

## **Product Data Sheet**

## **Anti-IRF3 Antibody**

Catalog #	Source	Reactivity	Applications		
CQA6569	Rabbit	H, M, R	WB, IH, IF/IC		
Description	R	abbit polyclonal antibody	to IRF3		
Immunogen	К	LH-conjugated synthetic p	eptide encompassing a sequence of human IRF3. The		
	e	xact sequence is proprieta	ry.		
Purification	TI	he antibody was purified k	oy immunogen affinity chromatography.		
Specificity	els of IRF3 protein.				
Clonality	P	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	aı	nd 0.01% sodium azide.			
Dilution	W	VB (1/500 - 1/1000), IH (1/50	0 - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	IF	RF3			
Alternative Names		Interferon regulatory factor 3; IRF-3			
Entrez Gene	rez Gene 3661 (Human); 54131 (Mouse)				
SwissProt	Q	(14653 (Human); P70671 (	Mouse)		
Storage/Stabi	i <b>lity</b> Sl	hipped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid		
	fr	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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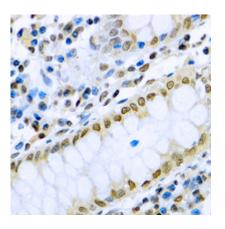
kDa A B

130 95

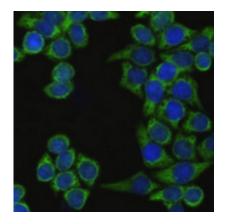
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Western blot analysis of IRF3 expression in Hela (A), NIH3T3 (B) whole cell lysates. (Predicted band size: 47 kD; Observed band size: 57 kD)



Immunohistochemical analysis of IRF3 staining in human colon 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 IRF3 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 488-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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