

## **Product Data Sheet**

## Anti-ZNF668 Antibody

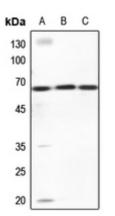
Catalog #	Source	Reactivity	Applications
CPA5146	Rabbit	H, M, R, B, Mk	WB, IH, IF/IC
Description	Rabb	it polyclonal antibody to	ZNF668
Immunogen	KLH-c	conjugated synthetic pep	tide encompassing a sequence within the center
	regio	n of human ZNF668. The	exact sequence is proprietary.
Purification	The a	ntibody was purified by i	mmunogen affinity chromatography.
Specificity	Reco	gnizes endogenous levels	of ZNF668 protein.
Clonality	Polyc	lonal	
Conjugation			
Form	Liquio	d in 0.42% Potassium pho	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and C	0.01% sodium azide.	
Dilution	WB (	1/500 - 1/1000) <i>,</i> IH (1/100	- 1/200), IF/IC (1/100 - 1/500)
Gene Symbol	ZNF6	68	
Alternative Na	ames Zinc f	inger protein 668	
Entrez Gene	7975	9 (Human); 244219 (Mou	ise)
SwissProt	Q96K	58 (Human); Q8K2R5 (M	ouse)
Storage/Stabi	lity Shipp	ed at 4°C. Upon delivery	aliquot 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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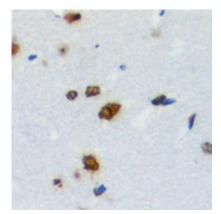
# Coherion



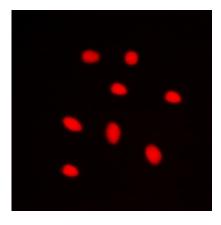
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Western blot analysis of ZNF668 expression in mouse lung (A), mouse kidney (B), rat kidney (C) whole cell lysates. (Predicted band size: 67 kD; Observed band size: 68 kD)



Immunohistochemical analysis of ZNF668 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 ZNF668 staining in HepG2 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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