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Datasheet for ABIN7150368 anti-Id2 antibody (AA 1-134)

2 Images



Overview

Quantity:	100 µg
Target:	ld2
Binding Specificity:	AA 1-134
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This Id2 antibody is un-conjugated
Application:	ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF)

Product Details

Immunogen:	Recombinant Human DNA-binding protein inhibitor ID-2 protein (1-134AA)
Isotype:	lgG
Cross-Reactivity:	Human
Purification:	>95%, Protein G purified

Target Details

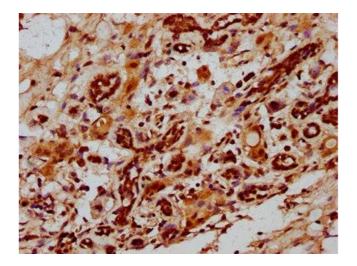
Target:	ld2
Alternative Name:	ID2 (Id2 Products)
Background:	Background: D (inhibitor of DNA binding) HLH proteins lack a basic DNA-binding domain but are
	able to form heterodimers with other HLH proteins, thereby inhibiting DNA binding. ID-2 may be

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	an inhibitor of tissue-specific gene expression.
	Aliases: bHLHb26 antibody, Cell growth inhibiting gene 8 antibody, class B basic helix loop helix
	protein 26 antibody, Class B basic helix-loop-helix protein 26 antibody, DNA binding protein
	inhibitor ID 2 antibody, DNA binding protein inhibitor ID2 antibody, DNA-binding protein inhibitor
	ID-2 antibody, GIG 8 antibody, GIG8 antibody, Helix loop helix protein ID2 antibody, ID2 antibody,
	ID2_HUMAN antibody, ID2A antibody, ID2H antibody, Inhibitor of differentiation 2 antibody,
	Inhibitor of DNA binding 2 antibody, Inhibitor of DNA binding 2, dominant negative helix loop
	helix protein antibody, MGC26389 antibody
UniProt:	Q02363

Application Details

Application Notes:	Recommended dilution: IHC:1:1000-1:2000, IF:1:200-1:500,
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.



Immunohistochemistry

Image 1. IHC image of ABIN7150368 diluted at 1:1000 and staining in paraffin-embedded human breast cancer 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.

Immunofluorescence

Image 2. Immunofluorescence staining of HepG2 cells with ABIN7150368 at 1:333, 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).

