

Product Datasheet

Anti-GAB1 (Phospho-Y627) Antibody (orb256545)

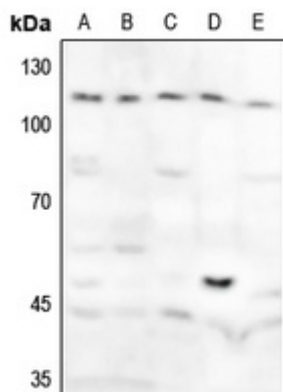
Description	Rabbit polyclonal antibody to GAB1 (phospho-Y627)
Species/Host	Rabbit
Reactivity	Human, Mouse, Rat
Conjugation	Unconjugated
Tested Applications	IH, WB
Immunogen	KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding Y627 of human GAB1 protein. The exact sequence is proprietary.
Target	GAB1
Preservatives	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.
Form/Appearance	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.
Storage	Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
Note	For research use only
Clonality	Polyclonal
Source	Rabbit
Uniprot ID	Q9QYY0, Q13480
Entrez	14388, 2549
Dilution Range	WB: 1:500-1000, IF/ICC: 1:100-500
Expiration Date	12 months from date of receipt.

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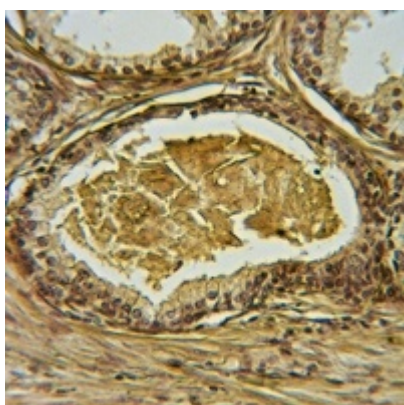
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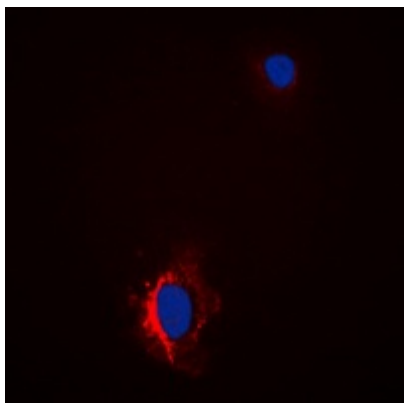
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Western blot analysis of GAB1 (Phospho-Y627) expression in HeLa (A), U2OS (B), mouse lung (C), mouse testis (D), rat lung (E) whole cell lysates. (Predicted band size: 76 kD; Observed band size: 110 kD)



Immunohistochemical analysis of GAB1 (Phospho-Y627) staining in human prostate formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (Phospho-H 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 GAB1 (Phospho-Y627) 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 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. DAPI was used to stain the cell nuclei (blue).

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