antibodies - online.com







anti-Keratin 36 antibody (Internal Region)

Images



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Target:

Quantity:	100 μL
Target:	Keratin 36 (KRT36)
Binding Specificity:	Internal Region
Reactivity:	Human, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This Keratin 36 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunocytochemistry (ICC), Immunofluorescence (IF)
Product Details	
Immunogen:	A synthesized peptide derived from human KRT36, corresponding to a region within the internal amino acids.
Isotype:	IgG
Specificity:	KRT36 Antibody detects endogenous levels of total KRT36.
Predicted Reactivity:	Pig,Bovine,Horse,Sheep,Dog
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).
Target Details	
-	V - V - 0.5 (VDT0.5)

Keratin 36 (KRT36)

Target Details

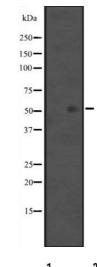
Alternative Name:	KRT36 (KRT36 Products)
Background:	Description: There are two types of hair/microfibrillar keratin, I (acidic) and II (neutral to basic). Gene: KRT36
Molecular Weight:	52 kDa
Gene ID:	8689
UniProt:	076013

Application Details

Application Notes:	WB 1:500-1:2000, IHC 1:50-1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only

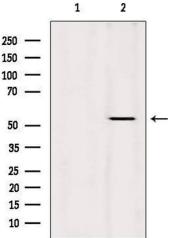
Handling

Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline, pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Expiry Date:	12 months



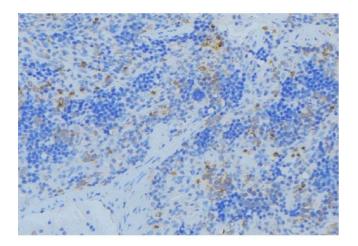
Western Blotting

Image 1. Western blot analysis of KRT36 expression in Hela cell lysate, The lane on the left is treated with the antigenspecific peptide.



Western Blotting

Image 2. Western blot analysis of extracts from Hela, using KRT36 Antibody. Lane 1 was treated with the blocking peptide.



Immunohistochemistry

Image 3. ABIN6273682 at 1/100 staining Human lymph node tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the antibody for 1.5 hours at 22¡ãC. An HRP conjugated goat anti-rabbit antibody was used as the secondary