

Storage/Stability

## **Product Data Sheet**

# Anti-G3BP1 (Phospho-S232) Antibody

freeze/thaw cycles.

Catalog #	Source	e Reactivity	Applications
CPA2456	Rabbit	: Н	WB, IH, IF/IC
Description		Rabbit polyclonal antibo	dy to G3BP1 (Phospho-S232)
Immunogen		KLH-conjugated syntheti	c phosphopeptide corresponding to residues surrounding
		S232 of human G3BP1 p	rotein. The exact sequence is proprietary.
Purification		The antibody was purific	ed by immunogen affinity chromatography.
Specificity		Recognizes endogenous	levels of G3BP1 protein only when phosphorylated at S232.
Clonality		Polyclonal	
Conjugation			
Form		Liquid in 0.42% Potassiu	m phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
		and 0.01% sodium azide	
Dilution		WB (1/500 - 1/1000), IH (	1/100 - 1/200), IF/IC (1/100 - 1/500)
Gene Symbol		G3BP1	
Alternative Na	ames	G3BP; Ras GTPase-activa	ating protein-binding protein 1; G3BP-1; ATP-dependent DNA
		helicase VIII; hDH VIII; G	AP SH3 domain-binding protein 1
Entrez Gene		10146 (Human)	
SwissProt		Q13283 (Human)	

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

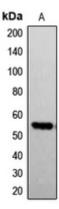
Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

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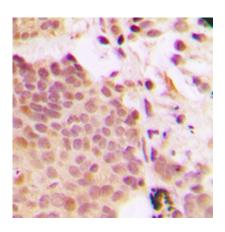
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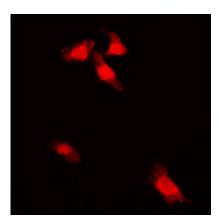
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Western blot analysis of G3BP1 (Phospho-S232) expression in A549 (A) whole cell lysates. (Predicted band size: 52 kD; Observed band size: 55 kD)



Immunohistochemical analysis of G3BP1 (Phospho-S232) staining in human breast 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 G3BP1 (Phospho-S232) staining in A549 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 humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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