

# **Product Data Sheet**

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

Catalog #	Source	Reactivity	Applications
CPA1139	Rabbit	H, M, R, D, Mk, P	WB, IH, IF/IC
Description	Ra	abbit polyclonal antibody to C	aspase 3
Immunogen	KL	LH-conjugated synthetic pepti	de encompassing a sequence within the center
	re	gion of human Caspase 3. The	e exact sequence is proprietary.
Purification	Tł	ne antibody was purified by in	nmunogen affinity chromatography.
Specificity	Re	ecognizes endogenous levels o	of Caspase 3 protein.
Clonality	Рс	olyclonal	
Conjugation			
Form	Lie	quid in 0.42% Potassium phos	phate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	ar	nd 0.01% sodium azide.	
Dilution	W	/B (1/500 - 1/1000), IH (1/50 - 1	/200), IF/IC (1/50 - 1/200)
Gene Symbol	CA	ASP3	
Alternative Na	ames CF	PP32; Caspase-3; CASP-3; Apo	pain; Cysteine protease CPP32; CPP-32; Protein
	Ya	ama; SREBP cleavage activity 1	l; SCA-1
Entrez Gene	83	36 (Human); 12367 (Mouse); 2	25402 (Rat)
SwissProt	PZ	42574 (Human); P70677 (Mou	ise); P55213 (Rat)
Storage/Stabi	<b>lity</b> Sh	nipped at 4°C. Upon delivery a	liquot and store at -20°C for one year. Avoid
	fre	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

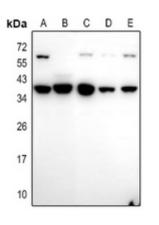
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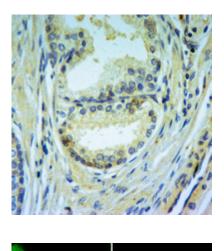




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Western blot analysis of Caspase 3 expression in SGC7901 (A), mouse kidney (B), A549 (C), Hela (D), LO2 (E) whole cell lysates. (Predicted band size: 31 kD; Observed band size: 35 kD)



Immunohistochemical analysis of Caspase 3 staining in human prostate 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.

Immunofluorescent analysis of Caspase 3 staining in HEK293T 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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