

Product Data Sheet

Anti-HDGF Antibody

Catalog #	Source	Reactivity	Applications		
CQA2001	Rabbit	Н, М	WB, IH, IF/IC, IP		
Description	I	Rabbit polyclonal ant	body to HDGF		
Immunogen	ł	Recombinant full len	th protein of human HDGF		
Purification	-	The antibody was pur	ified by immunogen affinity chromatography.		
Specificity	I	Recognizes endogenc	us levels of HDGF protein.		
Clonality	I	Polyclonal			
Conjugation					
Form	I	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ä	and 0.01% sodium az	de.		
Dilution	V	WB (1/500 - 1/2000), I	H (1/50 - 1/200), IF/IC (1/50 - 1/200), IP (1/20 - 1/50)		
Gene Symbol	ł	HDGF			
Alternative Na	ames l	HMG1L2; Hepatoma-	derived growth factor; HDGF; High mobility group protein		
	, -	1-like 2; HMG-1L2			
Entrez Gene	ŝ	3068 (Human); 15191 (Mouse)			
SwissProt	I	P51858 (Human); P51859 (Mouse)			
Storage/Stabi	lity S	Shipped at 4 $^\circ~$ C. Upo	n delivery aliquot and store at -20 $^\circ~$ C for one year. Avoid		
	f	freeze/thaw cycles.			

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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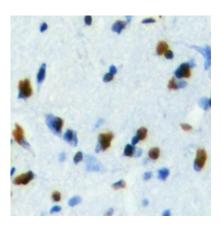
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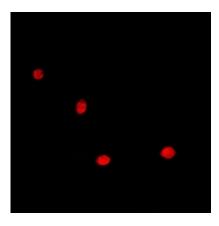
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Western blot analysis of HDGF expression in Hela (A), A549 (B), MCF7 (C) whole cell lysates. (Predicted band size: 25; 26; 28 kD; Observed band size: 37 kD)



Immunohistochemical analysis of HDGF staining in rat brain formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of HDGF staining in A549 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 $^{\circ}$ C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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