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anti-KTI12 antibody (AA 110-204)

3 Images



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Quantity:	100 μg
Target:	KTI12
Binding Specificity:	AA 110-204
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This KTI12 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF)

Product Details

Immunogen:	Recombinant Human Protein KTI12 homolog protein (110-204AA)
Isotype:	IgG
Cross-Reactivity:	Human, Rat
Purification:	>95%, Protein G purified

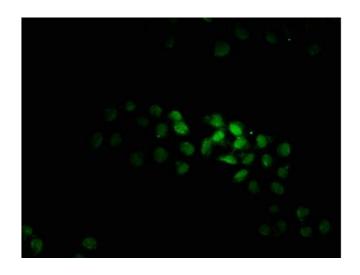
Target Details

Target:	KTI12
Alternative Name:	KTI12 (KTI12 Products)
Background:	KTI12 antibody, SBBI81 antibody, Protein KTI12 homolog antibody
UniProt:	Q96EK9

Application Details

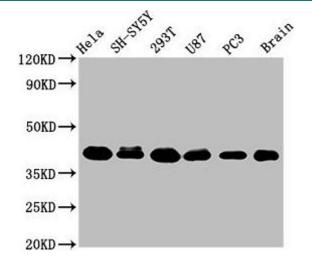
Application Notes:	Recommended dilution: WB:1:500-1:5000, IHC:1:200-1:500, IF:1:50-1:200,
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	Preservative: 0.03 % Proclin 300
	Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be
	handled by trained staff only.
Storage:	-20 °C,-80 °C
Storage Comment:	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.

Images



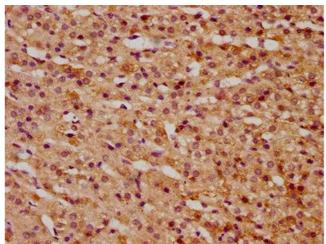
Immunofluorescence

Image 1. Immunofluorescence staining of Hela cells with ABIN7165725 at 1:100, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).



Western Blotting

Image 2. Western Blot Positive WB detected in: Hela whole cell lysate, SH-SY5Y whole cell lysate, 293T whole cell lysate, U87 whole cell lysate, PC-3 whole cell lysate, Rat brain tissue All lanes: KTI12 antibody at 3.2 μg/mL Secondary Goat polyclonal to rabbit lgG at 1/50000 dilution Predicted band size: 39 kDa Observed band size: 39 kDa



Immunohistochemistry

Image 3. IHC image of ABIN7165725 diluted at 1:300 and staining in paraffin-embedded human adrenal gland tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.