

# **Product Data Sheet**

### **Anti-Caspase 3 Antibody**

Catalog #	Source	Reactivity	Applications
CPA6370	Rabbit	H, M, R, B, D, Mk	WB, IH, IF/IC
Description	Rabb	it polyclonal antibody to Cas	pase 3
Immunogen	KLH-	conjugated synthetic peptide	encompassing a sequence within the center
	regio	n of human Caspase 3. The e	exact sequence is proprietary.
Purification	The	antibody was purified by imn	nunogen affinity chromatography.
Specificity	Reco	gnizes endogenous levels of	Caspase 3 protein.
Clonality	Poly	lonal	
Conjugation			
Form	Liqui	d in 0.42% Potassium phospl	nate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB (	1/500 - 1/1000), IH (1/50 - 1/2	00), IF/IC (1/50 - 1/200)
Gene Symbol	CASE	3	
Alternative Na	ames CPP3	2; Caspase-3; CASP-3; Apopa	in; Cysteine protease CPP32; CPP-32; Protein
	Yama	; SREBP cleavage activity 1;	5CA-1
Entrez Gene	836	Human); 12367 (Mouse); 25	402 (Rat)
SwissProt	P425	74 (Human); P70677 (Mouse	e); P55213 (Rat)
Storage/Stabi	lity Ship	oed at 4°C. Upon delivery alio	uot and store at -20°C for one year. Avoid
	freez	e/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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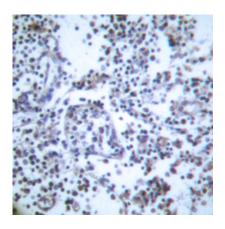
20

KDa A B C D E

For research purposes only, not for human use

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Western blot analysis of Caspase 3 expression in Raw264.7 (A), mouse brain (B), mouse lung (C), rat brain (D), rat lung (E) whole cell lysates. (Predicted band size: 31 kD; Observed band size: 35 kD)



Immunohistochemical analysis of Caspase 3 staining in human lymph node 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 Caspase 3 staining in NIH-3T3 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 Alexa Fluor 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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