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Datasheet for ABIN7168916 anti-HTRA1 antibody (AA 23-248)

3 Images



Overview

Quantity:	100 μL
Target:	HTRA1
Binding Specificity:	AA 23-248
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This HTRA1 antibody is un-conjugated
Application:	Immunohistochemistry (IHC), ELISA, Immunofluorescence (IF)

Product Details

Immunogen:	Recombinant Human Serine protease HTRA1 protein (23-248AA)
Isotype:	lgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

Target Details

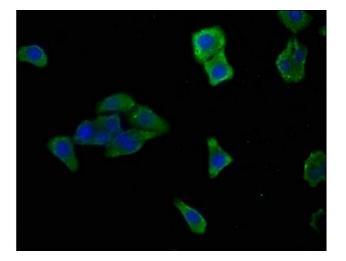
Target:	HTRA1
Alternative Name:	HTRA1 (HTRA1 Products)
Background:	Background: Serine protease with a variety of targets, including extracellular matrix proteins
	such as fibronectin. HTRA1-generated fibronectin fragments further induce synovial cells to up-

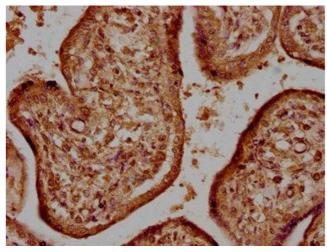
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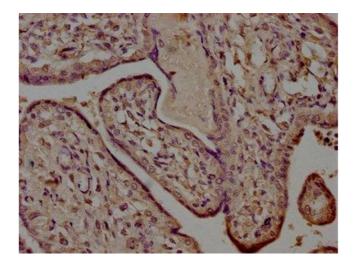
	regulate MMP1 and MMP3 production. May also degrade proteoglycans, such as aggrecan,
	decorin and fibromodulin. Through cleavage of proteoglycans, may release soluble FGF-
	glycosaminoglycan complexes that promote the range and intensity of FGF signals in the
	extracellular space. Regulates the availability of insulin-like growth factors (IGFs) by cleaving
	IGF-binding proteins. Inhibits signaling mediated by TGF-beta family members. This activity
	requires the integrity of the catalytic site, although it is unclear whether TGF-beta proteins are
	themselves degraded. By acting on TGF-beta signaling, may regulate many physiological
	processes, including retinal angiogenesis and neuronal survival and maturation during
	development. Intracellularly, degrades TSC2, leading to the activation of TSC2 downstream
	targets.
	Aliases: ARMD7 antibody, CARASIL antibody, High-temperature requirement A serine peptidase
	1 antibody, HtrA antibody, HtrA serine peptidase 1 antibody, HTRA1 antibody, HTRA1_HUMAN
	antibody, IGFBP5 protease antibody, L56 antibody, ORF480 antibody, Protease serine 11 (IGF
	binding) antibody, protease serine 11 antibody, PRSS11 antibody, Serine protease 11 antibody,
	Serine protease HTRA1 antibody, Serine protease HTRA1 precursor antibody
UniProt:	Q92743
Pathways:	Growth Factor Binding
Application Details	
Application Notes:	Recommended dilution: IHC:1:500-1:1000, IF:1:50-1:200,
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Ruffer:	Preservative: 0.03 % Proclin 300

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.

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Immunofluorescence

Image 1. Immunofluorescence staining of HepG2 cells with ABIN7168916 at 1:166, 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).

Immunohistochemistry

Image 2. IHC image of ABIN7168916 diluted at 1:500 and staining in paraffin-embedded human kidney 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 30 min 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.

Immunohistochemistry

Image 3. IHC image of ABIN7168916 diluted at 1:500 and staining in paraffin-embedded human placenta 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 30 min 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.

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