

Product Data Sheet

Anti-NF-kappaB p65 Antibody

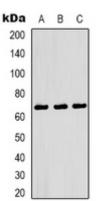
Catalog #	Source	Reactivity	Applications		
CPA9199	Mouse	H, M, R	WB, IH, IF/IC, IP		
Description	Μ	louse monoclonal antiboo	ly to NF-kappaB p65		
Immunogen	Re	ecombinant protein corre	sponding to human NF-kappaB p65.		
Purification					
Specificity	Re	ecognizes endogenous lev	els of NF-kappaB p65 protein.		
Clonality	Μ	Ionoclonal			
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 (1/2	00 - 1/500), IF/IC (1/100 - 1/200), IP (1/100 - 1/200)		
Gene Symbol	RI	ELA			
Alternative N	ames N	FKB3; Transcription factor	p65; Nuclear factor NF-kappa-B p65 subunit; Nuclear		
	fa	ctor of kappa light polype	ptide gene enhancer in B-cells 3		
Entrez Gene	59	970 (Human); 19697 (Mot	ıse)		
SwissProt	Q	04206 (Human); Q04207	(Mouse)		
Storage/Stabi	i lity Sł	nipped 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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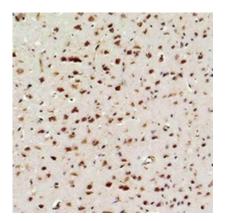
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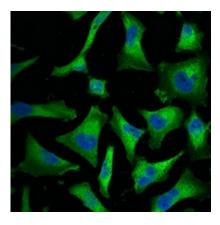
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Western blot analysis of NF-kappaB p65 expression in Hela (A), mouse spleen (B), rat heart (C) whole cell lysates. (Predicted band size: 60 kD; Observed band size: 65 kD)



Immunohistochemical analysis of NF-kappaB p65 staining in rat 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 NF-kappaB p65 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 FITC-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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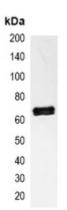
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Immunoprecipitation of NF-kappaB p65 from 0.5mg Hela whole cell extract lysate, using Anti-NF-kappaB p65 Antibody.



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