

Product Data Sheet

Anti-Connexin 46 Antibody

Catalog #	Source	Reactivity	Applications		
CPA3962	Rabbit	H, M, R, D	WB, IH, IF/IC		
Description	Rabbit polyclonal antibody to Connexin 46				
Immunogen	KLH	KLH-conjugated synthetic peptide encompassing a sequence within the center			
	regi	region of human Connexin 46. The exact sequence is proprietary.			
Purification	The	antibody was purified	by immunogen affinity chromatography.		
Specificity Recognizes endogenous levels of Connexin 46 protein.					
Clonality	Poly	Polyclonal			
Conjugation					
Form	Liqu	id in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB	(1/500 - 1/1000), IH (1/5	0 - 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	GJA	3			
Alternative Na	ames Gap	junction alpha-3 prote	in; Connexin-46; Cx46		
Entrez Gene	270	0 (Human); 14611 (Mo	use); 79217 (Rat)		
SwissProt	Q9Y	6H8 (Human); Q64448	(Mouse); P29414 (Rat)		
Storage/Stabi	lity Ship	ped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid		
	free	ze/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

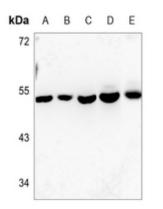
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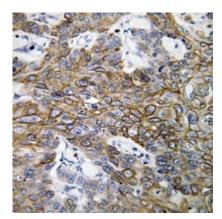




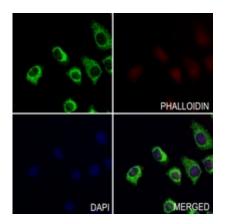
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Western blot analysis of Connexin 46 expression in A549 (A), U2OS (B), DLD (C), mouse heart (D), rat heart (E) whole cell lysates. (Predicted band size: 47 kD; Observed band size: 53 kD)



Immunohistochemical analysis of Connexin 46 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.



Immunofluorescent analysis of Connexin 46 staining in A549 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 AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin - AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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