

Product Data Sheet

Anti-TNF alpha Antibody

Catalog #	Source	Reactivity	Applications			
CPA9458	Rabbit	H, M, R	WB, IH			
Description	Rat	Rabbit polyclonal antibody to TNF alpha				
Immunogen	Rec	combinant protein corre	sponding to human TNF alpha.			
Purification	The	e antibody was purified l	by immunogen affinity chromatography.			
Specificity	Rec	ognizes endogenous lev	els of TNF alpha protein.			
Clonality	Pol	yclonal				
Conjugation						
Form	Liqu	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	and	0.01% sodium azide.				
Dilution	WB	s (1/1000 - 1/2000), IH (1/	50 - 1/200)			
Gene Symbol	TN	=				
Alternative Na	imes TNI	A; TNFSF2; Tumor necro	osis factor; Cachectin; TNF-alpha; Tumor necrosis factor			
	liga	nd superfamily member	2; TNF-a			
Entrez Gene	712	24 (Human); 21926 (Mou	ıse); 24835 (Rat)			
SwissProt	P01	1375 (Human); P06804 (Mouse); P16599 (Rat)			
Storage/Stabil	ity Shi	pped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid			
	free	eze/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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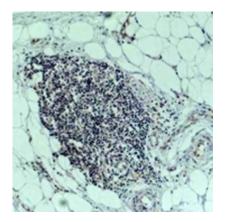
kDa A

70

For research purposes only, not for human use

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Western blot analysis of TNF alpha expression in HepG2 LPS-treated (A) whole cell lysates. (Predicted band size: 25 kD; Observed band size: 17; 25 kD)



Immunohistochemical analysis of TNF alpha staining in human breast cancer 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.

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