

Product Data Sheet

Anti-TWEAK Antibody

Catalog #	Source	Reactivity	Applications		
CPA2319	Rabbit	Н	WB, IH, IF/IC		
Description		Rabbit polyclonal antibody	r to TWEAK		
Immunogen		KLH-conjugated synthetic	peptide encompassing a sequence within the center		
		region of human TWEAK.	he exact sequence is proprietary.		
Purification		The antibody was purified	by immunogen affinity chromatography.		
Specificity		Recognizes endogenous levels of TWEAK protein.			
Clonality		Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/50 - 1/100), IF/IC (1/50 - 1/200)			
Gene Symbol		TNFSF12			
Alternative Names		APO3L; DR3LG; Tumor necrosis factor ligand superfamily member 12; APO3 ligand;			
		TNF-related weak inducer	of apoptosis; TWEAK		
Entrez Gene		407977, 8742 (Human)			
SwissProt		O43508 (Human)			
Storage/Stabi	lity	Shipped at 4°C. Upon deliv	ery aliquot and store at -20°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

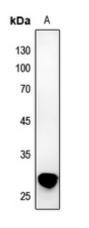
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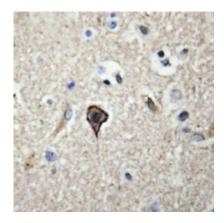
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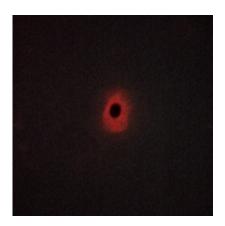
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Western blot analysis of TWEAK expression in Hela (A) whole cell lysates. (Predicted band size: 27 kD; Observed band size: 27 kD)



Immunohistochemical analysis of TWEAK staining in human 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 TWEAK staining in HeLa 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 hidified chamber. Cells were washed with PBST and incubated with a AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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