-Meara/Kobner/Weber-Cockayne types; K2C5_HUMAN; K5; Keratin 5 (epidermolysis bullosa simplex Dowling-Meara/Kobner/Weber-Cockayne types); keratin 5 (epidermolysis bullosa simplex, Dowling-Meara/Kobner/Weber-Cockayne types); Keratin 5; Keratin; keratin complex 2, basic, gene 5; Keratin Type II Cytoskeletal 5; keratin, type II cytoskeletal 5; Keratin-5; Keratin5; KRT 5; KRT 5A; KRT5; KRT5A; type II cytoskeletal 5; Type-II keratin Kb5.
Observed Band	62kD
Cell Pathway	nucleus,cytoplasm,mitochondrion,cytosol,intermediate filament,plasma membrane,membrane,keratin filament,extracellular exosome,
Tissue Specificity	Expressed in corneal epithelium (at protein level).

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Function

disease:Defects in KRT5 are a cause of epidermolysis bullosa simplex Dowling-Meara type (DM-EBS) [MIM:131760]. DM-EBS is a severe form of intraepidermal epidermolysis bullosa characterized by generalized herpetiform blistering, milia formation, dystrophic nails, and mucous membrane involvement.,disease:Defects in KRT5 are a cause of epidermolysis bullosa simplex Koebner type (K-EBS) [MIM:131900]. K-EBS is a form of intraepidermal epidermolysis bullosa characterized by generalized skin blistering. The phenotype is not fundamentally distinct from the Dowling-Meara type, although it is less severe.,disease:Defects in KRT5 are a cause of epidermolysis bullosa simplex Weber-Cockayne type (WC-EBS) [MIM:131800]. WC-EBS is a form of intraepidermal epidermolysis bullosa characterized by blistering limited to palmar and plantar areas of the skin.,disease:Defects in KRT5 are the cause of Dowling-D

Background

keratin 5(KRT5) Homo sapiens The protein encoded by this gene is a member of the keratin gene family. The type II cytokeratins consist of basic or neutral proteins which are arranged in pairs of heterotypic keratin chains coexpressed during differentiation of simple and stratified epithelial tissues. This type II cytokeratin is specifically expressed in the basal layer of the epidermis with family member KRT14. Mutations in these genes have been associated with a complex of diseases termed epidermolysis bullosa simplex. The type II cytokeratins are clustered in a region of chromosome 12q12-q13. [provided by RefSeq, Jul 2008],

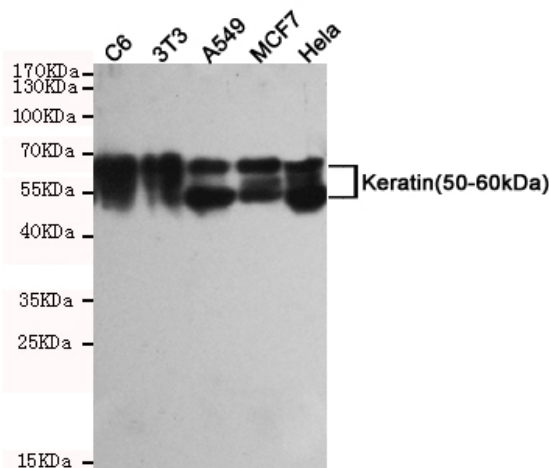
matters needing attention

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 detection of Keratin(50-60kDa) in C6,3T3,A549,MCF7 and HeLa cell lysates using Keratin(50-60kDa) mouse mAb (1:2000 diluted).Predicted band size50~60kDa.Observed band size:50~60kDa.

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