

## Cytokeratin 10 (ABT056) IHC kit

CatalogNo: IHCM6857

### Key Features

Host Species

Mouse

Reactivity
• Human,

Applications
• IHC

IsotypeIgG1,Kappa

#### **Recommended Dilution Ratios**

#### **Storage**

Storage\* 2°C to 8°C/1 year

#### **Basic Information**

| Clonality | Monoclonal |
|-----------|------------|
|-----------|------------|

Clone Number ABT056

#### Immunogen Information

| Immunogen   | Synthesized peptide derived from human CK10 AA range: 500-584   |
|-------------|---|
| Specificity | The antibody can specifically recognize human CK10 protein, and shows no cross reaction with CK4, 5, 6, 7, 8, 14, 15, 18, 19. |

#### Target Information

Gene name KRT10 KPP

#### Protein Name Cytokeratin-10

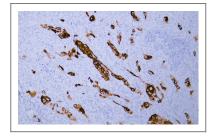
| Organism | Gene ID      | UniProt ID     |
|----------|--------------|----------------|
| Human    | <u>3858;</u> | <u>P13645;</u> |

### Cellular Cytoplasmic, Membranous Localization

# **Tissue specificity** Seen in all suprabasal cell layers including stratum corneum. Expressed on the surface of lung cell lines (PubMed:19627498).

**Function** Disease:Defects in KRT10 are a cause of bullous congenital ichthyosiform erythroderma (BCIE) [MIM:113800]; also known as epidermolytic hyperkeratosis (EHK) or bullous erythroderma ichthyosiformis congenita of Brocg, BCIE is an autosomal dominant skin disorder characterized by widespread blistering and an ichthyotic erythroderma at birth that persist into adulthood. Histologically there is a diffuse epidermolytic degeneration in the lower spinous layer of the epidermis. Within a few weeks from birth, erythroderma and blister formation diminish and hyperkeratoses develop., Disease: Defects in KRT10 are a cause of epidermal nevus epidermolytic hyperkeratotic type [MIM:600648]. Epidermal nevi affect about 1 in 1,000 people. They appear at or shortly after birth as localized lines of epidermal thickening. The extent of skin involvement varies widely., Disease: Defects in KRT10 are a cause of ichthyosis annular epidermolytic (AEI) [MIM:607602]; also known as cyclic ichthyosis with epidermolytic hyperkeratosis. AEI is a skin disorder resembling bullous congenital ichthyosiform erythroderma. Affected individuals present with bullous ichthyosis in early childhood and hyperkeratotic lichenified plagues in the flexural areas and extensor surfaces at later ages. The feature that distinguishes AEI from BCIE is dramatic episodes of flares of annular polycyclic plagues with scale, which coalesce to involve most of the body surface and can persist for several weeks or even months.,miscellaneous:There are two types of cytoskeletal and microfibrillar keratin: I (acidic; 40-55 kDa) and II (neutral to basic; 56-70 kDa).,online information:Keratin-10 entry,polymorphism:A number of alleles are known that mainly differ in the Gly-rich region (positions 490-560)., similarity: Belongs to the intermediate filament family., subunit: Heterotetramer of two type I and two type II keratins. keratin-10 is generally associated with keratin-1.,tissue specificity:Seen in all suprabasal cell layers including stratum corneum.,

#### Validation Data



Human cervix tissue was stained with Anti-Cytokeratin 10 (ABT056) Antibody



Human skin tissue was stained with Anti-Cytokeratin 10 (ABT056) Antibody

### **Contact information**

| Orders:    | order@immunoway.com                      |
|------------|--|
| Support:   | tech@immunoway.com                       |
| Telephone: | 408-747-0189 (USA) 400-8787-807(China)   |
| Website:   | http://www.immunoway.com                 |
| Address:   | 2200 Ringwood Ave San Jose, CA 95131 USA |



Please scan the QR code to access additional product information: **Cytokeratin 10** (ABT056) IHC kit

For Research Use Only. Not for Use in Diagnostic Procedures.

Antibody | ELISA Kits | Protein | Reagents