

Product Data Sheet

Anti-GBP3 Antibody

Catalog #	Source	Reactivity	Applications		
CPA1463	Rabbit	H, M, R	WB, IH, IF/IC		
Description	R	abbit polyclonal antibody t	o GBP3		
Immunogen	K	LH-conjugated synthetic pe	eptide encompassing a sequence within the C-term		
	re	egion of human GBP3. The	exact sequence is proprietary.		
Purification	T	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	R	ecognizes endogenous leve	els of GBP3 protein.		
Clonality	P	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	a	nd 0.01% sodium azide.			
Dilution	W	/B (1/500 - 1/1000), IH (1/50	- 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	G	BP3			
Alternative Na	<mark>ames</mark> G	uanylate-binding protein 3	; GTP-binding protein 3; GBP-3; Guanine		
	n	ucleotide-binding protein 3	3		
Entrez Gene	20	635 (Human)			
SwissProt	Q	9H0R5 (Human)			
Storage/Stabi	lity Sl	hipped at 4°C. Upon delive	ry 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

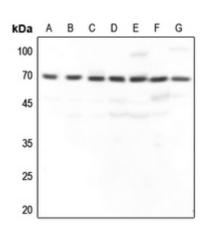
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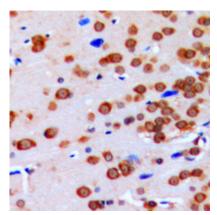
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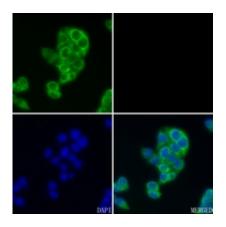
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Western blot analysis of GBP3 expression in HEK293T (A), Hela (B), H446 (C), mouse kidney (D), mouse lung (E), rat kidney (F), rat lung (G) whole cell lysates. (Predicted band size: 68 kD; Observed band size: 68 kD)



Immunohistochemical analysis of GBP3 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 GBP3 staining in MCF7 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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