

# **Product Data Sheet**

# Anti-EXT2 Antibody

Catalog #	Source	Reactivity	Applications			
CQA2778	Rabbit	Н, М	WB, IF/IC			
Description	Rabbit	polyclonal antibody to	o EXT2			
Immunogen	Recom	binant full length prot	ein of human EXT2			
Purification	The ar	tibody was purified by	/ immunogen affinity chromatography.			
Specificity	Recogi	nizes endogenous leve	ls of EXT2 protein.			
Clonality	Polyclo	Polyclonal				
Conjugation						
Form	Liquid	in 0.42% Potassium pl	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and 0.	01% sodium azide.				
Dilution	WB (1/	500 - 1/2000), IF/IC (1/	50 - 1/200)			
Gene Symbol	EXT2					
Alternative N	ames Exosto	Exostosin-2;				
	Glucur	onosyl-N-acetylglucos	aminyl-proteoglycan/N-acetylglucosaminyl-proteoglycan			
	4-alph	a-N-acetylglucosaminy	/ltransferase; Multiple exostoses protein 2; Putative			
	tumor	suppressor protein EX	T2			
Entrez Gene	2132 (	Human); 14043 (Mous	se)			
SwissProt	Q9306	3 (Human); P70428 (N	/louse)			
Storage/Stab	<b>ility</b> Shippe	ed at 4 $^{\circ}~$ C. Upon delive	ery aliquot and store at -20 $^\circ~$ C for one year. Avoid			
	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

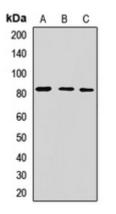
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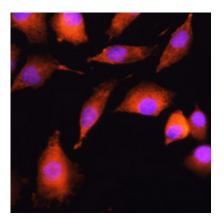


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Western blot analysis of EXT2 expression in Hela (A), SKOV3 (B), mouse liver (C) whole cell lysates. (Predicted band size: 82; 83; 85 kD; Observed band size: 82 kD)



Immunofluorescent analysis of EXT2 staining in L929 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 AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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