

# **Product Data Sheet**

### **Anti-Caspase 8 Antibody**

Catalog #	Source	Reactivity	Applications			
CPA1154	Rabbit	H, M, R	WB, IH			
Description	I	Rabbit polyclonal antibody to Caspase 8				
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center				
	I	region of human Caspase 8. The exact sequence is proprietary.				
Purification	-	The antibody was purified b	v immunogen affinity chromatography.			
Specificity	I	Recognizes endogenous leve	ls of Caspase 8 protein.			
Clonality	I	Polyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	ä	and 0.01% sodium azide.				
Dilution	,	WB (1/500 - 1/1000), IH (1/10	0 - 1/200)			
Gene Symbol	(	CASP8				
Alternative Na	ames	MCH5; Caspase-8; CASP-8; A	poptotic cysteine protease; Apoptotic protease Mch-5;			
	(	CAP4; FADD-homologous ICI	/ced-3-like protease; FADD-like ICE; FLICE; ICE-like			
	ä	apoptotic protease 5; MORT	1-associated ced-3 homolog; MACH			
Entrez Gene	8	841 (Human); 12370 (Mouse	)			
SwissProt	(	Q14790 (Human); O89110 (I	Mouse)			
Storage/Stabi	lity S	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid			
	1	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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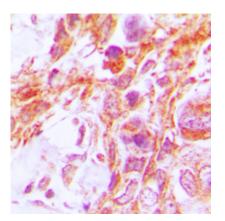
A B

С



# **Product Data Sheet**

Western blot analysis of Caspase 8 expression in DLD (A), mouse liver (B), Raw264.7 (C) whole cell lysates. (Predicted band size: 55 kD; Observed band size: 57 kD)



Immunohistochemical analysis of Caspase 8 staining in human lung 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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