

## **Product Data Sheet**

## **Anti-GBP1** Antibody

Catalog #	Source	Reactivity	Applications		
CPA4417	Rabbit	H, R, B	WB, IH, IF/IC		
Description	R	abbit polyclonal antibody t	o GBP1		
Immunogen	К	(LH-conjugated synthetic pe	ptide encompassing a sequence within the center		
	r	egion of human GBP1. The	exact sequence is proprietary.		
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	R	Recognizes endogenous leve	ls of GBP1 protein.		
Clonality	Р	Polyclonal			
Conjugation					
Form	L	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	nd 0.01% sodium azide.			
Dilution	V	VB (1/500 - 1/1000), IH (1/50	- 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	G	GBP1			
Alternative Na	a <mark>mes</mark> li	nterferon-induced guanylat	e-binding protein 1; GTP-binding protein 1; GBP-1;		
	F	luGBP-1; Guanine nucleotic	e-binding protein 1		
Entrez Gene	2	633 (Human)			
SwissProt	P	232455 (Human); Q01514 (N	/louse)		
Storage/Stabi	lity S	hipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid		
	f	reeze/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

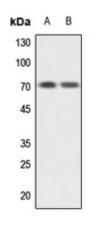
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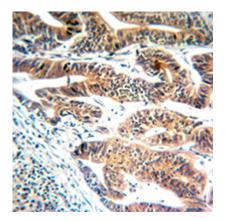
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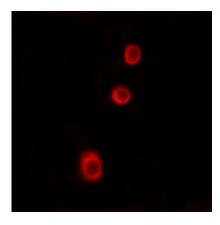
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Western blot analysis of GBP1 expression in HEK293T (A), H446 (B) whole cell lysates. (Predicted band size: 67 kD; Observed band size: 68 kD)



Immunohistochemical analysis of GBP1 staining in human colon 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 GBP1 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 AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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