

## **Product Data Sheet**

## Anti-FGF19 Antibody

Catalog #	Source	Reactivity	Applications	
CPA4644	Rabbit	Н	WB, IH, IF/IC	
Description	Ra	abbit polyclonal antibody	v to FGF19	
Immunogen	KL	H-conjugated synthetic	peptide encompassing a sequence within the C-term	
	re	gion of human FGF19. T	ne exact sequence is proprietary.	
Purification	Tł	ne antibody was purified	by immunogen affinity chromatography.	
Specificity	Re	ecognizes endogenous le	vels of FGF19 protein.	
Clonality	Pc	olyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	ar	nd 0.01% sodium azide.		
Dilution	W	/B (1/500 - 1/1000), IH (1/:	100 - 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	FC	GF19		
Alternative Na	ames Fi	broblast growth factor 1	9; FGF-19	
Entrez Gene		9965 (Human)		
SwissProt	09	95750 (Human)		
Storage/Stabi	<b>lity</b> Sh	nipped at 4°C. Upon deliv	ery aliquot 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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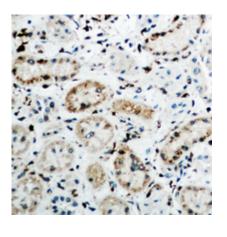
**kDa** A B 250

130

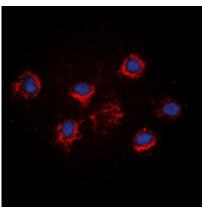
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Western blot analysis of FGF19 expression in HeLa (A), NCIH460 (B) whole cell lysates. (Predicted band size: 24 kD; Observed band size: 28 kD)



Immunohistochemical analysis of FGF19 staining in Human kidney 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 FGF19 staining in HeLa 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. DAPI was used to stain the cell nuclei (blue).

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