

## **Product Data Sheet**

### **Anti-Thrombospondin-5 Antibody**

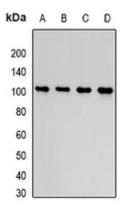
Catalog #	Source	Reactivity	Applications
CQA1275	Rabbit	H, M, R	WB, IF/IC
Description	Ra	abbit polyclonal antibody	to Thrombospondin-5
Immunogen	Re	ecombinant full length pr	otein of human Thrombospondin-5
Purification	Tł	ne antibody was purified	by immunogen affinity chromatography.
Specificity	Re	ecognizes endogenous lev	vels of Thrombospondin-5 protein.
Clonality	Po	olyclonal	
Conjugation			
Form	Li	quid in 0.42% Potassium	ohosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	ar	nd 0.01% sodium azide.	
Dilution	Ŵ	/B (1/500 - 1/2000), IF/IC (	1/50 - 1/200)
Gene Symbol	C	ОМР	
Alternative Na	ames Ca	artilage oligomeric matrix	protein; COMP; Thrombospondin-5; TSP5
Entrez Gene	13	311 (Human); 12845 (Mo	use); 25304 (Rat)
SwissProt	P4	49747 (Human); Q9R0G6	(Mouse); P35444 (Rat)
Storage/Stabi	<b>lity</b> Sł	nipped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid
	fr	eeze/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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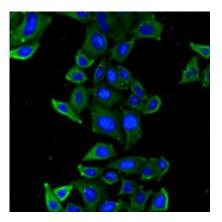




For research purposes only, not for human use

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Western blot analysis of Thrombospondin-5 expression in HepG2 (A), MCF7 (B), mouse heart (C), mouse ovary (D) whole cell lysates. (Predicted band size: 77; 82 kD; Observed band size: 110 kD)



Immunofluorescent analysis of Thrombospondin-5 staining in U2OS 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 °C in a humidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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